GLOBAL AIDS RESPONSE PROGRESS REPORTING 2014

Construction of Core Indicators for monitoring the 2011 United Nations Political Declaration on HIV and AIDS



Includes additional WHO/UNICEF Universal Access Health Sector Indicators







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Construction of Core Indicators for monitoring the 2011 United Nations Political Declaration on HIV and AIDS

Includes additional WHO/UNICEF Universal Access Health Sector Indicators

Please use the Global AIDS Response Progress Reporting website (www.unaids.org/AIDSReporting) to submit your indicator data by 31 March 2014.

Modelled HIV estimates using the updated Spectrum software are also due by 31 March 2014.

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Foreword

In 2013, the United Nations General Assembly conducted a midterm review of progress towards reaching the bold targets and commitments of the 2011 United Nations Political Declaration on HIV and AIDS. I congratulate the 26 countries which reported in 2013 that they succeeded in reducing by half the number of new HIV infections since 2001. Yet, the international community has committed to reach Millennium Development Goal 6 to "halt by 2015 and begin to reverse the spread of HIV and AIDS". This demands that we redouble our efforts in many more countries to identify gaps and take action rapidly to halt and reverse the HIV epidemic by the end of 2015.

These guidelines provide UN Member States with detailed information on how to conduct the next round of global AIDS response progress reporting in 2014. I encourage all countries to use this opportunity to consult with key country constituents, including civil society, on how to sharpen the national AIDS response. This round of reporting also provides the occasion to reprogramme efforts with development partners, including with the Global Fund to Fight AIDS, Tuberculosis and Malaria in preparation for its new funding model. It is only by demonstrating results and accelerating our progress that we can create the momentum needed to ensure that AIDS remains a priority in the post-2015 development agenda.

The response rate for global AIDS reporting has been exceptionally high: during the 2012 reporting round 186 out of 193 UN Member States (96%) submitted reports to UNAIDS. I encourage all Governments to sustain this effort and report on progress toward the commitments outlined in the 2011 United Nations Political Declaration on HIV and AIDS (General Assembly resolution 65/277).

Collecting and reporting high-quality results on the AIDS response are important elements of our agenda for shared responsibility and global solidarity. UNAIDS is determined to support you in this endeavour and has prepared these guidelines toward this end. I invite you to submit your monitoring data, HIV estimates and a narrative report for the year 2013 by 31 March 2014. The results of your next round of reporting will be used to inform several reports in 2014, including the *UNAIDS report on the global AIDS epidemic*.

If you have any questions or if you need additional support, do not hesitate to contact us at AIDSreporting@unaids.org.

I thank you for your continued engagement in the AIDS response. UNAIDS is available to provide any assistance you or your Government may require to monitor, review and report on progress towards the 2011 UN Political Declaration on HIV and AIDS and the achievement of Millennium Development Goals.

Michel Sidibé Executive Director UNAIDS

Acronyms

AIDS Acquired Immunodeficiency Syndrome

ANC Antenatal Clinic(s)
ART Antiretroviral therapy

BSS Behavioural Surveillance Survey
DHS Demographic and Health Survey

EID Early Infant Diagnosis

HIV Human Immunodeficiency Virus

IDU Injecting drug user/people who inject drugs (latter preferred language)

ILO International Labour Organization

MC Male circumcision

MTCT Mother-to-Child Transmission
MDG Millennium Development Goals
MICS Multiple Indicator Cluster Survey
MSM Men who have sex with men
M&E Monitoring and evaluation

NA Not Applicable

NAC National AIDS Committee(s) NAP National AIDS Programme

NASA National AIDS Spending Assessment NGO Nongovernmental Organization(s)

NSP National Strategic Plan

NSP Needle and Syringe Programmes

OECD Organisation for Economic Co-operation and Development

PLHIV People Living with HIV

PMTCT Prevention of Mother-to-Child Transmission

PRSP Poverty Reduction Strategy Paper

PWID People who inject drugs

STI Sexually Transmitted Infection(s)

TB Tuberculosis
UN United Nations

UNAIDS Joint United Nations Programme on HIV/AIDS
UNDAF United Nations Development Assistance Framework

UNFPA United Nations Population Fund

UNGASS United Nations General Assembly Special Session on HIV and AIDS

UNICEF United Nations Children's Fund WHO World Health Organization

2011 United Nations General Assembly Political Declaration on HIV and AIDS

Targets and elimination commitments







NT HIV ELIMINATE N DNG HIV INFECTI USERS AMONG CHIL





















Introduction

Purpose

The purpose of this document is to provide guidance to national AIDS programmes and partners actively involved in the country response to AIDS on use of core indicators to measure and report on the national response.

The "2011 UN Political Declaration on HIV and AIDS: Intensifying our Efforts to Eliminate HIV and AIDS" (General Assembly resolution 65/277), which was adopted at the United Nations General Assembly High Level Meeting on AIDS in June 2011, mandated UNAIDS to support countries to report on the commitments in the 2011 UN Political Declaration on HIV and AIDS.

The Global AIDS Response Progress Reporting (GARPR) indicators (before 2012 known as UNGASS indicators) were until 2012 reported at the global level every second year. However, from 2013 data is collected every year.

To assess progress made against the targets, the collection and reporting of indicator data is an important part. Countries are strongly encouraged to integrate these core indicators into their on-going monitoring and evaluation activities. These indicators are designed to help countries assess the current state of their national response and progress in achieving their national HIV targets. They will contribute to a better understanding of the global response to the HIV pandemic, including progress towards the global targets set in the 2011 UN Political Declaration on HIV and AIDS and the Millennium Development Goals.

These guidelines are designed to improve the quality and consistency of data collected at the country level, thus enhancing the accuracy of conclusions drawn at national, regional and global levels.

How to use these guidelines

These guidelines have been developed to help countries collect data and report on their national HIV response as effectively as possible. In the section "Core indicators for Global AIDS Response Progress Reporting" readers will find pages devoted to each indicator, giving reasons for inclusion and methods for collecting, constructing and measuring the indicator. The indicator's strengths and weaknesses are also discussed.

Help is available at every stage of the process. Key points and sources for additional information—including who to contact and how to reach them—is highlighted in this introductory section and pointed out with a orange arrow.

Background

We are rapidly approaching 2015, the end date of both the 2011 *Political Declaration on HIV and AIDS* and the Millennium Development Goals (MDGs), and this is an important opportunity to review progress and start preparing for the final reporting towards these targets.

The 2011 UN Political Declaration on HIV and AIDS builds on two previous political declarations: the 2001 *Declaration of Commitment on HIV/AIDS* and the 2006 *Political Declaration on HIV/AIDS*. At United Nations General Assembly Special Session on HIV/AIDS (UNGASS), in 2001, the declaration was adopted unanimously by the member states. This declaration reflected global consensus on a comprehensive framework to achieve Millennium Development Goal 6: halting and beginning to reverse the HIV epidemic by 2015. It recognized the need for multisectoral action on a range of fronts and addressed global, regional and country-level responses to prevent new HIV infections, expand health care access and mitigate the epidemic's impact. The 2006 declaration recognized the urgent need to achieve universal access to HIV treatment, prevention, care and support.

While the declarations have been adopted by governments, their vision extends far beyond the governmental sector to private industry and labour groups, faith-based organizations, nongovernmental organizations and other civil society entities, including organizations representing people living with HIV.

As indicated in the 2011 UN Political Declaration on HIV and AIDS, a successful AIDS response should be measured by the achievement of concrete, time-bound targets. It calls for careful monitoring of progress in implementing commitments and requires the United Nations Secretary-General to issue annual progress reports. These reports are designed to identify challenges and constraints and recommend action to accelerate achievement of the targets.

The guidelines in this document have been developed to enhance reporting of key indicators for the AIDS response. The reported data are used to monitor progress against the commitments and targets of the 2011 UN Political Declaration on HIV and AIDS and the AIDS related MDGs.

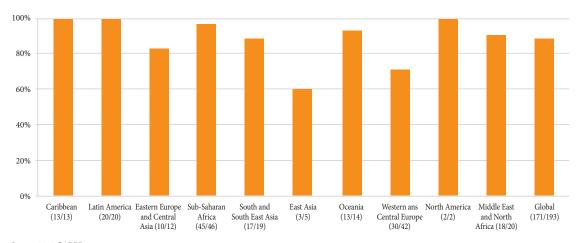
2011 UN Political Declaration on HIV and AIDS – Targets and elimination commitments

- 1. Reduce sexual transmission by 50% by 2015
- 2. Reduce transmission of HIV among people who inject drugs by 50% by 2015
- 3. Eliminate new HIV infections among children by 2015 and substantially reduce AIDS-related maternal deaths
- 4. Reach 15 million people living with HIV with lifesaving antiretroviral treatment by 2015
- 5. Reduce tuberculosis deaths in people living with HIV by 50% by 2015
- 6. Close the global AIDS resource gap by 2015 and reach annual global investment of US\$ 22-24 billion in low- and middle-income countries
- 7. Eliminate gender inequalities and gender-based abuse and violence and increase the capacity of women and girls to protect themselves from HIV
- 8. Eliminate stigma and discrimination against people living with and affected by HIV through promotion of laws and policies that ensure the full realization of all human rights and fundamental freedoms
- 9. Eliminate HIV-related restrictions on entry, stay and residence
- 10. Eliminate parallel systems for HIV-related services to strengthen integration of the AIDS response in global health and development efforts

Reporting history

UNAIDS has collected country progress reports from Member States for the purpose of monitoring the various political declarations every two years since 2004 and every year since 2013. Response rates increased from 102 (53%) Member States in 2004 to 186 (96%) in 2012. In 2013, the first year with yearly reporting, a slight decrease in response rates was noted with 171 Member States submitting reports (89%).

Proportion of countries that have participated in the 2013 Global AIDS Response Progress Reporting



Source: 2013 GARPR country reports (Countries reporting/total number of UN Member States in the region)

The information provided by country progress reports represents the most comprehensive data on both the status of, and response to the epidemic. The data from the previous reporting rounds are available online through AIDSinfo; aidsinfo.unaids.org. The full data base is available at www.aidsinfoonline.org, which can be used to produce charts, maps and tables. Unedited narrative country reports from the 2012 reporting are available at

www.unaids.org/en/dataanalysis/knowyourresponse/countryprogressreports/2012countries. Full NCPI reports are available at

www.unaids.org/en/dataanalysis/knowyourresponse/ncpi/2012countries.

Reporting format

When preparing Global AIDS Response Progress Reporting, countries should base their narrative reports on their national country reports. Where a recent national country report is available, this can be submitted as the narrative country progress report. A Country Progress Report template, with detailed instructions for completion of the different sections can be found in Appendix 1. The indicator data are considered an integral part of each Country Progress Report submission. Hence, both the narrative part of the Country Progress Report and the indicator data should be considered in the consultation and report preparation process as outlined in the section titled "Implementation of progress reporting at national level" on page 12 of these guidelines.



The deadline for report submission using the reporting website is 31 March 2014.

INTRODUCTION

Global AIDS Response Progress Reporting indicators are important for two reasons. First, they can help individual countries evaluate the effectiveness of their national response and second, when data from multiple countries are analysed collectively, the indicators can provide critical information on the effectiveness of the response at regional and global levels, and will be the basis for the regional and global analyses of progress towards the 2015 targets. At the same time this provides countries with insights into other national-level responses.

The changes in this round of reporting compared to the 2013 reporting round are summarized on page 19.

Countries should consider the applicability of each indicator to their epidemic. When countries choose not to report on a particular indicator, they should provide their reasons for choosing not to report as this enables differentiation between an absence of data and the inapplicability of specific indicators to particular country epidemics.

Most of the national indicators are applicable to all countries. The behaviour indicators for key populations at higher risk are relevant in countries with low-level and concentrated epidemics as well as countries with generalized epidemics. Similarly, countries with a concentrated epidemic are encouraged to collect data on sexual behaviours among young people as a means of tracking trend changes that could influence the national response in the future. However, a few are applicable to specific HIV epidemic contexts only.

UNAIDS strongly recommends that countries use these indicators within their national monitoring and evaluation systems. In accordance with specific needs, and if resources allow, countries may wish to include additional indicators in their national monitoring plans.

Five of the national indicators are also Millennium Development Goal indicators:

- percentage of young people who are living with HIV
- knowledge among young people about HIV
- condom use at last high-risk sex
- school attendance among orphans
- antiretroviral therapy coverage.

Data used by the UN Division of Statistics for reporting on the Millennium Development Goals are mainly sourced from data provided by Member States through Global AIDS Response Progress Reporting.



Full definitions for all indicators used for the Global AIDS Response Progress Reporting can be found in these guidelines. The indicators can also be found in the UNAIDS Indicator Registry at www.indicatorregistry.org. This online database provides complete definitions of the Global AIDS Response Progress Reporting indicators and clearly shows how these indicators relate to indicators used by WHO, UNAIDS, PEPFAR, the Global Fund and other key partners. The Indicator Registry also includes other HIV indicators used at country level. There are direct links from the online reporting tool to the indicators in the Indicator Registry. The indicators can also be exported from the Indicator Registry to Excel, Word or PDF.

National indicators for high-income countries

In adopting the 2011 Political Declaration on HIV and AIDS, high-income countries have committed themselves to reporting on progress made in their national responses to HIV. It is recognized that high-income countries may use relatively complex information systems and a variety of data sources which can make the calculation of a single national indicator challenging. However, this does not remove the need for high-income country data for monitoring global progress towards the targets of the Political Declaration on HIV and AIDS. European Union/European Economic Area (EU/EEA) countries have used innovative ways to link global HIV monitoring systems more closely to regional circumstances.

UNAIDS encourages other high-income countries to contact either the UNAIDS Strategic Information and Monitoring Division (AIDSreporting@unaids.org) if they require further technical advice regarding reporting on their domestic programmes, or the European Centre for Disease Prevention and Control (Dublindeclaration@ecdc.europa.eu) to learn more about the European approach of linking and harmonising global and regional monitoring systems.

Monitoring the Dublin Declaration on partnership to fight HIV and AIDS in Europe and Central Asia

In 2004, during the Irish Presidency of the Council of the European Union member states and neighbouring countries in Eastern Europe and Central Asia adopted the Dublin Declaration on Partnership to Fight HIV and AIDS in Europe and Central Asia. The European Centre for Disease Prevention and Control (ECDC) is coordinating reporting on this Declaration. UNAIDS and the ECDC are working closely to coordinate and harmonize global and regional monitoring systems. This should both improve the relevance of reporting for countries in the region and reduce their reporting burden. Data collection will take place at the same time as for the current Global AIDS Response Progress Reporting. In collaboration with UNAIDS, the ECDC will provide technical assistance to European countries preparing reports for both Global AIDS Response Progress Reporting 2014 and the Dublin Declaration 2014. For further information on the monitoring of the Dublin Declaration please contact Teymur Noori at the ECDC (teymur.noori@ecdc.europa.eu).

Implementation of progress reporting at national level

Indicator construction

For each indicator this manual provides the information needed to construct the indicator, including:

- summary of what it measures
- rationale for the indicator
- numerator, denominator and calculation
- recommended measurement tools
- measurement frequency
- strengths and weaknesses of the indicator (including summary interpretation of the indicator).

Measurement tools and data sources

The primary measurement tools vary by indicator and include:

- nationally representative, population-based sample surveys
- behavioural surveillance surveys
- specially designed surveys and questionnaires, including surveys of specific population groups (e.g. specific service coverage surveys)
- patient tracking systems
- health information systems
- sentinel surveillance
- national HIV estimates from Spectrum software
- the National Commitments and Policy Instrument (NCPI) questionnaire.

Existing data sources, including records and programme reviews from health facilities and schools as well as specific information from HIV surveillance activities and programmes, should be used to supplement the primary measurement tools.

Another source for denominators used in the GARPR reporting is the **Spectrum computer package** that allows countries to create population-level estimates of people in need of antiretroviral therapy, women in need of antiretroviral medicine and HIV-exposed children in need of virological testing.

In 2014 the process of completing the Spectrum file and submitting the GARPR data will be done simultaneously to ensure the results are harmonized. Initial Spectrum files should be provided to UNAIDS for review by 28 February so the files can be finalized before the GARPR and Spectrum submission due date of 31 March 2014. Instructions on how to update and submit Spectrum files will be sent to national HIV estimates teams in January 2014.

Spectrum files are created by a team of national experts who have been trained on how to populate and use the software. It is critical that the team completing the GARPR tool use the most-recent estimates developed by the national HIV estimates team.

Civil society organizations are valuable sources of data for many indicators, especially those that relate to interventions where nongovernmental, faith-based and community-based organizations play an active role. Examples include work with young people, key populations at higher risk and pregnant women.

In many countries, the bulk of the data required for the core national-level indicators may not be available from existing sources. Gathering such data is likely to require the adaptation of existing monitoring tools

or the addition of specific surveys. Countries that conduct regular, nationally representative, population based surveys such as the Demographic and Health Surveys or AIDS Indicator Surveys will collect important information, including behavioural data on young people. In countries where other types of population-based surveys are conducted, including those for purposes other than HIV, it is possible to adapt these surveys to collect data for selected core indicators.

Numerators and denominators

For each core indicator, detailed instructions for measuring the national response are provided. Most core national-level indicators use numerators and denominators to calculate the percentages that measure the current state of the national response. Countries are strongly encouraged to pay close attention to the dates attached to specific data when calculating an indicator. If data used for the numerator and denominator are collected at different times, the accuracy and validity of that information will be compromised.

The methods described have been designed to facilitate the construction of global estimates from national-level data. While these methods can be applied at the subnational level, simpler, faster and more flexible approaches that are tailored to local conditions may be more appropriate to guide decision-making below the national level.

A number of indicators related to coverage of services require a denominator that is based on the full population, i.e. not just those people that are seen at health care clinics. To calculate population-level indicators it is necessary to estimate the total number of people eligible for the service. For example, to estimate how close a country is to reaching 100% MTCT coverage it is necessary to estimate the total number of pregnant women living with HIV. UNAIDS recommends that countries use the Spectrum computer package to calculate the denominators needed for GARPR reporting.

Disaggregated data: sex and age

One of the key lessons learnt from previous rounds of reporting was the importance of obtaining disaggregated data, for example, breaking it down by sex and age. It is vital that countries collect data in their component parts and not simply in summary form. Without disaggregated data, it is difficult to monitor the breadth and depth of the response to the epidemic at both national and global levels. It is equally difficult to monitor access to activities, the equity of that access, the appropriateness of focusing on specific populations, and meaningful change over time.

Countries are strongly encouraged to make the collection of disaggregated data, especially by sex and age, one of the cornerstones of their monitoring and evaluation efforts. If possible, equity analyses should also be done. Key ministries should review their information systems, surveys and other instruments for collecting data to ensure that they capture disaggregated data at subnational levels, including facility and project levels. Special efforts should be made to follow disaggregated data up to the national level. In addition, the private sector and/or civil society organizations involved in the country's AIDS response must be advised of the importance of disaggregated data and make the collection and dissemination of the data a priority in their on-going operations.

The Global AIDS Response Progress Reporting website (www.unaids.org/AIDSreporting) clearly identifies the disaggregated data that are required to accurately report on the numerator and denominator for each indicator (see the preceding subsection entitled "Numerators and Denominators" for additional information). In general, where appropriate, all data should be disaggregated by sex and age. Where collecting disaggregated data has proved difficult, entry of partial data is possible, where necessary.

In situations where disaggregated data are not readily available, it may be possible to extract the information needed for core indicators from larger data sets, although the location of the data will vary from country to country. Countries should seek technical assistance from the United Nations System (including the UNAIDS, WHO and UNICEF country offices) and its partners for help with accessing the disaggregated data needed to properly complete the measurements of core indicators.

IMPLEMENTATION AT NATIONAL LEVEL

Governments are encouraged to look beyond their internal information resources to both collect and validate data. In many cases, civil society organizations may be able to provide valuable primary and secondary data.

Recent and representative survey data

Use the most-recently available nationally representative survey to calculate indicators that are based on general population surveys. This may mean that the data reported in this round will be the same as the data reported in the previous round, since such surveys are generally undertaken at five year intervals.

When more recent data is not available, survey data already reported in 2013 can be loaded in the on-line reporting tool at a click of a button and, therefore, do not need to be re-entered.

Ensuring that survey samples of key populations are truly representative is a great technical challenge.

Methods are being developed to try to achieve representative sampling of these populations (e.g. respondent-driven sampling). While these are being refined, it is recognized that countries may not be confident that samples used for surveys of key populations at higher risk of HIV exposure are representative. Countries are advised to use the most-recent survey of key populations that has been reviewed and endorsed by local technical experts, such as monitoring and evaluation technical working groups or national research councils. Countries are encouraged to report all recent, quality surveys of key populations, by site, with numerator, denominator and methods in the provided Excel spread sheets.

One of the challenges in developing burden of disease estimates and planning for programme needs is understanding the size of key populations. Countries are asked to report the size estimates for key populations, providing methods and any city/province-specific estimates calculated empirically. More details can be found on page 49. Some countries with empirical national size estimates for key populations are also able to aggregate prevention programme data. If a country can report against an indicator with national programme data, they may do so in the comment fields this year.



Countries needing additional information on implementation should seek technical assistance from their UNAIDS Strategic Information Advisers, UNICEF or WHO offices and HIV monitoring and evaluation working groups. Technical support is also available from the UNAIDS Regional Strategic Information Advisers based at the Regional Support Team and from the Strategic Information and Monitoring Division Team at the UNAIDS Secretariat who can be reached via email at AIDSreporting@unaids.org.

Interpretation and analysis

As each core indicator is discussed later in this manual, so too are their strengths and weaknesses. Countries should carefully review this section before they begin collecting and analysing data as it explains how to interpret each indicator and any potential issues related to it. The points raised in this section should be reviewed before finalization of the reporting and the writing of the narrative report to confirm the appropriateness of the findings for each indicator.

The sections on the strengths and weaknesses of each core indicator are designed to improve the accuracy and consistency of the data submitted to UNAIDS. Other points in this section provide additional information on the value of a particular indicator. The section acknowledges that variations may occur from country to country on issues as diverse as the relationship of costs to local income, standards for quality and variations in treatment regimens.

After compiling their data countries are strongly encouraged to continue analysing their findings. This will enable them to better understand their national response and identify opportunities to improve that response. Countries should be looking closely at the linkages between policy, implementation of HIV programmes, verifiable behaviour change and changes in the epidemic. For example, if a country has a policy on the reduction of mother-to-child transmission of HIV, does it also have field programmes that make prevention of mother-to-child transmission available to pregnant women? If these field programmes are in place, are women using them in sufficient numbers to have an impact on the number of HIV-infected infants born in that country?

These linkages exist in every facet of a national response and many of the most important ones are reflected in the core national-level indicators included in this manual. To effectively analyse these linkages, countries must draw on the widest range of data available, including quantitative and qualitative information from both the public and private sectors. An over-reliance on data of any one type or from any one source is less likely to provide the perspective or insights required to understand such linkages and to identify any existing or emerging trends.

Selection of indicators

Based on knowing the local HIV epidemic, countries should review all of the indicators to determine which ones are applicable in their situation. For example, a country with a concentrated epidemic among sex workers and men who have sex with men may not need to report on the core indicators related to people who inject drugs. However, they should regularly assess the situation to see whether injecting drug use is emerging as an issue that needs attention. They should calculate both the specific indicators for sex workers and men who have sex with men as well as broader indicators (e.g. young people's knowledge of HIV, higher-risk sex in women and men, and condom use during higher-risk sex), which are relevant in tracking the spread of HIV into the general population.

Similarly, countries with a generalized epidemic should include data on as many indicators as possible for key populations at higher risk. For example, a country with a higher-prevalence epidemic may also have a concentrated sub-epidemic among people who inject drugs. It would therefore be valuable to also calculate and report on the indicators that relate to the key populations at higher risk.

For each indicator, countries are requested to state the indicator relevance in the on-line reporting tool, depending on the epidemic situation in the country and if data is available. If it is felt that the area is relevant to the epidemic and response, but that the indicator itself is not relevant or appropriate for monitoring this issue, this should be stated in the online reporting tool comment boxes.

If a country is using an alternative indicator to effectively monitor the issue in question the comment boxes may be used to describe it (including a full definition and method of measurement), along with any available data for the indicator.

Geo-coding surveillance and monitoring and evaluation information

Through the Global AIDS Response Progress Reporting mechanism countries are asked to submit nationally representative data. However at the national level identifying geographic areas where localized HIV epidemics or specific populations most affected by the epidemic are not being reached by services is a key opportunity to strengthen the efficiency and effectiveness of national HIV responses. This is possible by attaching geographic information to indicator data. Geo-coding links surveillance and programmatic data from various sources to produce more detailed understandings of the HIV epidemic, facilitating implementation of focused and adapted interventions where they are most needed. To implement this approach, data collection must shift to sub-national levels that are programmatically relevant. Data collection is already expanding in many countries to lower geographical levels and among key populations. Confidentiality and ethical considerations must always be maintained in data collection, analysis and dissemination, to ensure geo-coded data are used to bring HIV-related services closer to the people who need them and not expose people to harm. Please see Appendix 7 for more information about geo-coding surveillance and M&E information.

Role of civil society

Civil society plays a key role in the response to the AIDS epidemic in countries around the world. The wide range of expertise within civil society organizations makes them ideal partners in the process of preparing Country Progress Reports. Specifically, civil society organizations are well positioned to provide quantitative and qualitative information to augment the data collected by governments. National AIDS councils/commissions committees or their equivalents should seek input from the full spectrum of civil society, including nongovernmental organizations, networks of people living with HIV, faith-based organizations, women, young people, trade unions and community-based organizations, for their reports

IMPLEMENTATION AT NATIONAL LEVEL

on the core national-level indicators underlying the 2011 UN Political Declaration on HIV and AIDS. The importance of securing input from the full spectrum of civil society, including people living with HIV, cannot be overstated. Civil society speaks with many voices and represents many different perspectives, all of which can be valuable in the monitoring and evaluation of a country's AIDS response.

National AIDS Committees or their equivalents should provide civil society organizations with easy access to their plans for data collection and denominator data. A straightforward mechanism for submitting and evaluating information should be developed. Mid-term review reports should include data from civil society service providers. As part of this effort, civil society organizations should also be invited to participate in workshops at the national level to determine how they can best support the country's reporting process. In every country civil society representatives should be given sufficient opportunity to review and comment on data before it is finalized and submitted. The report that is submitted to UNAIDS should be widely disseminated to ensure that civil society has ready access to it.

Country-level UNAIDS staff are available to assist with civil society input throughout the process. In particular, UNAIDS country-level staff should:

- brief civil society organizations on the indicators and the reporting process
- provide technical assistance on gathering, analysing and reporting data, including focused support to people living with HIV
- ensure the dissemination of reports including, whenever possible, reports in national languages.

Shadow reports by civil society will be accepted by UNAIDS as they were in previous rounds. It must be noted that shadow reports are not intended as a parallel reporting process for civil society. Wherever possible UNAIDS encourages civil society integration into national reporting processes, as described above. Shadow reports are intended to provide an alternative perspective where it is strongly felt that civil society was not adequately included in the national reporting process, where governments do not submit a report, or where data provided by government differs considerably from data collected by civil society monitoring government progress in service delivery.

Report contents

In 2014, countries are expected to submit data on all of the national indicators that are applicable to their response. National governments are responsible for reporting on national-level indicators with support from civil society and, where applicable, development partners. The procedures outlined in this manual should be used for collecting and calculating the necessary information for each indicator.

Countries are also requested, when possible, to submit copies of or links to primary reports from which data is drawn for the different indicators. These reports can be submitted through the online global reporting tool. This will facilitate the analyses of the data including trend analyses and comparisons between countries.

As discussed previously, and as required by the 2011 UN Political Declaration on HIV and AIDS, civil society, including people living with HIV, should be involved in the reporting process. The private sector at large should have a similar opportunity to participate in the reporting process. UNAIDS strongly recommends that national governments organize a workshop or forum to openly present and discuss the data before it is submitted. Joint United Nations Teams on AIDS are available in many countries to facilitate this discussion process.

The indicator data will be made available after a process of data cleaning, validation and reconciliation at www.AIDSinfo.com.

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If there are any questions, countries are advised to consult with UNAIDS locally or in Geneva at AIDSreporting@unaids.org. Updated information on Global AIDS Response Progress Reporting is available on the UNAIDS web site at: www.unaids.org/AIDSreporting.

Guidance on submission

- Countries needing additional information on the reporting tool and the submission mechanisms should seek technical assistance from their UNAIDS Strategic Information Advisers and HIV monitoring and evaluation working groups in country. The Strategic Information and Monitoring Division at the UNAIDS Secretariat is also available to provide support and can be reached via email at AIDSreporting@unaids.org.
- To facilitate contact with UNAIDS Geneva during the reporting process and followup, countries are requested to provide the name and contact details of the individual responsible for submitting the data as early as possible to AIDSreporting@unaids.org.

Reporting tool

Countries are asked to submit their data using the online global reporting tool found at http://AIDSreportingtool.unaids.org. Each country has an assigned national focal point that will be responsible for accessing this tool and entering their country information for submission. Countries may add/assign multiple rapporteurs in case data is provided from several sources and reporting structures.

As a new feature in the reporting tool, at the first log-on each country rapporteur is requested to create their username and password. Based on official communication with the country, one data editor is initially assigned per country, but the country rapporteur can extend these rights to others if he/she wishes to do so. Editors are able to add and make changes to the information to be submitted. As in the past years, the country rapporteur can also enable other people to view the data, allowing for broader country consultation. Viewers are able to see the information that will be submitted, yet make no changes to it. More details on this will be provided in the online presentations (www.unaids.org/AIDSreporting) on how to use the reporting tool.

As mentioned above, where countries do not submit data on an indicator, they should indicate whether this was due to an absence of appropriate data or because the indicator was not considered relevant to the epidemic. The comment boxes should be used for short explanatory notes stating how the numerator and denominator were calculated and assessing the accuracy of the composite and disaggregated data. For country level review, the data can also be printed out as one file if needed.

Progress in the reporting can be assessed in the main page, viewing the percentage or number of indicators being responded to. In addition to entering the current year data, countries may request to modify their past year's data if necessary. This will also be done through the online tool.

At the end, the data entry is finished by clicking the "submit" button. This closes the country's session in the online global reporting tool. The country will no longer be able to make editing changes or additions to its submission using this tool. UNAIDS will review the data and ask for clarifications if necessary. If there are queries to the data, the site will be opened again for the countries to edit their responses.

Problems with the online global reporting tool can be reported to AIDSreporting@unaids.org.

Joint reporting with WHO and UNICEF on health sector indicators

To minimize the reporting burden and facilitate the reporting process the Global AIDS Response Progress Reporting and the WHO and UNICEF health sector indicators will, as in the previous reporting round, be collected through the same online reporting tool.

The additional health sector indicators can be found in Part II of the guidelines.

For specific questions regarding these additional indicators, please e-mail: hivstrategicinfo@who.int.

Data submission

The indicator data should be submitted online by 31 March 2014. The national HIV estimates team should upload the Spectrum file to the designated folder (supplied by UNAIDS HIV Estimates team) by 31 March 2014. For any questions related to where to upload the file please contact estimates@unaids.org.

The indicator data should be entered online and the narrative report uploaded using the global reporting website (www.unaids.org/AIDSreporting). This will facilitate data processing and minimize errors.

Printed copies of reports may be sent to: Division Chief, Strategic Information and Monitoring Division, UNAIDS, 20 Avenue Appia CH-1211, Geneva 27, Switzerland.

The national-level reporting process: necessary actions

Complete reporting on the core indicators is essential if the reporting is to contribute to the global response to the epidemic. Countries are strongly encouraged to establish timetables and milestones for completing the necessary tasks. Listed below are necessary actions to facilitate completion of the report.

Under the direction of the National AIDS Committee or its equivalent, countries need to:

- 1. Identify the focal point for the reporting process and submit his/her name and contact details to UNAIDS Geneva through AIDSreporting@unaids.org;
- 2. Identify data needs in line with the national strategic plan requirements and these Global AIDS Response Progress Indicator guidelines; develop and disseminate a plan for data collection, including timelines and the roles of the National AIDS Committee or equivalent, other government agencies and civil society;
- 3. Identify relevant tools for data collection including meeting with national HIV estimates team;
- 4. Secure required funding for the entire process of collecting, analysing and reporting the data;
- 5. Collect and collate data in coordination with partner organizations from government, civil society and the international community;
- 6. Ensure that the HIV estimates team has submitted the draft Spectrum file to UNAIDS for review (latest 28 February);
- 7. Analyse data in coordination with partner organizations from government, civil society and the international community;
- 8. Calculate ART and PMTCT coverage estimates using denominators from updated and reviewed Spectrum files;
- 9. Allow stakeholders, including government agencies and civil society, to comment on the draft data;
- 10. Enter the data into the Global AIDS Response Progress Reporting website (http://AIDSreportingtool.unaids.org);
- 11. Upload the final Spectrum file to the designated national estimates folder;
- 12. Submit the indicator data before 31 March 2014;
- 13. Respond in a timely manner to queries on the submissions from UNAIDS, WHO or UNICEF.

It is important that the data that are reported are validated and reconciled between all partners in country. This process is supported in the online reporting tool through the ability to share the viewer credentials with national stakeholders. Several countries have reported that this feature enabled numerous civil society and other partners to view and provide inputs during the reporting process, hence allowing faster and wider stakeholder consultation and validation.

A summary checklist which may be used in the preparation and submission of the Country Progress Report is included as Appendix 4.

Summary of changes for 2014 Global AIDS Response progress reporting

2014 will be a "full" reporting year, i.e. including the core indicators, as well as the full National Commitments and Policy Instrument (NCPI) and a narrative country progress report.

Changes compared to the 2013 reporting round are summarized below:

- Indicators for sex-workers (Indicator 1.7, 1.8, 1.9 and 1.10) have, in addition to sex-disaggregation by female/male, also transgender as a possible disaggregation.
- Indicators for HIV testing among key populations (Indicators 1.9, 1.13 and 2.4) have the explanation of the denominator changed back to that of 2010 (see these indicators for details).
- The prevention of mother-to-child indicator (Indicator 3.1) has updated language to clarify the disaggregations and the links to Spectrum. Further, the indicator to measure coverage of PMTCT during breastfeeding has been added directly after this indicator (Indicator 3.1a, previously labelled Indicator 3.8).
- The indicator for ART coverage (Indicator 4.1) has a new denominator, now including all people living with HIV, not only these eligible for treatment. For a more detailed explanation for this change, please see the detailed indicator description for Indicator 4.1 (page 76). Further disaggregation of those newly initiated on ART (in the last 12 months) has been included (in previous rounds labelled 4.1a).
- The 12-month ART retention indicator (Indicator 4.2) has the possible disaggregations for pregnancy status and breastfeeding status at initiation included.
- The indicator for co-management of tuberculosis and HIV treatment (Indicator 5.1) has "adults" changed to "adults and children" in the numerator and "advanced" deleted from "advanced HIV infection".
- The AIDS spending indicator (Indicator 6.1) has split up bilateral donors into PEPFAR and other bilateral donors.
- A new indicator has been added related to Target 8: Discriminatory attitudes towards people living with HIV (Indicator 8.1). This indicator is new, so it is likely that most countries will not be able to report on the indicator during the 2014 reporting round. Instead countries are requested to report data from a previous version of one of the two questions of this indicator: "Would you buy fresh vegetables from a shopkeeper or vendor if you knew that this person had the AIDS virus?" This question has been routinely collected in DHS, and other surveys in many countries. In future reporting rounds, countries should report on the full indicator.
- Narrative reports are requested (please see Appendix 1 for more details).
- A full National Commitments and Policy Instrument (NCPI) is requested; the NCPI has been through slight revisions, please see Appendix 3 for the updated NCPI.

Key issues new in the 2013 reporting round that remain the same in the 2014 reporting round

- As in 2012 survey data that have not been updated since the last reporting round (i.e. 2012 or 2013 depending on when the last time reporting submitted) do not need to be re-entered (i.e. indicators 1.1, 1.2, 1.3, 1.4, 1.5, 1.6, 1.7, 1.8, 1.9, 1.10, 1.11, 1.12, 1.13, 1.14, 1.22, 2.2, 2.3, 2.4, 2.5, 7.1, 10.1, 10.2).
- The questions on key population size estimations from the 2012 reporting round are kept, but are now found among the indicators for Target 1 and Target 2 (page 49).
- The two indicators about prevalence of male circumcision and number of men circumcised that were added in 2013 for the 16 countries with high HIV-prevalence and low prevalence of male circumcision are still included (GARPR Indicators 1.22 and 1.23 and can be found in annex 6).
- Joint reporting of the Global AIDS Response Progress Reporting indicators and additional health sector indicators from WHO and UNICEF will continue. The additional health sector indicators can be found in Part II of these guidelines.

Core indicators for Global AIDS response progress reporting

Individual indicators may be used to track more than one target.

Individual indicators may be used to track more than one target.

Targets Indicators

largets	indica	ators			
Target 1. Reduce sexual transmission of HIV by 50% by 2015	1.1	Percentage of young women and men aged 15–24 who correctly identify ways of preventing the sexual transmission of HIV and who reject major misconceptions about HIV transmission*			
General population	1.2	Percentage of young women and men aged 15-24 who have had sexual intercourse before the age of 15			
	1.3	Percentage of adults aged 15–49 who have had sexual intercourse with more than one partner in the past 12 months			
	1.4	Percentage of adults aged 15–49 who had more than one sexual partner in the past 12 months who report the use of a condom during their last intercourse*			
	1.5	Percentage of women and men aged 15-49 who received an HIV test in the past 12 months and know their results			
	1.6	Percentage of young people aged 15-24 who are living with HIV*			
Sex workers	1.7	Percentage of sex workers reached with HIV prevention programmes			
	1.8	Percentage of sex workers reporting the use of a condom with their most recent client			
	1.9	Percentage of sex workers who have received an HIV test in the past 12 months and know their results			
	1.10	Percentage of sex workers who are living with HIV			
Men who have sex with men	1.11	Percentage of men who have sex with men reached with HIV prevention programmes			
	1.12	Percentage of men reporting the use of a condom the last time they had anal sex with a male partner			
	1.13	Percentage of men who have sex with men that have received an HIV test in the past 12 months and know their results			
	1.14	Percentage of men who have sex with men who are living with HIV			
Target 2. Reduce transmission of HIV among people	2.1	Number of syringes distributed per person who injects drugs per year by needle and syringe programmes			
who inject drugs by 50% by 2015	2.2	Percentage of people who inject drugs who report the use of a condom at last sexual intercourse			
	2.3	Percentage of people who inject drugs who reported using sterile injecting equipment the last time they injected			
	2.4	Percentage of people who inject drugs that have received an HIV test in the past 12 months and know their results			
	2.5	Percentage of people who inject drugs who are living with HIV			
Target 3. Eliminate new HIV infections among children by 2015 and substantially reduce AIDS-related maternal deaths**	3.1	Percentage of HIV-positive pregnant women who receive antiretrovirals to reduce the risk of mother-to-child transmission			
	3.1a	Percentage of women living with HIV receiving antiretroviral medicines for themselves or their infants during breastfeeding			
	3.2	Percentage of infants born to HIV-positive women receiving a virological test for HIV within 2 months of birth			
	3.3	Estimated percentage of child HIV infections from HIV-positive women delivering in the past 12 months			

Target 4. Reach 15 million people living with HIV with	4.1	Percentage of adults and children currently receiving antiretroviral therapy*		
lifesaving antiretroviral treatment by 2015	4.2	Percentage of adults and children with HIV known to be on treatment 12 months after initiation of antiretroviral therapy		
Target 5. Reduce tuberculosis deaths in people living with HIV by 50% by 2015	5.1	Percentage of estimated HIV-positive incident TB cases that received treatment for both TB and HIV		
Target 6. Close the global AIDS resource gap by 2015 and reach annual global investment of US\$ 22–24 billion in low- and middle-income countries	6.1	Domestic and international AIDS spending by categories and financing sources		
Target 7. Eliminating gender inequalities	7.1	Proportion of ever-married or partnered women aged 15-49 who experienced physical or sexual violence from a male intimate partner in the past 12 months		
		All indicators with sex-disaggregated data can be used to measure progress towards target 7		
Target 8. Eliminating stigma and discrimination	8.1	Discriminatory attitudes towards people living with HIV		
Target 9. Eliminate travel restrictions		Travel restriction data is collected directly by the Human Rights and Law Division at UNAIDS HQ, no reporting needed		
Target 10. Strengthening HIV integration	10.1	Current school attendance among orphans and non-orphans aged 10–14*		
	10.2	Proportion of the poorest households who received external economic support in the last 3 months		
Policy questions (relevant for all 10 targets)		National Commitments and Policy Instruments (NCPI)		

 $^{^{\}star} \quad \text{Millennium Development Goals indicator}$

*** The Global Plan towards the elimination of new HIV infections among children by 2015 and keeping their mothers alive defines this target as:

1. Reduce the number of new HIV infections among children by 90%

2. Reduce the number of AIDS-related maternal deaths by 50%

For further information see: http://www.unaids.org/en/media/unaids/contentassets/documents/unaidspublication/2011/20110609_JC2137_Global-Plan-Elimination-HIV-Children_en.pdf



TARGET 1. REDUCE SEXUAL TRANSMISSION OF HIV BY 50% BY 2015

General population

- 1.1 Percentage of young women and men aged 15–24 who both correctly identify ways of preventing the sexual transmission of HIV and who reject major misconceptions about HIV transmission*
- 1.2 Percentage of young women and men who have had sexual intercourse before the age of 15
- 1.3 Percentage of adults aged 15–49 who have had sexual intercourse with more than one partner in the past 12 months
- 1.4 Percentage of adults aged 15–49 who had more than one sexual partner in the past 12 months who report the use of a condom during their last intercourse*
- 1.5 Percentage of women and men aged 15-49 who received an HIV test in the past 12 months and know their results
- 1.6 Percentage of young people aged 15-24 who are living with HIV*

Sex workers

- 1.7 Percentage of sex workers reached with HIV prevention programmes
- 1.8 Percentage of sex workers reporting the use of a condom with their most recent client
- 1.9 Percentage of sex workers who have received an HIV test in the past 12 months and know their results
- 1.10 Percentage of sex workers who are living with HIV

Men who have sex with men

- 1.11 Percentage of men who have sex with men reached with HIV prevention programmes
- 1.12 Percentage of men reporting the use of a condom the last time they had anal sex with a male partner
- 1.13 Percentage of men who have sex with men that have received an HIV test in the past 12 months and know their results
- 1.14 Percentage of men who have sex with men who are living with HIV

^{*}Millennium Development Goals indicator

1.1 Young people: Knowledge about HIV prevention

Percentage of young people aged 15–24 who both correctly identify ways of preventing the sexual transmission of HIV and who reject major misconceptions about HIV transmission

What it measures

It measures progress towards universal knowledge of the essential facts about HIV transmission

Rationale

HIV epidemics are perpetuated through primarily sexual transmission of infection to successive generations of young people. Sound knowledge about HIV and AIDS is an essential prerequisite—albeit, often an insufficient condition—for adoption of behaviours that reduce the risk of HIV transmission.

Numerator: Number of respondents aged 15-24 years who gave the correct answer to all five

questions

Denominator: Number of all respondents aged 15–24

Calculation: Numerator / Denominator

Method of measurement:

Population-based surveys (Demographic and Health Survey, AIDS Indicator Survey, Multiple Indicator Cluster Survey or other representative survey)

This indicator is constructed from responses to the following set of prompted

questions:

1. Can the risk of HIV transmission be reduced by having sex with only one uninfected partner who has no other partners?

2. Can a person reduce the risk of getting HIV by using a condom every time

they have sex?

3. Can a healthy-looking person have HIV?

4. Can a person get HIV from mosquito bites?

5. Can a person get HIV by sharing food with someone who is infected?

Measurement frequency:

Preferred: every two years; minimum: every 3–5 years

Disaggregation: Sex

• Age (15-19 and 20-24)

Explanation of numerator

The first three questions should not be altered. Questions 4 and 5 ask about local misconceptions and may be replaced by the most common misconceptions in your country. Examples include: "Can a person get HIV by hugging or shaking hands with a person who is infected?" and "Can a person get HIV through supernatural means?"

Those who have never heard of HIV and AIDS should be excluded from the numerator but included in the denominator. An answer of "don't know" should be recorded as an incorrect answer.

Scores for each of the individual questions (based on the same denominator) are required as well as the score for the composite indicator.

Strengths and weaknesses

The belief that a healthy-looking person cannot be infected with HIV is a common misconception that can result in unprotected sexual intercourse with infected partners. Rejecting major misconceptions about modes of HIV transmission is as important as correct knowledge of true modes of transmission. For example, belief that HIV is transmitted through mosquito bites can weaken motivation to adopt safer sexual behaviour, while belief that HIV can be transmitted through sharing food reinforces the stigma faced by people living with HIV.

This indicator is particularly useful in countries where knowledge about HIV and AIDS is poor because it permits easy measurement of incremental improvements over time. However, it is also important in other countries as it can be used to ensure that pre-existing high levels of knowledge are maintained.

Further information

1.2 Sex before the age of 15

Percentage of young women and men aged 15-24 who have had sexual intercourse before the age of 15

What it measures

It measures progress in increasing the age at which young women and men aged 15-24 first have sex.

Rationale

A major goal in many countries is to delay the age at which young people first have sex and discourage premarital sexual activity because it reduces their potential exposure to HIV. There is also evidence to suggest that first having sex at a later age reduces susceptibility to infection per act of sex, at least for women.

Number of respondents (aged 15-24 years) who report the age at which they first Numerator:

had sexual intercourse as under 15 years

Number of all respondents aged 15-24 years Denominator:

Numerator / Denominator Calculation:

Population-based surveys (Demographic and Health Survey, AIDS Indicator Method of measurement:

Survey, Multiple Indicator Cluster Survey or other representative survey)

Respondents are asked whether or not they have ever had sexual intercourse and, if yes, they are asked: How old were you when you first had sexual intercourse for

the first time?

Measurement frequency:

Every 3-5 years

Sex Disaggregation:

Age (15-19 and 20-24)

Strengths and weaknesses

Countries where very few young people have sex before the age of 15 might opt to use an alternative indicator: percentage of young women and men aged 20-24 who report their age at sexual initiation as under 18 years. The advantage of using the reported age at which young people first had sexual intercourse (as opposed to the median age) is that the calculation is simple and allows easy comparison over time. The denominator is easily defined because all members of the survey sample contribute to this measure.

It is difficult to monitor change in this indicator over a short period because only individuals entering the group, i.e. those aged under 15 at the beginning of the period for which the trends are to be assessed, can influence the numerator. If the indicator is assessed every two to three years, it may be better to focus on changes in the levels for the 15-17 age group. If it is assessed every five years, the possibility exists of looking at the 15-19 age group.

In countries where HIV-prevention programmes encourage virginity or delaying of first sex, young people's responses to survey questions on this issue may be biased, including a deliberate misreporting of age at which they first had sex.

Further information

1.3 Multiple sexual partnerships

Percentage of women and men aged 15-49 who have had sexual intercourse with more than one partner in the past 12 months

What it measures

It measures progress in reducing the percentage of people who have multiple sexual partnerships.

Rationale

The spread of HIV largely depends upon unprotected sex among people with a high number of partnerships. Individuals who have multiple partners have a higher risk of HIV transmission than individuals that do not link into a wider sexual network.

Number of respondents aged 15-49 who have had sexual intercourse with more Numerator:

than one partner in the last 12 months

Number of all respondents aged 15-49 Denominator:

Numerator / Denominator Calculation:

Population-based surveys (Demographic and Health Survey, AIDS Indicator Method of measurement:

Survey, Multiple Indicator Cluster Survey or other representative survey)

Respondents' sexual histories are obtained. Analysis of sexual history is used to determine whether the respondent has had more than one partner in the

preceding 12 month period

Measurement frequency:

Every 3-5 years

Sex Disaggregation:

• Age (15-19, 20-24 and 25-49)

Strengths and weaknesses

This indicator gives a picture of levels of higher-risk sex. If people have only one sexual partner, the change will be captured by changes in this indicator. However, if people simply decrease the number of sexual partners they have, the indicator will not reflect a change, even though potentially this may have a significant impact on the epidemic spread of HIV and may be counted a programme success. Additional indicators may need to be selected to capture the reduction in multiple sexual partners in general.

Further information

1.4 Condom use at last sex among people with multiple sexual partnerships

Percentage of women and men aged 15-49 who had more than one partner in the past 12 months who used a condom during their last sexual intercourse

What it measures

It measures progress towards preventing exposure to HIV through unprotected among people with multiple sexual partners.

Rationale

Condom use is an important measure of protection against HIV, especially among people with multiple sexual partners.

Number of respondents (aged 15–49) who reported having had more than one Numerator:

sexual partner in the last 12 months who also reported that a condom was used

the last time they had sex

Number of respondents (15-49) who reported having had more than one sexual Denominator:

partner in the last 12 months.

