# **Program Quality Overview**

**PROGRAMMATIC M&E LFA TRAINING** 

OCTOBER 2019 - JANUARY 2020

GENEVA, SWITZERLAND



#### 1. Program Quality and the Role of the LFA

#### **Presentation outline:**







Planning and Implementation of Programmatic Spot Checks & Targeted HFA



**Expected outputs** 



Spot light- New Spot Check: AGYW



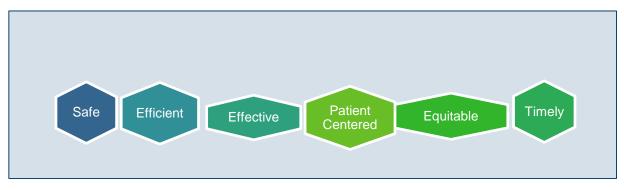


Case study – Review & Discussion

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#### What is Program Quality?

The degree to which health programs increase the likelihood of desired outcomes



- Poor quality health programs are an impediment to achieving Global Fund's objectives & waste resources