Numerator / Denominator Calculation:

Population-based surveys (Demographic Health Survey, AIDS Indicator Survey, Method of measurement:

Multiple Indicator Cluster Survey or other representative survey)

Respondents' sexual histories are obtained. Analysis of sexual history is used to determine whether the respondent has had more than one partner in the preceding 12 month period, and if so whether a condom was used the last time

the respondent had sexual intercourse

Measurement frequency:

3-5 years

Sex Disaggregation:

Age 15-19, 20-24 and 25-49 years

Strengths and weaknesses

This indicator shows the extent to which condoms are used by people who are likely to have higher-risk sex (i.e. change partners regularly). However, the broader significance of any given indicator value will depend upon the extent to which people engage in such relationships. Thus, levels and trends should be interpreted carefully using the data obtained on the percentages of people that have had more than one sexual partner within the last year.

The maximum protective effect of condoms is achieved when their use is consistent rather than occasional. The current indicator does not provide the level of consistent condom use. However, the alternative method of asking whether condoms were always/sometimes/never used in sexual encounters with non-regular partners in a specified period is subject to recall bias. Furthermore, the trend in condom use during the most recent sex act will generally reflect the trend in consistent condom use.

Further information

1.5 HIV testing in the general population

Percentage of women and men aged 15-49 who received an HIV test in the past 12 months and know their results

What it measures

It measures progress in implementing HIV testing and counselling.

Rationale

In order to protect themselves and to prevent infecting others, it is important for individuals to know their HIV status. Knowledge of one's status is also a critical factor in the decision to seek treatment.

Numerator: Number of respondents aged 15-49 who have been tested for HIV during the last

12 months and who know their results

Denominator: Number of all respondents aged 15-49

The denominator includes respondents who have never heard of HIV or AIDS

Calculation: Numerator / Denominator

Method of measurement:

Population-based surveys (Demographic and Health Survey, AIDS Indicator Survey, Multiple Indicator Cluster Survey or other representative survey)

Respondents are asked:

1. I don't want to know the results, but have you been tested for HIV in the last

12 months?

If yes:

2. I don't want to know the results, but did you get the results of that test?

Measurement frequency:

Every 3 to 5 years

Disaggregation: Sex

• Age (15-19, 20-24 and 25-49)

Strengths and weaknesses

The introductory statement by the interviewer "I don't want to know the results [of any testing], but..." allows for better reporting and reduces the risk of underreporting of HIV testing among people who do not wish to disclose their serostatus.

Knowledge of HIV test results in the past 12 months does not guarantee that a respondent knows their current HIV status. A respondent may have contracted HIV in the time since their last HIV test.

Further information

1.6 HIV prevalence in young people

Percentage of young people aged 15-24 who are living with HIV

What it measures

It measures progress towards reducing HIV infection.

Rationale

The goal in the response to HIV is to reduce HIV infection. However, given current inability to reliably measure HIV incidence in a cross-sectional survey, proxy measures of HIV incidence are required.

HIV prevalence at any given age is the difference between the cumulative numbers of people that have become infected with HIV up to this age minus the number who have died, expressed as a percentage of the total number alive at this age. At older ages, changes in HIV prevalence are slow to reflect changes in the rate of new infections (HIV incidence) because the average duration of infection is long. Declines in HIV prevalence can reflect saturation of infection among those individuals who are most vulnerable, and rising mortality, rather than behaviour change. Increases in HIV prevalence can reflect increasing numbers of individuals receiving antiretroviral therapy, and living longer. However at younger ages, trends in HIV prevalence are a better indication of recent trends in HIV incidence and risk behaviour since young people are likely to only recently have initiated sexual or injecting drug behaviours. In addition, young people who have recently been infected with HIV are not likely to have started antiretroviral therapy. Thus, reductions in HIV incidence associated with genuine behaviour change should first become detectable in trends in HIV prevalence figures for 15–24 years olds (or even earlier in 15–19-year-olds if this age breakdown is available). Where available, parallel behavioural surveillance survey data should be used to aid interpretation of trends in HIV prevalence.

Epidemic Type: Countries with generalized epidemics

Number of antenatal clinic attendees (aged 15–24) tested whose HIV test results

are positive

Denominator: Number of antenatal clinic attendees (aged 15–24) tested for their HIV infection

status

Calculation: Numerator / Denominator

Method of UNAIDS/WHO guidelines for HIV sentinel surveillance

measurement:

This indicator is calculated using data from pregnant women attending antenatal

clinics in HIV sentinel surveillance sites in the capital city, other urban areas and

rural areas

The sentinel surveillance sites used for the calculation of this indicator should

remain constant to allow for the tracking of changes over time

Measurement frequency:

Annual

Disaggregation: None

Strengths and weaknesses

In countries where the age at which young people first have sexual intercourse is late and/or levels of contraception use are high, HIV prevalence among pregnant women of 15–24 years of age will differ from that among all women in the age group. If fertility patterns are changing this trend might be biased if women living with HIV make different fertility choices.

This indicator (using data from antenatal clinics) gives a fairly good estimate of relatively recent trends in HIV infection in locations where the epidemic is heterosexually driven. It is less reliable as an indicator of HIV-epidemic trends in locations where most infections are primarily among key populations.

To supplement data from antenatal clinics, an increasing number of countries have included HIV testing in population-based surveys. If a country has produced HIV prevalence estimates from survey data, these estimates should be included in the comments box for this indicator in order to allow for comparisons between multiple surveys. Survey-based estimates should be disaggregated by sex.

The addition of new sentinel sites will increase the samples' representativeness and will therefore give a more robust point estimate of HIV prevalence. However, the addition of new sentinel sites reduces the comparability of values. As such it is important to use consistent sites when undertaking trend analyses.

As more children who were infected through mother to child transmission live into their reproductive years this indicator becomes more difficult to interpret. Countries should collect information on timing of infection for women with known HIV-positive sero-status to exclude these women from analyses of trends.

Further information

For further information, please consult the following links:

 $http://www.unaids.org/en/media/unaids/contentassets/documents/epidemiology/2013/gr2013/20131118_Methodology.pdf$

1.7 Sex workers: prevention programmes

Percentage of sex workers reached with HIV prevention programmes

What it measures

It measures progress in implementing basic elements of HIV prevention programmes for sex workers.

Rationale

Sex workers are often difficult to reach with HIV prevention programmes. However, in order to prevent the spread of HIV and AIDS among sex workers as well as into the general population, it is important that they access these services.

Note: Countries with generalized epidemics may also have a concentrated sub-epidemic among one or more key populations at higher risk. If so, they should calculate and report this indicator for those populations.

ADDITIONAL 2014 REPORTING: transgender women who sell sex have some of the highest-recorded HIV prevalence rates and therefore this new disaggregation has been included.

Numerator: Number sex workers who replied "yes" to both questions

Denominator: Total number of sex workers surveyed

Calculation: Numerator / Denominator

Method of measurement:

Behavioural surveillance or other special surveys

Sex workers are asked the following questions:

- 1. Do you know where you can go if you wish to receive an HIV test?
- 2. In the last twelve months, have you been given condoms? (e.g. through an outreach service, drop-in centre or sexual health clinic)

Scores for each of the individual questions—based on the same denominator—are required in addition to the score for the composite indicator

Whenever possible, data for sex workers should be collected through civil society organizations that have worked closely with this population in the field

Access to sex workers as well as the data collected from them must remain confidential

Measurement frequency:

Disaggregation:

Every two years

Sex (female, male, transgender)

■ Age (<25/25+)

Strengths and weaknesses

The data obtained may not be based on a representative national sample of the sex worker population being surveyed. If there are concerns that the data are not based on a representative sample, these concerns should be reflected in the interpretation of the survey data. Where different sources of data exist, the best available estimate should be used. Information on the sample size, the quality and reliability of the data, and any related issues should be included in the report submitted with this indicator.

The inclusion of these indicators for reporting purposes should not be interpreted to mean that these services alone are sufficient for HIV prevention programmes for the populations. The set of key interventions described above should be part of a comprehensive HIV prevention programme, which also includes elements such as provision of HIV prevention messages, (e.g. through outreach programmes and peer education), treatment of sexually transmitted diseases, and others. For further information on the elements of comprehensive HIV prevention programmes for sex workers please see the Practical guidelines for intensifying HIV prevention: towards universal access.

This indicator asks about services accessed in the past 12 months. If you have data available on another time period, such as the last three or six months or the last 30 days, please include this additional data in the comments section of the reporting tool.

To maximize the utility of these data, it is recommended that the same sample used for the calculation of this indicator be used for the calculation of the other indicators related to these populations.

Several countries have in previous reporting rounds reported HIV prevalence among subpopulations of transgender women through the additional comments field in the GARPR online reporting tool. This demonstrates that the data are feasible to obtain in different settings.

In addition to the above requested data, please report programme data if available for this indicator using the text box provided in the online reporting platform.

Further information

For further information, please consult the following references:

A framework for monitoring and evaluating HIV prevention programmes for most-at-risk populations. Geneva, UNAIDS, 2007.

Practical guidelines for intensifying HIV prevention: towards universal access. Geneva, UNAIDS, 2007.

Operational Guidelines for Monitoring and Evaluation of HIV Programmes for Sex Workers, Men who have Sex with Men, and Transgender People. MEASURE Evaluation (www.cpc.unc.edu/measure/publications/ms-11-49a).

1.8 Sex workers: condom use

Percentage of sex workers reporting the use of a condom with their most recent client

What it measures

It measures progress in preventing exposure to HIV among sex workers through unprotected sex with clients.

Rationale

Various factors increase the risk of exposure to HIV among sex workers, including multiple, non-regular partners and more frequent sexual intercourse. However, sex workers can substantially reduce the risk of HIV transmission, both from clients and to clients, through consistent and correct condom use.

Note: Countries with generalized epidemics may also have a concentrated sub-epidemic among sex workers. If so, it would be valuable for them to calculate and report on this indicator for this population.

ADDITIONAL 2014 REPORTING: transgender women who sell sex have some of the highest-recorded HIV prevalence rates and therefore this new disaggregation has been included.

Number of sex workers who reported that a condom was used with their

last client

Denominator: Number of sex workers who reported having commercial sex in the last

12 months

Calculation: Numerator / Denominator

Method of measurement:

Behavioural surveillance or other special surveys

Respondents are asked the following question:

Did you use a condom with your most recent client?

Whenever possible, data for sex workers should be collected through civil society

organizations that have worked closely with this population in the field

Access to sex workers as well as the data collected from them must remain

confidential

Measurement frequency:

Every two years

Disaggregation: Sex (female, male, transgender)

■ Age (<25/25+)

Strengths and weaknesses

Condoms are most effective when their use is consistent, rather than occasional. The current indicator will provide an overestimate of the level of consistent condom use. However, the alternative method of asking whether condoms are always/sometimes/never used in sexual encounters with clients in a specified period is subject to recall bias. Furthermore, the trend in condom use in the most recent sexual act will generally reflect the trend in consistent condom use.

This indicator asks about commercial sex in the past twelve months. If you have data available on another time period, such as the last three or six months, please include this additional data in the comments section of the reporting tool.

Surveying sex workers can be challenging. Consequently, data obtained may not be based on a representative national sample of the key populations at higher risk being surveyed. If there are concerns that the data are not based on a representative sample, these concerns should be reflected in the interpretation of the survey data. Where different sources of data exist, the best available estimate should be used. Information on the sample size, the quality and reliability of the data, and any related issues should be included in the report submitted with this indicator.

Several countries have in previous reporting rounds reported HIV prevalence among subpopulations of transgender women through the additional comments field in the GARPR online reporting tool. This demonstrates that the data are feasible to obtain in different settings.

To maximize the utility of these data, it is recommended that the same sample used for the calculation of this indicator be used for the calculation of the other indicators related to these populations.

Further information

For further information, please consult the following references:

A framework for monitoring and evaluating HIV prevention programmes for most-at-risk populations. Geneva, UNAIDS, 2007.

Practical guidelines for intensifying HIV prevention: towards universal access. Geneva, UNAIDS, 2007.

1.9 HIV testing in sex workers

Percentage of sex workers who received an HIV test in the past 12 months and know their results

What it measures

It measures progress in implementing HIV testing and counselling among sex workers.

Rationale

In order to protect themselves and to prevent infecting others, it is important for sex workers to know their HIV status. Knowledge of one's status is also a critical factor in the decision to seek treatment. Note: Countries with generalized epidemics may also have a concentrated sub-epidemic among one or more Key populations at higher risk. If so, they should calculate and report this indicator for those populations.

ADDITIONAL 2014 REPORTING: transgender women who sell sex have some of the highest-recorded HIV prevalence rates and therefore this new disaggregation has been included.

Numerator: Number of sex workers who have been tested for HIV during the last

12 months and who know their results

Denominator: Number of sex workers included in the sample

Calculation: Numerator / Denominator

Method of measurement:

Behavioural surveillance or other special surveys

Sex workers are asked the following questions:

1. Have you been tested for HIV in the last 12 months?

If yes:

2. I don't want to know the results, but did you receive the results of that test?

Whenever possible, data for sex workers should be collected through civil society

organizations that have worked closely with this population in the field

Access to sex workers as well as the data collected from them must remain

confidential

Measurement frequency:

Every two years

Disaggregation: Sex (female, male, transgender)

■ Age (<25/25+)

Strengths and weaknesses

The data obtained may not be based on a representative national sample of the sex workers being surveyed. If there are concerns that the data are not based on a representative sample, these concerns should be reflected in the interpretation of the survey data. Where different sources of data exist, the best available estimate should be used. Information on the sample size, the quality and reliability of the data, and any related issues should be included in the report submitted with this indicator.

Tracking sex workers over time to measure progress may be difficult due to mobility and the hard-to-reach nature of these populations with many groups being hidden populations. Thus, information about the nature of the sample should be reported in the narrative to facilitate interpretation and analysis over time.

Several countries have in previous reporting rounds reported HIV prevalence among subpopulations of transgender women through the additional comments field in the GARPR online reporting tool. This demonstrates that the data are feasible to obtain in different settings.

To maximize the utility of these data, it is recommended that the same sample used for the calculation of this indicator be used for the calculation of the other indicators related to these populations.

This indicator is most meaningful in settings where testing scale-up is relatively recent. People who tested more than 12 months ago and know they are positive will be considered "uncovered" by this indicator construction. Ideally, surveys should ask why respondents did not test in the past 12 months. If they report that they know their HIV status to be positive, they should not be included in the denominator. This indicator will be formally changed post-2015; we ask countries that can to report against this indicator while omitting known HIV-positive persons from the denominator and state that they've done this in the comment field.

Further information

For further information, please consult the following references:

A framework for monitoring and evaluating HIV prevention programmes for most-at-risk populations. Geneva, UNAIDS, 2007.

Practical guidelines for intensifying HIV prevention: towards universal access. Geneva, UNAIDS, 2007.

1.10 HIV prevalence in sex workers

Percentage of sex workers who are living with HIV

What it measures

It measures progress on reducing HIV prevalence among sex workers.

Rationale

Sex workers typically have higher HIV prevalence than the general population in both concentrated and generalized epidemics. In many cases, prevalence among these populations can be more than double the prevalence among the general population. Reducing prevalence among sex workers is a critical measure of a national-level response to HIV.

Countries with generalized epidemics may also have a concentrated sub-epidemic among sex workers. If so, it is valuable to calculate and report on this indicator for this population.

ADDITIONAL 2014 REPORTING: transgender women who sell sex have some of the highest-recorded HIV prevalence rates and therefore this new disaggregation has been included.

Numerator: Number of sex workers who test positive for HIV

Denominator: Number of sex workers tested for HIV

Calculation: Numerator / Denominator

Method of UNAIDS and WHO Working Group on Global HIV/AIDS and STI Surveillance:

measurement: Guidelines among populations most at risk for HIV (

WHO/UNAIDS, 2011)

This indicator is calculated using data from HIV tests conducted among

respondents in the primary sentinel site or sites

The sentinel surveillance sites used for the calculation of this indicator should

remain constant to allow for the tracking of changes over time

Measurement frequency:

Annual

Disaggregation: Sex (female, male, transgender)

■ Age (<25/25+)

Strengths and weaknesses

In theory, assessing progress in reducing the occurrence of new infections is best done through monitoring changes in incidence over time. However, in practice, prevalence data rather than incidence data are available. In analyzing prevalence data of sex workers for the assessment of prevention programme impact, it is desirable not to restrict analysis to young people but to report on those persons who are newly initiated to behaviours that put them at risk for infection (e.g. by restricting the analysis to people who have or participated in sex work for less than one year) This type of analysis also has the advantage of not being affected by the effect of ART in increasing survival and thereby increasing prevalence.

If prevalence estimates are available disaggregated by greater than and less than one year in sex work countries are strongly encouraged to report this disaggregation in their Country Progress Report, and to use the comments field in the reporting tool for this indicator to present disaggregated estimates.

Due to difficulties in accessing sex workers, biases in sero-surveillance data are likely to be far more significant than in data from a more general population, such as women attending antenatal clinics. If there are concerns about the data, these concerns should be reflected in the interpretation.

An understanding of how the sampled population(s) relate to any larger population(s) sharing similar risk behaviours is critical to the interpretation of this indicator. The period during which people belong to a key population is more closely associated with the risk of acquiring HIV than age. Therefore, it is desirable not to restrict analysis to young people but to report on other age groups as well.

Trends in HIV prevalence among sex workers in the capital city will provide a useful indication of HIV prevention programme performance in that city. However, it will not be representative of the situation in the country as a whole.

The addition of new sentinel sites will increase the samples representativeness and will therefore give a more robust point estimate of HIV prevalence. However, the addition of new sentinel sites reduces the comparability of values. As such it is important to use consistent sites when undertaking trend analyses.

Several countries have in previous reporting rounds reported HIV prevalence among subpopulations of transgender women through the additional comments field in the GARPR online reporting tool. This demonstrates that the data are feasible to obtain in different settings.

Further information

For further information, please consult the following links:

http://www.unaids.org/en/HIV_data/Methodology/default.asp

Revised guidelines on HIV surveillance for key populations at higher risk: WHO/UNAIDS Working Group on Global HIV/AIDS and STI Surveillance. Guidelines on surveillance among populations most at risk for HIV. Geneva, UNAIDS, 2011 (http://www.unaids.org/en/media/unaids) contentassets/documents/epidemiology/2011/20110518_Surveillance_among_most_at_risk.pdf

1.11 Men who have sex with men: prevention programmes

Percentage of men who have sex with men reached with HIV prevention programmes

What it measures

It measures progress in implementing basic elements of HIV prevention programmes for MSM.

Rationale

Men who have sex with men (MSM) are often difficult to reach with HIV prevention programmes. However, in order to prevent the spread of HIV and AIDS among MSM as well as into the general population, it is important that they access these services.

Note: Countries with generalized epidemics may also have a concentrated sub-epidemic among one or more key populations at higher risk. If so, they should calculate and report this indicator for those populations.

Number MSM who replied "yes" to both questions Numerator:

Total number of MSM surveyed Denominator:

Numerator / Denominator Calculation:

Method of measurement: Behavioural surveillance or other special surveys

Respondents are asked the following questions:

- 1. Do you know where you can go if you wish to receive an HIV test?
- 2. In the last twelve months, have you been given condoms? (e.g. through an outreach service, drop-in centre or sexual health clinic)

Scores for each of the individual questions—based on the same denominator are required in addition to the score for the composite indicator

Whenever possible, data for MSM should be collected through civil society organizations that have worked closely with this population in the field

Access to MSM as well as the data collected from them must remain confidential

Measurement frequency:

Every two years

Age (<25/25+) Disaggregation:

Strengths and weaknesses

The data obtained may not be based on a representative national sample of the MSM population being surveyed. If there are concerns that the data are not based on a representative sample, these concerns should be reflected in the interpretation of the survey data. Where different sources of data exist, the best available estimate should be used. Information on the sample size, the quality and reliability of the data, and any related issues should be included in the report submitted with this indicator.

The inclusion of these indicators for reporting purposes should not be interpreted to mean that these services alone are sufficient for HIV prevention programmes for the population. The set of key interventions described above should be part of a comprehensive HIV prevention programme, which also includes elements such as provision of HIV prevention messages, (e.g. through outreach programmes and peer education), treatment of sexually transmitted diseases, and others. For further information on the elements of comprehensive HIV prevention programmes for key populations at higher risk please see the *Practical guidelines for intensifying HIV prevention: towards universal access*.

This indicator asks about services accessed in the past 12 months. If you have data available on another time period, such as the last three or six months or the last 30 days, please include this additional data in the comments section of the reporting tool.

To maximize the utility of these data, it is recommended that the same sample used for the calculation of this indicator be used for the calculation of the other indicators related to these populations.

In addition to the above requested data, please report programme data if available for this indicator using the text box provided in the online reporting platform.

Further information

For further information, please consult the following references:

A framework for monitoring and evaluating HIV prevention programmes for most-at-risk populations. *Geneva, UNAIDS, 2007.*

Practical guidelines for intensifying HIV prevention: towards universal access. Geneva, UNAIDS, 2007.

1.12 Men who have sex with men: condom use

Percentage of men reporting the use of a condom the last time they had anal sex with a male partner

What it measures

It measures progress in preventing exposure to HIV among men who have unprotected anal sex with a male partner.

Rationale

Condoms can substantially reduce the risk of the sexual transmission of HIV. Consequently, consistent and correct condom use is important for men who have sex with men because of the high risk of HIV transmission during unprotected anal sex. In addition, men who have anal sex with other men may also have female partners, who could become infected as well. Condom use with their most recent male partner is considered a reliable indicator of longer-term behaviour.

Note: Countries with generalized epidemics may also have a concentrated sub-epidemic among men who have sex with men. If so, it would be valuable for them to calculate and report on this indicator for this population.

Numerator: Number of MSM who reported that a condom was used the last time they had

anal sex

Denominator: Number of MSM who reported having had anal sex with a male partner in the

last six months

Calculation: Numerator / Denominator

Method of measurement:

Behavioural surveillance or other special surveys

In a behavioural survey of a sample of men who have sex with men, respondents are asked about sexual partnerships in the preceding six months, about anal sex

within those partnerships and about condom use when they last had anal sex

Whenever possible, data for men who have sex with men should be collected through civil society organizations that have worked closely with this population

in the field

Access to MSM as well as the data collected from them must remain confidential

Measurement frequency:

Every two years

Disaggregation: ■ Age (<25/25+)

Strengths and weaknesses

For men who have sex with men, condom use at last anal sex with any partner gives a good indication of overall levels and trends of protected and unprotected sex in this population. This indicator does not give any idea of risk behaviour in sex with women among men who have sex with both women and men. In countries where men in the sub-population surveyed are likely to have partners of both sexes, condom use with female as well as male partners should be investigated. In these cases, data on condom use should always be presented separately for female and male partners.

This indicator asks about male to male sex in the past six months. If you have data available on another time period, such as the last three or twelve months, please include this additional data in the comments section of the reporting tool.

The data obtained may not be based on a representative national sample of the men who have sex with men being surveyed. If there are concerns that the data are not based on a representative sample, these concerns should be reflected in the interpretation of the survey data. Where different sources of data exist, the best available estimate should be used. Information on the sample size, the quality and reliability of the data, and any related issues should be included in the report submitted with this indicator.

To maximize the utility of these data, it is recommended that the same sample used for the calculation of this indicator be used for the calculation of the other indicators related to these populations.

Further information

For further information, please consult the following references:

A framework for monitoring and evaluating HIV prevention programmes for most-at-risk populations. Geneva, UNAIDS, 2007.

Practical guidelines for intensifying HIV prevention: towards universal access. Geneva, UNAIDS, 2007.

1.13 HIV testing in men who have sex with men

Percentage of men who have sex with men who received an HIV test in the past 12 months and know their results

What it measures

It measures progress in implementing HIV testing and counselling among men who have sex with men.

Rationale

In order to protect themselves and to prevent infecting others, it is important for men who have sex with men to know their HIV status. Knowledge of one's status is also a critical factor in the decision to seek treatment.

Note: Countries with generalized epidemics may also have a concentrated sub-epidemic among one or more key population at higher risk. If so, they should calculate and report this indicator for those populations.

Numerator: Number of men who have sex with men who have been tested for HIV during the

last 12 months and who know their results

Denominator: Number of men who have sex with men included in the sample

Calculation: Numerator / Denominator

Method of measurement:

Behavioural surveillance or other special surveys

Respondents are asked the following questions:

1. Have you been tested for HIV in the last 12 months?

If yes:

2. I don't want to know the results, but did you receive the results of that test?

Whenever possible, data for men who have sex with men should be collected through civil society organizations that have worked closely with this population

in the field

Access to MSM as well as the data collected from them must remain confidential

Measurement frequency:

Every two years

Disaggregation: Age (<25/25+)

Strengths and weaknesses

The data obtained may not be based on a representative national sample of the men who have sex with men being surveyed. If there are concerns that the data are not based on a representative sample, these concerns should be reflected in the interpretation of the survey data. Where different sources of data exist, the best available estimate should be used. Information on the sample size, the quality and reliability of the data, and any related issues should be included in the report submitted with this indicator.

Tracking men who have sex with men over time to measure progress may be difficult due to mobility and the often hard-to-reach nature of these populations. Thus, information about the nature of the sample should be reported in the narrative to facilitate interpretation and analysis over time.

To maximize the utility of these data, it is recommended that the same sample used for the calculation of this indicator be used for the calculation of the other indicators related to these populations.

This indicator is most meaningful in settings where testing scale-up is relatively recent. People who tested more than 12 months ago and know they are positive will be considered "uncovered" by this indicator construction. Ideally, surveys should ask why respondents did not test in the past 12 months. If they report that they know their HIV status to be positive, they should not be included in the denominator. This indicator will be formally changed post-2015; we will ask countries that can to report against this indicator while omitting known HIV-positive persons from the denominator and state that they've done this in the comment field.

Further information

For further information, please consult the following references:

A framework for monitoring and evaluating HIV prevention programmes for most-at-risk populations. Geneva, UNAIDS, 2007.

Practical guidelines for intensifying HIV prevention: towards universal access. Geneva, UNAIDS, 2007.

1.14 HIV prevalence in men who have sex with men

Percentage of men who have sex with men risk who are living with HIV

What it measures

It measures progress on reducing HIV prevalence among men who have sex with men.

Rationale

Men who have sex with men typically have the highest HIV prevalence in countries with either concentrated or generalized epidemics. In many cases, prevalence among these populations can be more than double the prevalence among the general population. Reducing prevalence among men who have sex with men is a critical measure of a national-level response to HIV.

Note: Countries with generalized epidemics may also have a concentrated sub-epidemic among one or more key population at higher risk. If so, it would be valuable for them to calculate and report on this indicator for those populations.

Numerator: Number of MSM who test positive for HIV

Denominator: Number of MSM tested for HIV

Calculation: Numerator / Denominator

Method of UNAIDS and WHO Working Group on Global HIV/AIDS and STI Surveillance: measurement: Guidelines among populations most at risk for HIV (WHO/UNAIDS, 2011)

This indicator is calculated using data from HIV tests conducted among

respondents in the primary sentinel site or sites

The sentinel surveillance sites used for the calculation of this indicator should

remain constant to allow for the tracking of changes over time

Measurement frequency:

Annual

Disaggregation: ■ Age (<25/25+)

Strengths and weaknesses

In theory, assessing progress in reducing the occurrence of new infections is best done through monitoring changes in incidence over time. However, in practice, prevalence data rather than incidence data are available.

In analyzing prevalence data of men who have sex with men for the assessment of prevention programme impact, it is desirable not to restrict analysis to young people but to report on those persons who are newly initiated to behaviours that put them at risk for infection (e.g. by restricting the analysis to people who first had sex with another man within the last year). This type of analysis also has the advantage of not being affected by the effect of ART in increasing survival and thereby increasing prevalence.

If prevalence estimates are available disaggregated by greater than and less than one year of sexual activity with other men countries are strongly encouraged to report this disaggregation in their Country Progress Report, and to use the comments field in the reporting tool for this indicator to present disaggregated estimates.

Due to difficulties in accessing men who have sex with men, biases in sero-surveillance data are likely to be far more significant than in data from a more general population, such as women attending antenatal clinics. If there are concerns about the data, these concerns should be reflected in the interpretation.

An understanding of how the sampled population(s) relate to any larger population(s) sharing similar risk behaviours is critical to the interpretation of this indicator. The period during which people belong to a key population is more closely associated with the risk of acquiring HIV than age. Therefore, it is desirable not to restrict analysis to young people but to report on other age groups as well.

Trends in HIV prevalence among men who have sex with men in the capital city will provide a useful indication of HIV-prevention programme performance in that city. However, it will not be representative of the situation in the country as a whole.

The addition of new sentinel sites will increase the samples representativeness and will therefore give a more robust point estimate of HIV prevalence. However, the addition of new sentinel sites reduces the comparability of values. As such it is important to use consistent sites when undertaking trend analyses.

Further information

For further information, please consult the following links:

http://www.unaids.org/en/HIV_data/Methodology/default.asp

Revised guidelines on HIV surveillance for key populations at higher risk: WHO/UNAIDS Working Group on Global HIV/AIDS and STI Surveillance. Guidelines on surveillance among populations most at risk for HIV. Geneva, UNAIDS, 2011 (http://www.unaids.org/en/media/unaids) contentassets/documents/epidemiology/2011/20110518_Surveillance_among_most_at_risk.pdf



TARGETS 1 AND 2. SIZE ESTIMATIONS FOR KEY POPULATIONS

Rationale

Programme planning for key populations can be much more efficient if there are accurate estimates of the size of these populations. The figures provide the MOH and UNAIDS with the ability to understand the scope of the potential HIV epidemic as well as the resources that will be needed to adequately meet the prevention needs of the at-risk populations.

1) Have you performed population size estimations for key populations?

Key population	Size estimation performed (yes/no)	If yes, when was the latest estimation performed? (year)	If yes, what was the size estimation?
a) Men who have sex with men			
b) People who inject drugs			
c) Sex workers			
d) Other key populations— please specify which key population in the comments box.			
e) Comments:			

To get a better understanding of the size estimates submitted, we request the following additional information about each estimate, to be included in the comment box:

- 1) the definition used of the population;
- 2) the method used to derive the size estimate;
- 3) site specific estimates for all available estimates.

In keeping with on-going effort to provide more granular data presentations, the latter will offer the opportunity for mapping denominator data with programme data if they are collected in the same survey areas.



TARGET 2. REDUCE TRANSMISSION OF HIV AMONG PEOPLE WHO INJECT DRUGS BY 50% BY 2015

- 2.1 Number of syringes distributed per person who injects drugs per year by needle and syringe programmes
- 2.2 Percentage of people who inject drugs who report the use of a condom at last sexual intercourse
- 2.3 Percentage of people who inject drugs who reported using sterile injecting equipment the last time they injected
- 2.4 Percentage of people who inject drugs that have received an HIV test in the past 12 months and know their results
- 2.5 Percentage of people who inject drugs who are living with HIV

2.1 People who inject drugs: prevention programmes

Number of needles and syringes distributed per person who injects drugs per year by needle and syringe programmes

What it measures

It measures progress in improving coverage of an essential HIV prevention service for people who inject drugs.

Rationale

Injecting drug use is the main route of transmission for approximately 10% of HIV infections globally and 30% of infections outside of sub-Saharan Africa. Preventing HIV transmission through injecting drug use is one of the key challenges to reducing the burden of HIV.

Needle and syringe programmes (NSPs) are one of nine interventions in the WHO, UNODC and UNAIDS comprehensive package for the prevention, treatment and care of HIV among people who inject drugs.

Needle and syringe programmes greatly affect HIV prevention for people who inject drugs. and there is a wealth of scientific evidence supporting its efficacy in preventing the spread of HIV (see http://www.who.int/hiv/topics/idu/needles/en/index.html).

Numerator: Number of needles and syringes distributed in past 12 months by NSPs

Denominator: Number of people who inject drugs in the country

Calculation: Numerator / Denominator

Method of Programme data used to count the number of needles and syringes distributed

measurement: (numerator)

Size estimation of the number of people who inject drugs in the country

(denominator)

Measurement frequency:

Every two years

Disaggregation: None

Strengths and weaknesses

Some difficulties regarding how to count needles and syringes are reported. Some commonly used syringes are 1 or 2ml needle and syringe units while others are syringes to which additional needles need to be fitted. In most cases only data on the number of syringes distributed via NSPs but not pharmacy sales will be available.

Estimating the size of PWID populations at country level is not without its challenges. Many different definitions of people who inject drugs exist in the literature and there are ranges of estimates. UNODC publishes size estimates of people who inject drugs in the World Drug Report. These estimates may be used. If there is a reason NOT to use them, please provide rationale in the comment box.

Countries can monitor this indicator against the following coverage levels:

- Low: <100 syringes per PWID per year
- Medium: >100-<200 syringes per PWID per year
- High: >200 syringes per PWID per year

These levels are based upon studies in developed country settings investigating the levels of syringe distribution and impact on HIV transmission. Note that the levels required for the prevention of hepatitis C are likely to be much higher than those presented here.

Further information

A full description of this indicator can be found in: WHO, UNODC and UNAIDS. *Technical guide for countries to set targets for universal access to HIV prevention, treatment and care for injecting drug users.* Geneva, World Health Organization, 2012

http://www.who.int/hiv/pub/idu/targets_universal_access/en/index.html.

For further information, please consult the following references:

Effectiveness of sterile needle and syringe programming in reducing HIV/AIDS among IDUs. Geneva, World Health Organization, 2004 (http://www.who.int/hiv/pub/idu/e4a-needle/en/index.html).

UNODC Global Assessment Programme on Drug Abuse. *Estimating prevalence: indirect methods for estimating the size of the drug problem. Vienna, UNODC, 2003.*

Hickman M et al. Estimating the prevalence of problematic drug use: a review of methods and their application. *Bulletin on Narcotics*, 2002, 54:15–32.

Most at risk populations sampling strategies and design tool. Atlanta, United States Department of Health and Human Services, Centers for Disease Control and Prevention, GAP Surveillance Team, 2009 (http://www.igh.org/surveillance).

http://www.idurefgroup.unsw.edu.au/IDURGWeb.nsf/page/publications (for more details on the Reference Group and to access reported country-level and global-level estimates of injecting drug use and HIV among injectors).

http://www.unaids.org/en/media/unaids/contentassets/documents/epidemiology/2011/2011_ Estimating_Populations_en.pdf (the WHO/UNAIDS working group on global HIV/AIDS and STI surveillance 2010 guidelines on estimating the size of populations most at risk to HIV).

WHO/UNAIDS Working Group on Global HIV/AIDS and STI Surveillance. *Guidelines on surveillance among populations most at risk for HIV.* Geneva, UNAIDS, 2011 (http://www.unaids.org/en/media/unaids) contentassets/documents/epidemiology/2011/20110518_Surveillance_among_most_at_risk.pdf.

2.2 People who inject drugs: condom use

Percentage of people who inject drugs reporting the use of a condom the last time they had sexual intercourse

What it measures

It measures progress in preventing sexual transmission of HIV among people who inject drugs.

Rationale

Safer injecting and sexual practices among people who inject drugs are essential, even in countries where other modes of HIV transmission predominate, because: (i) the risk of HIV transmission from contaminated injecting equipment is extremely high; and (ii) people who inject drugs can spread HIV (e.g. through sexual transmission) to the wider population.

Note: Countries with generalized epidemics may also have a concentrated sub-epidemic among people who inject drugs. If so, it would be valuable for them to calculate and report on this indicator for this population.

Numerator: Number of people who inject drugs who reported that a condom was used the

last time they had sex

Denominator: Number of people who inject drugs who report having injected drugs and having

had sexual intercourse in the last month

Calculation: Numerator / Denominator

Method of measurement:

Behavioural surveillance or other special surveys

people who inject drugs are asked the following sequence of questions:

- 1. Have you injected drugs at any time in the last month?
- 2. If yes: Have you had sexual intercourse in the last month?
- 3. If yes in answer to both 1 and 2: Did you use a condom when you last had sexual intercourse?

Whenever possible, data for people who inject drugs should be collected through civil society organizations that have worked closely with this population in the field

Access to survey respondents as well as the data collected from them must remain confidential

Measurement frequency:

Every two years

Disaggregation: Sex

■ Age (<25/25+)

Strengths and weaknesses

The data obtained may not be based on a representative national sample of the people who inject drugs being surveyed. If there are concerns that the data are not based on a representative sample, these concerns should be reflected in the interpretation of the survey data. Where different sources of data exist, the best available estimate should be used. Information on the sample size, the quality and reliability of the data, and any related issues should be included in the report submitted with this indicator.

The extent of injecting drug use-associated HIV transmission within a country depends on four factors: (i) the size, stage and pattern of dissemination of the national AIDS epidemic; (ii) the extent of injecting drug use; (iii) the degree to which people who inject drugs use contaminated injecting equipment; and (iv) the patterns of sexual mixing and condom use among people who inject drugs and between people who inject drugs and the wider population. This indicator provides partial information on the fourth factor.

To maximize the utility of these data, it is recommended that the same sample used for the calculation of this indicator be used for the calculation of the other indicators related to these populations.

Further information

For further information, please consult the following references:

WHO, UNODC and UNAIDS. Technical guide for countries to set targets for universal access to HIV prevention, treatment and care for injecting drug users. Geneva, World Health Organization, 2012 http://www.who.int/hiv/pub/idu/targets_universal_access/en/index.html.

A framework for monitoring and evaluating HIV prevention programmes for most-at-risk populations. Geneva, UNAIDS, 2007.

Practical guidelines for intensifying HIV prevention: towards universal access. Geneva, UNAIDS, 2007.

2.3 People who inject drugs: safe injecting practices

Percentage of people who inject drugs reporting the use of sterile injecting equipment the last time they injected

What it measures

It measures progress in preventing injecting drug use-associated HIV transmission.

Rationale

Safer injecting and sexual practices among people who inject drugs are essential, even in countries where other modes of HIV transmission predominate, because: (i) the risk of HIV transmission from contaminated injecting equipment is extremely high; and (ii) people who inject drugs can spread HIV (e.g., through sexual transmission) to the wider population.

Note: Countries with generalized epidemics may also have a concentrated sub-epidemic among people who inject drugs. If so, it would be valuable for them to calculate and report on this indicator for this population.

Number of people who inject drugs who report using sterile injecting equipment Numerator:

the last time they injected drugs

Number of people who inject drugs who report injecting drugs in the last month Denominator:

Numerator / Denominator Calculation:

Behavioural surveillance or other special surveys Method of measurement:

Respondents are asked the following questions:

1. Have you injected drugs at any time in the last month?

2. If yes: The last time you injected drugs, did you use a sterile needle and syringe?

Whenever possible, data for people who inject drugs should be collected through civil society organizations that have worked closely with this population in the field

Access to people who inject drugs as well as the data collected from them must

remain confidential Every two years

Measurement frequency:

Disaggregation:

Sex

■ Age (<25/25+)

Strengths and weaknesses

Surveying people who inject drugs can be challenging. Consequently, data obtained may not be based on a representative national sample of the people who inject drugs being surveyed. If there are concerns that the data are not based on a representative sample, these concerns should be reflected in the interpretation of the survey data. Where different sources of data exist, the best available estimate should be used. Information on the sample size, the quality and reliability of the data, and any related issues should be included in the report submitted with this indicator.

The extent of injecting drug use-associated HIV transmission within a country depends on four factors: (i) the size, stage and pattern of dissemination of the national AIDS epidemic; (ii) the extent of injecting drug use; (iii) the degree to which people who inject drugs use contaminated injecting equipment; and (iv) the patterns of sexual mixing and condom use among people who inject drugs and between people who inject drugs and the wider population. This indicator provides information on the third factor.

To maximize the utility of these data, it is recommended that the same sample used for the calculation of this indicator be used for the calculation of the other indicators related to these populations.

Further information

For further information, please consult the following references:

WHO, UNODC and UNAIDS. *Technical guide for countries to set targets for universal access to HIV prevention, treatment and care for injecting drug users*. Geneva, World Health Organization, 2012 http://www.who.int/hiv/pub/idu/targets_universal_access/en/index.html.

A framework for monitoring and evaluating HIV prevention programmes for most-at-risk populations. *Geneva, UNAIDS, 2007.*

Practical guidelines for intensifying HIV prevention: towards universal access. Geneva, UNAIDS, 2007.

2.4 HIV testing in people who inject drugs

Percentage of people who inject drugs who received an HIV test in the past 12 months and know their results

What it measures

It measures progress in implementing HIV testing and counselling among people who inject drugs.

Rationale

In order to protect themselves and to prevent infecting others, it is important people who inject drugs to know their HIV status. Knowledge of one's status is also a critical factor in the decision to seek treatment.

Note: Countries with generalized epidemics may also have a concentrated sub-epidemic among one or more key populations at higher risk. If so, they should calculate and report this indicator for those populations.

Numerator: Number of people who inject drugs respondents who have been tested for HIV

during the last 12 months and who know their results

Denominator: Number of people who inject drugs included in the sample

Calculation: Numerator / Denominator

Method of measurement:

Behavioural surveillance or other special surveys

Respondents are asked the following questions:

1. Have you been tested for HIV in the last 12 months?

If yes:

2. I don't want to know the results, but did you receive the results of that test?

Whenever possible, data for people who inject drugs should be collected through civil society organizations that have worked closely with this population in the field

Access to people who inject drugs as well as the data collected from them must

remain confidential

Measurement frequency:

Every two years

Disaggregation: Sex

■ Age (<25/25+)

Strengths and weaknesses

The data obtained may not be based on a representative national sample of the people who inject drugs being surveyed. If there are concerns that the data are not based on a representative sample, these concerns should be reflected in the interpretation of the survey data. Where different sources of data exist, the best available estimate should be used. Information on the sample size, the quality and reliability of the data, and any related issues should be included in the report submitted with this indicator.

Tracking people who inject drugs over time to measure progress may be difficult due to mobility and the hard-to-reach nature of these populations with many groups being hidden populations. Thus, information about the nature of the sample should be reported in the narrative to facilitate interpretation and analysis over time.

To maximize the utility of these data, it is recommended that the same sample used for the calculation of this indicator be used for the calculation of the other indicators related to these populations.

This indicator is most meaningful in settings where testing scale-up is relatively recent. People who tested more than 12 months ago and know they are positive will be considered "uncovered" by this indicator construction. Ideally, surveys should ask why respondents did not test in the past 12 months. If they report that they know their HIV status to be positive, they should not be included in the denominator. This indicator will be formally changed post-2015; we will ask countries that can to report against this indicator while omitting known HIV-positive persons from the denominator and state that they've done this in the comment field.

Further information

For further information, please consult the following references:

WHO/UNAIDS Working Group on Global HIV/AIDS and STI Surveillance. *Guidelines on surveillance among populations most at risk for HIV.* Geneva, UNAIDS, 2011 (http://www.unaids.org/en/media/unaids/contentassets/documents/epidemiology/2011/2011_Estimating_Populations_en.pdf).

Guidelines for using HIV testing technologies in surveillance: selection, evaluation and implementation. Geneva, World Health Organization, 2010 (http://www.who.int/hiv/pub/surveillance/hiv_testing_technologies_surveillance_.pdf).

WHO, UNODC and UNAIDS. *Technical guide for countries to set targets for universal access to HIV prevention, treatment and care for injecting drug users*. Geneva, World Health Organization, 2012 http://www.who.int/hiv/pub/idu/targets_universal_access/en/index.html.

A framework for monitoring and evaluating HIV prevention programmes for most-at-risk populations. Geneva, UNAIDS, 2007.

Practical guidelines for intensifying HIV prevention: towards universal access. Geneva, UNAIDS, 2007.

2.5 HIV prevalence in people who inject drugs

Percentage of people who inject drugs who are living with HIV

What it measures

It measures progress on reducing HIV prevalence among people who inject drugs.

Rationale

People who inject drugs typically have the highest HIV prevalence in countries with either concentrated or generalized epidemics. In many cases, prevalence among these populations can be more than double the prevalence among the general population. Reducing prevalence among people who inject drugs is a critical measure of a national-level response to HIV.

Countries with generalized epidemics may also have a concentrated sub-epidemic among people who inject drugs. If so, it is valuable for them to calculate and report on this indicator for those populations.

Numerator: Number of people who inject drugs who test positive for HIV

Denominator: Number of people who inject drugs tested for HIV

Calculation: Numerator / Denominator

Method of UNAIDS and WHO Working Group on Global HIV/AIDS and STI Surveillance: measurement: Guidelines among populations most at risk for HIV (WHO/UNAIDS, 2011)

This indicator is calculated using data from HIV tests conducted among respondents in the primary sentinel site or sites or in the context of a

surveillance survey

The sentinel surveillance sites used for the calculation of this indicator should

remain constant to allow for the tracking of changes over time

Measurement frequency:

Annual

Disaggregation: Sex

■ Age (<25/25+)

Strengths and weaknesses

In theory, assessing progress in reducing the occurrence of new infections is best done through monitoring changes in incidence over time. However, in practice, prevalence data rather than incidence data are available.

In analysing prevalence data of people who inject drugs for the assessment of prevention programme impact, it is desirable not to restrict analysis to young people but to report on those persons who are newly initiated to behaviours that put them at risk for infection (e.g. by restricting the analysis to people who have initiated injecting drug use within the last year). This type of analysis also has the advantage of not being affected by the effect of ART in increasing survival and thereby increasing prevalence.

If prevalence estimates are available disaggregated by greater than and less than one year of injecting drugs countries are strongly encouraged to report this disaggregation in their Country Progress Report, and to use the comments field for this indicator in the reporting tool to present disaggregated estimates.

Due to difficulties in accessing people who inject drugs, biases in sero-surveillance data are likely to be far more significant than in data from a more general population, such as women attending antenatal clinics. If there are concerns about the data, these concerns should be reflected in the interpretation.

An understanding of how the sampled population(s) relate to any larger population(s) sharing similar risk behaviours is critical to the interpretation of this indicator. The period during which people belong to a key population is more closely associated with the risk of acquiring HIV than age. Therefore, it is desirable not to restrict analysis to young people but to report on other age groups as well.

Trends in HIV prevalence among people who inject drugs in the capital city will provide a useful indication of HIV-prevention programme performance in that city. However, it will not be representative of the situation in the country as a whole.

The addition of new sentinel sites will increase the samples representativeness and will therefore give a more robust point estimate of HIV prevalence. However, the addition of new sentinel sites reduces the comparability of values. As such it is important to use consistent sites when undertaking trend analyses.

Further information

For further information, please consult the following links:

http://www.unaids.org/en/HIV_data/Methodology/default.asp

Revised guidelines on HIV surveillance for key populations at higher risk: WHO/UNAIDS Working Group on Global HIV/AIDS and STI Surveillance. *Guidelines on surveillance among populations most at risk for HIV.* Geneva, UNAIDS, 2011 (http://www.unaids.org/en/media/unaids)

contentassets/documents/epidemiology/2011/20110518_Surveillance_among_most_at_risk.pdf).



TARGET 3. ELIMINATE NEW HIV INFECTIONS AMONG CHILDREN BY 2015 AND SUBSTANTIALLY REDUCE AIDS-RELATED MATERNAL DEATHS

- 3.1 Percentage of HIV-positive pregnant women who receive antiretroviral medicine to reduce the risk of mother-to-child transmission
- 3.1a Percentage of women living with HIV who are provided with antiretroviral medicine for themselves or their infants during the breastfeeding period (formerly indicator 3.8)
- 3.2 Percentage of infants born to HIV-positive women receiving a virological test for HIV within 2 months of birth
- 3.3 Estimated percentage of child HIV infections from HIV-positive women delivering in the past 12 months

3.1 Prevention of mother-to-child transmission

Percentage of HIV-positive pregnant women who received antiretroviral medicine to reduce the risk of mother-to-child transmission

What it measures

This indicator measures progress in preventing mother-to-child transmission of HIV during pregnancy and delivery through the provision of antiretroviral medicine.

This indicator allows countries to monitor the coverage of antiretroviral medicines to HIV-positive pregnant women to reduce the risk for transmission of HIV to infants during pregnancy and delivery. When disaggregated by regimen, this indicator can show increased access to more effective antiretroviral drug regimens for pregnant women living with HIV. As the indicator usually measures antiretroviral drugs initially dispensed and not those consumed, it is not possible to determine adherence to the regimen in most cases.

The postpartum regimen, including ARV to reduce the risk of transmission during breastfeeding, is captured in indicator 3.1a. In addition, indicator 3.7 measures the percentage of infants born to HIV-infected women provided with ARV prophylaxis to reduce the risk of early mother-to-child transmission in the first 6 weeks.

Rationale

The risk of mother-to-child transmission can be significantly reduced by providing antiretroviral medicine (as lifelong therapy or as prophylaxis) for the mother during pregnancy and delivery, with antiretroviral prophylaxis for the infant, and antiretrovirals to the mother or child during breastfeeding (if breastfeeding), and use of safe delivery practices and safer infant feeding. The data will be used to track progress toward global and national goals towards elimination of mother-to-child transmission; to inform policy and strategic planning; for advocacy; and leveraging resources for accelerated scale up. It will help measure trends in coverage of antiretroviral prophylaxis and treatment, and when disaggregated by regimen type, will also assess progress in implementing more effective regimen and ART.

Numerator: Number of HIV-positive pregnant women who received antiretroviral medicine

during the past 12 months to reduce the risk of mother-to-child transmission during pregnancy and delivery. Global reports summarizing coverage of ARV for PMTCT will exclude women who received single dose nevirapine as it is considered a sub-optimal regimen. However the number of women who received

only a single dose of nevirapine should be reported by the country.

Denominator: Estimated number of HIV-positive women who delivered within the past 12 months

Calculation: Numerator / Denominator

Method of For the numerator: national programme records aggregated from programme measurement: monitoring tools, such as patient registers and summary reporting forms

For the denominator: estimation models such as Spectrum, or antenatal clinic surveillance surveys in combination with demographic data and appropriate

adjustments related to coverage of ANC surveys

Measurement Annual or more frequently, depending on a country's monitoring needs frequency:

Disaggregation: The numerator should be disaggregated by the six general regimens

described below

Explanation of numerator

The numerator should be disaggregated by the six categories below(the first three regimens are currently recommended by WHO) for HIV-positive pregnant women for the prevention of mother-to-child transmission:

1. Newly initiated on ART during the current pregnancy	
2. Already on ART before the current pregnancy	
3. Maternal triple ARV prophylaxis (prophylaxis component of WHO Option B)	
4. Maternal AZT (prophylaxis component during pregnancy and delivery of WHO Option A or WHO 2006 guidelines)	
5. Single dose nevirapine (with or without tail) ONLY	
6. Other (please comment: e.g. specify regimen, uncategorized, etc.)	

Disaggregation of regimen definitions

Categories	Further clarification	Common examples
The first two options include women receiving	A three-drug regimen intended to provide ART for life	Standard national treatment regimen, for example:
lifelong antiretroviral therapy (including Option B+)	Number of HIV-positive pregnant women identified in the reporting period newly initiated on ART for life	TDF+3TC+EFVAZT+3TC+NVP
newly initiated on treatment during the current pregnancy	2) Number of HIV-positive pregnant women identified in the reporting period who were already on ART at	
2) already on treatment before the pregnancy	their first ANC visit. If a woman is initiating ART for life during labour, she would be counted in category 1.	
	If the number of women on antiretroviral is not available by the timing of when they started ART the number can be included in the cell titled Total number of pregnant women on lifelong ART.	
3) Maternal triple ARV prophylaxis (prophylaxis component of WHO Option B during pregnancy and delivery)	A three-drug regimen provided for MTCT prophylaxis started during pregnancy or as late as during labour or delivery with the intention of stopping at the end of the breastfeeding period (or stopping at delivery if not breastfeeding) If a woman is receiving triple ARVs	TDF+3TC+EFVAZT+3TC+EFVAZT+3TC+LPV/r
	for the first time at labour or delivery then she should still be counted in this category if the facility is implementing Option B.	

TARGET 3: ELIMINATE NEW HIV INFECTIONS AMONG CHILDREN BY 2015 AND SUBSTANTIALLY REDUCE AIDS-RELATED MATERNAL DEATHS

4) Maternal AZT (prophylaxis component of WHO Option A during pregnancy and delivery)	A prophylactic regimen that uses AZT (or another NRTI) started as early as 14 weeks or as late as during labour or delivery to prevent HIV transmission If a woman is receiving ARVs for the first time at labour or delivery, then she should still be counted in this category if the facility is implementing Option A.	 AZT at any point before labour + intrapartum NVP AZT at any point before labour + intrapartum NVP +7 day post-partum tail of AZT/3TC
5) Single-dose nevirapine (sd-NVP) to the mother during pregnancy or delivery	 Nevirapine is the ONLY regimen provided to an HIV-positive pregnant woman during pregnancy, labour or delivery Do NOT count as sd-NVP if: Nevirapine is provided as part of Option A during pregnancy or An HIV+ pregnant woman is initiated on Option A, B, or B+ at labor and delivery 	 sd-NVP for mother ONLY at onset of labour sd-NVP + 7 day AZT/3TC tail ONLY sd-NVP for mother at onset of labour and sd-NVP for baby ONLY

The numerator must match the values included in Spectrum or an automated query will be sent requesting that the team make the values consistent.

GARPR	Spectrum
1) Newly initiated on treatment during the current pregnancy	ART started during current pregnancy
2) Already on treatment before the pregnancy	ART started before current pregnancy
3) Maternal triple ARV prophylaxis (prophylaxis component of WHO Option B during pregnancy and delivery)	Option B – Triple prophylaxis from 14 weeks
4) Maternal AZT (prophylaxis component of WHO Option A during pregnancy and delivery)	Option A—maternal AZT
5) Single-dose nevirapine (sd-NVP) to the mother during pregnancy or delivery	Single dose nevirapine
6) Other (usually limited to countries still providing maternal AZT started late in the pregnancy)	Maternal AZT according to 2006 WHO guidelines Spectrum requires data on historical regimens. This category is maintained to describe the regimens provided in previous years.

Explanation of denominator

Two methods can be used to estimate the denominator:

- 1. a projection model, such as Spectrum; use the output "number of pregnant woman needing PMTCT"; or
- 2. multiply the number of women who gave birth in the past 12 months (which can be obtained from estimates of the central statistics office or the United Nations Population Division or pregnancy registration systems with complete data) by the most recent national estimate of HIV prevalence in pregnant women (which can be derived from HIV sentinel surveillance in antenatal care clinics and appropriate adjustments related to coverage of ANC surveys.) if Spectrum projections are unavailable.

To ensure comparability the Spectrum output will be used for the denominator when global analyses are done.

Strengths and weaknesses

Countries are encouraged to track and report the actual number of women receiving the various regimens, so that the impact of antiretroviral drugs on mother-to-child transmission can be modelled on the basis of the efficacy of the regimens. If countries do not have a system for collecting and reporting data on the provision of different antiretroviral drug regimens for the prevention of mother-to-child transmission of HIV, they should establish such a system.

Further information

The prevention of mother-to-child transmission is a rapidly evolving programmatic area. Methods for monitoring coverage of this service are therefore also evolving. To access the most current information available please consult the following links:

www.who.int/hiv/pub/mtct/en/

www.who.int/hiv/pub/me/en/index.html

3.1a Prevention of mother-to-child transmission during breastfeeding

Percentage of women living with HIV who are provided with antiretroviral medicines for themselves or their infants during the breastfeeding period (formerly indicator 3.8)

What it measures

While indicator 3.1 captures whether programmes are reaching mothers during pregnancy and delivery, indicator 3.1a captures whether women are receiving prophylaxis for themselves or for their babies during the breastfeeding period.

Rationale

For women who are breastfeeding and not on antiretroviral therapy, the risk of transmitting HIV to the child during breastfeeding remains substantial. This risk can be reduced by providing prophylaxis to the mother or the baby during the entire duration of breastfeeding. The data will be used to track progress toward global and national goals towards elimination of mother-to-child transmission, to inform policy and strategic planning, for advocacy, and leveraging resources for accelerated scale up.

Numerator: Number of women living with HIV who were breastfeeding who received

antiretroviral prophylaxis for themselves or their infants to reduce the risk of mother-to-child transmission during breastfeeding during the past 12 months

Denominator: Estimated number of women living with HIV who were breastfeeding in the past

12 months

Calculation: Numerator / Denominator

Method of measurement:

For the numerator: national programme records aggregated from programme monitoring tools, such as patient registers and summary reporting forms. The data for the numerator can be collected at the infant's six-week Early Infant Diagnosis (EID) visit or DPT3 immunization visit (two to three months) and distinguished from ARV interventions given to prevent peripartum transmission. Data on whether maternal or infant antiretrovirals to reduce post-natal transmission were provided should be recorded for breastfeeding infants. HIV-infected pregnant women who are eligible for lifelong antiretroviral therapy, are receiving a treatment regimen and whose infants therefore benefit from the prophylactic effect of ART in reducing the risk of transmission through breastfeeding are also included in the numerator.

For the denominator: estimation models such as Spectrum, or antenatal clinic surveillance surveys in combination with demographic data and appropriate adjustments related to coverage of ANC surveys. The denominator should represent the number of women living with HIV who are breastfeeding. In settings where most HIV positive women breastfeed, the estimated number of HIV-positive pregnant women could be a proxy for the denominator (with some adjustment of infant deaths before the time point for measurement if available). In other settings, where a sizable population of HIV-exposed infants may not be breastfeeding, it will be necessary to estimate the number of HIV-exposed infants who are breastfeeding.

Measurement frequency:

Annual or more frequently, depending on a country's monitoring needs

Strengths and weaknesses

This indicator allows countries to monitor the coverage of programmes to reduce transmission to children during breastfeeding. As the indicator measures antiretroviral drugs dispensed and not those consumed, it is not possible to determine adherence to the regimen.

This indicator should not be confused with indicator 3.7 (Percentage of infants born to HIV-infected women provided with ARV prophylaxis to reduce the risk of early mother-to-child transmission in the first six weeks).

It is important to assess antiretroviral coverage throughout the breastfeeding period, but in many settings there is significant loss to follow-up after the six-week visit so it is difficult to get an accurate estimate of antiretroviral coverage at a later time point. In breastfeeding populations, effort should be made to ensure antiretroviral coverage during the breastfeeding period beyond six weeks or DPT3 as captured by this indicator.

If the data submitted for this indicator are not nationally representative, please state this in the comments field and describe the sample.

Further information

The prevention of mother-to-child transmission is a rapidly evolving programmatic area. Methods for monitoring coverage of this service are therefore also evolving. To access the most current information available please consult the following links:

www.who.int/hiv/pub/mtct/en

www.who.int/hiv/pub/me/en/index.html

3.2 Early infant diagnosis

Percentage of infants born to HIV-positive women receiving a virological test for HIV within 2 months of birth

What it measures

It measures progress in the extent to which infants born to HIV-positive women are tested within the first 2 months of life to determine their HIV status and eligibility for antiretroviral therapy disaggregated by test results.

Rationale

Infants infected with HIV during pregnancy, delivery or early postpartum often die before they are recognized as having HIV infection. WHO recommends national programmes to establish the capacity to provide early virological testing of infants for HIV at 6 weeks, or as soon as possible thereafter to guide clinical decision-making at the earliest possible stage. HIV disease progression is rapid in children; they need to be put on treatment as early as possible because without early treatment almost 50% of children would be dead by the second year.

Numerator: Number of infants who received an HIV test within two months of birth, during

the reporting period. Infants tested should only be counted once

Denominator: Number of HIV-positive pregnant women giving birth in the last 12 months

Calculation: Numerator / Denominator

Method of Early Infant Diagnosis (EID) testing laboratories for the numerator, and Spectrum estimates, central statistical offices, and/or sentinel surveillance for

the denominator

Measurement

frequency:

Annual or more frequently, depending on a country's monitoring needs

Explanation of numerator

To be collected from databases held at early infant diagnosis testing laboratories. The numerator should represent the number of infants who received virologic testing within two months of birth; it should not represent the number of samples tested at the laboratory. Data should be aggregated from the laboratory data bases. Where possible, double counting should be minimized when aggregating data to produce national-level data. It is expected that the number of infants receiving more than one virologic test in the first 2 months of life will be low. Efforts should be made to include all public, private and NGO-run health facilities that are providing HIV testing for HIV-exposed infants.

The test results should be reported as positive, negative, indeterminate or rejected for testing by the laboratory. When reporting this information only the most recent test result for an infant tested in the first 2 months of life should be included.

Explanation of denominator

This is a proxy measure for number of infants born to HIV-positive women.

Two methods can be used to estimate the denominator:

a) Using a projection model such as the one provided by Spectrum software use the output "the number of pregnant woman needing PMTCT" as a proxy,

or;

b) Multiplying the total number of women who gave birth in the last 12 months, (which can be obtained from central statistics office estimates of births or the UN Population Division estimates) by the most recent national estimate of HIV prevalence in pregnant women (which can be derived from HIV sentinel surveillance in ANC clinic and appropriate adjustments related to coverage of ANC surveys), if Spectrum projections are unavailable.

To ensure comparability the Spectrum output will be used for the denominator when global analyses are done.

Strengths and weaknesses

This indicator allows countries to monitor progress in providing early HIV virologic testing to HIV-exposed infants aged two months or less, critical for appropriate follow-up care and treatment. By limiting the age to two months of life or less, the chance of repeat tests for the same infant which can lead to double counting is also eliminated. The only three fields needed for this indicator: date of sample collection, age at collection (actual or calculated based upon date of birth), and results are systematically entered into central EID testing databases at testing laboratories.

Due to the small number of testing laboratories, and the electronic format of testing databases, this indicator should not have a heavy collection burden. Data quality at the laboratories is generally high, resulting in a robust indicator. The indicator does not capture the number of children with a definitive diagnosis (i.e. of HIV infection), or measure whether appropriate follow-up services were provided to the child based on interpretation of test results. It also does not measure the quality of testing nor the system in place for testing. A low value of the indicator could, however, signal systemic weaknesses, including poor country-level management of supplies of HIV virologic test kits, poor data collection, poor follow-up and mismanagement of testing samples.

Disaggregation by test results cannot be used as a proxy for overall MTCT transmission rates. If either the EID coverage of national need or the EID testing coverage in the first two months of life is low, low positivity rates among infants tested will not necessarily mean program success, as many other infants who are likely positive are not represented in this sample.

While early virological testing is a critical intervention for identifying infected infants, it is also important for countries to strengthen the quality of HIV-exposed infant follow-up and to train health providers to recognize signs and symptoms of early HIV infection among exposed infants, particularly where access to virological testing is limited. Inappropriate management of supplies can negatively affect the value of the indicator and significantly reduce access to HIV testing for infants born to HIV-positive women. Countries should ensure that appropriate systems and tools, particularly tools for LMIS, are in place to procure, distribute and manage supplies at facility, district and central level.

Further information

For further information, please consult the following reference and website:

WHO, UNICEF and UNAIDS. *Towards universal access: scaling up priority HIV/AIDS interventions in the health sector. Progress report, September 2010.* Geneva, World Health Organization, 2010 (http://www.who.int/hiv/pub/2009progressreport/en/index.html).

Next generation indicators reference guide. Washington, DC, United States President's Emergency Plan for AIDS Relief.2009.

Monitoring and evaluation toolkit. Part 2. Tools for monitoring programs for HIV, tuberculosis, malaria and health systems strengthening. Geneva, Global Fund to Fight AIDS, Tuberculosis and Malaria, 2009.

Measuring the impact of national PMTCT programmes: towards the elimination of new HIV infections among children by 2015 and keeping their mothers alive. A short guide on methods. Geneva, World Health Organization, 2012.

3.3 Mother-to-child transmission of HIV (modelled)

Estimated percentage of child HIV infections from HIV-positive women delivering in the past 12 months

What it measures

It measures progress towards eliminating mother-to-child HIV transmission.

Rationale

Efforts have been made to increase access to interventions that can significantly reduce mother-to-child transmission, including combination antiretroviral prophylactic and treatment regimens and strengthened infant-feeding counselling. It is important to assess the impact of PMTCT interventions in reducing new paediatric HIV infections through mother-to-child transmission.

The percentage of children who are HIV-positive should decrease as the coverage of interventions for PMTCT and the use of more effective regimens increases.

Numerator: The numerator is the estimated number of children who will be newly infected with

HIV due to mother-to-child transmission among children born in the previous

12 months to HIV-positive women

Denominator: Estimated number of HIV positive women who delivered in the previous

12 months

Calculation: Numerator / Denominator

Method of measurement:

The mother-to-child transmission probability differs with the antiretroviral drug regimen received and infant-feeding practices. The transmission can be calculated by using the Spectrum model. The Spectrum computer programme uses the information on:

- a. the distribution of HIV-positive pregnant women receiving different antiretroviral regimens prior to and during delivery (peripartum) by CD4 category of the mother
- b. the distribution of women and children receiving antiretrovirals after delivery (postpartum) by CD4 category of the mother.
- c. the percent of infants who are not breastfeeding in PMTCT programmes by age of the child
- d. mother-to-child transmission of HIV probabilities based on various categories of antiretroviral drug regimen and infant feeding practices

The estimated national transmission rate is reported in the PMTCT summary display in Spectrum. This variable can also be calculated using the variables in Spectrum on "New HIV infections" for children 0-14 years and dividing this by the variable "Women in need of PMTCT"

There is not enough information available about other HIV transmission routes for children to include such infections in the model. In addition other modes of transmission are believed to be a small fraction of the overall infections among children. The Spectrum output variable "New HIV infections for children 0-1 years" is not used because some infections due to breastfeeding will take place after age 1 year

Measurement frequency:

Annual

Disaggregation:

None

To ensure comparability the Spectrum output will be used for calculating this indicator when global analyses are done.

Strengths and weaknesses

Over time, this indicator assesses the ability of PMTCT programmes by estimating the impact of increases in the provision of antiretroviral drugs and the use of more efficacious regimens and optimal infant feeding practice. This indicator is generated from a model, which provides estimates of HIV infection in children. The estimated indicator is reliant on the assumptions and data used in the model. The indicator may not be a true measure of mother-to-child transmission. For example, in countries where other forms of PMTCT (e.g. Caesarean section) are widely practised, the indicator will overestimate mother-to-child transmission. It also relies on programme data that often captures antiretroviral drug regimens provided rather than taken, thus could underestimate mother-to-child transmission.

This indicator allows countries to assess the impact of PMTCT programmes by estimating the HIV transmission rate from HIV positive women to their children. It is difficult to follow up mother–children pairs, particularly at national level, because of the lag in reporting and the multiple health facility sites that mother-child pairs can visit for the wide range of PMTCT and child care interventions delivered over a timespan. In countries where data are available, facility attendance is high, and confirmatory tests are conducted systematically, efforts should be made to monitor the impact through directly assessing the percentage of children found to be HIV-positive among those born to HIV-positive mothers. All countries should make efforts to monitor the HIV status and survival of children born to HIV-positive women, gathered during follow-up health care visits.

Further information

http://www.who.int/hiv/pub/me/en/index.html



TARGET 4. REACH 15 MILLION PEOPLE LIVING WITH HIV WITH LIFESAVING ANTIRETROVIRAL TREATMENT BY 2015

- 4.1 Percentage of adults and children currently receiving antiretroviral therapy*
- 4.2 Percentage of adults and children with HIV known to be on treatment 12 months after initiation of antiretroviral therapy

^{*}Millennium Development Goals indicator

4.1 HIV treatment: antiretroviral therapy

Percentage of adults and children currently receiving antiretroviral therapy among all adults and children living with HIV

What it measures

Progress towards providing antiretroviral therapy to all people for treatment.

Rationale

Antiretroviral therapy (ART) has been shown to reduce HIV-related morbidity and mortality amongst those living with HIV and to reduce transmission of HIV. In recent years the guidelines on eligibility for antiretroviral therapy have changed a number of times. In addition national guidelines do not always match global guidelines. As a result, antiretroviral therapy coverage has been reported in numerous ways including being based on global guidelines, on national guidelines or both. When the guidelines are modified to include more people who are living with HIV, the coverage values for countries decrease. To avoid multiple antiretroviral therapy coverage values the number of people on antiretroviral therapy will be presented in relation to the total number of people living with HIV. The estimated coverage using all people living with HIV as a denominator is similar to the denominator been all people eligible for antiretroviral therapy under the 2013 antiretroviral therapy guidelines. Approximately 85% of people living with HIV are eligible under the 2013 WHO criteria for antiretroviral therapy provision.

Numerator: Number of adults and children currently receiving antiretroviral therapy in

accordance with the nationally approved treatment protocol (or WHO standards)

at the end of the reporting period

Denominator: Estimated number of adults and children living with HIV

National criteria for ART eligibility varies by country. To make this indicator comparable across countries global reports will present the ART coverage for

adults and children as a percentage of all people living with HIV.

Calculation: Numerator / Denominator

Additional Although coverage will be calculated using the total number of people living with information: HIV, please also provide the number eligible for ART under your national ART

criteria guidelines.

Method of Data should be collected continuously at the facility level. Data should be measurement: aggregated periodically. The most recent full year of data should be used for

annual reporting.

For the numerator: facility-based antiretroviral therapy registers and corresponding cross-sectional forms. For the denominator: HIV estimation

models such as Spectrum

Measurement Data should be collected continuously at the facility level. Data should be frequency: aggregated periodically, preferably monthly or quarterly. The most recent

monthly or quarterly data should be used for annual reporting

Disaggregation:

- Sex
- Age (less than 15 years, 15 years and older, 15-49, <1 year, 1-4 years, 5-9, 10-14, 15-19, 20-24, 25-49, 50+)
- Public/Private
- persons newly initiating antiretroviral therapy during the last reporting year (this indicator should be available from the same sources as the total number of people receiving antiretroviral therapy)

Explanation of numerator

The numerator can be generated by counting the number of adults and children who received antiretroviral therapy at the end of the reporting period.

The numerator should equal the number of adults and children who ever started antiretroviral therapy minus those patients who are not currently on treatment prior to the end of the reporting period.

Patients not currently on treatment at the end of the reporting period, in other words, those who are excluded from the numerator, are patients who died, stopped treatment or are lost to follow-up.

Some patients pick up several months of antiretroviral drugs at one visit, which could include antiretroviral medicine received for the last months of the reporting period, but not be recorded as visits for the last months in the patient register. Efforts should be made to account for these patients, as they need to be included in the numerator.

Antiretroviral medicines taken only for the purpose of prevention of mother-to-child transmission and postexposure prophylaxis are not included in this indicator. HIV-positive pregnant women who are on lifelong antiretroviral therapy are included in this indicator.

The number of adults and children currently receiving antiretroviral therapy can be obtained through data collected from facility-based antiretroviral therapy registers or drug supply management systems. These are then tallied and transferred to cross-sectional monthly or quarterly reports which can then be aggregated for national totals.

Patients receiving antiretroviral therapy in the private sector and public sector should be included in the numerator where data are available.

Explanation of denominator

The denominator is generated by estimating the number of people living with HIV. In previous years UNAIDS and WHO have reported on the percentage eligible based on the number eligible according to WHO criteria. In 2014 this will change to include all people living with HIV. This does not endorse the concept that all people living with HIV should receive antiretroviral therapy; instead this is a simpler measure that will not change over time and will result in coverage values that are consistent when compared globally and when calculated for national purposes.

Denominator estimates are most often based on the latest data available from sentinel surveillance used with a HIV modeling programme such as Spectrum. For further information on estimates of HIV need and the use of Spectrum please refer to the UNAIDS/WHO Reference Group on Estimates, Modelling and Projections methodology.¹⁵

Strengths and weaknesses

This indicator permits monitoring trends in coverage but does not attempt to distinguish between different forms of antiretroviral therapy or to measure the cost, quality or effectiveness of, or adherence to the treatment regimen provided. These will each vary within and between countries and are liable to change over time.

TARGET 4: REACH 15 MILLION PEOPLE LIVING WITH HIV WITH LIFESAVING ANTIRETROVIRAL TREATMENT BY 2015

The degree of utilization of antiretroviral therapy will depend on factors such as cost relative to local incomes, service delivery infrastructure and quality, availability and uptake of testing and counselling services, and perceptions of effectiveness and possible side effects of treatment.

The indicator measures the number of people provided with medication but does not measure whether the individual imbibed the medication thus it is not a measure of adherence.

Further information

http://www.who.int/hiv/topics/treatment/en/index.html

4.2 Twelve-month retention on antiretroviral therapy

Percentage of adults and children with HIV known to be on treatment 12 months after initiation of antiretroviral therapy

What it measures

It measures progress in increasing survival among infected adults and children by maintaining them on antiretroviral therapy.

Rationale

One of the goals of any antiretroviral therapy, programme is to increase survival among infected individuals. As antiretroviral therapy. is scaled up in countries around the world, it is also important to understand why and how many people drop out of treatment programmes. These data can be used to demonstrate the effectiveness of those programmes and highlight obstacles to expanding and improving them.

Number of adults and children who are still alive and on antiretroviral therapy at Numerator:

12 months after initiating treatment

Total number of adults and children who initiated antiretroviral therapy who Denominator:

> were expected to achieve 12-month outcomes within the reporting period, including those who have died since starting antiretroviral therapy., those who have stopped antiretroviral therapy, and those recorded as lost to follow-up at

month 12

Numerator / Denominator Calculation:

Programme monitoring tools; cohort/group analysis forms Method of

measurement: Antiretroviral therapy registers and antiretroviral therapy cohort analysis report

The reporting period is defined as any continuous 12-month period that has ended within a pre-defined number of months from the submission of the report. The pre-defined number of months can be determined by national reporting requirements. If the reporting period is January 1 to December 31, 2013, countries will calculate this indicator by using all patients who started antiretroviral therapy. any time during the 12-month period from January 1 to December 31, 2012. If the reporting period is July 1, 2012 to June 30, 2013, countries will include patients who started antiretroviral therapy from July 1, 2011 to June 30, 2012

A 12-month outcome is defined as the outcome (i.e., whether the patient is still alive and on antiretroviral therapy, dead or lost to follow-up) at 12 months after starting antiretroviral therapy. For example, patients who started antiretroviral therapy during the 12-month period from January 1 to December 31, 2011 will have reached their 12-month outcomes for the reporting period of January 1 to

December 31, 2013

Measurement frequency:

As patients start antiretroviral therapy, monthly cohort data should be collected continuously for these patients. Data for monthly cohorts that have completed at least 12 months of treatment should then be aggregated

Disaggregation:

- Sex
- Age (<15, 15+)
- Pregnancy status at start of therapy
- Breastfeeding status at start of therapy

Explanation of numerator

The numerator requires that adult and child patients must be alive and on antiretroviral therapy 12 months after their initiation of treatment. For a comprehensive understanding of survival, the following data must be collected:

- Number of adults and children in the antiretroviral therapy start-up groups initiating antiretroviral therapy at least 12 months prior to the end of the reporting period;
- Number of adults and children still alive and on antiretroviral therapy at 12 months after initiating treatment.

The numerator does not require patients to have been on antiretroviral therapy continuously for the 12-month period. Patients who may have missed one or two appointments or drug pick-ups, and temporarily stopped treatment during the 12 months since initiating treatment but are recorded as still being on treatment at month 12 are included in the numerator. On the contrary, those patients who have died, stopped treatment or been lost to follow-up at 12 months since starting treatment are not included in the numerator.

For example, for those patients who started antiretroviral therapy in May 2012, if at any point during the period May 2012 to May 2013 these patients die, are lost to follow-up (and do not return) or stop treatment (and do not restart), then at month 12 (May 2013), they are not on antiretroviral therapy, and not included in the numerator. Conversely, a patient who started antiretroviral therapy in May 2012 and who missed an appointment in June 2012, but is recorded as on antiretroviral therapy in May 2013 (at month 12) is on antiretroviral therapy and will be included in the numerator. What is important is that the patient who has started antiretroviral therapy in May 2012 is recorded as being alive and on antiretroviral therapy after 12 months, regardless of what happens from May 2012 to May 2013.

ART registries should include a number of variables describing the patients. For example the age of the patient at the start of ART. In addition many registries will include information indicating whether the patient was pregnant or was breastfeeding at the start of ART. ART retention for these sub-sets of women should be calculated to determine ART retention at 12 months for pregnant women and for breastfeeding women.

Explanation of denominator

The denominator is the total number of adults and children in the antiretroviral therapy start-up groups who initiated antiretroviral therapy at any point during the 12 months prior to the beginning of the reporting period, regardless of their 12-month outcome.

For example, for the reporting period January 1 to December 31, 2013, this will include all patients who started antiretroviral therapy during the 12-month period from January 1 to December 31, 2012. This includes all patients, both those on antiretroviral therapy as well as those who are dead, have stopped treatment or are lost to follow-up at month 12.

At the facility level, the number of adults and children on antiretroviral therapy at 12 months includes patients who have transferred in at any point from initiation of treatment to the end of the 12-month period and excludes patients who have transferred out during this same period to reflect the net current cohort at each facility. In other words, at the facility level, patients who have transferred out will not be counted either in the numerator or the denominator. Similarly, patients who have transferred in will be counted in both the numerator and denominator. At the national level, the number of transferred-in patients should match the number of transferred-out patients. Therefore, the net current cohort (the patients whose outcomes the facility is currently responsible for recording—the number of patients in the start-up group plus any transfers in, minus any transfers out) at 12 months should equal the number in the start-up cohort group 12 months prior.

Strengths and weaknesses

Using this denominator may underestimate true "survival", since a proportion of those lost to follow-up are alive. The number of people alive and on antiretroviral therapy (i.e. retention on antiretroviral therapy) in a treatment cohort is captured here.

Priority reporting is for aggregate survival reporting. If comprehensive cohort patient registries are available then it is encouraged for countries to track retention on treatment at 24, 36, and 48 months and yearly thereafter. This will enable comparison over time of survival on ART. As it stands, it is possible to identify whether survival at 12 months increases or decreases over time. However, it is not possible to attribute cause to these changes. For example, if survival at 12 months increases over time, this may reflect an improvement in care and treatment practices or earlier initiation of ART. The retention on antiretroviral therapy at 12 months therefore needs to be interpreted in view of the baseline characteristics of the cohort of patients at the start of antiretroviral therapy: mortality will be higher in sites where patients accessed antiretroviral therapy at a later stage of infection. Therefore, collection and reporting of survival over longer durations of treatment outcomes may provide a better picture of the long-term effectiveness of ART.

Further information

http://www.who.int/hiv/topics/treatment/en/index.html



TARGET 5. REDUCE TUBERCULOSIS DEATHS IN PEOPLE LIVING WITH HIV BY 50% BY 2015

5.1 Percentage of estimated HIV-positive incident TB cases that received treatment for both TB and HIV

5.1 Co-management of tuberculosis and HIV treatment

Percentage of estimated HIV-positive incident TB cases that received treatment for both TB and HIV

What it measures

It measures progress in detecting and treating TB in people living with HIV.

Rationale

Tuberculosis (TB) is a leading cause of morbidity and mortality in people living with HIV, including those on antiretroviral therapy. Intensified TB case-finding and access to quality diagnosis and treatment of TB in accordance with international/national guidelines is essential for improving the quality and quantity of life for people living with HIV. A measure of the percentage of HIV-positive TB cases that access appropriate treatment for their TB and HIV is important.

Number of adults and children with HIV infection who received antiretroviral Numerator:

> combination therapy in accordance with the nationally approved treatment protocol (or WHO/UNAIDS standards) and who were started on TB treatment (in accordance with national TB programme guidelines), within the reporting year

Estimated number of incident TB cases in people living with HIV Denominator:

Annual estimates of the number of incident TB cases in people living with HIV in

high TB burden countries are calculated by WHO and are available at:

http://www.who.int/tb/country/en

Numerator / Denominator Calculation:

Facility antiretroviral therapy registers and reports; programme monitoring tools Method of

measurement:

Programme data and estimates of incident TB cases in people living with HIV

Measurement

frequency:

Data should be collected continuously at the facility level. Data should be aggregated periodically, preferably monthly or quarterly, and reported annually.

The most recent year for which data and estimates are available should be

reported here

Strengths and weaknesses

Adequate detection and treatment of TB will prolong the lives of people living with HIV and reduce the community burden of TB. WHO provides annual estimates of the burden of TB among people living with HIV, based on the best available country estimates of HIV prevalence and TB incidence. All incident TB cases among people living with HIV should be started on TB treatment and depending on country specific eligibility criteria. Incident TB cases are defined as new cases that have occurred in that year, and this does not include latent cases. All people living with HIV who have TB should be given antiretroviral therapy as soon as possible. TB treatment should be started in accordance with national TB programme guidelines..

This indicator provides a measure of the extent to which collaboration between the national TB and HIV programmes is ensuring that people with HIV and TB disease are able to access appropriate treatment for both diseases. However, this indicator will also be affected by low uptake of HIV testing, poor access to HIV care services and ART, and poor access to TB diagnosis and treatment. Separate indicators exist for each of these factors and should be referred to when interpreting the results of this indicator.

It is important that those providing HIV care and antiretroviral therapy record TB diagnosis and treatment, as this information has important implications for antiretroviral therapy eligibility and choice of antiretroviral regimen. It is therefore recommended that the date of starting TB treatment is recorded in the ART register.

Further information

For further information, please consult the following reference:

Global tuberculosis control: surveillance, planning, financing. Geneva, World Health Organization, 2009 (http://www.who.int/tb/country/en).

A guide to monitoring and evaluation for collaborative TB/HIV activities. Geneva, World Health organization, 2009 (http://www.who.int/hiv/pub/tb/hiv_tb_monitoring_guide.pdf)



TARGET 6. CLOSE THE GLOBAL AIDS RESOURCE GAP BY 2015 AND REACH ANNUAL GLOBAL INVESTMENT OF US\$22–24 BILLION IN LOW- AND MIDDLE-INCOME COUNTRIES

6.1 Domestic and international AIDS spending by categories and financing sources

6.1 AIDS spending

Domestic and international AIDS spending by categories and financing sources

What it measures

It measures how funds are spent at the national level and where those funds are sourced in an accurate and consistent manner.

Rationale

As the national and international response to AIDS continues to scale up, it is increasingly important to accurately track in detail: i) how funds are spent at the national level and ii) where the funds originate. The data are used to measure annual global HIV expenditures, which is an important component of Monitoring the 2011 UN Political Declaration on HIV and AIDS. In addition, the data help national-level decision-makers monitor the scope and effectiveness of their programmes. When aggregated across multiple countries, the data also help the international community evaluate the status of the global response. This piece of strategic information supports the coordination role of the National AIDS Authority in each country and provides the basis for resource allocation and improved strategic planning processes.

Since different countries can choose among different methodologies and tools to monitor the flow of AIDS funding – i.e. National AIDS Spending Assessments (NASA), AIDS sub-account of the System of Health Accounts (SHA) or National Health Accounts (NHA) and ad hoc Resource Flows Surveys – the National Funding Matrix includes a spreadsheet that allows financial data from any of these three methodologies to be easily entered, reviewed and reported.

Measurement Tool: Prima

Primary tool/method:

1) NASA

Alternative tools/methods:

- 2) SHA/NHA—AIDS sub-accounts. There should not be any difference in the AIDS health spending measured by NASA or by the SHA/NHA subaccounts. However, some activities performed outside the health sector might not be included in Systemic/National Health Accounts
- 3) Resource Flows (RF) Survey. There has been an alignment process and countries that have been selected in the sample of this survey and have responded to the questionnaires may enter the information in the funding matrix at the aggregated level by main activities. Some activities performed outside the health sector might not be included in this RF Survey. In addition, some population-related actions should be excluded from the total for AIDS

The outputs from any of these measurement tools are to be used to complete the National Funding Matrix, which is to be submitted as part of the Country Progress Report (see Appendix 2).

Method of measurement:

The indicator on domestic and international AIDS spending is reported by completing the National Funding Matrix. Appendix 2 provides further instructions on how to submit the report of this indicator via the completed National Funding Matrix. The cover sheet as well as the information indicated in Appendix 2 needs to be submitted with the Country Progress Report.

Actual expenditures classified by eight AIDS Spending Categories and by financing source, including public expenditure from its own sources (i.e. government revenues such as taxes) and from international sources:

- 1. Prevention;
- 2. Care and treatment;
- 3. Orphans and vulnerable children;
- 4. Systems strengthening & programme coordination (this category was renamed from "Program management and administration strengthening" used in previous GARPR rounds)
- 5. Incentives for human resources;
- Social protection and social services (excluding orphans and vulnerable children);
- 7. Enabling environment and community development
- 8. Research (excluding operations research included under Systems Strengthening & Programme Coordination)

(There are multiple sub-categories in each AIDS Spending Category; see Appendix 2)

Three main groups of financing sources:

- 1. Domestic public;
- 2. International:
- 3. Domestic private (optional for global AIDS progress report reporting)

(There are multiple sub-categories for each source; see Appendix 2)

Measurement frequency:

2011, 2012 and 2013 (as available)

Calendar or fiscal year data (as available)

Countries that have submitted data for 2011 and/or 2012 already in their last report (GARPR 2013) do not need to fill this data again, unless data in the last report were missing, incomplete or there have been changed as more information has become available since. In this case, we will replace the data in our database reported for 2011 and/or 2012 in the last submission (GARPR 2013) with the newer data reported in the 2014 submission.

Strengths and weaknesses

The financial data entered in the National Funding Matrix must be actual expenditures, not budgets or commitments. They must also include AIDS expenditures that were made as part of broader systems of service provision. For example, the diagnosis and treatment of opportunistic infections would require a special costing estimate to track the specific resources allocated to AIDS-related diagnosis and treatment. Similarly, prevention activities in schools may benefit from a detailed estimation to calculate actual expenditures on AIDS activities. The AIDS expenditures might occur outside the health system given the nature of expanded responses to AIDS.

Completing the National Funding Matrix will provide a more detailed picture of the situation at the country level, which is useful for both national and global decision-making.

Further information

For further information, please consult the following references:

National AIDS Spending Assessment (NASA): classification, taxonomy and definitions. Geneva, UNAIDS, 2009 (http://www.unaids.org/en/dataanalysis/tools/nasapublications):

UNFPA/UNAIDS/NIDI. Details on resource flow surveys, survey instruments, countries sampled and more details on this tool are available at: www.resourceflows.org.

World Bank, WHO and USAID. *Guide to producing national health accounts*. Geneva, World Health Organization, This publication and other tools for national health accounts and AIDS sub-accounts can be found at: http://www.who.int/nha.

USAID, Health Systems 20/20 and UNAIDS. *Linking NASA and NHA concepts and mechanics*. Geneva, UNAIDS, 2009 (http://www.unaids.org/en/dataanalysis/tools/nasapublications).



TARGET 7: ELIMINATING GENDER INEQUALITIES

7.1 Proportion of ever-married or partnered women aged 15-49 who experienced physical or sexual violence from a male intimate partner in the past 12 months

7.1 Prevalence of recent intimate partner violence

Proportion of ever-married or partnered women aged 15-49 who experienced physical or sexual violence from a male intimate partner in the past 12 months.

What it measures

It measures progress in reducing prevalence of intimate partner violence against women (as an outcome itself and as a proxy for gender inequality).

An intimate partner is defined as a cohabiting partner, whether or not they had been married at the time. The violence could have occurred after they had separated.

Rationale

Globally, and particularly in sub-Saharan Africa, the observed high rates of HIV infection in women have brought into sharp focus the problem of violence against women. There is growing recognition that women and girls' risk of, and vulnerability to, HIV infection is shaped by deep-rooted and pervasive gender inequalities - violence against them in particular. Studies conducted in many countries indicate that a substantial proportion of women have experienced violence in some form or another at some point in their life. Studies from Rwanda, Tanzania, and South Africa show up to three-fold increases in risk of HIV among women who have experienced violence compared to those who have not.

Numerator: Women aged 15-49 who currently have or ever had an intimate partner, who

report experiencing physical or sexual violence by at least one of these partners in

the past 12 months

Denominator: Total women surveyed aged 15-49 who currently have or had an intimate partner

Calculation: Numerator / Denominator

Method of Population based surveys that are already being used within countries, such as WHO multi-country surveys, DHS/AIS (domestic violence module),

International Violence against Women Surveys (IVAWS)

Data collection on violence against women requires special methodologies that adhere to the ethical and safety standards to ensure that information is gathered in an ethical manner that does not pose a risk to study subjects, and in a way that

maximizes data validity and reliability

Measurement Frequency 3-5 years

Disaggregation: • Age (15-19, 20-24 and 25-49)

• HIV status (if available)

Explanation of numerator

Ever married or partnered women aged 15-49 include women who have ever been married or had an intimate partner. An intimate partner is defined as a cohabiting partner, whether or not they had been married at the time. These women are asked if they experienced physical or sexual violence from a male intimate partner in the past 12 months. Physical or sexual violence is determined by asking women if their partner did any of the following:

- Slapped her or threw something at her that could hurt her
- Pushed her or shoved her

- Hit her with a fist or something else that could hurt
- Kicked her, dragged her or beat her up
- Choked or burned her
- Threatened her with or used a gun, knife or other weapon against her
- Physically forced her to have sexual intercourse against her will
- Forced her to do something sexual she found degrading or humiliating
- Made her afraid of what he would do if she did not have sexual intercourse with him

Those reporting at least one incident corresponding to any one of these items the last 12 months are included in the numerator.

Explanation of denominator

Total women surveyed aged 15-49 who currently have or had an intimate partner.

Strengths and weaknesses

This indicator assesses progress in reducing the proportion of women who have experienced recent IPV, as an outcome in of itself. Further, the indicator should also be interpreted as a proxy for gender equality. A change in the prevalence level of recent violence over time will indicate a change in the level of gender equality—which is one of the structural factors driving the HIV epidemic. Gender equality has a clear, inverse relationship with IPV: In countries where IPV is high, gender equality, women's rates of education, and women's reproductive health and rights are low.

The indicator focuses on recent IPV, rather than ever experience of IPV, in order to enable monitoring and evaluating progress over time. Ever experience of IPV would show little change over time, no matter what the level of programming, since the numerator would include the same women for as long as they fell into the target age group. Sustained reductions in IPV are not possible without fundamental changes in unequal gender norms, gender relations at the household and community level, women's legal and customary rights, gender inequalities in access to health care, education, and economic and social resources, and male involvement in reproductive and child health. Thus, changes in this one IPV indicator will be a bellwether for changes in the status and treatment of women in all the different societal domains, which in turn directly and indirectly contributes to reduced risk of HIV.

Even after adhering to the WHO ethical and safety guidelines and providing a good setting in which to conduct interviews, there will always be some women who will not disclose this information. This means that estimates will likely be more conservative than the actual level of violence which has taken place in the surveyed population.

The complex relationship between violence against women and HIV has been conceptually illustrated in a comprehensive review of the current state of evidence and practice in developing and implementing interventions and strategies to address the intersection of violence against women and HIV. For over a decade, research world-wide has documented the undeniable link between violence against women (VAW) and HIV. Studies have demonstrated an association between VAW and HIV as both a contributing factor for infection as well as a consequence of infection. This relationship operates through a variety of direct and indirect mechanisms. For example:

- fear of violence may keep women from insisting on condom use by a male partner whom they suspect is HIV infected;
- fear of IPV may keep women from disclosing their HIV status or seeking treatment;
- forced vaginal penetration increases the likelihood of HIV transmission;
- rape is one manifestation of gender inequality and can result in HIV infection, although this represents a minority of cases; and
- rape, other sexual and physical abuse can result in psychological distress that is manifested in risky sexual behaviour, with the result of becoming infected with HIV.

Further information

Investing in gender equality: ending violence against women and girls. UNIFEM Brief, October 2010. Geneva, World Health Organization, 2010.

Addressing violence against women and HIV/AIDS: what works? Geneva, World Health Organization.

Dunkle KL, Head S, Garcia Moreno C. Current intervention strategies at the intersection of gender-based violence and HIV: a systematic review of the peer-reviewed literature describing evaluations of interventions addressing the interface between gender, violence and HIV. Geneva, World Health Organization, 2009.

Gender-based violence and HIV. Cambridge, MA, Program on International Health and Human Rights, Harvard School of Public Health, 2009.

Maman S et al. The intersections of HIV and violence: directions for future research and interventions. Social Science and Medicine, 2000, 50:459–478.



TARGET 8: ELIMINATING STIGMA AND DISCRIMINATION

8.1 Discriminatory attitudes towards people living with HIV

8.1 Discriminatory attitudes towards people living with HIV

Percentage of women and men aged 15–49 who report discriminatory attitudes towards people living with HIV

What it measures

It measures progress towards reducing discriminatory attitudes, and support for discriminatory policies.

Rationale

Discrimination is a human rights violation and is prohibited by international human rights law and most national constitutions. Discrimination in the context of HIV refers to unfair or unjust treatment (an act or an omission) of an individual based on his or her real or perceived HIV status. Discrimination exacerbates risks and deprives people of their rights and entitlements, fuelling the HIV epidemic.

This indicator is not a direct measure of discrimination but rather a measure of discriminatory attitudes which may result in discriminatory actions (or omissions). One item in this indicator measures the potential support by the respondents for discrimination that takes place at an institution while the other measures social distancing or behavioural expressions of prejudice. The composite indicator can be monitored as a measure of a key manifestation of HIV-related stigma and the potential for HIV-related discrimination within the general population. This indicator could provide further understanding and improve interventions in the area of HIV discrimination by: (1) showing change over time in the percentage of people with discriminatory attitudes, (2) allowing comparisons between national, provincial, state and more local administrations, and (3) pointing to priority areas for action.

Numerator: Number of respondents (aged 15–49 years) who respond "No" or "It depends" to

any of the two questions.

Denominator: Number of all respondents aged 15–49 years who have heard of HIV

Calculation: Numerator / Denominator

Method of measurement:

Population-based surveys (Demographic and Health Survey, AIDS Indicator Survey, Multiple Indicator Cluster Survey or other representative survey) This indicator is constructed from responses of respondents in a general

population survey who have heard of HIV to the following set of prompted

questions:

 Would you buy fresh vegetables from a shopkeeper or vendor if you knew that this person had HIV? (Yes; No; It depends; Don't know/ Not sure)1

 Do you think children living with HIV should be able to attend school with children who are HIV negative? (Yes; No; It depends; Don't know/ Not sure)

Measurement Frequency Every 3–5 years

Disaggregation: Responses for each of the individual questions (based on the same denominator)

are required as well as the consolidated response for the composite indicator.

Explanation of numerator

Those who have never heard of HIV and AIDS should be excluded from the numerator and denominator. Participants who respond "Don't Know/Not sure" and those who refuse to answer should also be excluded from the analyses.

It is important to assess the proportion of eligible survey participants who respond "Don't Know/Not sure" or who refuse to answer the questions. A high proportion of Don't Know/Not sure responses and refusals will reduce the precision of the results and may indicate problems with applicability of the question within the survey setting.

Strengths and weaknesses

This indicator directly measures discriminatory attitudes and support for discriminatory policies.

The question about buying vegetables is virtually identical to the question that has been used within DHS surveys for monitoring "accepting attitudes" towards people living with HIV, thereby enabling continued monitoring of trends. These measures improve upon the previously used measures for the "accepting attitudes" indicator as they are applicable in both high and low HIV prevalence settings, in both high and low income countries and are relevant across a wide cultural range. Individual measures and the composite indicator do not rely on the respondent having observed overt acts of discrimination against people living with HIV, which in many contexts are rare and difficult to both characterize and quantify. Rather, the individual measures and the composite indicator assess individuals' attitudes, which may have a more direct role in influencing behaviour.

The recommended questions assess agreement with hypothetical situations rather than measuring events of discrimination witnessed, and therefore social desirability bias may occur, leading to under-reporting of discriminatory attitudes. There is no mechanism for examining the frequency with which discrimination occurs, or the severity of the forms of discrimination.

In addition to conducting surveys that measure the prevalence of discriminatory attitudes in a community, where possible it would be ideal to collect qualitative data to inform the origins of discrimination. It would also be advisable to routinely collect data from people living with HIV about actual experiences of stigma and discrimination via the PLHIV Stigma Index process (www.stigmaindex.org) and compare findings with the data derived from the discriminatory attitudes indicator.

Further information

For further information on stigma and discrimination, and efforts to measure their prevalence, please see:

Thematic Segment on Non-Discrimination, 31st meeting of the UNAIDS Programme Coordinating Board. Background Note. (www.unaids.org/en/media/unaids/contentassets/documents/pcb/2012/20121111_PCB%2031_Non%20Discrimination_final_newcoverpage_en.pdf)

Stangl, A., Brady, L., Fritz, K. Technical Brief: Measuring HIV Stigma and Discrimination. Washington DC and London: International Center for Research on Women and London School of Tropical Medicine; STRIVE, 2012 (http://strive.lshtm.ac.uk/system/files/attachments/STRIVE_stigma%20 brief-A4.pdf).

Stangl, A., Lloyd, J., Brady, L. et al. A systematic review of interventions to reduce HIV-related stigma and discrimination from 2002 to 2013: how far have we come? Journal of the International AIDS Society. 2013, vol 16 Supplement (www.jiasociety.org/index.php/jias/issue/view/1464).

www.stigmaactionnetwork.org

For further information on DHS/AIS methodology and survey instruments, please visit: www.measuredhs.com Special Note for the 2014 Reporting Round:

 As this indicator is new, it is likely that many countries will not be able to report on the indicator during the 2014 reporting round;

- Instead countries are requested to report data from the previous version of question 1, 'Would you buy fresh vegetables from a shopkeeper or vendor if you knew that this person had the AIDS virus?'. This question has been routinely collected in DHS in many countries.
- In future reporting rounds, countries should report on the full indicator.

Special Note for the 2014 Reporting Round:

- As this indicator is new, it is likely that many countries will not be able to report on the indicator during the 2014 reporting round;
- Instead countries are requested to report data from the previous version of question 1, 'Would you buy fresh vegetables from a shopkeeper or vendor if you knew that this person had the AIDS virus?'. This question has been routinely collected in DHS in many countries.
- In future reporting rounds, countries should report on the full indicator.



TARGET 9: ELIMINATE TRAVEL RESTRICTIONS

Travel restriction data are collected directly by the Human Rights and Law Division at UNAIDS and no reporting is therefore needed.



TARGET 10: STRENGTHENING HIV INTEGRATION

- 10.1 Current school attendance among orphans and non-orphans aged 10–14*
- 10.2 Proportion of the poorest households who received external economic support in the last 3 months

 $^{{\}it *Millennium\ Development\ Goals\ indicator}$

10.1 Orphans school attendance

Current school attendance among orphans and non-orphans (10–14 years old, primary school age, secondary school age)

What it measures

It measures progress towards preventing relative disadvantage in school attendance among orphans versus non-orphans.

The indicator is split up in two parts so comparisons can be made between orphans and non-orphans:

Part A: current school attendance rate of orphans aged 10-14 primary school age, secondary school age.

Part B: current school attendance rate of children aged 10–14 primary school age, secondary school age both of whose parents are alive and who live with at least one parent.

Rationale

AIDS deaths in adults occur just at the time in their lives when they are forming families and bringing up children. Orphanhood is frequently accompanied by prejudice and increased poverty, factors that can jeopardize children's chances of completing school education and may lead to the adoption of survival strategies that increase vulnerability to HIV. It is important therefore to monitor the extent to which AIDS support programmes succeed in securing the educational opportunities of orphaned children.

Numerator: Part A: Number of children who have lost both parents and who attend school

aged 10-14, primary school age, secondary school age

Part B: Number of children both of whose parents are alive, who are living with at least one parent and who attend school aged 10-14, primary school age,

secondary school age

Denominator: Part A: Number of children who have lost both parents

Part B: Number of children both of whose parents are alive who are living with at

least one parent

Calculation: For both part A and B: Numerator / Denominator

Method of measurement:

Population-based survey (Demographic and Health Survey, AIDS Indicator Survey, Multiple Indicator Cluster Survey or other representative survey)

For every child aged 10-14, of primary school age, and secondary school age,

living in a household, a household member is asked:

1. Is this child's natural mother still alive? If yes, does she live in the household?

2. Is this child's natural father still alive? If yes, does he live in the household?

3. Did this child attend school at any time during the school year?

Measurement frequency:

Preferred: every two years Minimum: every 4–5 years

Disaggregation: Sex

Explanation of numerator

The definition of primary school age and secondary school age should be consistent with the UNESCO definition and as currently used for calculating other education-specific indicators such as net primary school enrolment/attendance rate and net secondary school enrolment/attendance rate for each country. The primary school age and secondary school age populations may vary slightly from country to country. Therefore this indicator uses the terms 'primary school age' and 'secondary school age' as currently applied in standard international measurements including in major survey programmes such as DHS or MICS to allow each country to apply its own national age ranges for primary and secondary school. The important point is to compare current school attendance of orphans and non-orphans across primary school and secondary school rather than by specific ages.

Strengths and weaknesses

The definitions of orphan/non-orphan used here—i.e., child aged 10–14 years as of the last birthday both of whose parents have died/are still alive—are chosen so that the maximum effect of disadvantage resulting from orphanhood can be identified and tracked over time. The age-range 10–14 years is used because younger orphans are more likely to have lost their parents recently so any detrimental effect on their education will have had little time to materialize. However, orphaned children are typically older than non-orphaned children (because the parents of younger children have often been HIV-infected for less time) and older children are more likely to have left school.

Typically, the data used to measure this indicator are taken from household-based surveys. Children not recorded in such surveys—e.g., those living in institutions or on the street—generally, are more disadvantaged and are more likely to be orphans. Thus, the indicator will tend to understate the relative disadvantage in educational attendance experienced by orphaned children.

This indicator does not distinguish children who lost their parents due to AIDS from those whose parents died of other causes. In countries with smaller epidemics or in the early stages of epidemics, most orphans will have lost their parents due to non-HIV-related causes. Any differences in the treatment of orphans according to the known or suspected cause of death of their parents could influence trends in the indicator. However, to date there is little evidence that such differences in treatment are common.

The indicator provides no information on actual numbers of orphaned children. The restrictions to double orphans and to 10–14 year-olds mean that estimates may be based on small numbers in countries with small or nascent epidemics.

Further information

For further information, please consult the following website:

http://www.unicef.org/aids/index_documents.html

10.2 External economic support to the poorest households

Proportion of the poorest households who received external economic support in the last 3 months

What it measures

It measures progress in providing external economic support to poorest households affected by HIV and AIDS.

Rationale

Economic support (with a focus on social assistance and livelihoods assistance) to poor and HIV-affected households remains a high priority in many comprehensive care and support programmes. This indicator reflects the growing international commitment to HIV-sensitive social protection. It recognizes that the household should be the primary unit of analysis since many of the care and support services are directed to the household level. Tracking coverage of households with orphans and within the poorest quintile remains a developmental priority.

Numerator: Number of the poorest households that received any form of external economic

support in the last 3 months

External economic support is defined as free economic help (cash grants, assistance for school fees, material support for education, income generation support in cash or kind, food assistance provided at the household level, or material or financial support for shelter) that comes from a source other than friends, family or neighbours unless they are working for a community-based group or organization. This source is most likely to be the national government or

a civil society organization

Denominator: Total number of poorest households

Poorest households are defined as a household in the bottom wealth quintile. Countries should use the exact indicator definition and method of measurement for standardized progress monitoring and reporting at national and global levels. This will allow monitoring of changes over time and comparisons across different countries. However, countries can add or exclude other categories locally (for example, other wealth quintiles) depending on the country needs with respect to

national programme planning and implementation

Calculation: Numerator / Denominator

Method of Population-based surveys such as Demographic and Health Survey, AIDS measurement: Indicator Survey, Multiple Indicator Cluster Survey or other nationally representative survey

An assessment of the household's wealth (through an assessment of asset ownership) is completed at the data analysis stage using the wealth quintile to identify the poorest 20% of households. However, since it is not possible to identify the poorest households at the time of data collection, questions on economic support should be asked to all households. Only those who fall in the

lowest wealth quintile will be included in the indicator

As part of a household survey, a household roster should be used to list all members of the household together with their ages, and identify all households with children less than 18 years of age, and with orphans, in the last year before the survey. Questions are then asked for each such household about the types of economic support received in the last 3 months, and the primary source of the help

The household heads or respondents are asked the following questions about the type of external economic support they have received in the last 3 months. Has your household received any of the following forms of external economic support in the last 3 months:

- a) Cash transfer (e.g., pensions, disability grant, child grant, to be adapted according to country context)
- b) Assistance for school fees
- c) Material support for education (e.g., uniforms, school books etc)
- d) Income generation support in cash or kind e.g. agricultural inputs
- e) Food assistance provided at the household or external institution (e.g., at school)
- f) Material or financial support for shelter
- g) Other form of economic support (specify)

An assessment of the household's wealth (through an assessment of asset ownership) is completed at the data analysis stage using the wealth quintile at which point it will possible to assess the extent to which the poorest households are receiving external support

Measurement frequency:

Every 4–5 years

Disaggregation:

It is recommended that the indicator is disaggregated by type of external economic support in order to track the different types of economic support provided – particularly to be able to distinguish between access to free social assistance such as cash transfers (often specifically for poor labour-constrained households) and livelihoods support, which is often targeted at poor households which are less labour-constrained. It is also recommended that the indicator is disaggregated by whether or not households have orphans as orphaning remains a major determinant of vulnerability, particularly in relation to access to services. Where possible, data should also be disaggregated by rural versus urban residence. For countries which opt to add data collection on households in other wealth quintiles in addition to those in the bottom quintile, the indicator can also be compared with other wealth quintiles to track whether external economic support is reaching the bottom quintile compared to wealthier quintiles

Strengths and weaknesses

This indicator reflects new evidence of the need for a greater focus on wealth dimensions of vulnerability and the fact that that targeting on the basis of extreme poverty in high prevalence contexts ensures good coverage of poor households affected by HIV. Proxy indicators of AIDS affectedness (such as "chronic illness") have often been poorly associated with HIV, have weak associations with adverse developmental outcomes, and have proven difficult to define in household questionnaires.

This indicator demonstrates changing levels of economic support for the poorest households. In high prevalence contexts, in particular, the majority are likely to be HIV affected. The indicator also demonstrates changes in the composition of external support (e.g. cash, food, livelihoods) received by poor households.

The indicator does not measure directly economic support to HIV infected and affected households, which is difficult to establish during a survey, but implicitly suggests that households living in the bottom wealth quintile in high prevalence contexts are more likely to be negatively impacted by HIV and AIDS and in need of economic assistance. In order to keep measurement as simple as possible, the indicator does not attempt to identify the different sources of support to households but this should be partly captured in National AIDS Spending Assessments (NASA).

The collection of data through population-based surveys, particularly DHS and MICS, means that the indicator does not capture the status of people living outside of households such as street children, children in institutions and internally displaced populations. Separate surveys are needed to track coverage for such vulnerable populations.

Further information

For further information, please consult the following website:

http://www.unicef.org/aids/index_documents.html

Government HIV and AIDS policies

National Commitments and Policy Instrument (NCPI)

What it measures

It measures progress in the development and implementation of national-level HIV and AIDS policies, strategies and laws.

Rationale

This indicator tracks progress made in implementing the laws, regulations and policies necessary for an effective response to HIV.

Method of measurement:

National Commitments and Policy Instrument (see Appendix 3)

The NCPI questionnaire is divided in two parts which cover the following areas:

Part A to be administered to government officials.

Part A covers:

I. Strategic plan

II. Political support and leadership

III. Human rights

IV. Prevention

V. Treatment, care and support

VI. Monitoring and evaluation

Part B to be administered to representatives from civil society organizations, bilateral agencies, and UN organizations.

Part B covers:

I. Civil society involvement²

II. Political support and leadership

III. Human rights

IV. Prevention

V. Treatment, care and support

Some questions occur in both Part A and Part B to ensure that the views of both the government and non-government respondents, whether in agreement or not, are obtained.

Measurement frequency:

Every two years. The NCPI is ideally completed in the last 3 months before submission (i.e. between January 2014 and March 2014 for the 2014 reporting round). As a variety of stakeholders need to be consulted, it is important to allow adequate time for the data gathering and data consolidation process.

Methods

Each section should be completed by (a) conducting a desk review of relevant documents and (b) interviewing key people most knowledgeable about the topic. It is important to submit a fully completed National Commitments and Policy Instrument (NCPI): check the relevant standardized responses as well as providing further information in the open text boxes where requested. This will facilitate a better understanding of the current country situation, provide examples of good practice for others to learn from, and pin-point some issues for further improvement. NCPI responses reflect the overall policy, strategy, legal and programme implementation environment of the HIV response. The open text boxes provide an opportunity to comment on issues that are perceived as important but insufficiently captured in the questions as asked e.g. important subnational variations; the level of implementation of strategies, policies, laws or regulations; explanatory notes; comments on the data sources etc. In general, draft strategies, policies, or laws are not considered 'in existence' (i.e., there is no opportunity yet to expect their influence on programme implementation) so questions about whether such a document exists should be answered with 'no'. It would, however, be useful to state that such documents are in draft form in the relevant open text box.

While the responsibility for submitting the consolidated NCPI data lies with the national government, the assistance of technical coordinators for data gathering, data consolidation and data validation is strongly advised. Accurate completion of the NCPI requires the involvement of a range of stakeholders including representatives of civil society organizations. It is strongly recommended (a) to organize an initial workshop with key stakeholders to agree on the NCPI data gathering process (including relevant documents for desk review, organizational representatives to be interviewed and process to be used for determining final responses) and the timeline; (b) to organize a final workshop with key stakeholders to present, discuss and validate the NCPI findings before official submission as part of the Global AIDS Progress Report. Agreement on the final NCPI data does not require that any discrepancies between overlapping questions in Part A and Part B be reconciled; it simply means that when there are different perspectives Part A respondents agree on their responses, Part B respondents agree on their responses, and both are submitted.

If not already the case, it is useful to collate all key documents (i.e. policies, strategies, laws, guidelines, reports, etc.) related to the HIV response in one place (such as a website), allowing easy access by all stakeholders. This will not only facilitate validation of NCPI responses but, even more importantly, increase awareness about and encourage use of these important documents in the implementation of the national HIV response going forward.

Strengths and weaknesses

The NCPI is the most comprehensive standardized questionnaire available to assess the policy, strategy, legal and programme implementation environment for the HIV response. Although the NCPI is generally referred to as an 'indicator' it is not used in that sense. The importance of the NCPI lies in the process of data collection and data reconciliation between different stakeholders, detailed analysis of the responses, and its use in strengthening the national HIV response. The NCPI process provides a unique opportunity for the variety of stakeholders to take stock of progress made and to discuss what still needs to be done to support an effective and efficient HIV response. When completed in a truly collaborative manner, inviting appropriate representation and respecting different views, the NCPI process can play an important role in strengthening in-country collaboration and increasing shared ownership of the HIV response.

It is important to analyse the data for each of the NCPI sections and include a write-up in the narrative section of the Country Progress Report in terms of progress made in (a) policy, strategy and law development and (b) implementation of these in support of the country's HIV response. Comments on the agreements or discrepancies between overlapping questions in Parts A and B should also be included, as well as a trend analysis on the key NCPI data from previous rounds of reporting, where available.

APPENDICES

Appendix 1. Country Progress Report template

Appendix 2. National Funding Matrix 2014

Appendix 3. National Commitments and Policy Instrument (NCPI)

Appendix 4. Sample checklist for Country Progress Report

Appendix 5. Selected bibliography

Appendix 6. Male circumcision indicators

Appendix 7. Geographic data collection in Surveillance, Monitoring and Evaluation

Appendix 1. Country Progress Report template

The following provides the full template of the narrative part of the Country Progress Report and detailed instructions for completion of the different sections included in it. It is highly recommended that the indicator data are submitted through the recommended online reporting tool.

COUNTRY PROGRESS REPORT [Country Name]

Submission date: fill in the date of the formal submission of the country report to UNAIDS.

Table of Contents

I. Status at a glance

Instructions: this section should provide the reader with a brief summary of

- (a) the inclusiveness of the stakeholders in the report writing process;
- (b) the status of the epidemic;
- (c) the policy and programmatic response;
- (d) Indicator data in an overview table.

II. Overview of the AIDS epidemic

[Instructions: This section should cover the detailed status of the HIV prevalence in the country during the period January 2012–December 2013 based on sentinel surveillance and specific studies. The source of information for all data provided should be included.

III. National response to the AIDS epidemic

Instructions: This section should reflect the change made in national commitment and programme implementation broken down by prevention, care, treatment and support; knowledge and behaviour change; and impact alleviation during the period January 2012–December 2013.

Countries should specifically address the linkages between the existing policy environment, implementation of HIV programmes, verifiable behaviour change and HIV prevalence as supported by the indicator data. Where relevant, these data should also be presented and analysed by sex and age groups. Countries should also use data from the National Commitments and Policy Instrument (NCPI) (see Appendix 3) to describe progress made in policy/strategy development and implementation, and include a trend analysis on the key NCPI data from previous rounds, where available. Countries are encouraged to report on additional data to support their analysis and interpretation of the reported data.

IV. Best practices

Instructions: This section should cover detailed examples of what is considered a best practice in-country in one or more of the key areas (such as political leadership; a supportive policy environment; scale-up of effective prevention programmes; scale-up of care, treatment and/or support programmes; monitoring and evaluation, capacity-building; infrastructure development. The purpose of this section is to share lessons learned with other countries.

V. Major challenges and remedial actions

Instructions: This section should focus on:

(a) progress made on key challenges reported in the 2012 Country Progress Report or during the Mid-Term Reviews of the Political Declaration many countries performed in 2013, if any;

- (b) challenges faced throughout the reporting period (2012–2013) that hindered the national response, in general, and the progress towards achieving targets, in particular;
- (c) concrete remedial actions that are planned to ensure achievement of agreed targets.

VI. Support from the country's development partners (if applicable)

Instructions: This section should focus on (a) key support received from, and (b) actions that need to be taken by development partners to ensure achievement of targets.

VII. Monitoring and evaluation environment

Instructions: This section should provide (a) an overview of the current monitoring and evaluation (M&E) system; (b) challenges faced in the implementation of a comprehensive M&E system; and (c) remedial actions planned to overcome the challenges, and (d) highlight, where relevant, the need for M&E technical assistance and capacity-building. Countries should base this section on the National Commitments and Policy Instrument (NCPI) (see Appendix 3).

ANNEXES

ANNEX 1: Consultation/preparation process for the country report on monitoring the progress towards the implementation of the Declaration of Commitment on HIV and AIDS

ANNEX 2: National Commitments and Policy Instrument (NCPI)

Please submit your complete Global AIDS Progress Report before 31 March 2014 using the recommended reporting tool.

Please direct all enquiries related to Global AIDS Reporting to the UNAIDS Secretariat at: AIDSreporting@unaids.org

Appendix 2. National Funding Matrix – 2014

Cover Sheet Please provide the following information when submitting the completed National Funding Matrix. Country: Contact Person at the National AIDS Authority/Committee (or equivalent): Title: Contact Information for the National AIDS Authority/Committee (or equivalent): Email: Telephone: _____ Fax: ____ Reporting Cycle 2011: calendar year _____ or fiscal year ____ Reporting Cycle 2012: calendar year _____ or fiscal year ____ Reporting Cycle 2013: calendar year _____ or fiscal year ____ For a fiscal year reporting cycle, please provide the start and end month/year: ____/ ___ to ____/ ___ Local Currency: Average exchange rate with US dollars during the reporting cycle: 2011: _____ / 2012: _____ / 2013: _____ Methodology: (Please confirm which methodology - National AIDS Spending Assessments, National Health Accounts or Resource Flows Surveys - supplied the data for the National Funding Matrix. In addition, please provide information on how and where to access the full report from whichever methodology was used to collect the data.) **Unaccounted Expenditures:** (Please specify if there were expenditures for activities in any of the AIDS Spending Categories or sub-categories that are not included in the National Funding Matrix and explain why these expenditures were not included.) Countries that have submitted data for 2011 and/or 2012 already in their last report (GARPR 2013) do not need to fill in data for these years again, unless data in the last report were missing, incomplete or there have been changes to the data as more information has become available since. In this case, we will replace the data in our database reported for these years in the last submission (GARPR 2013) with the newer data reported in the 2014 submission. 2011: 2012: Budget Support: Is budget support from an international source (e.g. a bilateral donor) included under the

2011: _____ Yes _____ No / 2012: _____ Yes ____ No / 2013: _____ Yes ____ No

Central/National and/or Subnational sub-categories under Public Sources of financing?

AIDS spending National Funding Matrix – 2014

Background

The AIDS Spending indicator is used to measure target # 6 of the 2011 UN Political Declaration on HIV and AIDS: "Reach a significant level of annual global expenditure (between \$22 billion and \$24 billion) in low and middle-income countries". AIDS Spending is reported completing the National Funding Matrix: AIDS Spending by category and by financing Source. The matrix is a spreadsheet that enables countries to record AIDS spending within eight categories across three funding sources. This indicator provides critical information that is valuable at both national and global levels of the AIDS response. The National Funding Matrix has been designed to be compatible with different data collection and tracking systems, i.e. National AIDS Spending Assessments (NASA), System of Health Accounts (SHA), National Health Accounts (NHA) and Resource Flows Surveys, so as to transfer information from these tools to the matrix. For countries using the NASA, the matrix is one of the outputs of this tool. (Countries interested in implementing the NASA are encouraged to contact UNAIDS for additional information on this tool.)

Structure of the matrix

The National Funding Matrix has two basic components:

- AIDS Spending Categories (How funds allocated to the national response are spent)
- Financing Sources (Where funds allocated to the national response are obtained)

There are eight AIDS Spending Categories: Prevention; Care and Treatment; Orphans and Vulnerable Children; Systems Strengthening and Programme Management; Incentives for Human Resources; Social Protection and Social Services (excluding Orphans and Vulnerable Children); Enabling Environment and Community Development; and Research.

Each spending category includes multiple sub-categories. Across the eight spending categories there are a total of 91 sub-categories. It is important to note that all of the spending categories and sub-categories are AIDS-specific; for example, expenditures listed under Enabling Environment and Community Development should only be those that are directly attributable to the AIDS response.

Prevention is the largest category with 22 sub-categories, ranging from voluntary counselling and testing to condom social marketing to blood safety; seven of the remaining eight spending categories have fewer than 10 sub-categories each. The purpose of the categories and sub-categories is to help national governments break out their spending as rationally and consistently as possible. As mentioned above, the matrix was designed to be compatible with common data collection and tracking systems in order to reduce the burden of reporting on national governments.

There are three major groups of Financing Sources: Domestic Public; International and Domestic Private (optional for the Global AIDS Progress reporting).

Similar to the spending categories, each financing source has multiple sub-categories. Public Sources has four sub-categories: Central/National, Subnational, Development Bank Reimbursable (loans) and All Other Public. International Sources has six subcategories: PEPFAR, Other Bilaterals, UN Agencies, Global Fund, Development Bank Grants (Non-reimbursable) and All Other International. Private Sources has two sub-categories: Corporations and Consumer/Out-of-Pocket. (Note: The data on Private Sources are optional for the Global AIDS Response Progress Reporting. However, countries are strongly encouraged to collect and report available data in this area.)

Instructions

■ The National AIDS Authority/Committee or equivalent should designate a technical coordinator to manage the collection and input of relevant data for the National Funding Matrix. It is recommended that this coordinator have good knowledge of tools and methodologies currently in use in the country for collecting this type of financial data (i.e. NASA, SHA/NHA, Resource Flows Survey). Also, it is encouraged that the coordinator contact other national resource tracking point persons,

- such as those in the Ministry of Health, who have been involved in reporting expenditures for HIV. The purpose of their involvement is to engender agreement on the national estimate for HIV expenditures and to avoid duplicate initiatives.
- Countries are requested to include as much detail in the National Funding Matrix as possible, including breakdowns by all applicable AIDS Spending and Funding Source Categories and sub-categories. Any categories or sub-categories that are not applicable in a country should be clearly identified; explanations for categories or sub-categories that do not include estimates for any other reason should be provided as part of the cover sheet to the matrix.
- The financial data in the matrix must be actual expenditures. They should not include budget figures that have not been validated as actual expenditures nor should the data reflect commitment or obligation figures. The actual expenditures must correspond to the calendar or fiscal years(s) of 2011, 2012 and/or 2013 (as available).
- The total for each line item should include funding from all sources listed for that item. In addition, there should be a sub-total for each of the eight AIDS Spending Categories, which captures all funding from all sources for all sub-categories in a given category.
- Amounts in each category or sub-category can be reported in local currency or in USD. If reported in
 national currency, the average exchange rate to US dollars for the calendar or fiscal year being reported
 needs to be stated on the cover sheet; (see the National Funding Matrix cover sheet on page 118).
- Spending categories and sub-categories are designed to be self-explanatory. Expenditures that do not clearly fit a specific sub-category should be listed in the Other/Not Classified Elsewhere sub-category that appears in each of the eight AIDS Spending Categories. (Detailed descriptions of the categories and sub-categories are available in the UNAIDS-published *National AIDS Spending Assessment (NASA): Classification taxonomy and Definitions*. See reference below.)
- Expenditures should only be counted in a single category or sub-category; they should never be double counted. For example, expenditures on activities for Orphans and Vulnerable Children should not be listed again under Social Protection and Social Services.
- Financing Sources categories and sub-categories are designed to be self-explanatory. Expenditures that do not clearly fit a specific sub-category should be listed in the "All Other" sub-category that appears under both Public and International Sources. Please note that the list of Financing Sources categories and sub-categories is not exhaustive; however, it is indicative of the main sources of financing.
- Financing in the Central/National and Subnational sub-categories under Public Sources should only include revenue generated by the government and allocated to the AIDS response. It should not include development assistance of any type from international sources; the only possible exception would be budget support from donor organizations that cannot be differentiated from domestic revenues. If the total amount of budget support can be identified, it should appear under the proper International Sources sub-category (e.g. "Other Bilaterals"). If any budget support is included in the Central/National and/or Subnational sub-categories, please indicate this fact on the cover sheet (see above).
- Financing provided by a Development Bank should be designated either as Reimbursable (e.g. loans), which appears under Public Sources, or Non-reimbursable (e.g. grants), which appears under International Sources. Countries that receive both loans and grants from Development Banks should be careful to allocate these funds to the correct categories.
- Financing provided by individual bilateral donors does not need to be disaggregated by donor agency in the funding matrix, with the exception of PEPFAR.
- Financing provided by international foundations should be listed in the "All Other" sub-category in the International category. Funds received from domestic foundations should be listed in the "All Other" sub-category in the "Public" category.
- Providing information on financing from Private Sources is optional. However, countries are strongly encouraged to collect and report available data in this area in order to provide a more complete picture of the funds available for the AIDS response.

- Key Populations at higher risk: all programmes targeting populations at higher risk, including risk-reduction activities, outreach (including by peers), voluntary and confidential HIV counselling and testing, and prevention of sexual transmission of HIV (including condoms, prevention and treatment of STIs) and programmes on developing and acquiring skills to negotiate safer behaviour, behaviour change and sustained engagement to prevent HIV infection should be coded and cross-classified under the corresponding AIDS Spending Category: ASC.01.08 Prevention programmes for sex workers and their clients: ASC.01.09, Programmes for men who have sex with men (MSM) or ASC.01.10 Harm-reduction programmes for injecting drug users (IDUs).
- All programmes targeting other specific populations (e.g. indigenous groups, migrants/mobile populations, military, police and other uniformed services, etc.), including risk-reduction activities, outreach (including by peers), voluntary and confidential HIV counselling and testing, and prevention of sexual transmission of HIV (including condoms, prevention and treatment of STIs) and programmes on developing and acquiring skills to negotiate safer behaviour, behaviour change and sustained engagement to prevent HIV infection should be coded and cross-classified under the corresponding AIDS Spending Category: ASC.01.04 Risk-reduction programmes for vulnerable and accessible populations.
- Programmes targeting the General Populations: all programmes targeting the general population, including risk-reduction activities, outreach (including by peers), voluntary and confidential HIV counselling and testing, and prevention of sexual transmission of HIV (including condoms, prevention and treatment of STIs) and programmes on developing and acquiring skills to negotiate safer behaviour, behaviour change and sustained engagement to prevent HIV infection should be coded and cross-classified under the corresponding AIDS Spending Category: ASC.01.01 Communication for social and behaviour change, ASC.01.02 Community mobilization, ASC.01.03 HIV testing and counselling (HCT), ASC.01.12 Condom social marketing, ASC.01.13 Public and commercial sector male condom provision and ASC.01.14 Public and commercial sector female condom provision.
- Incentives for Human resources: These expenditures are aimed at ensuring the availability of human resources for the AIDS response. Incentives for human resources refer to training, retention, deployment, and rewarding of quality performance of health care workers and managers for work in the HIV field. They only aim therefore at including the additional incentives for this purpose. The direct cost associated with human resources is included in the costs of each of the other spending categories. For example, the human resources are accounted for within the unitary costs of prevention and treatment interventions—*ASC.01 Prevention* and *ASC.02 Care and treatment*—and, where it concerns human resources required outside the point of care delivery, they are included in the programme costs as well—ASC.04 (Programme Management). Thus, the salary of a doctor should be accounted on the programmatic intervention where this doctor directly intervenes. Only the additional monetary incentive for the doctor, to work in a specific geographical area or for working on HIV, is to be classified under *ASC.05 Incentives for Human Resources*. Incentives for human resources covers mainly nurses and doctors.
- The Private Sources column for Corporations should list funds spent in-country by companies in the various AIDS Spending Categories and sub-categories; the adjacent Consumer/Out-of-pocket column should list funds spent by individuals and/or families in the AIDS Spending Categories and sub-categories. (Note: It is likely that most entries in the Consumer/Out-of-pocket column will be in the Care and Treatment and selected Prevention categories and sub-categories.)
- If a country has a National Health Account or Systemic Health Account (SHA) with an AIDS sub-account, it should implement a NASA-NHA/SHA Crosswalk in order to fill the Funding Matrix with the NHA or SHA AIDS sub-account results. The document *Linking NASA and NHA/SHA Concepts and Mechanics* is a comprehensive guide that shows how to crosswalk the spending categories from NHA/SHA-AIDS sub-account to the National Funding Matrix. Countries can contact the Strategic Information Adviser in their UNAIDS country office UNAIDS Investment and Efficiency Advisers in the region or the Strategic Information and Monitoring Division at UNAIDS headquarters in Geneva.
- If a country is working from a Resource Flows Survey, it may be able to correlate information from sub-totals in the survey to the eight AIDS Spending Categories in the National Funding Matrix.

- Electronic versions of the Notebook to Produce National AIDS Spending Assessment and the National AIDS Spending Assessment (NASA): Classification taxonomy and Definitions may be downloaded from the following page on the UNAIDS website: http://www.unaids.org/en/dataanalysis/tools/nasapublications/.
 An electronic version of the National Funding Matrix may be downloaded as an Excel file from the same website.
- The UNAIDS Secretariat strongly recommends the NAC or equivalent organize a one-day workshop of relevant stakeholders to review the National Funding Matrix before it is submitted as part of the Global AIDS Progress reporting process. Relevant stakeholders should include federal and provincial/regional/state government ministries and departments, local and international civil society organizations, multilateral agencies, bilateral donors, foundations and commercial sector entities, as well as representatives from other relevant resource tracking initiatives.

The National Funding Matrix is available on the Global AIDS Progress reporting tool (http://AIDSreportingtool.unaids.org).

Once the National Funding Matrix is filled, it has to be submitted through the Global AIDS Progress online reporting tool.

If you do not have access to the Global AIDS Progress reporting tool, please submit the National Funding Matrix by email to UNAIDS (AIDSreporting@unaids.org).

A DNS Grandling Colonicies	TOTAL (Local Currency) Public	Sub-Total Central / National	IsnoitsN -du2	Dev. Banks Reimbursable (e.g. Loans)	Social Security	All Other Public	lstorietnatini lstoT-du2	Silatetals a Aggag	PEPFAR Other Bilaterals	Other Bilaterals Multilaterals	NN Agencies	Global Fund	Dev. Bank Non- Reimburseable (e.g. Grants)	All Other Multilateral	rəfil Oflik Innetrastional	Private latoT-du2 thorq-ro4	institutions / Corporations	spunj plodesnoH	All Other Private
TOTAL (Local Currency)																			
1. Prevention (sub-total)																			
1.01 Communication for social and behavioural change (BCC)									+									+	
1.03 Voluntary counseling and testing (VCT)	+		+						+			\perp			+		+	+	
1.04 Risk-reduction and prevention activites for vulnerable and accessible populations																			
1.05. Prevention - Youth in school																			
1.06 Prevention - Youth out-of-school																			
1.07 Prevention of HIV transmission aimed at people living with HIV																			
1.08 Prevention programmes for sex workers and their clients																			
1.09 Programmes for men who have sex with men									-										
1.10 Harm-reduction programmes for injecting drug users			1						+									+	
1.11 Prevention programmes in the workplace			1						+									+	
1.12 Condom social marketing			\parallel						+									\dashv	
1.13 Public and commercial sector male condom provision									\dashv										
1.14 Public and commercial sector female condom provision									-										
1.15 Microbicides																			
1.16 Prevention, diagnosis and treatment of sexually transmitted infections (STI)																			
1.17 Prevention of mother-to-child transmission																			
1.18 Male Circumcision																			
1.19 Blood safety																			
1.20 Safe medical injections																			
1.21 Universal precautions																			
1.22 Post-exposure prophylaxis																			
1.23 Pre-exposure prophylaxis (new category for GARPR 2014)																			
1.98 Prevention activities not disaggregated by intervention																			
1.99 Prevention activities not elsewhere classified																			
2. Care and Treatment (sub-total)																			
2.01 Outpatient care (sub-total)																			
2.01.01 Provider- initiated testing and counseling																			
2.01.02 Opportunistic infection (OI) outpatient prophylaxis and treatment																			
2.01.03 Antiretroviral therapy																			
2.01.04 Nutritional support associated to ARV therapy			4																
2.01.05 Specific HIV-related laboratory monitoring									+										
2.01.06 Dental programmes for PLHIV			+	1					+									1	
2.01.07 Psychological treatment and support services			+	1					+					Ī			1	+	
2.01.08 Outpatient palliative care									+									+	
2.01.09 Home-based care	+		+	1					+			1		Ī	1		1	+	T
2.01.10 Traditional medicine and informal care and treatment services			+	1					+					Ī	1		1	+	T
2.01.98 Outpatient care services not disaggregated by intervention			\dashv	1					\dashv									+	
2.01.99 Outpatient Care services not elsewhere classified																			
2.02 In-patient care (sub-total)																			
2.02.01 Inpatient treatment of opportunistic infections (OI)																			
2.02.02 Inpatient palliative care																			
2.02.98 Inpatient care services not disaggregated by intervention																			
2.02.99 In-patient services not elsewhere classified																			
2.03 Patient transport and emergency rescue																			
2.98 Care and treatment services not disaggregated by intervention			_	$\overline{\downarrow}$					+					1			+	+	\exists
2.99 Care and treatment services not-elsewhere classified			$\frac{1}{2}$	1					+								-	-	

3. Orphans and Vulnerable Children (sub-total)
3.01 OVC Education
3.02 OVC Basic health care
3.03 OVC Family/home support
3.04 OVC Community support
3.05 OVC Social services and Administrative costs
3.06 OVC Institutional Care
3.98 OVC services not disaggregated by intervention
3.99 OVC services not-elsewhere classified
4. Systems Weapthening & Programme Coordination (sub-total) [renamed from "Program Anysteners and add-thinistration"
4.01 National planning, coordination and programme management
4.02 Administration and transaction costs associated with managing and disbursing funds
4.03 Monitoring and evaluation
4.04 Operations research
4.05 Scrological-surveillance (Serosurveillance)
4.06 HIV drug-resistance surveillance
4.07 Drug supply systems
4.08 Information technology
4.09 Patient tracking
4.10 Upgrading and construction of infrastructure
4.11 Mandatory HIV testing (not VCT)
4.98 Program Management and Administration Strengthening not disaggregated by type
4.99 Program Management and Administration Strengthening not-elsewhere classified
5. Incentives for human resources (sub-total)
5.01 Monetary incentives for human resources
5.02 Formative education to build-up an HIV workforce
S.03 Tarining
5.98 Incentives for Human Resources not specified by kind
5.99 Incentives for Human Resources not elsewhere classified
6. Social Protection and Social Services (excluding OVC) (sub-total)
6.01 Social protection through monetary benefits
6.02 Social protection through in-kind benefits
6.03 Social protection through provision of social services
6.04 HJV-specific income generation projects
6:98 Social protection services and social services and services are services and services and services are services and services and services are services are services and services are servic
699 Social protection services and social services and services are service
7. Enabling Environment (sub-total)
7.01 Advocacy
7.02 Human rights programmes
7.03 AIDS-specific institutional development
7.04 AIDS-specific programmes focused on women
7.05 Programmes to reduce Gender Based Violence
7.98 Enabling Environment and Community Development not disaggregated by type
7.99 Enabling Environment and Community Development not elsewhere classified
8. Research excluding operations research which is included under (sub-total)
8.01 Biomedical research
8.02 Clinical research
8.03 Epidemiological research
8.04 Social science research
8.05 Vacion-related research
8.98 Research not disaggregated by type
8.99 Research not elsewhere classified

Appendix 3. National Commitments and Policy Instrument (NCPI) 2014

COUNTRY:

Name of the National AIDS Committee Officer in charge of NCPI submission and who can be contacted for questions, if any:

Postal address:
Tel:
Fax:
E-mail:
Date of submission:

Instructions

The following instrument measures progress in the development and implementation of national HIV policies, strategies and laws. It is an integral part of the core indicators and is to be completed and submitted as part of the 2014 Country Progress Report.

This is the sixth version of the NCPI and the second revised version since the tool changed the name to National Commitments and Policy Instrument (NCPI), instead of the earlier National Composite and Policy Index (NCPI). The NCPI has since 2012 been updated where necessary to reflect new HIV programmatic guidance and includes the questions regarding HIV integration that were included in the Special 2013 GARPR questionnaire (can be found in Part AI). The majority of questions are identical to the previous rounds of reporting to allow for trend analyses. Countries are strongly advised to conduct a trend analysis and include a description of progress made in (a) policy, strategy and law development and (b) implementation of these in support of the country's HIV response. Comments on the agreements or discrepancies between overlapping questions in Parts A and B should also be included as well as a trend analysis on the key NCPI data, where available¹.

I. STRUCTURE OF THE QUESTIONNAIRE

The NCPI is divided into two parts.

Part A to be administered to government officials.

Part A covers:

- I. Strategic plan
- II. Political support and leadership
- III. Human Rights
- IV. Prevention
- V. Treatment, care and support
- VI. Monitoring and evaluation

¹ Compare NCPI in Guidelines on construction of core indicators from previous reporting rounds, for selecting questions for which trends can be analysed.

Part B to be administered to representatives from civil society organizations, bilateral agencies, and UN organizations.

Part B covers:

- I. Civil Society involvement
- II. Political support and leadership
- III. Human rights
- IV. Prevention
- V. Treatment, care and support

Some questions occur in both Part A and Part B to ensure that the views of both the national government and nongovernmental respondents, whether in agreement or not, are obtained.

For questions that pertain to key populations at higher risk for HIV (heretofore referred to as "key populations") and other vulnerable populations, the following definition is applied: Key populations are defined as most at risk for HIV within a defined epidemiological context, that have significantly higher levels of risk of acquiring and transmitting HIV, and with higher rates of mortality and/or morbidity; access or uptake of relevant services is often significantly lower than the rest of the population. Depending on the disease and the country context, some population groups may require explicit attention (for example, people who inject drugs, sex workers, men who have sex with men, transgender people and people living with HIV). Other populations that may be vulnerable to HIV are women and girls, clients of sex workers, prisoners, refugees, migrants or internally displaced populations, adolescents, and young people, vulnerable children and orphans, people with disabilities, ethnic minorities, people in low-income groups, people living in rural or geographically isolated settings or other group(s) specific to the country context.

It is important to submit a fully completed NCPI. Please check the relevant standardized responses as well as provide further information in the open text boxes where requested. This will facilitate a better understanding of the current country situation, provide examples of good practice for others to learn from, and pin-point some issues for further improvement. NCPI responses reflect the overall policy, strategy, legal and programme implementation environment of the HIV response. The open text boxes provide an opportunity to comment on anything that is perceived to be important but insufficiently captured by the standardized questions (e.g. important sub-national variations; the level of implementation of laws, policies or regulations; explanatory notes; comments on data sources etc). In general, draft strategies, policies, or laws are not considered 'in existence' (i.e. there is no opportunity yet to expect their influence on programme implementation) so questions about whether such a document exists should be answered with 'no'. It would, however, be useful to state that such documents are in draft form and any specifics about them in the relevant open text box.

The overall responsibility for collating and submitting the information requested in the NCPI lies with the national government, through officials from the National AIDS Committee (NAC) (or equivalent).

II. PROPOSED STEPS FOR DATA GATHERING AND DATA VALIDATION

The NCPI is ideally completed in the last 3 months before submission (i.e. between January 2014 and March 2014 for the 2014 reporting round). As a variety of stakeholders need to be consulted, it is important to allow adequate time for the data gathering and data consolidation process.

1. Designate two technical coordinators (one for part A; one for part B)

Technical coordinators should be given responsibility to undertake the desk review, to carry out interviews as needed, to bring together relevant stakeholders, and to facilitate collating and consolidating the NCPI data. Preferably, the technical coordinator for Part A is from the NAC (or equivalent) and for Part B is a person outside the government (mostly from civil society). They should ideally have understanding of the national policy and legal environment, a monitoring and evaluation background, and knowledge of the main actors in the national HIV response.

2. Agree with stakeholders on the NCPI data gathering and validation process

Accurate completion of the NCPI requires the involvement of a range of stakeholders including representatives of a variety of civil society organizations. It is strongly recommended to organize an initial workshop with key stakeholders to agree on the NCPI data-gathering process including relevant documents for desk review, organizational representatives to be interviewed, the process to be used for determining final responses, and the timeline.

3. Obtain data

The submitted NCPI data should represent the most recent stock-taking of the policy, strategic and legal environment. As the process involves a range of stakeholders and data need to be consolidated before official submission to UNAIDS, it is important to allow adequate time for completion.

Each section should include completion of the following tasks:

(i). Desk review of relevant documents.

If not already the case, it is useful to collate all key documents (i.e. policies, strategies, laws, guidelines, reports etc) related to the HIV response in one place which allows easy access by all stakeholders (such as a website). This will not only facilitate validation of NCPI responses but, even more importantly, increase awareness about and encourage use over time of these important documents in the implementation of the national HIV response.

- (ii). Interviewing (or other ways of obtaining the information efficiently) key people most knowledgeable about the specific topic including, but not restricted to the following:
 - For Strategic Plan and Political Support sections: the Director or Deputy Director of the National AIDS Programme or National AIDS Committee (or equivalent), the Heads of the AIDS Programme at provincial and at district levels (or equivalent decentralised levels).
 - For Monitoring and Evaluation section: Monitoring and Evaluation Officers of the National AIDS Committee (or equivalent), Ministry of Health, HIV focal points of other ministries, the national monitoring and evaluation technical working group.
 - For Human Rights questions: Ministry of Justice officials and human rights commissioners for questions in Part A; representatives of human rights and other civil society organizations/ networks, including representatives from networks of people living with HIV and from key populations and other vulnerable sub-populations, and legal aid centres/institutions working in the area of HIV for questions in Part B.
 - For Civil Society Participation section: key representatives of major civil society organizations working in the area of HIV. These specifically include representatives from networks of people living with HIV and from key populations and other vulnerable sub-populations.
 - For Prevention and Treatment, Care and Support sections: Ministries and major implementing agencies/organizations in those areas, including nongovernmental organizations and networks of people living with HIV.

Note that interviewees are requested to provide responses as representatives of their institutions or constituencies, not their own personal views.

4. Validate, analyse and interpret data

Once the NCPI is fully completed, the technical coordinators need to carefully review all responses to determine if additional consultations or review of more documents are needed.

It is important to analyse the data for each of the NCPI sections and include a write-up in the Country Progress Report in terms of progress made in policy/strategy development and implementation of programmes to tackle the country's HIV epidemic. Comments on the agreements/discrepancies between overlapping questions in Part A and Part B should also be included, as well as a trend analysis on the key NCPI data, where available.

It is strongly recommended to organize a final workshop with key stakeholders to present, discuss and validate the NCPI responses and the write-up of the findings before official submission. It is expected that representatives from civil society organizations working in the area of HIV are invited to participate. These specifically include representatives from networks of people living with HIV and from key populations and other vulnerable sub-populations. It is also important that persons with gender expertise and expertise with key populations be involved in the review and validation process. Ideally, the workshop would review the results from the last reporting round highlighting changes since that time and focus on validation of the NCPI data. Agreement on the final NCPI data does not require that discrepancies, if any, between overlapping questions in Part A and Part B be reconciled; it simply means that when there are different perspectives, that Part A respondents agree on their responses, Part B respondents agree on their responses, and that both are submitted. If there are no established mechanisms in place, the workshop can also provide an opportunity to discuss further collaboration between relevant stakeholders to address key gaps identified through the NCPI process.

5. Enter and submit data

Submit the final NCPI data before 31 March 2014, using the dedicated software provided on the Global AIDS Progress reporting website (www.unaids.org/AIDSReporting). If this is not possible, an electronic version of the completed questionnaire should be submitted as an appendix to the Country Progress Report before 15 March 2014 to allow time for the manual entry of data in Geneva.

National Commitments and Policy Instrument (NCPI) Data Gathering and Validation Process

Describe the process year for NCDI data gothering and validation.
Describe the process used for NCPI data gathering and validation:
Describe the process used for resolving disagreements, if any,
with respect to the responses to specific questions:
with respect to the responses to specific questions.
Highlight company of any miletal to the Construction of the same
Highlight concerns, if any, related to the final NCPI data submitted (such as data quality, potential misinterpretation of questions and the like):
(such as data quanty, potential infolict pretation of questions and the fixe).

NCPI Respondents

[Indicate information for **all** whose responses were compiled to fill out (parts of) the NCPI in the below table; add as many rows as needed]

NCPI - PART A [to be administered to government officials]

Organization	Names/Positions	Respondents to Part A [indicate which parts each respondent was queried on]				ed on]	
		A.I	A.II	A.III	A.IV	A.V	A.VI

Add details for all respondents.

NCPI - PART B

[to be administered to civil society organizations, bilateral agencies, and UN organizations]

Organization	Names/Positions	[indicat	Respondents to Part B [indicate which parts each respondent was queried on]			
		B.I	B.II	B.III	B.IV	B.V

Add details for all respondents.

National Commitments and Policy Instrument (NCPI)

Part A

[to be administered by government officials]

I. STRATEGIC PLAN		
1. Has the country developed a national multisectoral strategy	to respond to HIV?	
(Multisectoral strategies should include, but are not limited to, the ones listed under 1.2)	ose developed by Min	nistries such as the
	Yes	No
IF YES, what is the period covered [write in]:		
<i>IF YES</i> , briefly describe key developments/modifications betwee prior one.	en the current nationa	al strategy and the
<i>IF NO or NOT APPLICABLE</i> , briefly explain why.		
IEVES complete questions 1.1 through 1.10. IEVO as to security	n 2	
IF YES, complete questions 1.1 through 1.10; IF NO, go to question	11 2.	

1.1. Which government ministries or agencies have overall responsibility for the development and implementation of the national multi-sectoral strategy to respond to HIV?

Name of government ministries or agencies [write in]:

1.2. Which sectors are included in the multisectoral strategy with a specific HIV budget for their activities?

SECTORS	Included i	n Strategy	Earmarke	ed Budget
Education	Yes	No	Yes	No
Health	Yes	No	Yes	No
Labour	Yes	No	Yes	No
Military/Police	Yes	No	Yes	No
Social Welfare ²	Yes	No	Yes	No
Transportation	Yes	No	Yes	No
Women	Yes	No	Yes	No
Young People	Yes	No	Yes	No
Other [write in]:	Yes	No	Yes	No
	Yes	No	Yes	No

IF NO earmarked budget for some or all of the above sectors, explain what funding is used to ensure implementation of their HIV-specific activities?	

1.3. Does the multisectoral strategy address the following key populations/other vulnerable populations, settings and cross-cutting issues?

KEY POPULATIONS AND OTHER VULNERABLE POI	PULATIONS	
Discordant couples	Yes	No
Elderly persons	Yes	No
Men who have sex with men	Yes	No
Migrants/mobile populations	Yes	No
Orphans and other vulnerable children ³	Yes	No
People with disabilities	Yes	No
People who inject drugs	Yes	No
Sex workers	Yes	No
Transgender people	Yes	No
Women and girls	Yes	No
Young women/young men	Yes	No
Other specific vulnerable subpopulations ⁴	Yes	No
SETTINGS		

² This sector includes social protection

³ Orphans and other vulnerable children include children who have been abused, neglected or exploited, and children in homes/shelters/places of safety

⁴ Other specific vulnerable populations other than those listed above, that have been locally identified as being at higher risk of HIV infection (e.g. (in alphabetical order) bisexual people, clients of sex workers, indigenous people, internally displaced people, prisoners and refugees)

APPENDIX 3

Prisons	Yes	No
Schools	Yes	No
Workplace	Yes	No
CROSS-CUTTING ISSUES		
Addressing stigma and discrimination	Yes	No
Gender empowerment and/or gender equality	Yes	No
HIV and poverty	Yes	No
Human rights protection	Yes	No
Involvement of people living with HIV	Yes	No

IF NO, explain how key populations were identified?

$1.4. \quad \text{What are the identified key populations and vulnerable groups for HIV programmes in the country?}$

People living with HIV	Yes	No
Men who have sex with men	Yes	No
Migrants/mobile populations	Yes	No
Orphans and other vulnerable children	Yes	No
People with disabilities	Yes	No
People who inject drugs	Yes	No
Prison inmates	Yes	No
Sex workers	Yes	No
Transgender people	Yes	No
Women and girls	Yes	No
Young women/ young men	Yes	No
Other specific key populations/vulnerable subpopulations [write in]:	Yes	No

1.5. Does the country have a strategy for addressing HIV issues among its national uniformed services (such as military, police, peacekeepers, prison staff, etc)?

Yes No

1.6. Does the multisectoral strategy include an operational plan?

Yes No

	75 41 144 4 1		1	1 . 1 1
1.7.	Does the multisectoral	strategy or o	perational	blan include:

a) Formal programme goals?	Yes	No	N/A
b) Clear targets or milestones?	Yes	No	N/A
c) Detailed costs for each programmatic area?	Yes	No	N/A
d) An indication of funding sources to support programme implementation?	Yes	No	N/A
e) A monitoring and evaluation framework?	Yes	No	N/A

1.8.	Has the country ensured "full involvement and participation" of civil society ⁵ in	the
	development of the multisectoral strategy?	

Active	Moderate	No
involvement	involvement	involvement

IF NO or MODERATE INVOLVEMENT, briefly explain why this was the case:

Yes

No

1.9. Has the multisectoral strategy been endorsed by most external development partners

(bi-laterals, multi-laterals) ?

N/A

⁵ Civil society includes among others: networks and organisations of people living with HIV,women, young people, key affected groups (including men who have sex with men, transgender people, sex workers, people who inject drugs, migrants, refugees/displaced populations, prisoners); faith-based organizations; AIDS service organizations; community-based organizations; workers organizations, human rights organizations; etc. Note: The private sector is considered separately.

1.10. Have external development partners aligned and harmonized their HIV-related programmes to the national multisectoral strategy?

Yes, Yes, all partners some partners	No	N/A	
--------------------------------------	----	-----	--

SOME PARTNERS or NO, briefly explain for which areas there is no alignment/harmonization and why	IF

2.1. Has the country integrated HIV in the following specific development plans?

SPECIFIC DEVELOPMENT PLANS				
Common Country Assessment/UN Development Assistance Framework	Yes	No	N/A	
National Development Plan	Yes	No	N/A	
Poverty Reduction Strategy	Yes	No	N/A	
National Social Protection Strategic Plan	Yes	No	N/A	
Sector-wide approach	Yes	No	N/A	
Other [write in]:	Yes	No	N/A	
	Yes	No	N/A	

2.2. IF YES, are the following specific HIV-related areas included in one or more of the development plans?

HIV-RELATED AREA INCLUDED IN PLAN(S)			
Elimination of punitive laws	Yes	No	N/A
HIV impact alleviation (including palliative care for adults and children)	Yes	No	N/A
Reduction of gender inequalities as they relate to HIV prevention/treatment, care and/or support	Yes	No	N/A
Reduction of income inequalities as they relate to HIV prevention/ treatment, care and /or support	Yes	No	N/A
Reduction of stigma and discrimination	Yes	No	N/A
Treatment, care, and support (including social protection or other schemes)	Yes	No	N/A
Women's economic empowerment (e.g. access to credit, access to land, training)	Yes	No	N/A
Other [write in]:	Yes	No	N/A

3.	Has the country evaluated the impact of HIV on its socioeconomic development for planning
	purposes?

Yes	No	N/A
-----	----	-----

3.1. IF YES, on a scale of 0 to 5 (where 0 is "Low" and 5 is "High"), to what extent has the evaluation informed resource allocation decisions?

LOW		HIGH			
0	1	2	3	4	5

4. Does the country have a plan to strengthen health systems?

Yes?	
Please include information as to how this has impacted HIV-related infrastructure, human resources and capacities, and logistical systems to deliver medications?	

5. Are health facilities providing HIV services integrated with other health services?

Area	Many	Few	None
a) HIV counselling & testing with sexual & reproductive health			
b) HIV counselling & testing and tuberculosis			
c) HIV counselling & testing and general outpatient care			
d) HIV counselling & testing and chronic non-communicable diseases			
e) ART and tuberculosis			
f) ART and general outpatient care			
g) ART and chronic non-communicable diseases			
h) PMTCT with antenatal care/ maternal & child health			
i) Other comments on HIV integration:			

6. Overall, on a scale of 0 to 10 (where 0 is "Very Poor" and 10 is "Excellent"), how would you rate strategy planning efforts in your country's HIV programmes in 2013?

Very Poor									Excellent	
0	1	2	3	4	5	6	7	8	9	10

APPENDIX 3

Since 2011, what have been key achievements in this area:				
	L			
What challenges remain in this area:	7			

II. POLITICAL SUPPORT AND LEADERSHIP

Strong political support includes: government and political leaders who regularly speak out about HIV and AIDS and demonstrate leadership in different ways: allocation of national budgets to support HIV programmes; and, effective use of government and civil society organizations to support HIV programmes.

1.	Do the following high officials speak publicly and favoural forums at least twice a year?	oly about HIV efforts	s in major domestic
	A. Government ministers		
		Yes	No
	B. Other high officials at sub-national level		
		Yes	No
1.1	. In the last 12 months, have the head of government or othe demonstrated leadership in the response to HIV?	er high officials taker	n action that
	(For example, promised more resources to rectify identified we HIV as a human rights issue in a major domestic/internation an HIV clinic, etc.)		
		Yes	No
	riefly describe actions/examples of instances where the head of emonstrated leadership:	f government or othe	r high officials have
2.	Does the country have an officially recognized national mu (i.e., a National HIV Council or equivalent)?	ultisectoral HIV coo	rdination body
		Yes	No
II	FNO, briefly explain why not and how HIV programmes are b	eing managed:	

APPENDIX 3

2.1. IF YES:

Have terms of reference?	V	NT-
Have terms of reference?	Yes	No
Have active government leadership and participation?	Yes	No
Have an official chair person?	Yes	No
IF YES, what is his/her name and position title?		
Have a defined membership?	Yes	No
IF YES, how many members?		
Include civil society representatives?	Yes	No
IF YES, how many?		
Include people living with HIV?	Yes	No
IF YES, how many?		
Include the private sector?	Yes	No
Strengthen donor coordination to avoid parallel funding and duplication of effort in programming and reporting?	Yes	No

3. Does the country have a mechanism to promote interaction between government, civil society organizations, and the private sector for implementing HIV strategies/programmes?

Yes	No	N/A
-----	----	-----

IF YES, briefly describe the main achievements:					
What challenges remain in this area:					

			%
What kind of support does the National HIV Commiss organizations for the implementation of HIV-related ac			le to civil societ
Capacity-building		Yes	No
Coordination with other implementing partners		Yes	No
Information on priority needs		Yes	No
Procurement and distribution of medications or other suppl	ies	Yes	No
Technical guidance		Yes	No
Other [write in below]:		Yes	No
	determii	ne which, if any, a	re inconsistent
Has the country reviewed national policies and laws to with the National HIV Control policies?	determii		
	determi	Yes	No
with the National HIV Control policies?		Yes	No
. Has the country reviewed national policies and laws to with the National HIV Control policies? 1. IF YES, were policies and laws amended to be consistent.		Yes	No
with the National HIV Control policies?		Yes	No
with the National HIV Control policies?		Yes e National HIV (No Control policies
with the National HIV Control policies? 1. IF YES, were policies and laws amended to be consisten	t with th	Yes e National HIV (No Control policies
with the National HIV Control policies? 1. IF YES, were policies and laws amended to be consisten	t with th	Yes e National HIV (No Control policies
with the National HIV Control policies? 1. IF YES, were policies and laws amended to be consisten	t with th	Yes e National HIV (No Control policies
with the National HIV Control policies? 1. IF YES, were policies and laws amended to be consisten	t with th	Yes e National HIV (No Control policies
with the National HIV Control policies? 1. IF YES, were policies and laws amended to be consisten	t with th	Yes e National HIV (No Control policies
with the National HIV Control policies? 1. IF YES, were policies and laws amended to be consisten	t with th	Yes e National HIV (No Control policies
with the National HIV Control policies? 1. IF YES, were policies and laws amended to be consistent. IF YES, name and describe how the policies / laws were ame	nt with th	Yes e National HIV (Yes	No Control policies No
with the National HIV Control policies? 1. IF YES, were policies and laws amended to be consisten	nt with th	Yes e National HIV (Yes	No Control policies No
with the National HIV Control policies? 1. IF YES, were policies and laws amended to be consistent. IF YES, name and describe how the policies / laws were amended. Name and describe any inconsistencies that remain between	nt with th	Yes e National HIV (Yes	No Control policies No
with the National HIV Control policies? 1. IF YES, were policies and laws amended to be consistent. IF YES, name and describe how the policies / laws were amended. Name and describe any inconsistencies that remain between	nt with th	Yes e National HIV (Yes	No Control policies No

7. Overall, on a scale of 0 to 10 (where 0 is "Very Poor" and 10 is "Excellent"), how would you rate the political support for the HIV programme in 2013?

Very Poor										Excellent
0	1	2	3	4	5	6	7	8	9	10

Since 2011, what have been key achievements in this area:
What challenges remain in this area:

III. HUMAN RIGHTS

1.1. Does the country have non-discrimination laws or regulations which specify protections for specific key populations and other vulnerable groups? Circle yes if the policy specifies any of the following key populations and vulnerable groups:

KEY POPULATIONS AND VULNERABLE GROUPS		
People living with HIV	Yes	No
Men who have sex with men	Yes	No
Migrants/mobile populations	Yes	No
Orphans and other vulnerable children	Yes	No
People with disabilities	Yes	No
People who inject drugs	Yes	No
Prison inmates	Yes	No
Sex workers	Yes	No
Transgender people	Yes	No
Women and girls	Yes	No
Young women/young men	Yes	No
Other specific vulnerable subpopulations [write in]:	Yes	No

1.2. Does the country have a general (i.e., not specific to HIV-related discrimination) law on nondiscrimination?

103	No
aws:	
are implemented:	

Briefly comment on the degree to which they are currently implemented:

2.	Does the country have laws, regulations or policies that present obstacles ⁶ to effective HIV
	prevention, treatment, care and support for key populations and vulnerable groups?

Yes	No
-----	----

IF YES, for which key populations and vulnerable groups?		
People living with HIV	Yes	No
Elderly persons	Yes	No
Men who have sex with men	Yes	No
Migrants/mobile populations	Yes	No
Orphans and other vulnerable children	Yes	No
People with disabilities	Yes	No
People who inject drugs	Yes	No
Prison inmates	Yes	No
Sex workers	Yes	No
Transgender people	Yes	No
Women and girls	Yes	No
Young women/young men	Yes	No
Other specific vulnerable populations ⁷ [write in below]:	Yes	No

Briefly describe the content of these laws, regulations or policies:	
Briefly comment on how they pose barriers:	

⁶ These are not necessarily HIV-specific policies or laws. They include policies, laws or regulations which may deter people from or make it difficult for them to access prevention, treatment, care and support services. Examples cited in country reports in the past have include: "laws that criminalize same sex relationships", "laws that criminalize possession of condoms or drug paraphernalia"; "loitering laws"; "laws that preclude importation of generic medicines"; "policies that preclude distribution or possession of condoms in prisons"; "policies that preclude non-citizens from accessing ART"; "criminalization of HIV transmission and exposure", "inheritance laws/rights for women", "laws that prohibit provision of sexual and reproductive health information and services to young people", etc.

⁷ Other specific vulnerable populations other than above, may be defined as having been locally identified as being at higher risk of HIV infection (e.g. (in alphabetical order) bisexual people, clients of sex workers, indigenous people, internally displaced people, prisoners, and refugees)

IV. PREVENTION

1.	Does the country have a policy or strategy that promotes information, education and commu-
	nication (IEC) on HIV to the general population?

Yes No

Delay sexual debut	Yes	No
Engage in safe(r) sex	Yes	No
Fight against violence against women	Yes	No
Greater acceptance and involvement of people living with HIV	Yes	No
Greater involvement of men in reproductive health programmes	Yes	No
Know your HIV status	Yes	No
Males to get circumcised under medical supervision	Yes	No
Prevent mother-to-child transmission of HIV	Yes	No
Promote greater equality between men and women	Yes	No
Reduce the number of sexual partners	Yes	No
Use clean needles and syringes	Yes	No
Use condoms consistently	Yes	No
Other [write in below]:	Yes	No

1.2. In the last year, did the country implement an activity or programme to promote accountry implement and activity or programme to promote accountry in the programme activity in the programme activit	urate
reporting on HIV by the media?	

Yes	No

2. Does the country have a policy or strategy to promote life-skills based HIV education for young people?

Yes	No

2.1.

Is HIV education part of the curriculum in:							
Primary schools?	Yes	No					
Secondary schools?	Yes	No					
Teacher training?	Yes	No					

2.2. Do	es the strategy include						
a)	age-appropriate sexual	and reprod	uctive heal	th element.	s?		
					Yes		No
<i>b</i>)	gender-sensitive sexual	and reprod	uctive heal	th elements	s?		
					Yes		No
2.3. Do	es the country have an HI	IV educatio	n strategy i	for out-of-s	school young	g people?	
					Yes		No
	es the country have a polition and other preventive						
					Yes		No
Briefl	y describe the content of th	nis policy or	strategy:				
3.1. IF	YES, which populations ar	nd what eler	nents of HI	V preventi	on does the 1	policy/strate	egy address?
□ Che	ck which specific populat	ions and ele	ements are	included in	the policy/s	strategy	
		1018	3.403.49	Sex	Customers	Prison	Other

	IDU ⁸	MSM ⁹	Sex workers	Customers of Sex Workers	Prison inmates	Other populations ¹⁰ [write in]
Condom promotion						
Drug substitution therapy						
HIV testing and counseling						
Needle & syringe exchange						
Reproductive health, including sexually transmitted infections prevention and treatment						
Stigma and discrimination reduction						
Targeted information on risk reduction and HIV education						
Vulnerability reduction (e.g. income generation)						

 $⁸ ext{ IDU} = People who inject drugs$

⁹ MSM = men who have sex with men

¹⁰ Other vulnerable population other than those listed above, that have been locally identified as being at higher risk of HIV infection (e.g. (in alphabetical order) bisexual people, clients of sex workers, indigenous people, internally displaced people, prisoners, and refugees)

3.2.	Overall, on a scale of 0 to 10 (where 0 is "Very Po	oor"	and 10 is "	Excellent"),	how would y	you rate
	policy efforts in support of HIV prevention in 20	013?				

Very Poor										Excellent	
0	1	2	3	4	5	6	7	8	9	10	

Since 2011, what have been key achievements in this area:		
What challenges remain in this area:		
4. Has the country identified specific needs for HIV prevention	on programmes?	
, 1	on programmes.	
	Yes	No
		No
IF YES, how were these specific needs determined?		No
		No
IF YES, how were these specific needs determined?		No
IF YES, how were these specific needs determined?		No
IF YES, how were these specific needs determined?		No
IF YES, how were these specific needs determined?		No

4.1. To what extent has HIV prevention been implemented?

The majority of people in need have access to	Strongly disagree	Disagree	Agree	Strongly agree	N/A
Blood safety	1	2	3	4	N/A
Condom promotion	1	2	3	4	N/A
Economic support e.g. cash transfers	1	2	3	4	N/A
Harm reduction for people who inject drugs	1	2	3	4	N/A
HIV prevention for out-of-school young people	1	2	3	4	N/A
HIV prevention in the workplace	1	2	3	4	N/A
HIV testing and counseling	1	2	3	4	N/A
IEC11 on risk reduction	1	2	3	4	N/A
IEC on stigma and discrimination reduction	1	2	3	4	N/A
Prevention of mother-to-child transmission of HIV	1	2	3	4	N/A
Prevention for people living with HIV ¹²	1	2	3	4	N/A
Reproductive health services including sexually transmitted infections prevention and treatment	1	2	3	4	N/A
Risk reduction for intimate partners of key populations	1	2	3	4	N/A
Risk reduction for men who have sex with men	1	2	3	4	N/A
Risk reduction for sex workers	1	2	3	4	N/A
Reduction of Gender based violence	1	2	3	4	N/A
School-based HIV education for young people	1	2	3	4	N/A
Treatment as prevention	1	2	3	4	N/A
Universal precautions in health care settings	1	2	3	4	N/A
Other[write in]:	1	2	3	4	N/A

5. Overall, on a scale of 0 to 10 (where 0 is "Very Poor" and 10 is "Excellent"), how would you rate the efforts in implementation of HIV prevention programmes in 2013?

Very Poor										Excellent
0	1	2	3	4	5	6	7	8	9	10

¹¹ IEC = information, education, communication.

¹² Positive Prevention places PLHIV at the centre of managing their health and wellbeing. It recognises and emphasizes the leadership roles of PLHIV in responding holistically to HIV prevention and treatment needs.

V. TREATMENT, CARE AND SUPPORT

1. Has the country identified the essential elements of a comprehensive package of HIV treatment, care and support services?

|--|

<i>If YES</i> , Briefly identify the elements and what has been prioritized:						
Briefly identify how HIV treatment, care and support services are being scaled-up?						

1.1. To what extent have the following HIV treatment, care and support services been implemented?

The majority of people in need have access to	Strongly disagree	Disagree	Agree	Strongly agree	N/A
Antiretroviral therapy	1	2	3	4	N/A
ART for TB patients	1	2	3	4	N/A
Cotrimoxazole prophylaxis in people living with HIV	1	2	3	4	N/A
Early infant diagnosis	1	2	3	4	N/A
Economic support	1	2	3	4	N/A
Family based care and support	1	2	3	4	N/A
HIV care and support in the workplace (including alternative working arrangements)	1	2	3	4	N/A
HIV testing and counselling for people with TB	1	2	3	4	N/A
HIV treatment services in the workplace or treatment referral systems through the workplace	1	2	3	4	N/A
Nutritional care	1	2	3	4	N/A
Paediatric AIDS treatment	1	2	3	4	N/A
Palliative care for children and adults	1	2	3	4	N/A

APPENDIX 3

The majority of people in need have access to	Strongly disagree	Disagree	Agree	Strongly agree	N/A
Post-delivery ART provision to women	1	2	3	4	N/A
Post-exposure prophylaxis for non-occupational exposure (e.g., sexual assault)	1	2	3	4	N/A
Post-exposure prophylaxis for occupational exposures to HIV	1	2	3	4	N/A
Psychosocial support for people living with HIV and their families	1	2	3	4	N/A
Sexually transmitted infection management	1	2	3	4	N/A
TB infection control in HIV treatment and care facilities	1	2	3	4	N/A
TB preventive therapy for people living with HIV	1	2	3	4	N/A
TB screening for people living with HIV	1	2	3	4	N/A
Treatment of common HIV-related infections	1	2	3	4	N/A
Other[write in]:	1	2	3	4	N/A

2. Does the government have a policy or strategy in place to provide social and economic support to people infected/affected by HIV?

Yes No	Yes	No
--------	-----	----

Please clarify which social and economic support is provided ¹³ :							

3. Does the country have a policy or strategy for developing/using generic medications or parallel importing of medications for HIV?

Yes	No
140	1 10

¹³ Core types of economic assistance may include: (1) economic assistance (cash in exchange for labour, assistance for the elderly, disability grant, child grant); (2) educational support (assistance for school fees, material support related to schooling); (3) support for housing or shelter; (4) food assistance (food assistance at school or early learning centre/at home/in the community/at health facility, food vouchers, food in exchange for labour); (5) support for income generation in the form of agricultural inputs such as seeds, fertilizers or livestocks).

medica	.101131									
					Yes		No		1	N/A
F YES , for	· which co	mmodities	?							
		e of 0 to 10 implement								
ery Poor										Excell
0	1	2	3	4	5	6	7	8	9	10
nce 2011	, what hav	ve been key	achieven	ients in th	nis area:					
/hat chall	enges rem	nain in this	area:							

6. Does the		have a po	olicy or s	trategy t	o address	the need	ds of orp	hans and	l other v	ulnerable
					Yes		No	0		N/A
6.1. IF YES,	is there ar	ı operati	onal defi	nition fo	or orphan	s and vu	lnerable	children	in the co	ountry?
							Ye	s		No
6.2. IF YES,	does the co	ountry ha	ave a nati	onal acti	on plan s _j	pecificall	y for orp	hans and	vulneral	ole children?
							Ye	s		No
	, on a scale									
0	1	2	3	4	5	6	7	8	9	10
Since 2011	, what have	e been ke	y achieve	ements in	n this area	:				
What chal	enges rema	ain in thi	s area:							

VI. MONITORING AND EVALUATION

1.	Does the country	y have one national	Monitoring and	Evaluation (M&E) plan for HIV?
----	------------------	---------------------	----------------	---------------------	--------------------

		Yes	In Progress	No
Briefly describe any cha	allenges in developm	ent or implementation	on:	
	anongeo m de verop m		U111	
1.1. IF YES, years cover	ed [write in]:			
1.2. IF YES, have key pa	etnore aligned and	harmanizad thair M	[&E roquiromants (in	cluding indi
cators) with the nat		narmomzed their w	i&E requirements (in	cluding mai-
	Yes,	Yes,		
	all partners	some partners	No	N/A
			, 1	
Briefly describe what th	ne issues are:			

2. Does the national Monitoring and Evaluation plan include?

A data collection strategy	Yes	No
IF YES, does it address:		
Behavioural surveys	Yes	No
Evaluation / research studies	Yes	No
HIV Drug resistance surveillance	Yes	No
HIV surveillance	Yes	No
Routine programme monitoring	Yes	No

APPENDIX 3

A data analysis strategy	Yes	No
A data dissemination and use strategy	Yes	No
A well-defined standardised set of indicators that includes sex and age disaggregation (where appropriate)	Yes	No
Guidelines on tools for data collection	Yes	No

3. Is there a budget for implementation of the M&E plan?

Yes	In Progress	No
-----	-------------	----

3.1. IF YES, what percentage of the total HIV programme funding is budgeted for M&E activities?

%

4. Is there a functional national M&E Unit?

Yes	In Progress	No
-----	-------------	----

Briefly describe any obstacles:		

4.1. Where is the national M&E Unit based?

In the Ministry of Health?	Yes	No
In the National HIV Commission (or equivalent)?	Yes	No
Elsewhere [write in]?	Yes	No

4.2. How many	and what type of	professional staff a	re working in the na	ational M&E Unit?
---------------	------------------	----------------------	----------------------	-------------------

POSITION [write in position titles in spaces below]	Fullt	ime	Part tim	ne Since when
Permanent Staff [Add as many as needed]				
	Fullt	ime	Part tim	ne Since when
Temporary Staff [Add as many as needed]				
.3. Are there mechanisms in place to ensure that all ke the M&E Unit for inclusion in the national M&E sys				I
			Yes	No
Briefly describe the data-sharing mechanisms:				
Ziton, accorde une annu cinaring incommission				
What are the major challenges in this area:				

IF YES, briefly describe the national database and who manages it. 1. IF YES, does it include information about the content, key populations and geographical cover of HIV services, as well as their implementing organizations? Yes, all of the Yes, but only some of the No, none of the some of the No, none of the some of the No, none	. Is there a national M&E Committe activities?	e or W	orking Group that	meets regularly to	coordinate M&E
I.I. IF YES, does it include information about the content, key populations and geographical covor of HIV services, as well as their implementing organizations? Yes, all of the above Some of the above the	. Is there a central national database	with l	HIV- related data?		
At national level At national level At subnational level At what level(s)? [write in]				Yes	No
The services, as well as their implementing organizations? Yes, all of the above	IF YES, briefly describe the national da	tabase	and who manages	it.	
The services, as well as their implementing organizations? Yes, all of the above					
The services, as well as their implementing organizations? Yes, all of the above					
The services, as well as their implementing organizations? Yes, all of the above					
The services, as well as their implementing organizations? Yes, all of the above					
Yes, all of the above Yes, but only some of the above No, none of the above No, none of the above 1. Is there a functional Health Information System 14? At national level At subnational level Yes No If YES, at what level(s)? [write in] 1. Are there reliable estimates of current needs and of future needs of the number of adults and children requiring antiretroviral therapy?					ographical cover
2. Is there a functional Health Information System 14? At national level At subnational level Yes No If YES, at what level(s)? [write in] 1. Are there reliable estimates of current needs and of future needs of the number of adults and children requiring antiretroviral therapy?	of firv services, as well as their im	pieiliei T	nting organization		T
2. Is there a functional Health Information System ¹⁴ ? At national level At subnational level Yes No IF YES, at what level(s)? [write in] 1. Are there reliable estimates of current needs and of future needs of the number of adults and children requiring antiretroviral therapy?				some of the	No, none of the above
2. Is there a functional Health Information System ¹⁴ ? At national level At subnational level Yes No IF YES, at what level(s)? [write in] 1. Are there reliable estimates of current needs and of future needs of the number of adults and children requiring antiretroviral therapy?	IE VEC but only some of the shows wh	iah aa	a a ata da aa it in alud	-2	
At national level At subnational level Yes No IF YES, at what level(s)? [write in] 1. Are there reliable estimates of current needs and of future needs of the number of adults and children requiring antiretroviral therapy?	11. 123, but only some of the above, wh	iicii as _l	pects does it ilicidate		
At national level At subnational level Yes No IF YES, at what level(s)? [write in] 1. Are there reliable estimates of current needs and of future needs of the number of adults and children requiring antiretroviral therapy?					
At national level At subnational level Yes No IF YES, at what level(s)? [write in] 1. Are there reliable estimates of current needs and of future needs of the number of adults and children requiring antiretroviral therapy?					
At national level At subnational level Yes No IF YES, at what level(s)? [write in] 1. Are there reliable estimates of current needs and of future needs of the number of adults and children requiring antiretroviral therapy?					
At national level At subnational level Yes No IF YES, at what level(s)? [write in] 1. Are there reliable estimates of current needs and of future needs of the number of adults and children requiring antiretroviral therapy?					
At subnational level IF YES, at what level(s)? [write in] 1. Are there reliable estimates of current needs and of future needs of the number of adults and children requiring antiretroviral therapy?	2. Is there a functional Health Inform	nation	System ¹⁴ ?		
1. Are there reliable estimates of current needs and of future needs of the number of adults and children requiring antiretroviral therapy?	At national level			Yes	No
1. Are there reliable estimates of current needs and of future needs of the number of adults and children requiring antiretroviral therapy?	At subnational level			Yes	No
children requiring antiretroviral therapy?	IF YES, at what level(s)? [write in]				
children requiring antiretroviral therapy?					
Estimates Estimates				needs of the numbe	r of adults and
of Current and of Current No					

Future Needs

Needs Only

¹⁴ Such as regularly reporting data from health facilities which are aggregated at district level and sent to national level; data are analysed and used at different levels)?

Yes	No
Yes	No
Yes	No
Yes	No
	Yes

	Yes	No
. How are M&E data used?		
For programme improvement?	Yes	No
In developing / revising the national HIV response?	Yes	No
For resource allocation?	Yes	No
Other [write in]:		
Briefly provide specific examples of how M&E data are used, a	illu the main chancinges,	II airy.
0. In the last year, was training in M&E conducted		
O. In the last year, was training in M&E conducted At national level?	Yes	No
· · · · · · · · · · · · · · · · · · ·	Yes	No
At national level?	Yes	No No
At national level? IF YES, what was the number trained:		
At national level? IF YES, what was the number trained: At subnational level?		
At national level? IF YES, what was the number trained: At subnational level? IF YES, what was the number trained	Yes	No

11. Overall, on a scale of 0 to 10 (where 0 is "Very Poor" and 10 is "Excellent"), how would you rate the HIV-related monitoring and evaluation (M&E) in 2013?

Very Poor										Excellent
0	1	2	3	4	5	6	7	8	9	10

Since 2011, what have been key achievements in this area:
What challenges remain in this area:

National Commitments and Policy Instrument (NCPI)

Part B

[to be administered to representatives from civil society organizations, bilateral agencies, and UN organizations]

I. CIVIL SOCIETY¹⁵ INVOLVEMENT

1. To what extent (on a scale of 0 to 5 where 0 is "Low" and 5 is "High") has civil society contributed to strengthening the political commitment of top leaders and national strategy/policy formulations?

LOW					HIGH
0	1	2	3	4	5

Comments and examples:			

2. To what extent (on a scale of 0 to 5 where 0 is "Low" and 5 is "High") have civil society representatives been involved in the planning and budgeting process for the National Strategic Plan on HIV or for the most current activity plan (e.g. attending planning meetings and reviewing drafts)?

LOW					HIGH
0	1	2	3	4	5

Comments and examples:		

¹⁵ Civil society includes among others: networks and organisations of people living with HIV, women, young people, key affected groups (including men who have sex with men, transgender people, sex workers, people who inject drugs, migrants, refugees/displaced populations, prisoners); faith-based organizations; AIDS service organizations; community-based organizations; ; workers organizations, human rights organizations; etc. Note: The private sector is considered separately.

3. To what extent (on a scale of 0 to 5 where 0 is "Low" and 5 is "High") are the services provided by civil society in areas of HIV prevention, treatment, care and support included in:

a. The national HIV strategy?

LOW					HIGH
0	1	2	3	4	5

b. The national HIV budget?

LOW					HIGH
0	1	2	3	4	5

c. The national HIV reports?

LOW					HIGH
0	1	2	3	4	5

Comments and examples:		

4. To what extent (on a scale of 0 to 5 where 0 is "Low" and 5 is "High") is civil society included in the monitoring and evaluation (M&E) of the HIV response?

a. Developing the national M&E plan?

LOW					HIGH
0	1	2	3	4	5

b. Participating in the national M&E committee / working group responsible for coordination of M&E activities?

LOW					HIGH
0	1	2	3	4	5

c. Participate in using data for decision-making?

LOW					HIGH
0	1	2	3	4	5

APPENDIX 3

Comments and	examples:				
in HIV effort	s inclusive of div		s (e.g. organizati	ons and networ	iety representation ks of people living anizations)?
LOW		·		- -	HIGH
0	1	2	3	4	5
Comments and o	examples:				
To what exter	nt (on a scale of (0 to 5 where 0 is "I	Low" and 5 is "Hi	igh") is civil soci	iety able to access:
a. Adequat	te financial supp	ort to implement i	its HIV activities	?	
LOW					HIGH
0	1	2	3	4	5
b. Adequat	te technical supp	ort to implement	its HIV activities	?	
LOW					HIGH
0	1	2	3	4	5
Comments and o	examples:				

7. What percentage of the following HIV programmes/services is estimated to be provided by civil society?

Prevention for key-populations				
People living with HIV	<25%	25-50%	51-75%	>75%
Men who have sex with men	<25%	25-50%	51-75%	>75%
People who inject drugs	<25%	25-50%	51-75%	>75%
Sex workers	<25%	25-50%	51-75%	>75%
Transgender people	<25%	25-50%	51-75%	>75%
Palliative care	< 25%	25-50%	51-75%	> 75%
Testing and Counselling	<25%	25-50%	51-75%	>75%
Know your Rights/ Legal services	<25%	25-50%	51-75%	>75%
Reduction of Stigma and Discrimination	<25%	25-50%	51-75%	>75%
Clinical services (ART/OI)*	<25%	25-50%	51-75%	>75%
Home-based care	<25%	25-50%	51–75%	>75%
Programmes for OVC**	<25%	25-50%	51-75%	>75%

 $^{^*\}quad ART = Antiretroviral\ Therapy;\ OI = Opportunistic\ infections$

8. Overall, on a scale of 0 to 10 (where 0 is "Very Poor" and 10 is "Excellent"), how would you rate the efforts to increase civil society participation in 2013?

Very Poor										Excellent
0	1	2	3	4	5	6	7	8	9	10

Since 2011, what have been key achievements in this area:
What challenges remain in this area:

^{**} OVC = Orphans and other vulnerable children

II. POLITICAL SUPPORT AND LEADERSHIP

1.	Has the Government, through political and financial supp key populations and/or other vulnerable sub-populations programme implementation?		•
		Yes	No
I	FYES, describe some examples of when and how this has happ	pened:	

III. HUMAN RIGHTS

1.1 Does the country have non-discrimination laws or regulations which specify protections for specific key populations and other vulnerable subpopulations? Circle yes if the policy specifies any of the following key populations:

KEY POPULATIONS AND VULNERABLE SUBPOPULATI	ONS	
People living with HIV	Yes	No
Men who have sex with men	Yes	No
Migrants/mobile populations	Yes	No
Orphans and other vulnerable children	Yes	No
People with disabilities	Yes	No
People who inject drugs	Yes	No
Prison inmates	Yes	No
Sex workers	Yes	No
Transgender people	Yes	No
Women and girls	Yes	No
Young women/young men	Yes	No
Other specific vulnerable subpopulations [write in]:	Yes	No

1.2. Does the country hav	e a general (i.e.	, not specific to	HIV-related	discrimination)	law on non-
discrimination?					

Yes	No

<i>IF YES</i> to Question 1.1 or 1.2, briefly describe the contents of these laws:
Briefly explain what mechanisms are in place to ensure that these laws are implemented:

Briefly comment on the degree to which they are currently implemented:			

2. Does the country have laws, regulations or policies that present obstacles¹⁶ to effective HIV prevention, treatment, care and support for key populations and other vulnerable subpopulations?

2.1. IF YES, for which sub-populations?

KEY POPULATIONS AND VULNERABLE SUBPOPULATIONS		
People living with HIV	Yes	No
Men who have sex with men	Yes	No
Migrants/mobile populations	Yes	No
Orphans and other vulnerable children	Yes	No
People with disabilities	Yes	No
People who inject drugs	Yes	No
Prison inmates	Yes	No
Sex workers	Yes	No
Transgender people	Yes	No
Women and girls	Yes	No
Young women/young men	Yes	No
Other specific vulnerable populations ¹⁷ [write in]:	Yes	No

¹⁶ These are not necessarily HIV-specific policies or laws. They include policies, laws, or regulations which may deter people from or make it difficult for them to access prevention, treatment, care and support services. Examples cited in country reports in the past have include: "laws that criminalize same sex relationships", "laws that criminalize possession of condoms or drug paraphernalia"; "loitering laws"; "laws that preclude importation of generic medicines"; "policies that preclude distribution or possession of condoms in prisons"; "policies that preclude non-citizens from accessing ART"; "criminalization of HIV transmission and exposure", "inheritance laws/rights for women", "laws that prohibit provision of sexual and reproductive health information and services to young people", etc

¹⁷ Other specific vulnerable populations other than above, may be defined as having been locally identified as being at higher risk of HIV infection (e.g. (in alphabetical order) bisexual people, clients of sex workers, indigenous people, internally displaced people, prisoners, and refugees)

Briefly describe the content of these laws, regulations or policies	:			
Briefly comment on how they pose barriers:				
3. Does the country have a policy, law or regulation to reduce example, victims of sexual assault or women living with HI		men, including for		
	Yes	No		
Briefly describe the content of the policy, law or regulation and t	he populations inclu-	ded.		
4. Is the promotion and protection of human rights explicitly m	nentioned in any HIV	policy or strategy?		
	Yes	No		
IF YES, briefly describe how human rights are mentioned in this HIV policy or strategy:				

				Yes		No
F YES, briefly describe this	mechanism:					
,						
Does the country have a						
services are provided free applicable).	e-of-charge to	all people, to	some peop	le or not at al	l (circle "yes'	' or "no'
			Prov	vided		
	Provided charge to a			-charge	Provi but o	
	in the co	ountry		e people country	at a cost	
Antiretroviral treatment	Yes	No	Yes	No	Yes	No
HIV prevention services ¹⁸	Yes	No	Yes	No	Yes	No
HIV-related care and upport interventions	Yes	No	Yes	No	Yes	No
f applicable, which populate	ions have beer	n identified as	s priority, ar	nd for which s	ervices?	
Does the country have a prevention, treatment, c			re equal acc	cess for wome	en and men t	o HIV
•			re equal acc		en and men t	
•			re equal acc	cess for wome	en and men t	o HIV
•	are and suppo	ort? policy or str	rategy to en	Yes sure access to	HIV prever	No ntion,

¹⁸ Such as blood safety, condom promotion, harm reduction for people who inject drugs, HIV prevention for out-of-school young people, HIV prevention in the workplace, HIV testing and counseling, IEC on risk reduction, IEC on stigma and discrimination reduction, prevention of motherto-child transmission of HIV, prevention for people living with HIV, reproductive health services including sexually transmitted infections prevention and treatment, risk reduction for intimate partners of key populations, risk reduction for men who have sex with men, risk reduction for sex workers, school-based HIV education for young people, universal precautions in health care settings.

	Yes	No
YES, Briefly describe the content of this policy/s	trategy and the populations inclu	ded:
IF YES, does this policy/strategy include different key populations and/or other vulnerations.		re equal access
	Yes	No
YES, briefly explain the different types of approa	ches to ensure equal access for di	fferent populati
· - · · - · -		ployment purp
Does the country have a policy or law prohibitin (recruitment, assignment/relocation, appointm		ployment purp
· - · · - · -	ent, promotion, termination)?	
· - · · - · -		ployment purp No
(recruitment, assignment/relocation, appointm	ent, promotion, termination)? Yes	
(recruitment, assignment/relocation, appointm	ent, promotion, termination)? Yes	
· - · · -	ent, promotion, termination)? Yes	
(recruitment, assignment/relocation, appointm	ent, promotion, termination)? Yes	
(recruitment, assignment/relocation, appointm	ent, promotion, termination)? Yes	
(recruitment, assignment/relocation, appointm	ent, promotion, termination)? Yes	
(recruitment, assignment/relocation, appointm	ent, promotion, termination)? Yes	
(recruitment, assignment/relocation, appointm	ent, promotion, termination)? Yes	
(recruitment, assignment/relocation, appointment) YES, briefly describe the content of the policy or	ent, promotion, termination)? Yes law:	No
(recruitment, assignment/relocation, appointment) YES, briefly describe the content of the policy or	ent, promotion, termination)? Yes law:	No
YES, briefly describe the content of the policy or Does the country have the following human right.	ent, promotion, termination)? Yes law:	No No nt mechanisms
YES, briefly describe the content of the policy or Does the country have the following human ri a. Existence of independent national institu	ent, promotion, termination)? Yes law: ghts monitoring and enforcementions for the promotion and prot	No nt mechanisms
YES, briefly describe the content of the policy or Does the country have the following human ri a. Existence of independent national institurights, including human rights commission	ent, promotion, termination)? Yes law: ghts monitoring and enforcementions for the promotion and protons, law reform commissions, wa	No nt mechanisms
YES, briefly describe the content of the policy or Does the country have the following human ri a. Existence of independent national institu	ent, promotion, termination)? Yes law: ghts monitoring and enforcementions for the promotion and protons, law reform commissions, wa	No nt mechanisms
YES, briefly describe the content of the policy or Does the country have the following human ri a. Existence of independent national institurights, including human rights commission	ent, promotion, termination)? Yes law: ghts monitoring and enforcementions for the promotion and protons, law reform commissions, wa	No nt mechanisms
TYES, briefly describe the content of the policy or Does the country have the following human rights, including human rights commission	ent, promotion, termination)? Yes law: tions for the promotion and protions, law reform commissions, was ed issues within their work	No nt mechanisms fection of huma tchdogs, and

IF YES on any of the above questions, describe some example	les:		
 In the last 2 years, have there been the following trainin 	g and/or capac	city-building	activities:
a. Programmes to educate, raise awareness among pe concerning their rights (in the context of HIV) ¹⁹ ?	-		
	Ye	es	No
b. Programmes for members of the judiciary and law issues that may come up in the context of their wor	•	on HIV and	human rights
	Ye	es	No
a. Legal aid systems for HIV casework			
	Ye	es	No
b. Private sector law firms or university-based centres services to people living with HIV	s to provide fre	ee or reduced	l-cost legal
	Ye	es	No
3. Are there programmes in place to reduce HIV-related s	stigma and disc	crimination?	
	Ye	es	No
IF YES, what types of programmes?			
Programmes for health care workers		Yes	No
Programmes for the media		Yes	No
Programmes in the work place		Yes	No
Other [write in]:		Yes	No

¹⁹ Including, for example, Know-your-rights campaigns – campaigns that empower those affected by HIV to know their rights and the laws in context of the epidemic (see UNAIDS Guidance Note: Addressing HIV-related law at National Level, Working Paper, 30 April 2008)

²⁰ Including, for example, judges, magistrates, prosecutors, police, human rights commissioners and employment tribunal/labour court judges or commissioners

14.	Overall, on a scale of 0 to 10 (where 0 is "Very Poor" and 10 is "Excellent"), how would you rate
	the policies, laws and regulations in place to promote and protect human rights in relation to HIV
	in 2013?

Very Poor										Excellent
0	1	2	3	4	5	6	7	8	9	10

Since 2011,	what have	e been ke	y achieve	ements in	this area:					
What challe	enges rem	ain in thi	s area:							
15. Overall,	on a scale	e of 0 to 1	0 (where	e O is "Ver	v Door"	and 10 is	"Excelle	ent"), hov	w would	you rate
the effor										
				hts related						Evcellent
Very Poor	t to imple	ement hu	man rigl	hts related	d policies	, laws an	d regula	tions in 2	2013?	Excellent
										Excellent 10
Very Poor 0	t to imple	2	man rigl	hts related	d policies 5	6	d regula	tions in 2	2013?	
Very Poor	t to imple	2	man rigl	hts related	d policies 5	6	d regula	tions in 2	2013?	
Very Poor 0	t to imple	2	man rigl	hts related	d policies 5	6	d regula	tions in 2	2013?	
Very Poor 0	t to imple	2	man rigl	hts related	d policies 5	6	d regula	tions in 2	2013?	
Very Poor 0	t to imple	2	man rigl	hts related	d policies 5	6	d regula	tions in 2	2013?	
Very Poor 0	t to imple	2	man rigl	hts related	d policies 5	6	d regula	tions in 2	2013?	
Very Poor 0 Since 2011,	1 what have	2 e been ke	3 y achieve	hts related	d policies 5	6	d regula	tions in 2	2013?	
Very Poor 0	1 what have	2 e been ke	3 y achieve	hts related	d policies 5	6	d regula	tions in 2	2013?	
Very Poor 0 Since 2011,	1 what have	2 e been ke	3 y achieve	hts related	d policies 5	6	d regula	tions in 2	2013?	
Very Poor 0 Since 2011,	1 what have	2 e been ke	3 y achieve	hts related	d policies 5	6	d regula	tions in 2	2013?	
Very Poor 0 Since 2011,	1 what have	2 e been ke	3 y achieve	hts related	d policies 5	6	d regula	tions in 2	2013?	

IV. PREVENTION

1. Has the country identified the specific needs for HIV prevention programmes?

	Yes	No
--	-----	----

IF YES, how were these specific needs determined?
IF YES, what are these specific needs?

1.1 To what extent has HIV prevention been implemented?

	The m	ajority of p	eople in nee	d have acce	ss to
HIV prevention component	Strongly disagree	Disagree	Agree	Strongly agree	N/A
Blood safety	1	2	3	4	N/A
Condom promotion	1	2	3	4	N/A
Harm reduction for people who inject drugs	1	2	3	4	N/A
HIV prevention for out-of-school young people	1	2	3	4	N/A
HIV prevention in the workplace	1	2	3	4	N/A
HIV testing and counseling	1	2	3	4	N/A
IEC ²¹ on risk reduction	1	2	3	4	N/A
IEC on stigma and discrimination reduction	1	2	3	4	N/A
Prevention of mother-to-child transmission of HIV	1	2	3	4	N/A

²¹ IEC = information, education, communication

	The m	ajority of p	eople in nee	d have acce	ss to
HIV prevention component	Strongly disagree	Disagree	Agree	Strongly agree	N/A
Prevention for people living with HIV	1	2	3	4	N/A
Reproductive health services including sexually transmitted infections prevention and treatment	1	2	3	4	N/A
Risk reduction for intimate partners of key populations	1	2	3	4	N/A
Risk reduction for men who have sex with men	1	2	3	4	N/A
Risk reduction for sex workers	1	2	3	4	N/A
School-based HIV education for young people	1	2	3	4	N/A
Universal precautions in health care settings	1	2	3	4	N/A
Other[write in]:	1	2	3	4	N/A

2. Overall, on a scale of 0 to 10 (where 0 is "Very Poor" and 10 is "Excellent"), how would you rate the efforts in the implementation of HIV prevention programmes in 2013?

Very Poor										Excellent
0	1	2	3	4	5	6	7	8	9	10

Since 2011, what have been key achievements in this area:
What challenges remain in this area:

V. TREATMENT, CARE AND SUPPORT

1. Has the country identified the essential elements of a comprehensive package of HIV treatment, care and support services?

Yes	No

IF YES, Briefly identify the elements and what has been prioritized:
Briefly identify how HIV treatment, care and support services are being scaled-up?
briefly laterity flow 111 v treatment, care and support services are being scared-up:

1.1. To what extent have the following HIV treatment, care and support services been implemented?

	The majority of people in need have access to							
HIV treatment, care and support service	Strongly disagree	Disagree	Agree	Strongly agree	N/A			
Antiretroviral therapy	1	2	3	4	N/A			
ART for TB patients	1	2	3	4	N/A			
Cotrimoxazole prophylaxis in people living with HIV	1	2	3	4	N/A			
Early infant diagnosis	1	2	3	4	N/A			
HIV care and support in the workplace (including alternative working arrangements)	1	2	3	4	N/A			
HIV testing and counselling for people with TB	1	2	3	4	N/A			
HIV treatment services in the workplace or treatment referral systems through the workplace	1	2	3	4	N/A			
Nutritional care	1	2	3	4	N/A			
Paediatric AIDS treatment	1	2	3	4	N/A			
Post-delivery ART provision to women	1	2	3	4	N/A			

	The majority of people in need have access to							
HIV treatment, care and support service	Strongly disagree	Disagree	Agree	Strongly agree	N/A			
Post-exposure prophylaxis for non-occupational exposure (e.g., sexual assault)	1	2	3	4	N/A			
Post-exposure prophylaxis for occupational exposures to HIV	1	2	3	4	N/A			
Psychosocial support for people living with HIV and their families	1	2	3	4	N/A			
Sexually transmitted infection management	1	2	3	4	N/A			
TB infection control in HIV treatment and care facilities	1	2	3	4	N/A			
TB preventive therapy for people living with HIV	1	2	3	4	N/A			
TB screening for people living with HIV	1	2	3	4	N/A			
Treatment of common HIV-related infections	1	2	3	4	N/A			
Other[write in]:	1	2	3	4	N/A			

1.2. Overall, on a scale of 0 to 10 (where 0 is "Very Poor" and 10 is "Excellent"), how would you rate the efforts in the implementation of HIV treatment, care and support programmes in 2013?

Very Poor										Excellent
0	1	2	3	4	5	6	7	8	9	10

Since 2011, what have been key achievements in this area:
What challenges remain in this area:

2.		e country ble childr		olicy or s	strategy to	o addres	s the nee	ds of orp	hans and	other	
								Υe	es		No
2.1.	IF YES,	is there a	n operati	onal defi	inition fo	r orphai	ıs and vu	lnerable	children	in the co	untry?
								Ye	es		No
2.2.	IF YES,	does the c	ountry ha	ive a nati	onal actio	on plan sj	pecifically	y for orpl	nans and v	vulnerabl	e children?
								Υe	es		No
										I	
3.		on a scal									
Ve	ery Poor										Excellent
	0	1	2	3	4	5	6	7	8	9	10
Si	nce 2011,	, what hav	e been ke	ey achieve	ements in	this area	ı:				
W	hat chall	enges rem	ain in th	is area:							

Appendix 4. Sample checklist for Country Progress Report

Reporting process established, including timelines and milestones, and roles of NAC, government agencies, UN agencies, civil society and other relevant partners.
Funding secured for all aspects of the reporting process.
Data collection, vetting and analysis process established, including:
• Identification of relevant tools (including Spectrum) and sources for data collection for each indicator
 Timeline for data collection in line with other data collection efforts, including those via funding agencies such as the Global Fund, PEPFAR and UN agencies
 Reporting timeline for facility-based indicators for national level aggregation
 Data vetting and triangulation workshops with the aim of reaching consensus on the correct value for each indicator
Protocols established for data processing and management, including:
Basic data cleaning and validation
 One database for analysis and reporting purposes
Relevant data analysed in coordination with partner organizations from government, civil society and the international community
Consensus reached with stakeholders, including government agencies and civil society, on the final report to be submitted
Data entered into and narrative report attached to the online reporting tool by 31 March 2014
Data queries answered (sent from AIDSreporting@unaids.org or directly in the online reporting tool).

Appendix 5. Selected bibliography

UNAIDS (2010) 12 Components M&E System Assessment - Guidelines to support preparation, implementation and follow up activities. Geneva: UNAIDS

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UNAIDS (2008) Organizing Framework for a Functional National HIV Monitoring and Evaluation System. Geneva: UNAIDS

UNAIDS (2013) UNAIDS Report on the Global AIDS Epidemic. Geneva: UNAIDS

UNAIDS (2011) Securing the future today – Synthesis of Strategic Information on HIV and Young People; Geneva, UNAIDS

UNAIDS (2010) Strategic Guidance for the Evaluation of HIV prevention programmes. Geneva: UNAIDS

WHO (2013) Consolidated guidelines on the use of antiretroviral drugs for treating and preventing HIV infection. Geneva: WHO

UNICEF (2013). The State of the World's Children Report. New York: UNICEF

UNICEF (2009) Country-led Monitoring and Evaluation Systems. New York; UNICEF

Appendix 6. Male circumcision indicators

These two indicators are only required from 16 countries with high HIV prevalence, low levels of male circumcision and generalized heterosexual epidemics i.e. Botswana, Ethiopia, Central African Republic, Kenya, Lesotho, Malawi, Mozambique, Namibia, Rwanda, South Africa, South Sudan, Swaziland, Uganda, United Republic of Tanzania, Zambia and Zimbabwe.

1.22 Proportion of males circumcised

Percentage of men 15-49 that are circumcised

What it measures

It measures progress towards increased coverage of male circumcision.

Rationale

There is compelling evidence that male circumcision reduces the risk of heterosexually acquired HIV infection in men by approximately 60%. Three randomized controlled trials have shown that male circumcision provided by well-trained health professionals in properly equipped settings is safe and can reduce the risk of HIV acquisition. WHO/UNAIDS recommendations emphasize that male circumcision should be considered an efficacious intervention for HIV prevention in countries and regions with heterosexual epidemics, high HIV and low male circumcision prevalence.

Numerator: Number of male respondents aged 15-49 years who report that they

are circumcised.

Denominator: Number of all male respondents aged 15–49 years

Calculation: Numerator / Denominator

Method of Population-based surveys (Demographic and Health Survey, AIDS Indicator measurement: Survey, Multiple Indicator Cluster Surveys or other representative survey)

Measurement frequency:

Every 3–5 years

Disaggregation: • Age 15-19, 20-24 and 25-49 years

Source/practitioner of circumcision procedure: formal healthcare system or traditional

Strengths and weaknesses

Changing rates of male circumcision may or may not be the result of a programme. For example, changing societal norms not due to a programme may be leading to changing rates of male circumcision. This indicator measures total change in the population, whatever the reason(s).

Existing population-based surveys (such as DHS) may not accurately measure true male circumcision status because of a lack of knowledge of what male circumcision is, confusion about circumcision status, or perceived social desirability of circumcision status. Other approaches to determining circumcision status might be used, e.g. the use of pictures or drawings (drawings may be more culturally appropriate), prompts or even direct examination. Modelling the potential impact of changing rates of male circumcision on HIV incidence requires accurate knowledge of male circumcision status over time.

APPENDIX 6

Further information

For further information on Male Circumcision indicators, see *A guide to indicators for male circumcision programmes in the formal health care system, WHO, UNAIDS, 2009* http://whqlibdoc.who.int/publications/2009/9789241598262_eng.pdf

1.23 Number of male circumcisions performed

Number of male circumcisions performed according to national standards during the last 12 months

What it measures

It measures progress in scaling up male circumcision services.

Rationale

There is compelling evidence that male circumcision reduces the risk of heterosexually acquired HIV infection in men by approximately 60%. Three randomized controlled trials have shown that male circumcision provided by well-trained health professionals in properly equipped settings is safe and can reduce the risk of HIV acquisition. WHO/UNAIDS recommendations emphasize that male circumcision should be considered an efficacious intervention for HIV prevention in countries and regions with heterosexual epidemics, high HIV and low male circumcision prevalence.

Numerator: Number of males circumcised during the past 12 months according

to national standards

Denominator: Not applicable

Method of

Health facility recording and reporting forms

measurement:

Measurement frequency:

Yearly

Disaggregation: Age: <1, 1-9, 10-14, 15-19, 20-24, 25-49, and 50+ years

Strengths and weaknesses

The total number of male circumcisions carried out indicates either change in the supply of services or change in demand. Comparing the results against previous values shows where male circumcision services have been newly instituted or where male circumcision volume has changed.

Further disaggregations are recommended at country level:

- i) HIV positive by test(s) on site; HIV negative by test(s) on site; HIV indeterminate result by test(s) on site; Unknown/refused HIV test;
- ii) Type and location of health facility
- iii) Cadre of provider

When the number of male circumcisions is disaggregated by HIV status and age it will be possible to determine the impact of male circumcision programmes on HIV incidence using models. If a country has prioritized particular age groups this disaggregation will help determine whether age-specific communication strategies are creating demand. Further if the data are available by type and location of health-care facility where the circumcision was performed resource allocation needs can be assessed. Finally by disaggregating these data by the cadre of health-care provider will determine if task-shifting efforts are succeeding and determine resource allocation.

Some programmes will work closely with voluntary HIV counselling and testing services to provide HIV testing. A patient desiring male circumcision may have been recently tested, in which event an on-site HIV test may be unnecessary. In these cases, a written 'verified result' may be requested at the facility to verify

HIV status. There is no specific length of time before male circumcision that the test should have been done, but within three months is suggested (the purpose of testing is not to identify every man who might be infected but to provide HIV testing to men seeking health care and to identify HIV-positive men who, if they choose to be circumcised, are likely to be at higher risk of surgical complications, i.e. men who are chronically infected and with low CD4 counts).

Further information

For further information on Male Circumcision indicators, see *A guide to indicators for male circumcision programmes in the formal health care system, WHO, UNAIDS, 2009* http://whqlibdoc.who.int/publications/2009/9789241598262_eng.pdf Appendix 4.

Appendix 7. Geographic data collection in Surveillance, Monitoring and Evaluation

There is a programmatically relevant geographic aspect to virtually every element of surveillance and Monitoring and Evaluation (M&E) systems. Many input, output, and outcome indicators can be represented in a Geographic Information System (GIS) for analysis and presentation. As a result, geographic data can be leveraged for epidemic assessment, monitoring and evaluation. Epidemiology has an obvious geographic dimension, and the range of prevention, care and treatment services also come together in specific places. Geography therefore has a key role in integrating data from surveillance and different programmatic streams. A standards-based approach to spatial data can support country systems where operational guidance directs programmes to take account of geographic aspects of interventions. Such an approach will also promote consistency of geospatial data between data sources, facilitating sharing and use of geospatial data by countries and partners, as well as the bringing together of all available data sources to inform analysis and decision-making at local level.

Spatial data inputs to surveillance and M&E

To facilitate data integration and analysis, geographic markers for data should be maintained with indicators at the appropriate level of precision and using standardized geographic references and naming conventions. The appropriate level of precision may be as general as a health district, province or even a national administrative boundary. However, attaching geographic information to the more granular data that compose aggregate indicators can enable a wide array of analysis, such as geographic coverage of services, spatial distribution of human resource and expenditures, and the estimation of change over time for small areas.

For many surveillance applications, geographic representativeness dictates the scale at which data can be used. For example, population-based surveys are typically representative of populations at the province level. ANC sentinel surveillance data is usually linked to specific health facilities or more rarely to a cluster of rural health facilities. The geographic localization of the ANC site or cluster of ANC sites should be attached to the he HIV prevalence data from these sites. As surveillance systems transition to using HIV prevalence data generated by PMTCT programmes, the prevalence data should be accompanied by the geo-location of the PMTCT sites. Sentinel surveillance data for key populations (e.g. collected through integrated bio-behavioural surveys – IBBS) can either be located to the central facility at which the surveillance is conducted (e.g. when using respondent-driven sampling or clinic-based surveillance), or to the actual location where respondents are encountered during the surveillance (e.g. when using time-location sampling).

For indicator data that characterize a health facility, the finest geographic representation is a point based on the latitude and longitude of the facility, which is information that should be maintained as part of a Ministry of Health's master list of health facilities, where that exists, or possibly in a GIS unit within a central statistical agency. Key monitoring data that should be geographically tagged include people tested for HIV, new HIV diagnoses, pregnant women tested for HIV, people initiated on ART, people on ART, pregnant women receiving antiretrovirals and early infant diagnoses.

Many community-based activities whether for key populations or for supporting treatment programmes may also be located with latitude and longitude, although the geography of non-facility based activities in the continuum of response can be diffuse or complex.

Spatial data standards and metadata

Most countries have National Spatial Data Infrastructure (NSDI) initiatives or an explicit spatial data component in a larger national information and communication infrastructure. NSDI includes the technology, policies, standards, human resources and related activities necessary to acquire, process, distribute, use, maintain and preserve spatial data." A data management plan for spatial data is recommended to reduce duplication and to support country ownership and sustainability by ensuring that these data become part of the NSDI of the country.

To the extent possible, databases should include international naming standards in addition to any local naming standards and place codes. For country-specific data where metadata standards are specified by NSDI policy, spatial data can follow the FGDC Metadata Standard and include any additional metadata elements enumerated in the local standard. Adherence to spatial data standards is necessary for alignment with national programmes and systems.

Unique identification of individuals

Place can uniquely identify individuals, especially when linked with other data elements Care must be taken in determining whether the release of specific spatial data could be inappropriately leveraged with other data to violate confidentiality. Extreme care should be taken when developing maps of stigmatized key populations or the places where key populations congregate.

Geospatial tools

A variety of commercial and free and open-source tools to support geographic mapping are available. Elementary spatial analysis can be conducted in spreadsheets or using digital globes. More advanced spatial analysis, management of spatial data, and displays of spatial data can be accomplished using a GIS. The right tool should be matched with the right data, the right analysis and at the right scale. Skills that potentially already exist in countries to conduct geospatial analysis should be sought. Analysis using geographic mapping tools can be complemented through participatory methodologies involving community stakeholders.

A GUIDE ON INDICATORS FOR MONITORING AND REPORTING ON THE HEALTH SECTOR RESPONSE TO HIV AND AIDS

2014







List of Abbreviations

3TC ABC	lamivudine abacavir	NRTI	nucleoside reverse transcriptase inhibitors
ANC	antenatal care	OST	opioid substitution therapy
ART		PCR	polymerase chain reaction
ARV	antiretroviral therapy	PEPFAR	United States President's
	antiretroviral drug	ILIIAN	Emergency Plan for AIDS Relief
AZT CTX	zidovudine co-trimoxazole	PITC	provider-initiated testing and counselling
DHS	Demographic and Health Survey	PLHIV	people living with HIV
DPT3	third dose of diphtheria, pertussis and tetanus vaccine	PMTCT	prevention of mother-to-child transmission
d4T	stavudine	RPR	rapid plasma reagin
EBF	exclusive breastfeeding	RF	replacement feeding
EIA	enzyme immunoassay	SAM	Service Availability Mapping
EFV	efavirenz	SPA	Service Provision Assessment
FTC	emtricitabine	SRH	Sexual and Reproductive Health
GARPR	Global AIDS Response	STI	sexually transmitted infections
	Progress Reporting	SW	sex workers
HIV	human immunodeficiency virus	TB	tuberculosis
HIVDR	HIV drug resistance	T&C	testing and counselling
HTC	HIV testing and counselling	TDF	tenofovir
IDP	internally displaced persons	TPHA	treponema pallidum
IF	infant feeding	ППА	haemagglutination assay
IPT	isoniazid preventive therapy. Also can be termed TBPT	TPPA	treponema pallidum particle agglutination assay
L&D	(TB preventive therapy) labour and delivery	UNAIDS	United Nations Joint Programme on HIV/AIDS
LMIS	logistics management information system	UNGASS	United Nations General Assembly Special Session on HIV/AIDS
LPV/r	lopinavir	UNPD	United Nations Population Division
M&E	monitoring & evaluation	UNICEF	United Nations Children's Fund
MC	male circumcision	UNODC	United Nations Office on Drugs and
MDG	Millennium Development Goal		Crime
MF	mixed feeding	VCT	voluntary counselling and testing
MTCT	mother-to-child transmission	VDRL	venereal disease research
NAP	National AIDS Programme		laboratory test
NSP	needle and syringe programme	VL	viral load
NNRTI	non-nucleoside reverse transcriptase inhibitors		

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I. Introduction

As countries scale up their national HIV and AIDS programmes towards the goal of universal access (UA) to prevention, treatment, care and support, it is increasingly important to strengthen strategic information on the epidemic and national responses to inform policies and programmes, improve the effectiveness of interventions and promote accountability.

At the international level, WHO is committed since the 59th World Health Assembly in 2006 to monitor and report annually on global progress in countries' health sector responses towards universal access to HIV prevention, treatment, care and support.²² WHO is working with UNICEF and UNAIDS to harmonize the global monitoring and reporting on the health sector response to HIV and AIDS towards universal access. This joint work of the UN partners aims to harmonize data collection and minimize the reporting burden on countries.

In order to collect data from countries, WHO, UNAIDS and UNICEF have developed a Joint Online Reporting Tool. The reporting tool and guidance on the Global AIDS Response Progress Reporting indicators and the UA health sector indicators are available at http://AIDSreporting.unaids.org.

This part of the guide describes in detail the additional health sector indicators that are not described in the UNAIDS Global AIDS Response Progress Reporting. It can also be considered for use to monitor the health sector response at the national level, in addition with other information, to review progress. In summary:

- Global Reporting: This part of the guide complements the UNAIDS Global AIDS Response
 Progress Reporting 2014: guidelines. Construction of core indicators for monitoring the 2011
 Political Declaration on HIV and AIDS. The overall recommended country reporting process is
 described in detail in the global reporting guidelines.
 - This section aims to support and facilitate data collection using the Joint Online Reporting Tool with a focus on the additional indicators of the 2014 health sector reporting requested which are not part of the GARPR indicators. The online data collection tool, disseminated to all countries, is the main tool to enable annual global reporting on the health sector progress towards universal access to HIV prevention, care, and treatment.
- National Monitoring: This guide can also be used for national monitoring of the health sector's response to HIV/AIDS. It can be adapted to the epidemic context of each country. For example, countries should select indicators that would support monitoring of their own nationally-set targets. They may also add or remove some of the indicators depending on the importance of intervention areas to their country epidemic.

^{22.} HIV/AIDS. WHO's contribution to universal access to HIV/AIDS prevention, treatment and care: report by the Secretariat. Geneva, WHO, 2006.

Indicator descriptions in this guide The indicator descriptions follow this format:

The indicator number is the number in the 2014 GARPR/UA reporting tool

This is the number used in last year's tool

	X. INDICATOR TITLE (# y.y)
Rationale	Why this indicator is important
What it measures	What the indicator measures
Numerator	Definition of the numerator
Denominator	Definition of the denominator (sources of information must be specified: for some indicators, estimates only are possible and/ or required)
How to Measure and Measurement Tools	What is included in the numerator and denominator Method of measurement Tools used for measurement
Disaggregation	Recommended disaggregation. Even if not included for breakdown in the Reporting Tool, disaggregation is recommended to be collected for national monitoring and reporting as appropriate.
Strengths and weaknesses	Description of the strengths and weaknesses of the indicator
Additional considerations	Other points for countries to note
Data utilization	How this indicator can be used and some implications
Data Quality Control and Notes for the Reporting Tool	Additional information on issues to consider when filling in the reporting tool. Includes elements of: • Double Reporting: What to pay attention to in order to assess possible double reporting. • National Representativeness: What to pay attention to in order to assess the national representativeness of the value reported. • Denominator Issues: Issues to note about the denominator • Triangulation Options: Other data sources that can be reviewed to assess the validity of the indicator value
Other References	References related to the indicator, e.g.: PMTCT: Indicator in the updated Monitoring and Evaluating the PMTCT of HIV A guide for national programmes (2011) HIV/TB: Indicators in the updated A guide to monitoring and evaluation for collaborative TB/HIV M&E activities (2009) People who inject drugs: Technical Guide for countries to set targets for universal access to HIV prevention, treatment and care for injecting drug users (2009)

Technical Support and Contact for Questions

WHO, UNICEF and UNAIDS are committed to support countries improve their strategic information system, including and not limited to the review of health sector M&E systems; data quality and validation; evaluating impact; surveillance; operational research; and training in various aspects of strategic information.

Please do not hesitate to contact WHO at hivstrategicinfo@who.int for any questions or requests, or to send any comments and suggestions for improving this guidance.

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II. INDICATOR DESCRIPTIONS

The present table gives an overview of the indicators described in the Global AIDS Response Progress Reporting 2014 guidelines and those described in this guide for the 2014 health sector reporting for universal access (UA 2014).

GARPR	UA 2014		
		Target '	1. Reduce sexual transmission of HIV by 50% by 2015
		Indicate	ors for the general population
x		1.1	Young People: Knowledge about HIV Prevention
х		1.2	Sex Before the Age of 15
х		1.3	Multiple sexual partners
х		1.4	Condom Use During Higher-Risk Sex
х		1.5	HIV Testing in the General Population
х		1.6	Reduction in HIV Prevalence
		Indicate	ors for sex workers
x		1.7	Sex Workers: Prevention programmes
x	x	1.8	Sex Workers: Condom Use
X	x	1.9	Sex Workers: HIV Testing
X	x	1.10	Sex Workers: HIV Prevalence
		Indicate	ors for men who have sex with men
X		1.11	Men who have sex with men: Prevention programmes
X	x	1.12	Men who have sex with men: Condom Use
X	x	1.13	Men who have sex with men: HIV Testing
x	x	1.14	Men who have sex with men: HIV Prevalence
		Testing	and Counselling
	x	1.16	HIV Testing in 15+ (from programme records)
	x	1.16.1	Rapid test kits stock-outs (new)
		Sexuali	ly Transmitted Infections
	x	1.17	Sexually Transmitted Infections (STIs)
		1.17.1	Percentage of women accessing antenatal care (ANC) services who were tested for syphilis
		1.17.2	Percentage of antenatal care attendees who were positive for syphilis

GARPR	UA 2014		
		1.17.3	Percentage of antenatal care attendees positive for syphilis who received treatment
		1.17.4	Percentage of sex workers with active syphilis
		1.17.5	Percentage of men who have sex with men with active syphilis
		1.17.6	Number of adults reported with syphilis (primary/secondary and latent/unknown) in the past 12 months
		1.17.7	Number of reported congenital syphilis cases (live births and stillbirth) in the past 12 months
		1.17.8	Number of men reported with gonorrhoea in the past 12 months
		1.17.9	Number of men reported with urethral discharge in the past 12 months
		1.17.10	Number of adults reported with genital ulcer disease in the past 12 months
	X	1.18	Percentage (%) of pregnant women with a positive syphilis serology whose sexual contacts were identified and treated for syphilis (PAHO only)
		Male cir	cumcision
х	X	1.22	Male circumcision, prevalence
x	x	1.23	Number of men circumcised last year
		Target 2 by 50%	. Reduce transmission of HIV among people who inject drugs by 2015
X	x	2.1	People who inject drugs: Number of needles/person who injects drugs
X	x	2.2	People who inject drugs: Condom Use
х	X	2.3	People who inject drugs: Safe Injecting Practices
x	x	2.4	People who inject drugs: HIV Testing
x	x	2.5	People who inject drugs: HIV Prevalence
	x	2.6	People on opioid substitution therapy
	X	2.7	NSP and OST sites
		_	Eliminate mother-to-child transmission of HIV by 2015 and tially reduce AIDS-related maternal deaths
x	x	3.1	Prevention of Mother-to-Child Transmission
х	X	3.2	Early Infant Diagnosis
x	x	3.3	Mother-to-Child transmission rate (modelled)

GARPR	UA 2014		
	х	3.4	Pregnant women who were tested for HIV and received their results
	x	3.5	Percentage of pregnant women attending antenatal care whose male partner was tested for HIV in the last 12 months
	x	3.6	Percentage of HIV-infected pregnant women assessed for ART eligibility through either clinical staging or CD4 testing
	X	3.7	Percentage of infants born to HIV-infected women provided with ARV prophylaxis to reduce the risk of early mother-to-child-transmission in the first 6 weeks
	X	3.9	Percentage of infants born to HIV-infected women started on co-trimoxazole (CTX) prophylaxis within two months of birth
	X	3.10	Distribution of feeding practices for infants born to HIV-infected women at DPT3 visit
	X	3.11	Number of pregnant women attending ANC at least once during the reporting period
	x	3.11.1	Percentage of HIV-positive pregnant women who had their pregnancy terminated (EURO8)
	x	3.11.2	Percentage of HIV-positive pregnant women who delivered during the reporting year (EURO9)
	x	3.12	PAHO-specific indicators
	x	3.13	EURO-specific PMTCT Indicator (pregnant women who inject drugs)
	X	3.13.1	Percentage of HIV-positive pregnant women who were injecting drug users (PWIDs) (EURO11)
	x	3.13.2	Percentage of HIV-positive pregnant PWID women who received OST during pregnancy (EURO12)
	X	3.13	Percentage of HIV-positive pregnant PWID women who received ARVs to reduce the risk of mother-to-child transmission during pregnancy (EURO13)
		•	I. Have 15 million people living with HIV on antiretroviral nt by 2015
х	x	4.1	ART coverage (adults and children), including number of eligible adults and children who newly enrolled on antiretroviral therapy during the reporting period (2013)
x	x	4.2a	HIV Treatment: 12 months retention
	x	4.2b	HIV Treatment: 24 months retention
	x	4.2c	HIV Treatment: 60 months retention

GARPR	UA 2014		
	х	4.2.1	Percentage of injecting drug users with HIV still alive and known to be on treatment 12 months, 24 months and 60 months after initiation of antiretroviral therapy (EURO4)
	X	4.3	Health facilities that offer antiretroviral therapy
	X	4.4	ARV stock-outs
	X	4.5	Late HIV diagnoses (PAHO only)
	X	4.6	HIV care
		4.7	Viral Load
		Target 5	i. Reduce tuberculosis deaths in people living with HIV by 50%
x	x	5.1	Co-Management of Tuberculosis and HIV Treatment
	x	5.2	Percentage of people living with HIV (PLHIV) newly enrolled in care who are detected having active TB disease (new)
	х	5.3	Percentage of adults and children newly enrolled in HIV care (starting isoniazid preventive therapy (IPT))
	х	5.4	Percentage of adults and children enrolled in HIV care who had TB status assessed and recorded during their last visit
		_	5. Reach a significant level of annual global expenditure –24 billion) in low and middle-income countries
x		6.1	AIDS Spending - Domestic and international AIDS spending by categories and financing sources
		Target 7	'. Eliminating gender inequalities
x		7.1	Prevalence of Recent Intimate Partner Violence (IPV)
		Target 8	3. Eliminating stigma and discrimination
			development, Stigma indicator for general population planned to for 2014 reporting
		Target 9	. Eliminate Travel restrictions
			estriction data collected by Human Rights and Law Division at HQ, no data collected needed
		Target 1	0. Strengthening HIV integration
х		10.1	Orphans and non-orphans school attendance
x		10.2	Economic support for eligible households

GARPR	UA 2014		
		Policy questions, relevant questions for all 10 targets.	
x		P.1 Special 2014 GARPR questionnaire	
	x	P.1b WHO policy questions	
		P.1c European NCPI supplement	
		Appendices	
	х	Appendix 1. HIV/Hepatitis Indicators (EURO)	
		Number of adults and children currently in HIV care who were screened for hepatitis B (EUR15)	
	х	Percentage of HIV-positive hepatitis B cases eligible for hepatitis B treatment who received treatment for both hepatitis B and HIV (EUR16)	
	x	Number of adults and children currently in HIV care who were screened for hepatitis C (EUR17)	
	x	Percentage of HIV-positive hepatitis C cases eligible for hepatitis C treatment who received treatment for hepatitis C (EUR18)	
	х	Appendix 2. HIV/AIDS cases (PAHO)	
		P.1d Number of HIV cases diagnosed in 2012 and reported, by sex for 2012 (PAHO)	
	x	P.1e Number of AIDS cases diagnosed in 2012 and reported, by sex for 2012 (PAHO)	
		Key population size estimations	

Note on Defining "Health Facility"

A frequently asked question is what we are defining as a health facility. For the purposes of this reporting process, we are excluding health facilities that provide specialized care which would never provide any HIV services (e.g. an eye clinic). If you have difficulties trying to define what is counted as a health facility for this exercise, please provide any comments you have in the Comment box or e-mail WHO at hivstrategicinfo@who.int.

Target 1: Reduce Sexual transmission of HIV by 50% by 2015 Testing and Counselling

	n and men aged 15 and older who received HIV testing and counselling of the and know their results
Rationale	Knowledge of HIV status is critical for access to HIV treatment, care and support, and prevention. There are different models for delivery of the testing and counselling services such as client-initiated testing and counselling (CITC) and provider-initiated testing and counselling (PITC). The essential elements of TC are that those who are tested are appropriately counselled and know the results.
What it measures	Number of people aged 15 and older who received HIV T&C through any method or setting (excluding mandatory T&C) in the past 12 months and know their results
	(Note: Although not required for the purposes of this indicator the denominator may be gauged by using the general population as the denominator in generalized epidemics, and the key populations at higher risk and other groups for low-level and concentrated epidemics. These data can be reviewed along with an estimate of what percentage of the HIV+ population already know their status, and what the recommended HIV testing policy or frequency is.
	Programmatic progress for testing and counselling. Tracking the number of individuals who are tested and counselled and know their status provides an indication of uptake of T&C in the country.
How to Measure and Measurement Tools	Programme service statistics compiled from routine reports of the number of people tested and know the results from all service points, including clinics, hospitals, VCT sites, other NGO sites and outreach points, mobile testing, home and community testing, testing delivered in the workplace, schools, testing as part of special campaigns, and all other forms of testing (excluding mandatory T&C) which are often aggregated at the district levels and subsequently at the national level. This indicator is not measured through population-based surveys.
Disaggregation	Sex: male, female, pregnancy
	Serostatus: HIV positive, HIV negative
	If possible:
	Age: 15-19, 20-24, 25+ Test: New test , Repeated test
	HIV transmission mode: injecting drug use, sex between men,
	heterosexual contact, mother-to-child transmission, other and unknown (European Region only)

Strengths and weaknesses	This indicator permits comparison of trends of the quantity of TC services delivered and the strength of scaling up TC services over time. This indicator may provide information on the number of times T&C occurred, and not necessarily the number of people who received T&C services unless countries have a mechanism to avoid double-counting of repeat testers. The indicator does not provide information on whether those who were tested were adequately referred to and receiving follow-up services to benefit from knowing their status.
Data utilization	To review the number of tests conducted in the country, data can be compared with previous years to look at trends while considering the percentage of the population that may have already been tested recently. It can be useful to explore any patterns in testing, for example whether there were more tests conducted in a particular season or month when there were campaigns, or whether many more people are being tested in particular health facilities or in the communities.
Additional considerations for countries	In some countries, a significant proportion of testing and counselling services are provided by community-based organizations or unregistered organizations, which often may not be included as part of national statistics. These organizations should be encouraged to register with national authorities so all data on testing and counselling could be reflected in the national statistics.
Data Quality Control and Notes for Reporting	Double Reporting: Countries will need to estimate the extent of repeat testers in order to determine the true number of persons tested over the period. If countries have a mechanism to make such a meaningful assessment (e.g. record of the number of repeat tests or re-testers within a year), please do so and note how this was done. Otherwise, please report the total number of tests reported and clarify that repeat tests are likely included.
	National Representativeness: Try to ensure information from non-governmental and private facilities is also available at the central level. If significant information is missing, note it down in the comments section.
	Denominator Issues: Although not required for the purposes of this indictor the validity of the numerator may be gauged by comparing the general population as the denominator in generalized epidemics, and the size of the key populations at higher risk and other groups for low-level and concentrated epidemics.
	Triangulation Options: In generalized epidemics, data from population-based surveys asking for the number (and calculating the percentage) of people tested can be compared to with this indicator value to assess and discuss any major differences.

1.16.1 Percentage of health facilities dispensing HIV rapid test kits that experienced a stock- out in the last 12 months (new)		
Numerator	Number of health facilities dispensing rapid test kits that experienced a stock-out in the last 12 months.	
Denominator	Total number of health facilities dispensing rapid test kits	

Sexually Transmitted Infections

1.17.1 STIs: Percentage of women accessing antenatal care (ANC) services who were tested for syphilis	
Rationale	Testing pregnant women for syphilis early in pregnancy is important both for their health and the health of the fetus, and for second generation surveillance purposes. It also contributes to monitoring of the quality of ANC services.
What it measures	Coverage of syphilis testing in women attending first ANC services
Numerator	Number of women attending ANC services who were tested for syphilis
Denominator	Number of women attending ANC services
How to Measure and Measurement Tools	How to measure: All pregnant women should be tested ("screened") for syphilis at their first antenatal care visit. Ideally countries will report on testing at any visit as well as testing at first visit. Countries unable to distinguish first visit from testing at any visit should still report data on this indicator, but should make sure that it is clearly reported as data for "any visit". This indicator should be measured annually. Either non-treponemal tests that measure reaginic antibody (e.g., VDRL or RPR) or treponemal tests that measure treponemal antibody (e.g., TPHA, TPPA, EIA or rapid treponemal tests) may be used for screening. For this indicator simply being tested by either type of test is sufficient, although being tested with both is preferred. Please indicate in the "Comments" section what test type is generally used in your country. Measurement tools: Ideally national programme records aggregated from health facility data should be used. However, if national programme data are not available, data from sentinel surveillance or special studies
	can be reported only if the data are felt to be representative of the national situation. Please specify the source and coverage of your data (for example, national programme data from all 12 provinces) in the "Comments" section.
Disaggregation	Tested at any visit, tested at first visit
Additional considerations	Countries may wish to also monitor the week of pregnancy that each woman is tested. Preventing congenital syphilis requires testing early in pregnancy, as stillbirth may occur in the second trimester. Knowing that women are being tested late in pregnancy will indicate either that women are not accessing ANC early or that testing is not occurring early in pregnancy.
	Programmes that test pregnant women for syphilis and those that test pregnant women for HIV should work together to enhance the effectiveness of their individual programme work.

Data utilization	Global: Examine trends over time to assess progress towards target levels of testing coverage required for elimination of mother-to-child transmission of syphilis. Knowledge of testing policies and practices should be used to assist with interpretation of trends in coverage. Local: Data can be used to identify clinics not fully implementing national policy.
Data Quality Control and Notes for the Reporting Tool	Please comment on if the data you are providing is routine programme data, and if it is felt to be representative of the entire country.
Other References	Recommended indicator in "National-Level Monitoring of the Achievement of Universal Access to Reproductive Health: Conceptual and practical considerations and related indicators" and "Methods for Surveillance and Monitoring of Congenital Syphilis Elimination within Existing Systems".

1.17.2 STIs: Percentage	of antenatal care attendees who were positive for syphilis
Rationale	Syphilis infection in antenatal care attendees can be used to guide STI prevention programme needs, and may provide early warning of potential changes in HIV transmission in the general population.
What it measures	The percentage of pregnant women attending antenatal clinics with a positive (reactive) syphilis serology
Numerator	Number of antenatal care attendees who tested positive for syphilis
Denominator	Number of antenatal care attendees who were tested for syphilis
How to Measure and Measurement Tools	How to measure: Syphilis positivity can be measured using either non-treponemal tests (e.g., RPR or VDRL), treponemal tests (e.g. TPHA, TPPA, EIA, or a variety of available rapid tests), or ideally a combination of both. A reactive non-treponemal test, particularly if the titre is high, is suggestive of active infection, whereas positivity with a treponemal test indicates any previous infection even if treated successfully. For the purposes of this indicator (intended to measure seropositivity), it is acceptable to report positivity based on a single test result. If both treponemal and non-treponemal test results on an individual patient are available, then syphilis positivity should be defined as having positive results on both tests. Use of rapid treponemal test has allowed syphilis testing to occur in settings without laboratory capacity, greatly increasing the number of women who can be tested and treated for syphilis in pregnancy. Data should be collected annually.
	Measurement tools: National programme records aggregated from health facility data, sentinel surveillance, or special surveys, using serologic tests to detect reaginic and/or treponemal antibody may be used. Please specify the source and coverage of your data (for example, sentinel surveillance of all ANC attendees in 2 of 10 provinces) as well as what test type is generally used in your country in the "Comments" section (e.g., non-treponemal (RPR, VDRL), treponemal (rapid tests, TPPA), patients positive on both, or unknown).
Disaggregation	Age groups: Total, 1524 years, 25 years and over

Strengths and weaknesses	Strengths: Data on syphilis positivity in pregnant women are available in most countries through routine health system reporting. Weaknesses: Differences in test type used or changes in testing practices may affect data. Knowledge of testing practices within the country (e.g., proportion of treponemal vs. non-treponemal testing used) should be used to assist with interpretation of disease trends.
Additional considerations	Countries are encouraged to use unique identifiers or registers that separate first and subsequent tests so that the data reflect syphilis true prevalence or incidence rather than test positivity. Since most countries will have data from a variety of test types, sub-analysis (disaggregation) in 15 to 24 year old women may increase the likelihood that test positivity reflects recent infection.
Data utilization	Global/regional: Estimate perinatal mortality and morbidity caused by syphilis that could be averted with effective programmes to eliminate MTCT of syphilis. Identify areas at greatest need of comprehensive congenital syphilis prevention interventions. Local: Follow trends over time to assess changes in burden of disease and STI prevention programme needs. All levels: Compare data on trends of syphilis and HIV to look for early warning of increased risk of HIV transmission.
Data Quality Control and Notes for the Reporting Tool	Please comment on if the data you are providing is routine programme data, if it is felt to be representative of the entire country, and what test type was used to define positivity (e.g., non-treponemal, treponemal, patients positive on both, or mixed/unknown).
Other References	Recommended indicator in "National-Level Monitoring of the Achievement of Universal Access to Reproductive Health: Conceptual and practical considerations and related indicators" and "Methods for Surveillance and Monitoring of Congenital Syphilis Elimination within Existing Systems".

1.17.3 STIs: Percentage of antenatal care attendees positive for syphilis who received treatment	
Rationale	Treatment of antenatal care attendees positive for syphilis is a direct measure of the elimination of mother-to-child transmission of syphilis programme efforts and efforts to strengthen primary HIV prevention.
What it measures	Percentage of antenatal care attendees during a specified period with a positive syphilis serology who were treated adequately.
Numerator	Number of antenatal care attendees with a positive syphilis serology who received at least one dose of benzathine penicillin 2.4 mU IM
Denominator	Number of antenatal care attendees with a positive syphilis serology

How to Measure and Measurement Tools	How to measure: Data should be collected annually. Seropositivity on either treponemal or non-treponemal test is sufficient for being considered positive for syphilis for this indicator.
	Measurement tools: Ideally national programme records aggregated from health facility data should be used. However, if national programme data are not available, data from sentinel surveillance or special studies can be reported if it is felt to be representative of the national situation. Please specify the source and coverage of your data (for example, national programme data from all 12 provinces) in the "Comments" section.
Disaggregation	None
Strengths and weaknesses	Strengths: Data on treatment of syphilis in antenatal care attendees is often routinely monitored in health facilities.
	Weaknesses: Collection of treatment data may require collaboration with MCH programmes to ensure that it is available at a national level.
Additional considerations	For purposes of this indicator, documentation of a single dose of penicillin is sufficient. Treatment of a pregnant woman positive for syphilis with a single injection of 2.4 mU benzathine penicillin prior to 24 weeks gestational age is sufficient to prevent transmission of syphilis from mother to infant. However, three injections spaced at weekly intervals are recommended to treat latent syphilis and prevent tertiary syphilis in the mother.
Data utilization	Global/regional/local: Estimate programme effectiveness in reducing syphilis-associated perinatal morbidity and mortality.
	Local: Identify areas in need of assistance with programme implementation or additional resources.
	All levels: Knowledge of treatment policies and practices should be used to assist with interpretation of trends in treatment.
Data Quality Control and Notes for the Reporting Tool	If the data you are providing does not cover the entire country, please comment.
Other References	Recommended indicator in "National-Level Monitoring of the Achievement of Universal Access to Reproductive Health: Conceptual and practical considerations and related indicators"; recommended indicator in "Methods for Surveillance and Monitoring of Congenital Syphilis Elimination within Existing Systems".

1.17.4 STIs: Percentage of sex workers (SWs) with active syphilis	
Rationale	Testing sex workers (SWs) for syphilis is important for their health, and for second generation surveillance purposes.
What it measures	Progress in decreasing high-risk sexual behaviour, and intervention efforts to control syphilis among sex workers.

Numerator	Number of sex workers who tested positive for active syphilis
Denominator	Number of sex workers who were tested for active syphilis
How to Measure and Measurement Tools	Measurement tools: Data from routine health information systems, sentinel surveillance or special surveys may be used.
	How to measure: The traditional approach to determining seroprevalence has been to screen with a non-treponemal test that measures reaginic antibody (e.g., VDRL or RPR) and confirm positive results with a treponemal test that measures treponemal antibody (e.g., TPHA, TPPA, EIA, or rapid treponemal test). Newer, rapid treponemal tests are comparatively easy to use, a feature which encourages the use of these tests for screening, ideally paired with a non-treponemal test that detects reaginic antibody. Whichever approach is used, the proposed indicator requires both a positive non-treponemal test AND a positive treponemal test to give a proxy for active infection.
	Just a non-treponemal test, or just a treponemal test, while useful in some situations for therapeutic purposes, is not sufficiently specific for surveillance of sex workers. The requirement for both a positive non-treponemal test and a positive treponemal test in sex workers differs from the indicator on syphilis testing in antenatal care attendees because sex workers are more likely to have a history of previous infection. A positive treponemal test measures lifetime exposure, whereas the non-treponemal test is a better indicator of active infection.
Disaggregation	Sex: total, male, female
Strengths and weaknesses	Strengths: Requiring testing by both tests enhances specificity of the reported numbers of positive tests. In addition, requiring testing by both tests will increase the likelihood of identifying active disease. Weaknesses: Requiring testing by both tests increases the difficulty of acquiring data for this indicator.
Additional considerations	Quality assurance and quality control should be an integral part of syphilis testing to ensure reliable results.
Data utilization	Look at trends in comparable groups over time. Compare with data on trends of syphilis and HIV where available.
Data Quality Control and Notes for the Reporting Tool	Please describe in "Comments" what type of sex workers the data represent and what setting the data were collected in. It is important NOT to count multiple tests run on the same patient. That is, if a person has been tested more than once in the past 12 months, they should not be counted more than once.

1.17.5 STIs: Percentage	of men who have sex with men with active syphilis
Rationale	Testing of syphilis among men who have sex with men is important for their health, and for second generation surveillance purposes.
What it measures	Progress in decreasing high-risk sexual behaviour, and intervention efforts to control syphilis among men who have sex with men.
Numerator	Number of men who have sex with men who tested positive for syphilis
Denominator	Number of men who have sex with men who were tested for syphilis
How to Measure and Measurement Tools	Measurement tools: Routine health information systems, sentinel surveillance or special surveys. How to measure: The traditional approach to determining seroprevalence has been to screen with a non-treponemal test that measures reaginic antibody (e.g., VDRL or RPR) and confirm positive results with a treponemal test that measures treponemal antibody (e.g., TPHA, TPPA, EIA, or rapid treponemal test). Newer, rapid treponemal tests are comparatively easy to use, which encourages the use of these tests for screening, ideally paired with a non-treponemal test that detects reaginic antibody. Whichever approach is used, the proposed indicator requires both a positive non-treponemal test AND a positive treponemal test to give a proxy for active infection. Just a non-treponemal test, or just a treponemal test, while useful in some situations for therapeutic purposes, is not sufficiently specific for surveillance of men who have sex with men. The requirement for both a positive non-treponemal test and a positive treponemal test in men who have sex with men differs from the indicator on syphilis testing in antenatal care attendees because men who have sex with men are more likely to have a history of previous infection. A positive treponemal test measures
	lifetime exposure, whereas the non-treponemal test is a better indicator of active infection.
Disaggregation	None
Strengths and weaknesses	Strengths: Requiring testing by both tests enhances specificity of the reported numbers of positive tests. In addition, requiring testing by both tests will increase the likelihood of identifying active disease. Weaknesses: Requiring testing by both tests increases the difficulty of acquiring data for this indicator.
Additional considerations	Quality assurance and quality control should be an integral part of syphilis testing to ensure reliable results.
Data utilization	Look at trends in comparable groups over time. Compare with data on trends of syphilis and HIV where available.
Data Quality Control and Notes for the Reporting Tool	It is important NOT to count multiple tests run on the same patient. That is, if a person has been tested more than once in the past 12 months, they should not be counted more than once. Please describe in "Comments" what setting the data were collected in.

1.17.6 STIs: Number of adults reported with syphilis (primary/secondary and latent/unknown) in the past 12 months	
Rationale	Infection with an acute bacterial STI such as primary/secondary syphilis is a marker of unprotected sexual intercourse and facilitates HIV transmission and acquisition. Therefore, surveillance for primary/secondary syphilis contributes to second-generation HIV surveillance through providing early warning of the epidemic potential of HIV from sexual transmission and on-going high-risk sexual activity that may need more aggressive programme interventions to reduce risk. Furthermore, untreated syphilis causes stillbirths and neonatal disease, and can progress to debilitating or fatal outcomes in adults.
What it measures	Progress in reducing unprotected sex in the general population.
Numerator	Number of adults reported with syphilis during the reporting period
Denominator	Number of individuals aged 15 and older
How to Measure and Measurement Tools	Routine health information systems
Disaggregation	Sex, Primary/secondary vs. latent/unknown: Total, Total Female, Total Male, Female primary/secondary, Male primary/secondary
Strengths and weaknesses	Although WHO has provided a global case definition, actual case definition may vary between and within countries. Furthermore, diagnostic capacity may vary between and within countries. Although underreporting of this indicator may occur, in the absence of changes in case definition or major changes in screening practices, these data can generally be used for following trends over time within a country.
Additional considerations	It is important that countries when reporting on syphilis communicate on the extent to which the data are felt to be representative of the national population. If a country is unable to report on the denominator, WHO will use denominator per UNPD.
Data utilization	Look at trends in comparable groups over time.
Data Quality Control and Notes for the Reporting Tool	Recommended indicator in: "Strategies and laboratory methods for strengthening surveillance of sexually transmitted infection 2012"

1.17.7 STIs: Number of reported congenital syphilis cases (live births and stillbirth) in the past 12 months	
Rationale	Untreated syphilis infection in pregnancy can not only increase risk of HIV transmission and acquisition in the mother and the infant, but also lead to stillbirth, neonatal death, and congenital disease (collectively defined as "congenital syphilis"). Given the high efficacy, simplicity, and low cost of syphilis testing and treatment, global and regional initiatives to eliminate mother-to-child transmission (MTCT) of syphilis have been launched. The rate of congenital syphilis is a measure of the impact of programmatic interventions to eliminate MTCT of syphilis.
What it measures	Progress in elimination of MTCT of syphilis.
Numerator	Number of reported congenital syphilis cases (live births and stillbirths) in the past 12 months
Denominator	Number of live births
How to Measure and Measurement Tools	Routine health information systems
Disaggregation	None
Strengths and weaknesses	Diagnosis of congenital syphilis is most reliable when using specific diagnostic tests that are seldom available even in developed countries. Therefore, in most countries diagnosis of congenital syphilis relies on clinical history and examination, making surveillance challenging. Although WHO has a global case definition for surveillance purposes, actual case definition may vary between and within countries and regions.
Additional considerations	It is important that countries when reporting on syphilis communicate on the extent to which the data are felt to be representative of the national population. If a country is unable to report on the denominator, WHO will use denominator per UNPD.
Data utilization	Given the difficulties in diagnosing congenital syphilis, and depending on the case definition used, either underreporting or overreporting can be a problem. The likely magnitude of such reporting errors should always be considered when looking at rates of congenital syphilis. However, with use of a consistent case definition, trends over time may be useful.
Data Quality Control and Notes for the Reporting Tool	Countries should comment on any major differences between the national case definition and the global surveillance case definition (available on page 15 of: http://www.who.int/reproductivehealth/publications/rtis/9789241505895/en/index.html). Recommended indicator in "Methods for Surveillance and Monitoring of Congenital Syphilis Elimination within Existing Systems".

1.17.8 STIs: Number of	1.17.8 STIs: Number of men reported with gonorrhoea in the past 12 months	
Rationale	Infection with an acute bacterial STI such as gonorrhoea is a marker of unprotected sexual intercourse and facilitates HIV transmission and acquisition. Therefore, surveillance for gonorrhoea contributes to second-generation HIV surveillance through providing early warning of the epidemic potential of HIV from sexual transmission and on-going high-risk sexual activity that may need more aggressive programme interventions to reduce risk. Furthermore, untreated gonorrhoea can result in pelvic inflammatory disease, ectopic pregnancy, infertility, blindness, and disseminated disease. Increasing resistance to currently recommended treatment options may render this infection untreatable.	
What it measures	Progress in reducing unprotected sex in men.	
Numerator	Number of men reported with gonorrhoea during the reporting period	
Denominator	Number of males aged 15 and older	
How to Measure and Measurement Tools	Routine health information systems	
Disaggregation	None	
Strengths and weaknesses	Although WHO has provided a global case definition, actual case definition may vary between and within countries. Furthermore, diagnostic capacity may vary between and within countries. Although underreporting of this indicator may occur, in the absence of changes in case definition or major changes in screening practices, these data can generally be used for following trends over time within a country.	
Additional considerations	It is important that countries when reporting on gonorrhoea communicate on the extent to which the data are felt to be representative of the national population. Data on gonorrhoea among women, although useful for monitoring purposes at a local and national level, are not requested at the global level because the majority of women infected with <i>Neisseria gonorrhoeae</i> are asymptomatic and sensitive diagnostic tests for gonorrhoea in women are not widely available in developing countries. Therefore data on gonorrhoea among women are felt to be too dependent on diagnostic resources and screening practices to be monitored appropriately at the global level. If a country is unable to report on the denominator, WHO will use	
Data utilization	denominator per UNPD.	
Data utilization	Look at trends in comparable groups over time.	
Data Quality Control and Notes for the Reporting Tool	Recommended indicator in: "Strategies and laboratory methods for strengthening surveillance of sexually transmitted infection 2012"	

1.17.9 STIs: Number of	men reported with urethral discharge in the past 12 months
Rationale	Urethral discharge in men is an STI syndrome generally most commonly caused by <i>Neisseria gonorrhoeae</i> or <i>Chlamydia trachomatis</i> . Presentation with an acute STI syndrome such as urethral discharge is a marker of unprotected sexual intercourse and urethral discharge facilitates HIV transmission and acquisition. Therefore, surveillance for urethral discharge contributes to second-generation HIV surveillance through providing early warning of the epidemic potential of HIV from sexual transmission and on-going high-risk sexual activity that may need more aggressive programme interventions to reduce risk. Furthermore, untreated urethral discharge can result in infertility, blindness, and disseminated disease. Increasing resistance to currently recommended treatment options for <i>Neisseria gonorrhoeae</i> may render this infection untreatable.
What it measures	Progress in reducing unprotected sex in men.
Numerator	Number of men reported with urethral discharge during the reporting period
Denominator	Number of males aged 15 and older
How to Measure and Measurement Tools	Routine health information systems.
Disaggregation	None
Strengths and weaknesses	Although WHO has provided a global case definition, actual case definition may vary between and within countries. Furthermore, clinical diagnostic capacity may vary between and within countries. Although underreporting of this indicator may occur, in the absence of changes in case definition or major changes in screening practices, these data can generally be used for following trends over time within a country.
Additional considerations	It is important that countries when reporting on urethral discharge communicate on the extent to which the data are felt to be representative of the national population. Following trends in urethral discharge is a feasible means to monitor incident STI in a population. Data on vaginal discharge among women, although useful for monitoring purposes at a local and national level, are not requested at the global level because in many settings the majority of vaginal discharge cases are not due to sexually transmitted infections. Countries should conduct periodic assessments of the etiology of urethral discharge syndrome in order to understand the predominant causes of urethral discharge and therefore appropriate therapy. If a country is unable to report on the denominator, WHO will use denominator per UNPD.
Data utilization	Look at trends in comparable groups over time.
Data Quality Control and Notes for the Reporting Tool	Recommended indicator in: "Strategies and laboratory methods for strengthening surveillance of sexually transmitted infection 2012"

1.17.10 STIs: Number of adults reported with genital ulcer disease in the past 12 months	
Rationale	Genital ulcer disease is an STI syndrome generally most commonly caused by syphilis, chancroid, or herpes simplex virus. Presentation with an acute STI syndrome such genital ulcer disease is a marker of unprotected sexual intercourse and facilitates HIV transmission and acquisition. Therefore, surveillance for genital ulcer disease contributes to second-generation HIV surveillance through providing early warning of the epidemic potential of HIV from sexual transmission and on-going high-risk sexual activity that may need more aggressive programme interventions to reduce risk. Furthermore, untreated genital ulcer diseases can cause stillbirths and neonatal disease, and can progress to debilitating or fatal outcomes in adults.
What it measures	Progress in reducing unprotected sex in the general population.
Numerator	Number of adults reported with genital ulcer disease during the reporting period
Denominator	Number of individuals aged 15 and older
How to Measure and Measurement Tools	Routine health information systems
Disaggregation	Sex: total, men, women
Strengths and weaknesses	Although WHO has provided a global case definition, actual case definition may vary between and within countries. Furthermore, clinical diagnostic capacity may vary between and within countries. Although underreporting of this indicator may occur, in the absence of changes in case definition or major changes in screening practices, these data can generally be used for following trends over time within a country.
Additional considerations	It is important that countries when reporting on genital ulcer disease communicate on the extent to which the data are felt to be representative of the national population.
	Countries should conduct periodic assessments of the etiology of genital ulcer disease in order to ensure appropriate drug selection for syndromic management and to understand the extent to which genital ulcer disease reflects incident infection due to recurrent HSV infection versus acute infection with syphilis, chancroid, or HSV.
	If a country is unable to report on the denominator, WHO will use denominator per UNPD.
Data utilization	Look at trends in comparable groups over time.
Data Quality Control and Notes for the Reporting Tool	Recommended indicator in: "Strategies and laboratory methods for strengthening surveillance of sexually transmitted infection 2012"

1.18 Percentage (%) of pregnant women with a positive syphilis serology whose sexual contacts were identified and treated for syphilis	
Numerator	The number of pregnant women who tested positive for syphilis and whose sexual contacts were identified and treated.
	This numerator calls for providing counselling for each pregnant woman and identifying all her sexual contacts. Only if all her reported sexual partners are being treated, can that woman be included in this numerator.
Denominator	Number of pregnant women who tested positive for syphilis during pregnancy.

Target 2: Reduce transmission of HIV among people who inject drugs by 50% by 2015

2.6 Number of people on opioid substitution therapy (OST)	
Rationale	Opioid substitution therapy represents a commitment to treat opioid dependence and to reduce the frequency of injecting, preferably to zero. OST is the most effective public health tool for reducing injecting drug use among opioid injectors. OST also provides a crucial support for the treatment of other health conditions, including HIV, TB and viral hepatitis.
What it measures	National commitment and progress towards the treatment of opioid dependence and reduction of HIV transmission probabilities among people who inject drugs.
How to Measure and Measurement Tools	Programme data, census data
Disaggregation	Administrative units: urban, rural
Strengths and weaknesses	Number of people on OST should be readily available and valid since they are typically licensed by the relevant authorities.
Additional considerations	Please refer to the WHO/UNODC/UNAIDS Technical Guide for countries to set targets for universal access to HIV prevention, treatment and care for injecting drug users (http://www.who.int/hiv/topics/idu/en/index.html) for a proposed complete set of globally agreed indicators for people who inject drugs.
Data utilization	Try to assess whether sufficient OSTs are available for the number and distribution of people who are dependent on opioids in the country.
Other References	WHO/UNODC/UNAIDS Technical Guide for countries to set targets for universal access to HIV prevention, treatment and care for injecting drug users (http://www.who.int/hiv/topics/idu/en/index.html)

2.7 Number of NSP and OST sites: - Number of needle and syringe programme (NSP) sites	
Rationale	Needle and syringe distribution programmes are among the most effective interventions for preventing transmission of HIV among people who inject drugs. Sufficient access to clean needles for the injecting population is measured with this indicator.
What it measures	Number of NSP sites (including pharmacy sites providing at no cost needles and syringes). Availability of sites that can provide clean needles and syringes to injection drug users.
How to Measure and Measurement Tools	National programme data

Disaggregation	Administrative unit
	Urban, rural
Strengths and weaknesses	Many NSPs are not "official" and therefore not counted among national programme data
Additional considerations	Needle and syringe programmes (NSPs) are any programmes that include access to clean equipment and safe disposal through fixed or mobile exchange programmes and/or through pharmacies where equipment is available free of charge. In many countries pharmacy sales of injecting equipment are an important and sometimes the most significant source of clean injecting equipment accessible to drug users. However, pharmacies that sell needles and syringes are typically not counted in a retrievable database as part of a public health or harm reduction programme. If they are available, they should be counted and highlighted, if possible. Pharmacies that distribute needles and syringes free of cost typically do maintain records of needles distributed as part of the programme and should be included.
	Please refer to the WHO/UNODC/UNAIDS Technical Guide for countries to set targets for universal access to HIV prevention, treatment and care for injecting drug users (http://www.who.int/hiv/topics/idu/en/index.html) for a proposed complete set of globally agreed indicators for people who inject drugs.
Data utilization	Get an idea of the availability of NSP sites, and trends over time. Also try to analyse data based on geographical location of the NSP sites and geographical distribution and population density of people who inject drugs in the country. Try to assess whether sufficient NSPs are available for the number and distribution of people who inject drugs in the country.
Data Quality Control and Notes for the Reporting Tool	National Representativeness: Many NSP sites are not "official" and may be run by NGOs, which the government may not have information on. Please try to assess the national representativeness of the number you are reporting.
Other References	WHO/UNODC/UNAIDS Technical Guide for countries to set targets for universal access to HIV prevention, treatment and care for injecting drug users (http://www.who.int/hiv/topics/idu/en/index.html)

	2.7 Number of NSP and OST sites: - Number of opioid substitution therapy (OSP) sites	
Rationale	Opioid substitution therapy represents a commitment to treat opiate users and to reduce the frequency of injection, preferably to zero. OST is the single most effective public health tool for reducing injection drug use.	
What it measures	National commitment and progress towards the treatment of opiate users and reduction of HIV transmission probabilities among people who inject drugs. The number of OST sites and the availability of sites that can provide OST to injecting drug users.	
How to Measure and Measurement Tools	National programme data	
Disaggregation	Administrative unit Urban, rural	
Strengths and weaknesses	OST sites should be readily available and valid since they are typically licensed by the relevant authorities. However, the number of sites does not indicate the number of slots that may be available.	
	Obtaining subgroup population size estimates will be difficult and add extra uncertainty.	
Additional considerations	Please refer to the WHO/UNODC/UNAIDS Technical Guide for countries to set targets for universal access to HIV prevention, treatment and care for injecting drug users (http://www.who.int/hiv/topics/idu/en/index.html) for a complete set of globally agreed indicators for people who inject drugs.	
Data utilization	Get an idea of the availability of OST sites and trends over time in relation to the population size of opiate injectors in the country. Also try to analyse data based on geographical location of the OST sites and geographical distribution and population density of people who inject opioid drugs in the country. If possible, try to interpret this indicator considering information available on the number of OST slots in various sites. Try to assess whether sufficient OSTs are available for the number and distribution of opiate injectors in the country.	
Data Quality Control and Notes for the Reporting Tool	National Representativeness: Many OST sites are not "official" and may be run by NGOs, which the government may not have information on. Please try to assess the national representativeness of the number you are reporting.	
Other References	WHO/UNODC/UNAIDS Technical Guide for countries to set targets for universal access to HIV prevention, treatment and care for injecting drug users (http://www.who.int/hiv/pub/idu/idu_target_setting_guide.pdf)	

Target 3: Eliminate mother-to-child transmission on HIV by 2015 and substantially reduce AIDS-related maternal deaths

(tested for HIV an	egnant women who know their HIV status and received their results - during pregnancy, during labour and delivery, best-partum period (<72 hours), including those with previously known
Rationale	Identification of a pregnant woman's HIV serological status provides an entry point for other services for PMTCT and to tailor prevention, care and treatment to her needs.
What it measures	This indicator assesses efforts to identify the HIV serological status of pregnant women in the previous 12 months.
Numerator	Number of pregnant women of known HIV status. This is compiled from the number of women of unknown HIV serological status attending antenatal care, labour and delivery and postpartum services, who have been tested for HIV and know their results and women with known HIV infection attending antenatal care for a new pregnancy in the past 12 months.
	Pregnant women with known HIV infection: women who were tested and confirmed to be HIV-positive at any time before the current pregnancy, who are attending antenatal care for a new pregnancy. These women may not need to be retested if there is documented proof of their positive status ²³ , and in line with national guidelines on testing pregnant women. These women do, however, need services for PMTCT and are counted in the numerator.
	Pregnant (and postpartum) women of unknown serological status: women who were not tested during antenatal care or at labour and delivery for this pregnancy or do not have documented proof of having been tested during this pregnancy.
	The numerator is the sum of categories a-c below:
	(a-1) pregnant women who have an HIV test and receive their result during antenatal care;
	(a-2) pregnant women with known HIV infection attending antenatal care for a new pregnancy;
	(b) pregnant women of unknown HIV serological status attending labour and delivery who were tested and received results; and
	(c) women of unknown HIV serological status attending postpartum services within 72 hours of delivery who were tested and received results.
	Categories a-1, b and c include all women who were tested and received results, irrespective of the HIV test result. Category a-2 includes women with previously known HIV-positive status.
	Data reported from facilities may be disaggregated into:
	(a) women with known (positive) HIV infection at antenatal care;
	(b) women newly identified as HIV positive; and(c) women testing HIV negative (the remainder).See below for Disaggregation for Global Reporting.
Denominator	Estimated number of pregnant women in the past 12 months

^{23.} Documentation of HIV infection (care and treatment card, maternal card from previous pregnancy or other reliable written documentation of HIV status) is generally required in most settings. Without proof of existing HIV infection, women are usually considered as being of 'unknown' status and are often retested. National guidelines should be consulted.

How to Measure and Measurement Tools	The numerator is calculated from national programme records aggregated from facility registers for antenatal care, labour and delivery and postpartum care. In countries with high rates of facility attendance for labour and delivery, data can be collected from labour and delivery registers only, as the results of HIV testing will be available for most pregnant women from this one source. Health facility registers should record known HIV infection in pregnant women coming to antenatal care clinics for a new pregnancy, so that they receive services for PMTCT. All public, private and nongovernmental organization-run health facilities that are providing testing and counselling for pregnant women should be included. The denominator is derived from a population estimate of the number of pregnant women giving birth in the past 12 months. This can be obtained from estimates of births from the central statistics office or from the United Nations Population Division or pregnancy registration systems with complete data.
Disaggregation	Pregnancy stages: ANC, L&D, postpartum Receipt of results: tested, tested and received results HIV serostatus: number HIV+
Strengths and weaknesses	This indicator enables a country to monitor trends in HIV testing among pregnant women. The points at which drop-outs occur during the testing and counselling process and the reasons why they occur are not captured by this indicator. This indicator does not measure the quality of the testing or counselling. It also does not capture the number of women who received pre-test counselling.
Additional considerations for countries	Health facility registers should reflect known HIV infection among HIV-infected pregnant women coming to the ANC for a new pregnancy (even if they are not tested at that site), such as through a code, circle, or other method, in order for them to receive subsequent PMTCT interventions. Not all categories will be applicable or significant to all settings (e.g. women of unknown status tested within 72 hours postpartum). Countries may want to prioritize investment of resources (revision of tools, time, money) for measuring the categories that are appropriate to their country context. It may be important for programme managers to use additional sub-national and facility level indicators to measure trends and progress in the testing and counselling process, such as uptake of testing and receipt of results. It is also important to know the number of women whose HIV status has been identified at each service, i.e. % ANC attendees whose HIV status is known; % L&D attendees whose HIV status is known, etc. This indicator could be triangulated and validated using population-based surveys, such as the DHS, which generally occurs every five years, or the AIDS Indicator Survey, a population-based survey that can be done on a more periodic basis.

Data utilization	Look at trends over time. If disaggregated data is available by region, see whether any lower performing areas can be identified. Review if data is available on % of ANC attendees who know their status (including those with previously confirmed HIV status and those tested) and % of L&D attendees who know their status.
Data Quality Control and Notes for the Reporting Tool	Double Reporting: There is a risk of double counting with this indicator, as a pregnant woman can be tested a few times during ANC, L&D, or postpartum. This is particularly true where women get re-tested in different facilities, or where they come to the L&D without documentation of their test. While not feasible to avoid double counting entirely, countries should ensure a data collection and reporting system is in place to minimize it, such as using patient held and facility held ANC records to document that testing took place.
	Please do not add <i>all</i> the number of women tested from ANC <i>and</i> L&D to get the <i>total</i> number of women tested. We are interested in knowing the number of <i>women</i> tested, and not the total number of tests (i.e. if a women is tested at ANC and again at L&D, try to only count her once). It is important to include those with previously known HIV infection in the numerator – even if they do not receive an HIV test, their HIV infection is identified for subsequent PMTCT interventions.
	Number tested, as well as tested and received results: If available, please report the number of pregnant women tested, as well as the number of pregnant women tested and received results (latter should not exceed the former).
	If your data collection system does not currently separate those with known and unknown HIV status and you are unable to provide the specific disaggregated data, please review the data available, and derive the best data for the number of <i>pregnant women whose HIV status has been identified</i> during pregnancy, L&D, or during the post-partum period within 72 hours.
	Please provide any details that would help to interpret your data in the Comment section.
	Please comment on the source of your denominator.
Other References	PMTCT M&E Core Indicator #3

3.5 Percentage of pregnant women attending antenatal care (ANC) whose male partner was tested for HIV in the last 12 months	
Rationale	Male involvement is a critical element in providing family-focused services to HIV-infected pregnant mothers, their infants and family members. It is also important in the prevention of HIV infection and can help couples who are seronegative to remain seronegative. Partner testing is the first step in involving the male partner, regardless of the couple's HIV status.
What it measures	The percentage of pregnant women attending antenatal care whose male partner was tested during their female partner's pregnancy in the past 12 months.
Numerator	Number of pregnant women attending antenatal care whose male partner was tested in the last 12 months
Denominator	Number of pregnant women attending antenatal care
How to Measure and Measurement Tools	The numerator can be calculated from national programme records compiled from facility registers. Male partners can be tested with the woman at the first antenatal care visit or at a follow-up visit or tested alone on a separate visit, such as a day reserved for male partner testing. Data can be aggregated from antenatal care or testing and counselling register, depending on the context. All public, private and nongovernmental organization-run health facilities that provide antenatal care services should be included. If feasible, programmes may consider collecting data on whether or not
	the male and female partner disclosed their HIV status to each other in the presence of a clinic staff member.
Strengths and weaknesses	This indicator allows countries to monitor efforts at increasing testing of male partners of pregnant women attending ANC services. It does not measure whether the male partner received his result or any follow-up services. The indicator does not take into account ANC clients that have more than one partner or that may change partners over time. It also may not include partners that received HIV testing at non-ANC settings and which are not linked to ANC (e.g. general VCT or provider initiated testing). Not all sites may be collecting data on male partner testing or routinely aggregating and reporting the data. Measuring this indicator may require additional investment and resources to revise data collection tools and summary reporting forms.
Additional considerations	Although testing male partners is an important tool for increasing male involvement and preventing infection during pregnancy, it is also a critical entry point into on-going and family-focused care for the man. Health providers should ensure and document that appropriate follow-up services are provided to all male partners who test HIV-positive, as part of a comprehensive care and treatment programme.

Data utilization	Interpret based on country context and applicability. Discuss how to increase coverage.
Data Quality Control and Notes for the Reporting Tool	Please provide any comments that would help to interpret the representativeness of the data. If the number of discordant couples is easily available, please provide data in the comments section with any supporting comments.
Other References	PMTCT Additional Indicator # A-3

3.6 Percentage of HIV-infected pregnant women assessed for ART eligibility through either clinical staging or CD4 testing	
Rationale	HIV-infected pregnant women who meet the clinical and (when available) immunological criteria for antiretroviral therapy should receive it. Antiretroviral therapy preserves maternal health and reduces the risk for mother-to-child transmission. Services for the prevention of mother-to-child transmission of HIV should undertake such assessments. Women who are not yet eligible for antiretroviral therapy should receive antiretroviral drug prophylaxis for PMTCT according to the national guidelines and recommendations.
What it measures	Coverage of eligibility assessment for antiretroviral therapy among HIV-infected pregnant women, either clinically by WHO clinical staging criteria or immunologically by CD4 testing. Assessments can be made on site or by referral.
Numerator	Number of HIV-infected pregnant women assessed for eligibility for antiretroviral therapy by either clinical staging or CD4 testing, on site or by referral, in the past 12 months. 'On site' means that the service is offered in a health facility structure or compound. For instance, HIV clinical staging may be available in the antenatal care unit, while blood draw for CD4 testing is available at the HIV care and treatment unit in the same health facility. Both these services are considered to be on site. Referral can be made on site or off site and is defined as sending a patient to a different service unit, health provider or health facility. Often, patients return to the original health facility, service unit or provider, where the services received at the referral site are fed back to the original site, and the patient continues with follow-up care. Referral facilities should document the services provided and patient outcomes. This indicator should be disaggregated by type of assessment (clinical staging or CD4 testing). Women who were assessed by CD4 testing and clinical staging should be counted only once as having been assessed by CD4 testing.
Denominator	Estimated number of HIV-infected pregnant women in the past 12 months

How to Measure and Measurement Tools

The numerator is calculated from national programme records aggregated from facility registers.

Assessment can be conducted in antenatal care clinics and HIV care and treatment units, on site or by referral. Data should be aggregated from the appropriate register, with consideration of which registers capture the data, where the assessment actually took place, possible double-counting or under-counting and the need for accurate data for the national level.

All public, private and nongovernmental organization-run health facilities that assess eligibility of HIV-infected pregnant women for antiretroviral therapy, either on site or by referral, should be included

Two methods can be used to calculate the denominator:

a projection model such as that provided by Spectrum software: use the output "number of pregnant woman needing prevention of mother-to-child transmission of HIV"; or

multiply the number of women who gave birth in the past 12 months (which can be obtained from estimates of the central statistics office or the United Nations Population Division or pregnancy registration systems with complete data) by the most recent national estimate of HIV prevalence in pregnant women (which can be derived from HIV sentinel surveillance in antenatal care clinics), if Spectrum projections are unavailable.

Disaggregation

Method of ART eligibility assessment: Clinical staging, CD4 testing

Strengths and weaknesses

The strength of this indicator is that it enables countries to monitor the extent to which HIV-infected pregnant women are receiving an intervention that is critical for accessing ART for their own health.

It does not capture whether HIV-infected pregnant women who were eligible for ART actually received it.

Although each category is mutually exclusive, there is a risk of double counting this indicator where HIV-infected pregnant women have been assessed both clinically and immunologically, as well as where women are assessed in different units or in a different facility. Countries should ensure systems are in place to minimize the risk of double counting.

This indicator does not capture women who may have been identified HIV-positive at labour and delivery and subsequently assessed for ART eligibility.

Additional considerations

It is recommended that countries disaggregate by eligibility status for additional information on national trends in the percentage of pregnant women who are eligible for ART.

In settings where HIV-infected pregnant women are referred out to another health facility or another service unit within the same health facility, health providers should make an effort to document referrals made and services received for these women in the ANC/PMTCT register for better patient tracking and monitoring of HIV-infected pregnant women.

Data utilization	The goal would be to aim for 100%; once 100% is reached routinely, this indicator may become obsolete. Explore further information on disaggregated data on whether eligibility was assessed through clinical staging or CD4 tests and any data available on how long it takes to receive a CD4 test result in various places.
Data Quality Control and Notes for the Reporting Tool	Please provide any comments that would help to interpret the data.
Other References	PMTCT M&E Core Indicator #4

3.7 Percentage of infants born to HIV-infected women provided with antiretroviral (ARV) prophylaxis to reduce the risk of early mother-to-child transmission in the first 6 weeks (i.e. early postpartum transmission around 6 weeks of age)	
Rationale	The risk for mother-to-child transmission can be significantly reduced by the complementary approaches of providing antiretroviral drugs (as treatment or as prophylaxis) for the mother during pregnancy and delivery, with antiretroviral prophylaxis for the infant, and antiretrovirals to the mother or child during breastfeeding (if breastfeeding), and use of safe delivery practices and safer infant feeding.
What it measures	Progress in the prevention of early postpartum mother-to-child transmission by the provision of antiretroviral prophylaxis for HIV-exposed infants
Numerator	Number of infants born to HIV-infected women during the past 12 months who received antiretroviral prophylaxis to reduce early mother-to-child transmission (i.e. early postpartum, in the first 6 weeks).
Denominator	Estimated number of live births to pregnant HIV-infected women in the past 12 months

How to Measure and Measurement Tools

The numerator is calculated from national programme records aggregated from facility registers.

Antiretroviral drugs can be given to HIV-exposed infants shortly after delivery, at facilities for labour and delivery for infants born at facilities, at outpatient postnatal care or child clinics for infants born at home and brought to the facility, or at HIV care and treatment or other sites, depending on the country.

Three methods for calculating the numerator can be considered:

- Counting at the point of antiretroviral drug provision: In settings with low facility delivery rates, data for the numerator should be compiled from the sites where antiretroviral drugs are dispensed and where the data are recorded. There is a risk of double-counting when antiretroviral drugs are provided during more than one visit or at different health facilities. Countries should establish data collection and reporting systems to minimize double-counting.
- Counting around time of delivery: In settings where a high proportion of women give birth in health facilities, countries can estimate the numerator from only the labour and delivery register by counting the number of HIV-exposed infants who received a specific antiretroviral drug regimen before discharge from the labour and delivery ward. This may be the most reliable and accurate method for calculating this indicator in settings with a high proportion of facility deliveries and low follow-up, as the corresponding antiretroviral drug regimen dispensed is counted at the time of provision to the infant.
- Counting at postnatal or child health sites: Countries can also count and aggregate the number of HIV-exposed infants who received antiretroviral prophylaxis recorded at postnatal or child health clinics if attendance is high and the exposure status of the child is likely to be known (e.g. from postnatal registers, stand-alone registers or integrated HIV-exposed infant registers).

All public, private and nongovernmental organization-run health facilities that provide antiretroviral drugs to HIV-exposed infants for the prevention of mother-to-child transmission of HIV should be included.

Two methods can be used to estimate the denominator:

- a projection model, such as that provided by Spectrum software; use the output "number of pregnant woman needing prevention of mother-tochild transmission of HIV" as a proxy; or
- multiply the number of women who gave birth in the past 12 months (which can be obtained from estimates by central statistics office or the United Nations Population Division or pregnancy registration systems with complete data) by the most recent national estimate of HIV prevalence in pregnant women (which can be derived from HIV sentinel surveillance in antenatal care clinics), if Spectrum projections are unavailable.
- If there are data on the number of live births, they should be adjusted to derive a better proxy.

Disaggregation

None requested

Strengths and weaknesses	This indicator allows countries to monitor the coverage of antiretrovirals regimens dispensed or initiated among HIV-exposed infants to reduce the risk of early maternal HIV transmission. The indicator measures the extent to which ARVs were dispensed for infants as prophylaxis. It does not capture whether the ARVs were consumed; thus it is not possible to determine adherence to the ARV regimen, nor whether ARV regimens were completed.
Additional considerations	Countries that have developed mechanisms for reaching HIV-exposed infants at the community level with ARVs will want to ensure a system of data collection is in place for reporting infants receiving ARV regimens at the community level.
Data utilization	Compare the indicator value with coverage of the maternal ARV regimen (Indicator I-10) and discuss what the data may mean in the country context. Some countries may want to explore further and do a linked review of the infant ARV prophylaxis regimen vis-à-vis the maternal ARV regimen can be assessed.
Data Quality Control and Notes for the Tool	Please provide any comments that would help to interpret the data.
Other References	PMTCT M&E Core Indicator #6

	3.9 Percentage of infants born to HIV-infected women started on co-trimoxazole (CTX) prophylaxis within two months of birth	
Rationale	Co-trimoxazole prophylaxis is a simple, cost-effective intervention to prevent <i>Pneumocystis jiroveci</i> pneumonia in HIV-infected infants. This infection is the leading cause of serious respiratory disease in these infants in resource-constrained countries and often occurs before HIV infection can be diagnosed. Owing to resource and logistical constraints in diagnosing HIV infection in young infants, all infants born to HIV-infected women should receive co-trimoxazole prophylaxis, starting 4–6 weeks after birth and continuing until HIV infection has been excluded and the infant is no longer at risk of acquiring HIV through breastfeeding.	
What it measures	The provision and coverage of co-trimoxazole prophylaxis for HIV-exposed infants in line with international guidelines ²⁴	
Numerator	Number of infants born to HIV-infected women started on co-trimoxazole prophylaxis within 2 months of birth in the past 12 months	
Denominator	Estimated number of HIV-infected pregnant women who gave birth in the past 12 months	

^{24.} WHO. Guidelines on co-trimoxazole prophylaxis for HIV-related infections among children, adolescents and adults: Recommendations for a public health approach. Geneva, World Health Organization, 2006.

How to Measure and Measurement Tools

The numerator is calculated from national programme records aggregated from facility registers.

Data should be aggregated from the appropriate facility registers, such as a stand-alone or integrated HIV-exposed infant register. The register used may depend on where services are offered. For example, where HIV-exposed infants are followed by health workers in HIV care and treatment facilities, countries could aggregate information from a register based at that site. All public, private and nongovernmental organization-run health facilities that provide co-trimoxazole prophylaxis for HIV-exposed infants should be included.

Two methods can be used to estimate the denominator:

a projection model such as that provided by Spectrum software; use the output "number of pregnant woman needing PMTCT (prevention of mother-to-child transmission of HIV)" as a proxy; or

multiply the total number of women who gave birth in the past 12 months (which can be obtained from central statistics offices or the United Nations Population Division or pregnancy registration systems with complete data) by the most recent national estimate of HIV prevalence in pregnant women³ (which can be derived from HIV sentinel surveillance in antenatal care clinic), if Spectrum projections are unavailable.

If there are data on the number of live births, they should be adjusted to derive a better proxy.

Disaggregation

None requested

Strengths and weaknesses

This indicator allows countries to monitor progress in the early follow-up of exposed infants by measuring provision of co-trimoxazole in line with international guidelines. It can also be used as a proxy indicator for early follow-up visits of exposed infants within the recommended first 4-6 weeks of life. The indicator captures only those infants who return for HIV-exposed infant follow-up services within two months of birth. It does not measure actual coverage of co-trimoxazole prophylaxis for HIV-exposed infants as some infants may have been started on treatment after 2 months. A low value of the indicator could signal potential bottlenecks in the system, including poor management of CTX supplies in the country, poor data collection, and inadequate distribution systems.

Additional considerations

Countries may also wish to document provision of CTX for HIV-exposed infants older than 2 months as a way to monitor overall progress of the programme, identify existing challenges with early initiation of CTX, and to monitor consumption for procurement needs.

Inappropriate management of supplies can negatively affect the value of the indicator and significantly reduce access to CTX for HIV-exposed infants. Countries should ensure appropriate systems and tools, particularly tools for LMIS, are in place to adequately procure, distribute, and manage supplies at facility, district and central levels.

Data utilization	Data can also be reviewed as an indication of the number of exposed infants who are seen at a facility within 2 months of birth. If indicator value is low, explore reasons why (e.g. whether exposed-infants are not attending facilities within 2 months, or if there are stock-outs of CTX, etc.).
Data Quality Control and Notes for the Reporting Tool	National Representativeness: If this indicator is obtained from a sub-set of facilities, comments should be added regarding the representativeness. Triangulation Options: pharmacy registers If the data reported represents CTX provided in infants beyond 2 months of age, please note it in the Comments section.
Other References	PMTCT M&E Core Indicator #8

	eding practices (exclusive breastfeeding, replacement feeding, mixed infants born to HIV-infected women at DPT3 visit
Rationale	HIV can be transmitted during breastfeeding even in settings where 100% of HIV-infected pregnant women receive either lifelong antiretroviral therapy or antiretroviral medicines as prophylaxis for the prevention of mother-to child transmission of HIV. Mixed feeding before 6 months of age increases the risk for HIV transmission when compared with exclusive breastfeeding. WHO therefore recommends that when mothers known to be HIV-infected breastfeed, they should be given ARVs to reduce the risk of transmission and also exclusively breastfeed for the first 6 months, introduce complementary feeds from 6 months and continue breastfeeding until 12 months of age.
	Coverage with the third dose of diphtheria, pertussis and tetanus vaccine close to the recommended age of 14 weeks is relatively high in most countries. It is proposed to collect data at this time because most infants are seen then and it is mid-way between birth and the time at which exclusive breastfeeding would stop, making it comparable to the way that exclusive breastfeeding is usually reported for the general population in demographic and health surveys.
What it measures	Feeding ²⁵ of HIV-exposed infants, derived from 24-h recall, measured at the time of the third dose of diphtheria, pertussis and tetanus vaccine (DPT3), which is often around 3 months of age or at the closest visit after 3 months.

^{25.} The infant feeding practices measured with this indicator are defined as follows:

Exclusive breastfeeding: An infant receives only breast milk and no other liquids or solids, not even water, with the exception of drops or syrups consisting of vitamins, mineral supplements or medicines, for up to 6 months. Breast milk is defined as including milk from a wet nurse and a mother's expressed milk.

Replacement feeding (no breast milk at all): Feeding an infant who is not receiving any breast milk a diet that provides all the necessary nutrients until the child is fully fed on family foods. During the first 6 months, the food should be a suitable breast-milk substitute, which is usually a commercial infant formula, as home modified animal milk is no longer recommended for feeding infants during the entire first 6 months of life, except as an emergency measure.

Mixed feeding (-partial breastfeeding): Feeding both breast milk and other foods or liquids to infants for 0–6 months.

I12a number of HIV-exposed infants who were exclusively breastfeeding at or around the DPT3 visit; I12b number of HIV-exposed infants who received replacement feeding at or around the DPT3 visit; and I12c number of HIV-exposed infants who received mixed feeding at or around the DPT3 visit. The numerators capture feeding practices only for known HIV-exposed infants who visit a health facility. Denominator The denominator is the same for all three indicators: the number of HIV exposed infants whose feeding practice has been assessed at a DPT3 visit. Infants will be aged around 3 months or more. The numerators are calculated from national programme records aggregated from facility registers. Ideally, data from appropriate sites and registers such as a stand-alone or integrated HIV-exposed infant registers should be aggregated, depending on where the services are and where data are recorded. At each visit, the health-care provider should enquire about infant-feeding practices during the previous 24 hours, by asking: "What did you give your infant to eat or drink yesterday during the day and during the night?" After each response, the health provider should ask: "Anything else?" The response will be recorded as exclusive breastfeeding, replacement feeding or mixed feeding. While this information is collected and recorded on the child health card at every visit, providers should record it in the register only once, during the third visit for diphtheria, pertussis and tetanu vaccination. This record will be used for compilation and reporting to national level. In settings where HIV-exposed infants are seen in HIV care and treatment facilities, data should be collected at a visit when the infant is around 3 months. The denominator is calculated from the total number of exposed infants whose feeding was assessed. Exposed infants who did not attend facilities are not included in the denominator. All public, private and nongovernmental organization-run health facilities that provide HIV-exposed infant follow-up		
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practices during the previous 24 hours, by asking: "What did you give your infant to eat or drink yesterday during the day and during the night?" After each response, the health provider should ask: "Anything else?" The response will be recorded as exclusive breastfeeding, replacement feeding or mixed feeding. While this information is collected and recorded on the child health card at every visit, providers should record it in the register only once, during the third visit for diphtheria, pertussis and tetanu vaccination. This record will be used for compilation and reporting to national level. In settings where HIV-exposed infants are seen in HIV care and treatment facilities, data should be collected at a visit when the infant is around 3 months. The denominator is calculated from the total number of exposed infants whose feeding was assessed. Exposed infants who did not attend facilities are not included in the denominator. All public, private and nongovernmental organization-run health facilities that provide HIV-exposed infant follow-up services should be included.		integrated HIV-exposed infant registers should be aggregated, depending
whose feeding was assessed. Exposed infants who did not attend facilities are not included in the denominator. All public, private and nongovernmental organization-run health facilities that provide HIV-exposed infant follow-up services should be included.		practices during the previous 24 hours, by asking: "What did you give your infant to eat or drink yesterday during the day and during the night?" After each response, the health provider should ask: "Anything else?" The response will be recorded as exclusive breastfeeding, replacement feeding or mixed feeding. While this information is collected and recorded on the child health card at every visit, providers should record it in the register only once, during the third visit for diphtheria, pertussis and tetanus vaccination. This record will be used for compilation and reporting to national level. In settings where HIV-exposed infants are seen in HIV care and treatment facilities, data should be collected at a visit when the infant
that provide HIV-exposed infant follow-up services should be included.		whose feeding was assessed. Exposed infants who did not attend facilities
Disaggregation Report distribution of IF practice: EBF, RF, MF/Other; Uncategorized/other		
	Disaggregation	Report distribution of IF practice: EBF, RF, MF/Other; Uncategorized/other

Strengths The indicators measure important progress in safer infant feeding practices and weaknesses among HIV-infected women and their exposed infants. They can also be used to indicate the quality of infant feeding counselling (with low rates of mixed feeding likely to indicate adequate infant feeding counselling and support), and can also be used to model the impact of the intervention in a country (see Core Indicator 12 in the PMTCT M&E guide, or GARPR 3.3 - modelled MTCT rate). It should be noted that the indicator says nothing about the quality of replacement feeding given, nor the impact of the feeding practices on child survival. The information can be compared with population surveys (e.g. DHS), which monitor infant feeding practices in the general population. The indicators may not reflect the actual distribution of infant feeding practices of HIV-exposed infants at the national level, as it does not include HIV-exposed infants who may have already died, infants whose exposure status is unknown, nor HIV-exposed infants whose mothers did not attend a facility with their infant for DPT3 or for another reason at or around 3 months. Additional To fully understand the extent and type of infant feeding practices, considerations countries may consider carrying out special studies with a cohort of HIV-infected women who choose to replacement feed and exclusively breastfeed. As well as measuring infant feeding practices, these studies could examine the reasons why women who have chosen either breastfeeding or replacement feeding are or are not practicing the chosen option exclusively, and whether the AFASS criteria were present. It could also examine the types of foods and liquids given to infants in addition to breast milk or formula before six months, and issues around cessation of exclusive breastfeeding at six months and complementary feeding after that time. Another issue to be examined is the impact of early infant diagnosis on infant feeding practices, and if the impact is negative, what can be done to better support mothers at this time. In countries where exposed infant follow-up has been integrated into community outreach services, programmes should consider identifying a system for collecting data at the community level for this indicator. Countries may wish to consider collecting this information at other time points, for example at both 6 weeks and 6 months. They may also wish to calculate the indicators using different denominators, such as the estimated number of HIV-exposed infants who should have been followed-up. Data utilization Review the distribution of infant feeding practice and discuss strategies to move towards safer practices. **Data Quality Control** Please provide any relevant information that would allow to better interpret and Notes for the the data reported. **Reporting Tool** If this data is not available, please provide an estimate of the distribution of IF practice among HIV+ women in the country in the Comments section.

PMTCT M&E Core Indicator #10

Other References

3.11 Number of pregnant women attending ANC at least once during the reporting period	
Notes for the Reporting Tool	Please report the number of ANC attendees with at least one visit during the reporting period.
	Please note that this counts the number of people, and not the number of attendances, meaning that a pregnant woman making 3 ANC visits will only be counted once.
	If the number does not represent the national number (e.g. if you only have data from 65% of the districts or facilities; or if the number represents multiple visits instead of "at least one visit"), please comment on the representativeness of the number you are reporting.

EURO only

3.11.1 Percentage of HIV-positive pregnant women who had their pregnancy terminated (EURO8)	
Rationale	Pregnancy termination is common in eastern European countries. HIV positive pregnant women who terminated their pregnancy do not need to take ARV drugs to prevent mother-to-child transmission. This indicator helps to assess access to effective contraceptive methods among HIV positive women, quality of counselling on reproductive health and family planning and reflects common medical practices.
What is measured	This indicator measures termination of pregnancy among pregnant HIV positive women.
Numerator	Number of pregnancy terminations among HIV-positive pregnant women during the reporting year.
Denominator	Number of diagnosed HIV-positive women who had pregnancy registered during the reporting year.
How to Measure and Measurement Tools	The numerator is calculated from national programme records aggregated from health care facility registers.
Disaggregation	None requested.
Strength and weaknesses	Prevention of unintended pregnancies among HIV positive women and improved access to family planning and effective contraception is one of the key elements of a comprehensive PMTCT strategy. This indicator helps for better planning reproductive health services for HIV positive women.

Data utilization	Look at trends over time. Although disaggregation is not required for this indicator, disaggregated data by geographical regions in the country (if available) is useful for in-country analysis allowing identification of lower performing areas. This indicator will have impact on other indicators, including ARV coverage.
Data quality control and notes for the reporting tool	It is important to comment how the indicator was calculated. Variation could happen due to different HIV testing policies among pregnant women between countries, for example:
	HIV testing is offered to all pregnant women, including those who terminate pregnancy
	HIV testing is offered only for women who will continue pregnancy, excluding those who opt for termination

3.11.2 Percentage of HIV-positive pregnant women who delivered during the reporting year (EURO9)	
Rationale	The number and percentage of HIV-positive pregnant women who delivered during the reporting period provides the basis for calculating reported rates of mother-to-child transmission of HIV. Further, elective Caesarean section is an intervention that reduces the risk of mother-to-child transmission. This indicator will help to monitor access to PMTCT interventions and calculate mother-to-child transmission rate and provides information about current health system practices.
What is measured	This indicator measures the proportion of HIV-positive women who delivered during the reporting year.
Numerator	Number of HIV-positive pregnant women who delivered during the reporting year.
Denominator	Number of HIV-positive pregnant women who had pregnancy registered during the reporting year.
How to Measure and Measurement Tools	The numerator is calculated from national programme records aggregated from health care facility registers.
Disaggregation	Type of delivery: Elective Caesarian section (defined as Caesarian section conducted prior to uterus contractions started and foetal membranes ruptured) versus other types of delivery (spontaneous, induced, acute Caesarean section etc.)
Strength and weaknesses	Elective Caesarean section among HIV positive women has been one of the key interventions prior to the use of ART for PMTCT. If ART is used and viral load at 36 weeks of gestation is less than 1000 copies/ml there is limited benefit of these interventions. Still there are countries in the Region that do not have access to routine monitoring of viral load. With unknown viral load status, elective Caesarean section is an important intervention for PMTCT.
Data utilization	This indicator will help to calculate mother-to-child transmission rates.

3.12 PAHO Specific Indicators

3.121 Number of infants born to HIV positive mothers ("HIV-exposed infants") born in 2012 (or latest data available)

What it measures Reported number of infants born to HIV-positive mothers within a defined calendar year (2012).

3.12.2 Number of infants, born in 2012 (or latest data available) to HIV positive mothers, classified as indeterminate (i.e.: all lost to follow up, death before definitive diagnosis, indeterminate lab results)

What it measures

Number of infants born during the defined calendar year (2012) to
HIV-positive women, who did not complete diagnostic testing to evaluate
their HIV status due to their being lost to follow-up, to their death, or to their
transfer to another facility and/or were not tested.

3.12.3 Number of infants born in 2012 (or latest data available) to HIV + mothers that are diagnosed as positive for HIV

What it measures

Number of infants born to HIV-positive mothers in 2012, who were diagnosed as HIV positive.

3.12.4 Number of infants born to HIV + mothers in 2012 (or latest data available) that are diagnosed as negative for HIV

What it measures

Number of infants born to HIV-positive mothers in 2012, who were diagnosed as HIV negative.

3.13 EURO-specific PMTCT Indicator (pregnant women who inject drugs)

3.13.1 Percentage of HIV (EURO11)		
Rationale	HIV positive pregnant women who are injecting drugs remain the hardest to reach population by PMTCT interventions. Significant proportion of children who are HIV positive or/and abandoned were born to HIV positive women using drugs. It is a strategic focus for the Region to improve access of PWID women to PMTCT interventions and services.	
What is measured	This indicator measures proportion of HIV-positive pregnant women who were actively injecting drugs during pregnancy.	
Numerator	Number of HIV-positive pregnant women who were injecting drug users (PWID).	
Denominator	Number of diagnosed HIV-positive women who had pregnancy registered during the reporting year.	
How to Measure and Measurement Tools	The numerator is calculated from national programme records aggregated from health care facility registers.	
Disaggregation	None requested.	
Strength and weaknesses	Injecting drug use and substance use has always been associated with stigma and discrimination. Therefore some PWID women might not report their drug use to healthcare providers, or be less inclined to seek health care services and the true prevalence of drug injection could be underestimated while some PWID women might not access the services they need.	
Data utilization	This indicator will help to monitor trends of PWID among pregnant HIV positive women and better plan interventions to address it.	
Data quality control and notes for the reporting tool	It is important to put a note if numerator and denominator has all delivered PWID women, or also include those who terminated there pregnancies.	

3.13.2 Percentage of HIV-positive pregnant PWID women who received OST during pregnancy (EURO12)	
Rationale	HIV positive pregnant women who are injecting drugs remain the hardest to reach population by PMTCT interventions. Opioid substitution therapy (OST) is a critical intervention to improve access of PWID women to PMTCT services.
What is measured	This indicator measures proportion of HIV-positive pregnant drug dependent women who were receiving OST (methadone, buprenorphin) during pregnancy.
Numerator	Number of HIV-positive pregnant PWID women who received OST during pregnancy.

Denominator	Number of diagnosed HIV-positive PWID women who had pregnancy registered during the reporting year.
How to Measure and Measurement Tools	The numerator is calculated from national programme records aggregated from health care facility registers.
Disaggregation	None requested.
Strength and weaknesses	OST has been documented as an effective intervention to improve pregnancy outcome, including reduced rates of neonatal morbidity and mortality. Due to stigma and discrimination of PWID women, some of them could under report their injecting drug use, which may in turn have an impact on the indicator and overestimate coverage with OST.
Data utilization	This indicator will help to monitor trends and access of PWID pregnant HIV positive women to OST.
Data quality control and notes for the reporting tool	It is important to put a note clarifying if the numerator and denominator includes delivering PWID women only, or also include those who terminated their pregnancies.

3.13.3 Percentage of HIV-positive pregnant PWID women who received ARVs to reduce the risk of mother-to-child transmission during pregnancy (EURO13)	
Rationale	HIV positive pregnant women who are injecting drugs remain the hardest to reach population by PMTCT interventions. Antiretroviral Treatment (ART) is a critical intervention to reduce the risk of mother-to-child transmission in HIV-positive pregnant PWID women.
What is measured	This indicator measures the proportion of HIV-positive pregnant women who inject drugs who were receiving ARVs during pregnancy.
Numerator	Number of HIV-positive pregnant PWID women who received ARVs during pregnancy.
Denominator	Number of diagnosed HIV-positive PWID women who had pregnancy registered during the reporting year.
How to Measure and Measurement Tools	The numerator is calculated from national programme records aggregated from health care facility registers.
Disaggregation	None requested.
Strength and weaknesses	ANC data are often incomplete and might influence the indicator. Due to stigma and discrimination of PWID women, some of them could under report their injecting drug use, which may in turn have an impact on the indicator. The indicator does not assess adherence.
Data utilization	This indicator will help to monitor trends and access of PWID pregnant HIV positive women to ARVs.
Data quality control and notes for the reporting tool	It is important to put a note clarifying if the numerator and denominator includes delivering PWID women only, or also include those who terminated their pregnancies.

Target 4: Have 15 million people living with HIV on antiretroviral treatment by 2015

4.1. Percentage of adults and children currently receiving antiretroviral therapy

Note that the above indicator is described in the first part of the Guidelines whereas the following indicator on people newly initiating ART is additional to section 4.1 and not included in the GARPR Guidelines:

4.1 – additional: HIV treatment: Antiretroviral therapy Number of eligible adults and children who newly initiated antiretroviral therapy (ART) during the reporting period (2013)	
Rationale	In addition to coverage it is important to monitor ART initiation. Comparing the evolution of the number of people on ART at the end of the years (indicator G2) does not inform about the number newly initiated, especially since ART attrition is high in the first year and thus the patients newly initiating during the reporting year are not all continuing at the end of the year. Therefore this indicator captures the number of patients newly initiated on ART during a reporting year.
What it measures	Number of eligible adults and children who newly initiated antiretroviral therapy during the reporting period (2013) Yearly evolution of the number of patients newly enrolled in antiretroviral therapy
How to Measure and Measurement Tools	Facility ART registers and drug supply management forms. By counting the number of patients who are newly enrolled in ART within the reporting period. Patients with records that transfer in from another facility or who temporarily stopped therapy and have started again in the reporting period should not be counted (risk of double counting). ARV drugs taken for purpose of PMTCT (except ART for the mother's own health) and post-exposure prophylaxis are not included in this indicator.
Disaggregation	By male and female By age groups: <1, 1-4, 5-14, 15+ By public and private By mode of transmission, injecting status, OST recipient status, imprisonment status (European Region only) These and other disaggregations to be included if available in the Comments box
Strengths and weaknesses	This indicator permits monitoring trends in initiation but does not attempt to distinguish between different forms of antiretroviral therapy or to measure the cost, quality or effectiveness of treatment provided. These will each vary within and between countries and are liable to change over time. The degree of initiation of ART will depend on factors such as cost relative to local incomes, service delivery infrastructure and quality, availability and uptake of voluntary counselling and testing services, and perceptions of effectiveness and possible side effects of treatment.
Additional considerations	This indicator should be analysed in view of the 'waiting list' i.e. patients eligible for ART and not initiated.

Data utilization	In addition to the number of old patients retained on ART (retention on ART) the number of patients newly initiated is necessary for accurate planning of resources and drug stocks (avoiding shortage and wastage)
Data Quality Control and Notes for the Reporting Tool	Double Reporting: If patients transferred in and out are not correctly registered and if patients followed in different ART sites are not identified, there is a risk for double reporting which could lead to an overestimation of ART initiation. If this is the case, please comment.
	Similarly if patients temporarily stopping ART and restarting are coded as new patients, this will overestimate the true number of patients newly initiated.
	National Representativeness: the numerator is a national cumulative indicator, usually produced by all health facilities, otherwise it may estimate ART initiation. Please comment on your data as necessary.
	Triangulation Options: Pharmacy report, comparing the number of new patients in the pharmacy register and the ART register
Other References	PEPFAR indicator and guidelines

- 4.2 Percentage of adults and children with HIV still alive and known to be on antiretroviral therapy
 - (a) 12 months after initiating treatment among patients initiating antiretroviral therapy during 2013
 - (b) 24 months after initiating treatment among patients initiating antiretroviral therapy during 2013
 - (c) 60 months after initiating treatment among patients initiating antiretroviral therapy during 2013

3	
Rationale	Antiretroviral is a life-long intervention. Measuring retention on ART is critical for determining the effectiveness of programmes, inferring their impact and to highlight obstacles to expanding and improving them.
What it measures	This indicator measures the retention on ART related to the increase in survival and willingness to continue ART. It should be produced at 12 months and for longer duration of follow-up; the 24 and 60 months retention are described here (the 12 months retention is included in the GARPR indicator guidance). It completes programme coverage as a measure of the effectiveness.
Numerator	Number of adults and children who are still alive and on ART at b) 24 months, c) 60 months, after initiating treatment (among those who initiated ART in b) 2011 and c) 2008).

Denominator (b) at 24 months: Total number of adults and children who initiated ART in 2011 (or another specified period), who were expected to achieve 24-month outcomes within the 2013 reporting period (or 24 months after the specified initiation period), including those who have died since starting ART, those who have stopped ART, and those recorded as lost to follow-up at month 24. (c) at 60 months: Total number of adults and children who initiated ART in 2008 (or another specified period), who were expected to achieve 60-month outcomes within the 2013 reporting period (or 60 months after the specified initiation period, including those who have died since starting ART, those who have stopped ART, and those recorded as lost to follow-up at month 60. How to Measure and **Numerator and denominator:** Programme monitoring tools; ART register; **Measurement Tools** cohort analysis forms. In measuring retention, it is important to carefully select the patients according to the period they have initiated ART and to check their outcomes when they reached the expected duration of follow-up. Assessing outcomes at 24 months should include all patients started 2 years ago and at 60 months, all patients started 5 years ago. If the data available does not really fit this standard yearly period, it is important to specify the period the patients have initiated ART. Disaggregation Among the people who started (denominator), in addition to reporting the (1) number of people alive and on treatment (numerator), it is also important to report the number (2) lost to follow-up, (3) stopped therapy, and (4) died. These 4 outcomes should sum to the number of people who started ART. When generating information at site level, patients transferred in should be included in the statistics and patients transferred out should be excluded. From the compilation of site reports, if the number of patients transferred in and transferred out is summed at the national level, these statistics should be reported for 12-month analysis. **Strengths** The continuation of ART is mostly related to survival (but also willingness and weaknesses to continue). Survival might reflect the services offered but also depends on the baseline characteristics of the patients started on ART. Clinical, immunological and virological staging are independent predictors of survival under ART. Baseline characteristics of the cohort of patients should help in interpreting the results and, in particular, comparing ART sites.

Additional considerations	If data on 24-month or 60-month retention are not available for patients that initiated antiretroviral therapy in 2011 or 2008, respectively, but available for patients that initiated antiretroviral therapy during an earlier time period (e.g. 2010 or 2009, or 2007 or 2006), please specify the period in the comment field: e.g. "Started antiretroviral therapy between [month]/[year] and [month]/[year]".
	The numerator does not require patients to have been on antiretroviral therapy continuously for the 24 month or 60 month period. For example, patients who may have missed one or two appointments or drug pick-ups, and temporarily stopped treatment since initiating treatment but are recorded as still being on treatment at month 24 or 60 are included in the numerator. On the contrary, those patients who have died, stopped treatment or been lost to follow-up at 24 or 60 months since starting treatment are not included in the numerator.
	In countries where this indicator is not produced in all ART sites but in a sub-set of facilities, data should be interpreted keeping in mind the representativeness, and this should be stated in the <i>Comments</i> box.
Data utilization	Note any particularly low retention and assess reasons behind it, by analysing the distribution of those who are not on ART: dead, stopped, loss to follow up. If data is available, try to assess the lost-to-follow-up population to see if they are likely to be dead, stopped, or transferred out. Compare cohorts.
Data Quality Control and Notes for the Reporting Tool	National Representativeness: If this indicator is only produced in a sub-set of facilities, comment should be added on the source of information and whether the information is representative of all ART sites.

EURO only

4.2.1 Percentage of injecting drug users with HIV still alive and known to be on treatment a) 12 months, b) 24 months and c) 60 months after initiation of antiretroviral therapy (EUR4)	
Rationale	ART is a lifelong therapy that increases survival and reduces transmission. In WHO European Region, where injecting drug users (PWID) are most affected by the HIV/AIDS epidemic, access to and retention in ART is among key the interventions in health sector response.
What it measures	This indicator measures the retention on ART related to the increase in survival and willingness to continue ART. It should be produced at 12 months and then yearly after the beginning of ART. It completes program coverage by a measure of the effectiveness.
Numerator	Number of PWID who are still alive and on ART a) 12 months, b) 24 months, c) 60 months after initiating treatment.

Denominator

- a) At 12 months: Total number of injecting drug users who initiated ART in 2012 and so, who were expected to achieve 12-month outcomes within the reporting period (2013), including those who have died since starting ART, those who have stopped ART, and those recorded as lost to follow-up at month 12.
- b) at 24 months: Total number of injecting drug users who initiated ART in 2011 and so, who were expected to achieve 24-month outcomes within the reporting period (2013), including those who have died since starting ART, those who have stopped ART, and those recorded as lost to follow-up at month 24.
- c) at 60 months: Total number of injecting drug users who initiated ART in 2008 and so, who were expected to achieve 60-month outcomes within the reporting period (2013), including those who have died since starting ART, those who have stopped ART, and those recorded as lost to follow-up at month 60.

How to Measure and Measurement Tools

Numerator and denominator: Programme monitoring tools; ART register and cohort analysis report form.

In measuring retention for the 3 different intervals, it is important to carefully select the PWID patients according the period they have started therapy and to check the outcomes when they reached the expected duration of follow-up.

Assessing outcomes at 12 months should include all PWID patients who started therapy in the last year, at 24 months, all PWID patients who started 2 years ago and at 60 months, all PWID patients who started 5 years ago. If the data available do not fit this standard yearly period it is important to specify the period used for calculation and when the patients initiated treatment.

PWID patients must be alive and on antiretroviral therapy at 12/24/60 months after their initiation of treatment. The numerator does not require patients to have been on antiretroviral therapy continuously for the 12/24/60-months period. PWID patients who may have missed one or two appointments or drug pick-ups, and temporarily stopped treatment during the 12/24/60 months since initiating treatment but are recorded as still being on treatment at month 12/24/60 are included in the numerator. On the contrary, those patients who have died, stopped treatment or been lost to follow-up at 12/24/60 months since starting treatment are not included in the numerator.

When generating information at site level, patients transferred in should be included in the statistics and patients transferred out should be excluded. From the compilation of site reports, if the number of patients transferred in and transferred out is summed at national level, these statistics should be reported for 12 months analysis.

Disaggregation

As much as possible, this indicator is to be disaggregated by sex, by age (<15, 15+), by 1st line and 2nd line regimens at the end point.

Strengths and weaknesses	The continuation of ART is mostly related to survival (but also willingness to continue treatment). Survival might reflect the services offered but also depends on the baseline characteristics of the PWID patients started on ART. Clinical, immunological and virological staging are independent predictors of survival under ART. For injecting drug users, various underlying health conditions may additionally affect survival rates. Baseline characteristics of the cohort of patients should help in interpreting the results and in comparing ART sites.
Additional considerations	In countries where this indicator is not produced in all ART sites but in a sub-set of facilities, data should be interpreted keeping in mind the representativeness.
Data utilization	Note any particularly low coverage and use the data to assess the reasons behind it. Try to get data on the distribution of those who are no longer on ART: dead, stopped, loss to follow up. If data are available, try to assess loss to follow-up population to see if they are likely to be dead, stopped, or transferred out. Compare cohorts.
Data Quality Control and Notes for the Reporting Tool	National Representativeness: If this indicator is only produced in a sub-set of facilities, comment should be added on the source of information, sample size and whether the information is representative of all ART sites.

4.3.a Number of health facilities that offer antiretroviral therapy (ART)	
Rationale	Antiretroviral therapy is a cornerstone of effective HIV treatment, and measuring the percentage of health facilities that offer ART provides valuable information about ART availability.
What it measures	Number of health facilities that offer ART (i.e., prescribe and/or provide clinical follow-up).
	Capacity of health facilities to provide antiretroviral therapy (ART), expressed as percentage of health facilities that offer ART (i.e., prescribe and/or provide clinical follow-up). Health facilities include public and private facilities, health centres and clinics (including TB centres), as well as health facilities that are run by faith-based or nongovernmental organizations.

How to Measure and Measurement Tools	The numerator is calculated by summing of the number of facilities reporting availability of ART services. Information on the availability of specific services is usually kept at the national or sub-national level. National AIDS Programmes should have a record of all health facilities offering ART services. A health facility census or survey can also provide this information, along with more in-depth information on available services, provided the information is collected from a representative sample of health facilities in the country. Responses to a series of questions establish whether providers in that facility provide ART services directly (i.e., prescribe ART and/or provide clinical follow-up for ART patients) or refer patients to other health facilities for these services. In addition, facility records documenting the current status of service provision should be consulted. One potential limitation to facility surveys or censuses is that they are usually only conducted once every few years. Countries should regularly update their programme records on health facilities offering ART services, and supplement these data with those obtained through a health facility survey or census every few years. For health facility surveys or censuses, tools such as the Service Provision Assessment (SPA) or the Service Availability Mapping (SAM) can be used.
Disaggregation	Sector: public, private By type: hospital, health centre, ANC facility, TB facility, STI services.
Strengths and weaknesses	This indicator provides valuable information about the availability of ART services in health facilities, but it does not capture information about the quality of services provided. Antiretroviral therapy itself is complex, and it should be delivered as part of a package of care interventions, including the provision of co-trimoxazole prophylaxis, the management of opportunistic infections and comorbidities, nutritional support and palliative care. Simple monitoring of ART availability does not ensure that all ART-related services are adequately provided to those who need them. Nevertheless, it is important to know what percentage of health facilities provide ART services in order to plan for service expansion as needed to meet universal access targets.
Additional considerations	One strategy to scale up ART services is to make ART available in more health facilities. This may be achieved by decentralizing ART services from tertiary facilities (e.g., hospitals) to primary or secondary-level health facilities. Greater availability of ART services provides crucial support to the goal of universal access to HIV treatment. Depending on the country's epidemic type, the denominator may not be as relevant if the HIV programme strategy aims to target a limited number of
Data utilization	sites to offer ART in. To look at progress in the percentage of health facilities which provide antiretroviral therapy. Analyzing the data geographically and by type of health facilities, and triangulating the data with estimates of HIV density can provide insight into where there is a need to increase availability of ART services.

Data Quality Control and Notes for the Reporting Tool	Please comment on whether the data reported is from a national facility listing or census, or from a survey. If data from the private or other sectors is missing, please comment.
	If it is possible to easily report any additional information on the geographical distribution of facilities offering ART (e.g. urban/rural, % facilities with ART in areas with a high concentration of people living with HIV), please provide extra details.
Other References	Additional Recommended Indicators for NAP #5

4.3.b Health facilities Number of health facilities that offer paediatric antiretroviral therapy (ART)		
Rationale	Antiretroviral therapy is a cornerstone of effective HIV treatment, and measuring the percentage of health facilities that offer paediatric ART provides valuable information about capacity to address HIV care in children.	
What it measures	Number of health facilities that offer paediatric ART.	
	Capacity of health facilities to provide paediatric antiretroviral therapy (ART), expressed as percentage of health facilities that offer paediatric ART. Health facilities include public and private facilities, health centres and clinics (including TB centres), as well as health facilities that are run by faith-based or nongovernmental organizations.	
How to Measure and Measurement Tools	The numerator is calculated by summing the number of facilities reporting availability of paediatric ART services. Information on the availability of specific services is usually kept at the national or subnational level. National AIDS Programmes should have a record of all health facilities offering ART services. A health facility census or survey can also provide this information, along with more in-depth information on available services, provided the information is collected from a representative sample of health facilities in the country. Responses to a series of questions establish whether providers in that facility provide paediatric ART services directly or refer patients to other health facilities for these services.	
	In addition, facility records documenting the current status of service provision should be consulted. One potential limitation to facility surveys or censuses is that they are usually only conducted once every few years. Countries should regularly update their programme records on health facilities offering paediatric ART services, and supplement these data with those obtained through a health facility survey or census every few years. For health facility surveys or censuses, tools such as the Service Provision Assessment (SPA) or the Service Availability Mapping (SAM) can be used.	

A denominator is not requested in the UA reporting tool but some countries trying to expand paediatric ART nationally can consider Total number of health facilities, excluding specialized facilities where paediatric ART services are/will never be relevant, which can be calculated by summing the total number of health facilities included in the sample. Information for construction of the denominator may come from programme records, facility listings, and/or national strategy or planning documents. It should exclude specialized facilities where paediatric ART services are/will never be relevant, (e.g., facilities where paediatric ART services are/will never be relevant, (e.g., facilities specializing in eye care where ART will never be introduced) Disaggregation Sector: public, private This indicator provides valuable information about the availability of paediatric ART services in health facilities, but it does not capture information about the quality of services provided. Antiretroviral therapy itself is complex, and it should be delivered as part of a package of care interventions, including the provision of co-trimoxazole prophylaxis, the management of opportunistic infections and comorbidities, nutritional support and palliative care. Simple monitoring of ART availability does not ensure that all ART-related services are adequately provided to those who need them. Nevertheless, it is important to know what percentage of health facilities, nutritional access targets. One potential limitation to facility surveys or censuses is that they are usually only conducted once every few years and may not capture the latest information especially in setting with recent intensified scale-up. Additional considerations One strategy to scale up ART services is to make ART including paediatric ART services available in more health facilities. This may be achieved by decentralizing ART services from tertiary facilities (e.g. hospitals) to primary or secondary-level health facilities. Greater availability of paediatri		
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Other References UNAIDS Additional Recommended Indicators for NAP #5	and Notes for	listing or census, or from a survey. If a survey, please remember to report the year of the survey. If data from the private or other sectors is missing, please comment. If it is possible to easily report any additional information on the geographical distribution of facilities offering paediatric ART (e.g. urban/rural, %facilities with ART in areas with a high concentration of people
	Other References	UNAIDS Additional Recommended Indicators for NAP #5

	alth facilities dispensing ARVs that experienced a stock-out of at least / in the last 12 months
Rationale	As countries scale-up ART services, it is important to ensure that ARVs are available to those who need them. ART is a long-term treatment strategy for people living with advanced HIV infection, and treatment interruptions may lead to treatment failure and HIV drug resistance. Efficient supply management is needed to ensure an uninterrupted supply of ARVs.
What it measures	This indicator measures a key aspect of antiretroviral (ARV) drug supply management: whether health facilities dispensing ARV drugs have run out of stock of at least one required ARV in the last 12 months.
Numerator	Number of health facilities dispensing ARVs that experienced a stock-out of one or more required ARV drug in the last 12 months.
Denominator	Total number of health facilities dispensing ARVs.
How to Measure and Measurement Tools	This information is collected at central level, where health facilities submit their inventory control reports or requisition forms for ARVs. These forms have information on patients on ART, consumption data, and stock on hand with stock out information if any.
	This indicator requires the following tools:
	 stock inventory control reports from health facilities indicating also the stock level of each item in the report;
	 requisition forms submitted from facilities during a defined period of time (e.g. last order period, last quarter, last year) for ARVs; and
	 list of ARVs that each facility is expected to dispense, if not already included in the inventory control reports or requisition forms.
	All the above work if the national logistics management information systems (LMIS) is operational. If the national LMIS is not operational, or health facility surveys such as the Service Provision Assessment (SPA) or the Service Availability Mapping (SAM) may be used provided they include questions on ARV stock-outs.
	If there is one national logistics management information system (LMIS) with details on ARV availability at the health facility level, information should be extracted from this system to construct this indicator. Alternatively, the information may need to be collected through a special survey or site visits. If there are only a limited number of health facilities where ARVs are dispensed in the country, all health facilities dispensing ARVs should be included in the survey or site visits. If the number of health facilities dispensing ARVs is large, it may be necessary to select a representative sample from the total number of health facilities dispensing ARVs (the full list should be available at the national level). When sampling, it is important to ensure that the sample includes facilities at different levels (such as central, district, and peripheral levels). In countries where ARV drugs are dispensed at pharmacies or other non-health facility delivery points, stockouts should also be monitored in these venues; feasibility will depend on the coverage of the Logistics Management Information System.

Disaggregation	Sector: public, private	
Strengths and weaknesses	This indicator captures a crucial component of the ART programme: whether or not there is a continuous, uninterrupted supply of ARV drugs at the health facility level.	
	This indicator does not, however, provide information on why stock-out problems occur; which ARV drug(s) are/were out of stock; or how long the stock-out lasted for a particular ARV drug. It also does not provide information on the quality of ARV drug storage, delivery, and distribution.	
Additional considerations	In some situations, simply monitoring stock-outs could be misleading because a facility may keep reserve stock but maintains a policy of not issuing the reserve stock. These facilities would not be counted as having experienced a stock-out using this indicator definition, even though a patient would not be receiving a required ARV drug for treatment. In settings where reserve stock is not issued during ARV stock-outs, it is preferable to collect information on a functional stock-out (i.e., the inability to access or make use of a required ARV drug).	
Data utilization	If stock-outs exist, assess whether the problem lies in the national distribution system or if it is a financial flow problem or a global ARV shortage problem. Find out whether the reason is due to projections of supply order or the distribution system or any other issue. Use this as an opportunity to see whether LMIS is functioning.	
Data Quality Control and Notes for the Reporting Tool	Comment on whether the data is based on national data or survey data from a sample of facilities. Please provide any other comments that would help the interpretation of data (e.g. if only public or private sector data is included, and whether it may be an over- or underestimate).	
Other References	Harmonized monitoring and evaluation indicators for procurement and supply management systems.	

PAHO only

4.5 Late HIV diagnoses: Percentage of HIV positive persons with first CD4 cell count < 200 cells/μL in 2013	
Rationale	As countries scale-up HIV services, it is important to monitor whether people are diagnosed at an earlier stage (or what percentage is still diagnosed at a late stage).
What it measures	This indicator measures the proportion of people with a CD4 cell count <200 cells/µl out of those who had a first CD4 count during the reporting period.
Numerator	Number of HIV-positive people with first CD4 cell count <200 cells/µl in 2013
Denominator	Total number of HIV-positive people with first CD4 cell count in 2013

4.6	HIV CARE: HIV treatment: Antiretroviral therapy		
4.6.a	Total number of people enrolled in HIV care at the end of the reporting period		
4.6.b	Number of adults and children newly enrolled in HIV care during the reporting period (2013)		
Ration	ale	In addition to HIV testing it is important to monitor linking to HIV care and treatment. Comparing the evolution of the number of people tested for HIV at the end of the years does not inform about the number new people enrolled in HIV care especially since loses in HIV continuum of care cascade may be high with high attrition and lost to follow up. Therefore this indicator captures the number of patients that are either on HIV care waiting for ART initiation or on ART treatment during a reporting year.	
What i	t measures	Number of adults and children who are being followed up by health services for HIV care, including those in antiretroviral therapy during the reporting period (2013). People in HIV care include those seen at the HIV clinic at least once during the reporting year. Yearly evolution of the number of HIV+ patients enrolled in the health services for HIV.	
	o Measure and rement Tools	Health facility services that received patients for ART assessment needs and ART registers. By counting the number of patients who are linked to care and ART within the reporting period. Transfer-in patients, those who temporarily stopped therapy but continue to be monitored, pregnant women taking ARVs for PMTCT purpose should be included as linked to care but caution is required to avoid double counting	
Disago	gregation	By gender : Male/ Female By age groups: <15, 15+ By mode of transmission (European Region only)	
Streng and we	nths eaknesses	This indicator permits monitoring trends of total patients linked to HIV health services but does not attempt to distinguish between HIV care and ART or to measure the cost, quality or effectiveness of treatment provided. The degree of ART initiation will depend on factors such as new policies, cost relative to local incomes, service delivery infrastructure and quality, availability and uptake of voluntary counselling and testing services, and perceptions of effectiveness and possible side effects of treatment.	
Additio	onal Ierations	This indicator should be analysed in view of the 'waiting list' i.e. patients eligible for ART and not initiated.	
Data u	tilization	In addition to the number of people on ART, the number of patients on care is necessary for accurate planning of resources and drug stocks (avoiding shortage and wastage)	

Data Quality Control and Notes for the Reporting Tool	Double Reporting: If patients transferred in and out are not correctly registered and if patients followed in different ART sites are not identified, there is a risk for double reporting which could lead to an overestimation of ART initiation. If this is the case, please comment.
	Similarly if patients temporarily stopping ART and restarting are coded as new patients, this will overestimate the true number of patients newly initiated.
	National Representativeness: the numerator is a national cumulative indicator, usually produced by all health facilities,. Please comment on your data as necessary.
	Triangulation Options: Pharmacy report, comparing the number of people being tested, the number of patients in the pharmacy register and the ART register

4.7 Viral Load suppression

Other References

- 4.7 a) percentage of people on ART tested for viral load (VL) who have an undetectable viral load in reporting period (2013)
- 4.7 b) percentage of people on ART tested for viral load (VL) with VL level below ≤ 1,000 copies after 12 months of therapy (2013)

PEPFAR indicator and guidelines

Rationale

Viral load is the recommended measure of ART efficacy and also provides an indication of treatment adherence and the risk of HIV transmission at the individual and population levels

Effective ART reduces transmission of HIV. Various study results provide strong support for the premise that treatment of the HIV-infected individual can significantly reduce sexual transmission of HIV. Thus suppression of viral load to undetectable levels should greatly reduce the risk of transmission to the uninfected partner. ART also prevents perinatal transmission of HIV. ART is considered effective when it consistently suppresses plasma viral load to undetectable levels.

Persons receiving antiretroviral therapy (ART) frequently develop treatment resistance. A key determinant of treatment failure is increase in viral load. Measurement of viral suppression (VL ≤1000 copies/ml) is key programmatic indicator related to effective treatment.

What it measures

Viral load is a measure of the effect of ART on viral replication. A viral load threshold of >1,000 copies/ml defines treatment failure according to the WHO 2013 ART guidelines.

A supressed viral load under the limits of detection is termed: undetectable viral load, usually between 50 to 400 copies/ml.

The viral load of patients in care may be used as a quality of care indicator for the population engaged in care. If measured over time, it should reflect access to healthcare, acceptance and adherence to antiretroviral therapy, and adequate clinical monitoring of VL. For a particular healthcare system it can be used as a rough proxy measure of access to antiretrovirals, level of antiretroviral medication adherence, patient compliance with disease monitoring, and quality of care delivered to a patient population.

Numerator	4.7. a	(cross sectional data)
		number of people on ART tested for viral load in the reporting period with suppressed viral load (i.e. ≤ 1000 copies)
	4.7. b	(cohort data)
		Number of people tested after 12 months therapy for VL and have suppression ($VL \le 1000$ copies) during the reporting period
Denominator	4.7.a	(cross sectional data)
		number of people on ART tested for viral load in the reporting period
	4.7. b	(cohort data)
		Number of people tested after 12 months therapy for VL during the reporting period
How to Measure and Measurement Tools	Where viral load testing is done routinely, results will be recorded in patient files or in laboratory systems. Viral load test results may also be recorded electronically and reported as part of cohort monitoring studies as the percentage of patients who are virologically suppressed at defined time points.	
Disaggregation	By age groups: <15, 15+	
	By gen	der: Male/ Female
Strengths and weaknesses	Strengths: viral load measurements provide information on adherence, treatment efficacy, and transmission risk at the individual and programme level	
	limited i may no load, as This ap all ART outcom determi	esses: viral load monitoring capacity is scaling up but remains in low-income settings. Summary data from the viral load indicator to be representative of the broader ART treatment population viral is results may only be attainable from a non-representative subset. Polies in particular if viral load testing in not performed routinely for patients, but only selectively for those with questionable treatment ess. Cut-off VL values for treatment failure are not universally ned. Values to define supressed undetectable viral load varies ing on the sensitivity of the assays used.
Additional considerations		ve reasons, this indicator is only applicable if VL is performed y (rather than on a "as needed" basis).
		ortant to restrict this indicator to <i>people</i> on ART (and not include all erformed) in order to exclude re-testing in the reporting period.
	approa a highe	ettings use dried blood spots for viral load measures; this ch is currently poorly accurate at lower thresholds and therefore r threshold for defining virological failure needs to be applied copies/ml).

Data utilization	Viral load testing can help programmes to plan for second-line drug needs (in the case of treatment failure) and potential interventions to limit HIV transmission. The percentage of patients with undetectable viral load is a proxy measure of the program's success.
Data Quality Control and Notes for the Reporting Tool	Patient monitoring system may yield both cross sectional and cohort data. Cohort data may also stem from special studies. If laboratory data is used, data needs to be adjusted to avoid double counting of patients with more than one VL test in the reporting period.
Other References	Early warning indicators for HIVDR

Target 5: Reduce tuberculosis deaths in people living with HIV (PLHIV) by 50% by 2015

5.2 Percentage of adults and children living with HIV newly enrolled in care who are detected having active TB disease (new)	
Rationale	The primary aim of the ICF activities is early detection of TB among PLHIV and provision of prompt TB treatment and ART which if optimally implemented along with provision of early ART, Isoniazid Preventive Therapy and Airborne Infection Control practices, reduce TB burden among the PLHIV.
	While ICF is to be implemented in all the PLHIV attending HIV-care and treatment facilities during every visit, it is critically important among the PLHIV newly enrolled in HIV care and treatment, as risk of undetected TB among them is greater than those already on ART. Hence this indicator measures both burden of active TB disease among PLHIV newly enrolled in HIV care as well as the access to TB diagnosis
What it measures	Total TB cases detected among HIV positive patients who are newly enrolled in HIV care (Pre-ART or ART) during the reporting period
Numerator	Total number of adults and children newly enrolled in HIV care who are diagnosed as having active TB disease during the reporting period
Denominator	Total number of adults and children newly enrolled in pre-ART care or on ART during the reporting period
How to Measure and Measurement Tools	The outcome of TB status assessment among PLHIV attending ART facilities (whether TB is diagnosed) is to be recorded in ART card ("investigations" column on "encounters" section) and the pre-ART and ART register (monthly /quarterly follow-up sections respectively).
	At the end of reporting period count the total number of TB cases detected among PLHIV newly enrolled in HIV care i.e. those enrolled in pre-ART register plus ART register , during the reporting period. Care should be taken to avoid double counting the same individual on both registers. Direct registration into ART register may be part of country adaptation and TB patients detected among those enrolled in this register during reporting period may be enumerated and considered in the numerator . The denominator consists of total number of PLHIV newly enrolled in HIV care during the same period
	This indicator is to be collected from pre-ART or ART registers and summarized on cross-sectional quarterly reports. It could also be assessed from a systematic sample of patient HIV care/ART cards during annual patient monitoring reviews.
Disaggregation	Data for this indicator should be disaggregated by sex and age (<15 years/15+)

Strengths and weaknesses	It measures burden of TB among PLHIV newly enrolled in care. It also measures the level of access to TB diagnosis in newly enrolled cases, indirectly. However it does not consider burden of TB/HIV cases who are detected through HIV testing of TB patients but may not enrolled in HIV care. Actual burden of TB among PLHIV may still be higher since some patients may remain asymptomatic or have disseminated form of TB which may be missed during evaluation if not for high level of suspicion or thorough investigation using sensitive tools	
Additional Considerations	Data are to be collected continuously and reported to sub-national or national level as part of routine cross-sectional reporting quarterly. It should also be submitted annually to WHO.	
Data Quality Control and Notes for the Reporting Tool	This is a "new" indicator and so countries are asked to provide comments on problems with reporting – particularly if they are unable to report.	

5.3 Percentage of adults and children newly enrolled in HIV care starting isoniazid preventive therapy (IPT)	
Rationale	To ensure that eligible HIV-positive individuals are given treatment for latent TB infection and thus to reduce the incidence of TB in people living with HIV.
What it measures	Number of adults and children newly-enrolled in HIV care who started on treatment for latent TB infection, isoniazid preventative therapy (IPT) expressed as a proportion of the total number of adults and children newly-enrolled in HIV care over a given time period.
Numerator	Number of adults and children <u>newly</u> enrolled (i.e. started) in HIV care (pre-ART and ART) who <u>also</u> start (i.e. given at least one dose) isoniazid preventive therapy treatment during the reporting period <i>HIV care</i> includes pre-ART and ART.
Denominator	Number of adults and children newly enrolled (i.e. started) in HIV care during the reporting period.
How to Measure and Measurement Tools	HIV treatment card and modified HIV care register. The data needed for this indicator is collected from pre ART and ART registers at the HIV care service sites, depending on where isoniazid preventive therapy (IPT) is to be administered. HIV-positive clients should be screened for TB. Those clients found not to have evidence of active TB will be offered IPT according to nationally determined guidelines. All those accepting IPT and receiving at least the first dose of treatment should be recorded. This information is being recorded in an extra column in the HIV care registers. Accurately predicting drug requirements for supply management requires the collection of more detailed information.

Disaggregation	None
Strengths and weaknesses	Treatment of latent TB infection will reduce the incidence of developing TB disease in People living with HIV who are infected with TB but who have no active TB disease. To include clients who are given at least one dose is relatively easy, even in resource-limited settings. This information is the minimum necessary to ensure that TB preventive therapy is being offered to HIV-positive clients without evidence of active TB. However, unless further data are collected, this indicator provides no information about how many clients adhere to or complete the TB preventive therapy course. Much greater resources are required to collect more complete data on adherence or completion, but programmes may wish to undertake periodic studies to establish, for example, adherence rates, and the accuracy of the screening questionnaire.
Additional considerations	A pharmacy based TB preventive therapy (INH) register should record client attendance to collect further drug supplies (usually monthly). From this register, facilities would be able to report the number of new, and continuing cases and treatment completion on a quarterly basis. If such information is collected routinely, the indicator of choice would be 'the number of HIV-positive clients completing treatment of latent TB infection, as a proportion of the total number of HIV-positive clients started on such treatment".
	From pilot testing sites it is apparent that many clients who test HIV-positive can be expected to start TB preventive therapy; some will not meet the eligibility criteria, some will decline and some will drop out during the screening process. The proportion likely to start TB preventive therapy depends on the screening algorithm used (for example, using tuberculin skin test as a screening tool reduces the number that are eligible) and also on the type of facility at which HIV diagnosis is made.
Data utilization	If low value, explore reasons why and compare disaggregated data with the national average to identify places needing special attention and reasons for suboptimal coverage. Explore further available data on completion of TBPT/IPT.
Data Quality Control and Notes for the Reporting Tool	Please provide any comments on whether the data you provide covers the entire country, or is from a selected sample (if so, please provide details on what the data represents, as well as any assumptions made to extrapolate the data to a national figure)
Other References	° A guide to monitoring and evaluation for collaborative TB/HIV activities

5.4 Percentage of adults and children enrolled in HIV care who had TB status assessed and recorded during their last visit		
Rationale	This is a process indicator for an activity intended to reduce the impact of TB among people living with HIV. It will demonstrate the level of implementation of the recommendation that people living with HIV are screened for TB at diagnosis and at follow-up visits using their last visit as proxy measure.	
What it measures	Number of adults and children enrolled in HIV care who had TB status assessed and recorded during their last visit.	
Numerator	Number of adults and children in HIV care, who had their TB status assessed and recorded during their last visit. HIV care includes pre-ART and ART.	
Denominator	Total number of adults and children in HIV care in the reporting period.	
How to Measure and Measurement Tools	WHO recommends the use of a simplified screening algorithm for intensified TB case findings that includes 4 clinical symptoms: (1) current cough, (2) fever, (3) weight loss and (4) night sweats.	
	Using this simplified algorithm assessment of TB status at every visit during the reporting period ('Yes' if 'no signs', 'suspect' or 'on treatment' and 'No' if TB status not assessed) should be recorded on the patient HIV care/ART card, and transferred onto the pre-ART or ART registers as appropriate at all facilities providing routine HIV care. Enrolled in care includes all those continuing in care and those newly enrolled during the reporting period This data should be analysed and reported together with other cross sectional data at national level.	
	The numerator is taken from the pre ART and ART registers by counting the number of patients who had their TB status assessed during the reporting period. For patients who started on ART during the reporting period, care should be taken to count them in the ART register and not in the pre-ART register.	
	The denominator for pre-ART patients will be those seen for care during the reporting period. The denominator for ART patients will be those current on ART during the reporting period.	
	The denominator is taken from the pre-ART and ART registers by counting the number of patients with a visit during the reporting period. This is then recorded on the cross sectional reporting form.	
	TB and HIV programmes should collaborate to ensure that agreed criteria for identifying a TB suspect and methods of TB screening are used that are consistent with TB control programme protocols. ²⁶	
Disaggregation	None	

^{26.} A suggested method of conducting the screening would be to ask HIV-positive clients whether they are currently on TB treatment. If not, they are then asked about the key symptoms of TB disease (e.g. cough > 2 weeks, persistent fever, night sweats, unexplained weight loss and lymphadenopathy). A simple checklist could be used and any positive response would indicate that the individual may be a TB suspect. If on questioning they are defined as a TB suspect (as per national protocols) they should be investigated for TB (or referred to TB service for investigation) and treated appropriately. Those found not to have TB should be offered six months of isoniazid preventive therapy (IPT))

Strengths and weaknesses	TB status assessment among people living with HIV, followed by prompt referral for diagnosis and treatment, increases the chances of survival, improves quality of life and reduces transmission of TB in the community. TB status assessment identifies HIV-positive clients who show no evidence of active TB and would benefit from treatment with isoniazid for latent TB infection.
	The indicator does not measure the quality of intensified TB case-finding nor does it reveal whether those identified as suspects are investigated further or effectively for TB. However, it does emphasize the importance of intensified TB case-finding for people living with HIV at diagnosis and at every contact they have with HIV treatment and care services.
	Programmes should aim for a high value for this indicator (close to 100%) but should interpret it in conjunction with values of indicators related to the % of people in HIV care who are: a) on TB treatment and b) who were given treatment for latent TB infection, to ensure that appropriate action follows the screening process. A low value will demonstrate that Objective B - reducing the impact of TB among people living with HIV - is unlikely to be met.
Data utilization	See section on Strengths and Weaknesses for interpretation of data and further areas to explore. If low value, review disaggregated data and explore reasons why.
Data Quality Control and Notes for the Reporting Tool	Please provide any comments on how this data was collected and any assumptions made in establishing a national estimate.
Other References	A guide to monitoring and evaluation for collaborative TB/HIV activities

Policy questions

Toney quodiono		
P.1b WHO Policy Questions		
HIV testing and counselling	Does the current national HIV testing and counselling policy/guidelines: for each: (1) yes (2) no (3) specify: - address testing of children? - address testing of adolescents? - address testing of "mature minors"? - address HIV testing for key and vulnerable populations? - recommend provider initiated testing and counselling (PITC) in all patient encounters? - recommend PITC for all pregnant women? - recommend PITC for most-at-risk and vulnerable populations? - support rapid testing with same day result provision? - support HIV testing and counselling (HTC) provided by community services? - support HIV rapid testing (point of care) done by lay or community workers?	

Antiretroviral therapy	What is the status of ARV guidelines revision? Provide year of last completed revision: Adult ART guidelines: year PMTCT guidelines: year Paediatric ART guidelines: year
	Have recommendations of the WHO 2013 Guidelines on the use of ARVs for the Prevention and Treatment of HIV been adapted in a national process? for each (1) yes, completed (2) on-going, (3) no, (4) specify Adult ART guidelines: PMTCT guidelines: Paediatric ART guidelines:
	What are the national ART target(s): Target number of people on ART and year: PMTCT ART coverage target: % by year
	If national guidelines recommend a CD4 threshold of 500, is there prioritization given to persons with a CD4 ≤ 350 or to those with advanced clinical disease? (1) Yes, (2) No, (3) Not applicable (e.g. country has not yet adopted CD4 threshold of 500), (4) if yes, please specify
	What are ART initiation criteria in infants and children? - Age cut-off to treat all children irrespective of symptoms: (1) < 5 years, (2) < 2 years, (3) other (specify) - CD4 thresholds in children aged 5 years and older who are asymptomatic: (1) regardless of CD4 count, (2) ≤ 500, (3) ≤ 350, (4) other (specify)
	Do national guidelines recommend ART for all HIV-infected patients with active TB? (1) Yes (2) No (3) Other (specify)
	Do national guidelines recommend ART for the HIV positive partner in sero-discordant couples? (1) Yes, (2) No, (3) specify
	Does the country use fixed-dose ART combinations in preference? (1) Yes, (2) No, (3) Other (specify)
	Regimen: Is TDF/3TC(FTC)/EFV the preferred 1st line ARV combination for adults and adolescents in the national guidelines? (1) Yes, (2) No
	Is there a policy to phase out D4T? (1) Yes (2) No (3) Other (specify)

Is AZT/3TC(FTC)/ATV/r(LPV/r) the preferred 2nd line ARV combination for adults and adolescents in national guidelines? (1) Yes, (2) No, (3) Other (specify_____) What is the preferred NRTI for children less than 3 years of age? (1) Abacavir (ABC), (2) Zidovudine (AZT), (3) Stavudine (d4T), (4) Other (specify Are LPV/r based-regimens preferred for all infants and children < 36 months (irrespective of NNRTI exposure) in the national guidelines? (1) Yes, all; (2) No, but recommended for NNRTI-exposed infants only, (3) Not recommended Is Efavirenz (EFV) recommended as the preferred NNRTI for children aged 3 years and older) (1) Yes, (2) No, (3) Other (specify_____) What is the recommended NRTI backbone for children aged 3-10 years? (1) TDF + 3TC (or FTC); (2) AZT + 3TC (or FTC); (3) ABC + 3TC (or FTC); (4) Other (specify_____ What is the recommended NRTI backbone for adolescents > 35kg and at least 10 years of age? (1) TDF + 3TC (or FTC); (2) AZT + 3TC (or FTC); (3) ABC + 3TC (or FTC); (4) Other (specify_____ Monitoring: Does the country use point-of-care CD4 technology? (1) What proportion of district hospitals have CD4 point of care? Provide an estimate ----- % (2) What proportion of primary health care facilities have access to CD4 point of care for testing their patients, whether on-site or nearby referral)? Provide an estimate -----% Service delivery: Which of the following service provision modalities are included in the ART national policy? ART provision in TB clinics by TB providers: specify: TB treatment in ART settings by ART providers ART provision in MNCH clinics by MNCH providers ART provision in settings providing opioid substitution therapy Community health workers engaged in ART patient support Hepatitis C diagnosis and management as part of HIV care Hepatitis B and Hepatitis C testing in ART clinics Hepatitis B vaccination provided at ART clinics Hepatitis C treatment provided in ART clinics Which of the following co-infection policies are in place? (Questions to be asked for adults and children): Isoniazid preventive therapy (IPT) for people living with HIV Intensified TB case finding in PLHIV TB Infection control for PLHIV

Co-trimoxazole prophylaxis

Prevention of mother-Do you have national plan for the elimination of MTCT of HIV? to-child transmission (1) Yes, if yes specify the MTCT transmission rate target(s) and year: _____ (2) no (3) if yes specify the elimination target(s) and year: _____ Do you have a national plan for elimination of MTCT of syphilis? (1) Yes, integrated with HIV or other elimination initiative (2) Yes, stand-alone (not integrated with HIV or other elimination initiative) (3) No national plan What is the current nationally recommended PMTCT option? Option A: Option B: if yes since Option B+: if yes since _____ If currently implementing Option A, is transition to option B/B+ planned? (1) yes, (2) no, (3) if yes in what year: ____ What is the current nationally recommended first line ART regimen for pregnant and breastfeeding women with HIV? (1) TDF/3TC(FTC)/EFV (2) other, please specify _____ What is the current nationally recommended PMTCT regimen, and duration, for exposed infants? Current nationally recommended PMTCT regimen Duration Is there a national recommendation on infant feeding for HIV-exposed infants? 1) Yes – breastfeeding (duration _____ months or unspecified ____) 2) Yes - replacement feeding 3) Yes – both recommended, left to individual choice or different settings 4) No If breastfeeding is recommended for HIV positive women and exposed infants, is the duration specified? ___ Yes No If Yes, please specify the duration in months: ()

Sexually transmitted Are there national STI treatment guidelines or recommendations? infections (STI) If so, what year were they last updated? Yes, If yes, year updated (____) No **Key populations** Which of the following key population or vulnerable groups are explicitly addressed in the national HIV policy or national plans? (tick box): men who have sex with men, transgender persons, sex workers. people who inject drugs, prisoners, adolescent key populations Which of the following components of the comprehensive package of HIV prevention, treatment and care interventions for sex workers are implemented in the country? (1) Comprehensive condom programming (yes/no) (2) HIV testing and counselling (yes/no) (3) Antiretroviral therapy and care (yes/no) (4a) Symptomatic STI treatment (yes/no) (4b) Asymptomatic STI treatment (yes/no) (4c) Periodic presumptive STI treatment (yes/no) (5) Comprehensive package of interventions for SW who inject drugs (yes/no) Empowerment of sex workers (participation in planning and implementation of HIV/AIDS/STI prevention and care activities) (yes/no) Which of the following components of the comprehensive package of HIV prevention, treatment and care interventions for men who have sex with men are implemented in the country? (1) Comprehensive condom programming (yes/no) (2) HIV testing and counselling (yes/no) (3) Antiretroviral therapy and care(yes/no) (4) Sexually transmitted infection (STI) prevention and treatment (yes/no) Comprehensive package of interventions for men who have sex with men who inject drugs (yes/no)

Male circumcision (only for 14 countries)	What is the current timeframe and target number of voluntary medical male circumcisions to achieve? Target: Year: What is the status of operational planning and monitoring? (tick boxes): Operational plan for 2014 exists Annual MC programme performance review conducted: if yes please specify in what year MC is integrated in infant care programmes: MC is linked with adolescent sexual reproductive health (SRH) What are the recommended medical male circumcisions methods? Conventional surgical methods (dorsal slit, forceps guided, sleeve resection) A prequalified device method has been approved for use, please specify:
Surveillance	Does the country carry out sentinel surveillance in special populations? if yes every years; number of sites, last survey in year (i) ANC attendees (ii a) sex workers? (ii b) people who inject drugs? (ii c) men who have sex with men? (ii d) transgender (iii) Other specific populations (please specify)
Monitoring and evaluation	What is the current status of planning for M&E of the HIV/AIDS health sector response? A national M&E plan exists: last update in year An review of the M&E system was conducted: year of last review, specify A review of the M&E system is planned:, in year, specify:
HIV Drug Resistance	Is national HIV drug resistance strategy in place? (1) yes, (2) no, (3) if yes, embedded in national HIV strategy? Has the country carried out HIV Drug Resistance (HIVDR) surveillance according to the following WHO protocols? for each: if yes, last in year, next in year 1. Transmitted drug resistance surveys 2. Pretreatment drug resistance surveys 3. Acquired drug resistance surveys 4. Paediatric drug resistance surveys 5. Monitoring of Early warning indicators for HIV drug resistance, if yes, number of ART clinics participating:

Toxicity monitoring surveillance	What is the status of national ARV toxicity surveillance? please specify: A national policy/strategy on ARV toxicity surveillance exists Toxicity surveillance activities are at pilot stage Toxicity surveillance activities are part of a national programme Toxicity surveillance data are integral part of M&E reporting within ART programme				
Strategic planning and review	What is the status of national HIV/AIDS Programme development (that includes HIV in the health sector)? The HIV national (health sector) strategic plan is in place, valid from: (year) to (year) The next HIV (health sector) programme review is planned for year: () Does the current national HIV [health sector] strategy address the following elements: a) achieving universal access to ART b) collaboration between HIV and other services including reproductive health c) strengthening health systems d) reducing inequities				
Reproductive Health and Research	In your country, do you have service delivery points providing appropriate medical and psychological care and support for women and men who have been raped & experienced incest? Appropriate medical and psychological care and support includes and is in accordance with the recommendations of the WHO clinical and policy guidelines - Responding to intimate partner violence and sexual violence against women (2013): Provision of first-line support or what is known as psychological first aid Provision of emergency contraception to women who seek services within 5 days Offer safe abortion if a woman is pregnant as a result of rape, in accordance with the national law Provision of STI and HIV post-exposure prophylaxis (within 72 hours of a sexual assault) as needed yes no				

Appendix 1. HIV/Hepatitis Indicators (EURO)

EUR15 Number of adults and children in HIV care who were screened for hepatitis B						
Rationale	HIV patients are often co-infected with HBV, notably in the WHO Europea Region, due to the same modes of transmission of HIV and HBV. Screeni of HBV informs physician strategy on patient management (recommendin vaccination against hepatitis B of uninfected and not vaccinated patients, or further evaluation and treatment of Hepatitis B). This is part of a comprehensive approach to the management of PLHIV promoted in the WHO European Region.					
What is measured	This indicator measures the number of people living with HIV enrolled in HIV care who were screened for HBsAg with the purpose of addressing patient's health needs regarding hepatitis B.					
Numerator	Number of HIV-positive adults and children in HIV care who were screened for hepatitis B using HBsAg tests during the reporting year.					
How to Measure and Measurement Tools	Calculated from clinical records of health care facilities which provide HIV/ AIDS treatment and care.					
Disaggregation	None requested.					
Strengths and weaknesses	The strength of this indicator is that it allows countries to monitor the extent to which HIV infected patients are being screened for hepatitis B – an intervention that is critical for assessing further needs related to the management of hepatitis B. Presence of HBsAg for a minimum of 6 months indicates chronic hepatitis B and informs clinicians on the need for further clinical and laboratory					
	evaluation and treatment. Knowing HIV/Hepatitis B status allows prescribing ARVs which are effective against both HBV and HIV infections.					
Additional considerations	Additional information regarding the number of adults and children in HIV care and screened for hepatitis B who were diagnosed with hepatitis B during the reporting period is also requested as part of this indicator. This data allows evaluating access to treatment among those who need it					
Data utilization	Look at trends over time. Useful information for clinical management and quality control in patient management.					
Data quality control and notes for the reporting tool	National Representativeness: if this indicator is only produced in a sub-set of facilities, comment should be added on the source of information, sample size and whether the information is representative of all sites where HIV/AIDS treatment and care delivered.					

EUR16 Percentage of HIV-positive hepatitis B cases eligible for hepatitis B treatment who received treatment for both hepatitis B and HIV				
Rationale	HIV patients are often co-infected with HBV due to the same modes of transmission of HIV and HBV. Co-infection rates are particularly high in the WHO European Region where a large proportion of HIV infections are related to injecting drug use. Treatment of hepatitis B in PLHIV has an impact on patients' quality of life, life expectancy and mortality. Some antiretroviral drugs are effective against both HIV and HBV viruses, which simplifies treatment of coinfected patients.			
What is measured	This indicator measures the number of HBV/HIV co-infected patients receiving treatment for both hepatitis B and HIV with effective ARVs for both viruses among patients enrolled in HIV care who were evaluated on hepatitis disease progression and found eligible for treatment.			
Numerator	Number of HIV-positive hepatitis B cases eligible for hepatitis B and HIV treatment who received treatment for both hepatitis B and HIV with effective ARVs for both viruses during the reporting year.			
Denominator	Number of HIV-positive hepatitis B cases who were eligible for both hepatitis B and HIV treatment during the reporting year.			
How to Measure and Measurement Tools	The numerator and denominator are calculated from clinical records of health care facilities providing HIV/AIDS treatment and care.			
Disaggregation	None requested.			
Strengths and weaknesses	The strength of this indicator is that it provides information on hepatitis B disease burden in PLHIV. It also allows monitoring access to hepatitis B treatment for PLHIV co-infected with HBV who are eligible for treatment.			
Data utilization	Look at trends over time. Useful information for clinical management and quality control in patient management.			
Data quality control and notes for the reporting tool	National Representativeness: if this indicator is only produced in a sub-set of facilities, comments should be added on the source of information, sample size and whether the information is representative of all sites where HIV/ AIDS treatment and care delivered.			

EUR17 Number of adults and children in HIV care who were screened for hepatitis C				
Rationale	HIV patients are often co-infected with HCV, notably in the WHO European Region, due to the same modes of transmission of HIV and HCV. Screening of HCV informs physician strategy on patient management (further evaluation and treatment of Hepatitis C if indicated or counselling on how to minimize risk of HCV infection in the future). This is part of a comprehensive approach to the management of PLHIV promoted in the WHO European Region.			

What is measured	This indicator measures the number of people living with HIV enrolled in HIV care who were screened for HCV a/b with the purpose of addressing patient's health needs regarding hepatitis C.						
Numerator	Number of HIV positive adults and children in HIV care who were screened for nepatitis C using HCV a/b tests during the reporting year.						
How to Measure and Measurement Tools	Calculated from clinical records of health care facilities which provide HIV/						
Disaggregation	None requested.						
Strengths and weaknesses	The strength of this indicator is that it allows countries to monitor the extent to which HIV infected patients are being screened for hepatitis B – an intervention that is critical for assessing further needs related to the management of hepatitis C. Presence of HCV a/b provides information on HIV/HCV co-infection rates, informs clinicians on need for further clinical and laboratory evaluation and treatment.						
Additional considerations	Additional information regarding the number of adults and children in HIV care and screened for hepatitis C who were diagnosed with hepatitis C during the reporting year is also requested as part of this indicator. This data allows evaluating access to treatment among those who need it						
Data utilization	Look at trends over time. Useful information for clinical management and quality control in patient management.						
Data quality control and notes for the reporting tool	National Representativeness: if this indicator is only produced in a sub-set of facilities, comment should be added on the source of information, sample size and whether the information is representative of all sites where HIV/ AIDS treatment and care delivered.						

EUR18 Percentage of HIV-positive hepatitis C cases eligible for hepatitis C treatment who received treatment for hepatitis C				
Rationale	HIV patients are often co-infected with HCV due to the same modes of transmission of HIV and HCV. Co-infection rates are particularly high in the WHO European Region where a large proportion of HIV infections are related to injecting drug use. Treatment of hepatitis C in PLHIV has an impact on patients' quality of life, life expectancy, and mortality.			
What is measured	This indicator measures number of HCV/HIV co-infected patients receiving hepatitis C treatment among patients enrolled in HIV care who were screened, evaluated on hepatitis disease progression and found eligible for treatment.			
Numerator	Number of HIV positive hepatitis C cases eligible for hepatitis C treatment who received hepatitis C treatment during reporting year			

Denominator	Number of HIV positive hepatitis C cases who were eligible for hepatitis C treatment during the reporting year					
How to Measure and Measurement Tools	The numerator and denominator are calculated from clinical records of health care facilities providing HIV/AIDS treatment and care.					
Disaggregation	None requested.					
Strengths and weaknesses	The strength of this indicator is that it provides information on hepatitis C disease burden in PLHIV. It also allows monitoring access to hepatitis C treatment for PLHIV co-infected with HCV who are eligible for treatment.					
Data utilization	Look at trends over time. Useful information for clinical management and quality control in patient management.					
Data quality control and notes for the reporting tool	National Representativeness: if this indicator is only produced in a sub-set of facilities, comments should be added on the source of information, sample size and whether the information is representative of all sites where HIV/AIDS treatment and care delivered.					

Appendix 2. Diagnosis of HIV/AIDS cases (PAHO)

P.1d Number of HIV cases diagnosed in 2012 , by sex for 2012				
What it measures	Number of HIV cases diagnosed in 2012 , by sex for 2012			

P.1e Number of AIDS	Number of AIDS cases diagnosed in 2012 and reported, by sex for 2012				
What it measures	Number of AIDS cases diagnosed in 2012 and reported, by sex for 2012				

79

Votes		
	 	

Notes				
				
				

Votes		





The purpose of these guidelines is to provide countries with technical guidance on how to measure the core indicators for the monitoring of the 2011 UN Political Declaration on HIV and AIDS. These guidelines provide technical guidance on the detailed specifications of the core indicators, on the information required and the basis of their construction, and on their interpretation. The guidelines also aim to maximize the validity, internal consistency and comparability across countries and over time of the indicator estimates obtained. In particular, the guidelines aim to ensure consistency in the types of data and methods of calculation employed.

UNAIDS Joint United Nations Programme on HIV/AIDS

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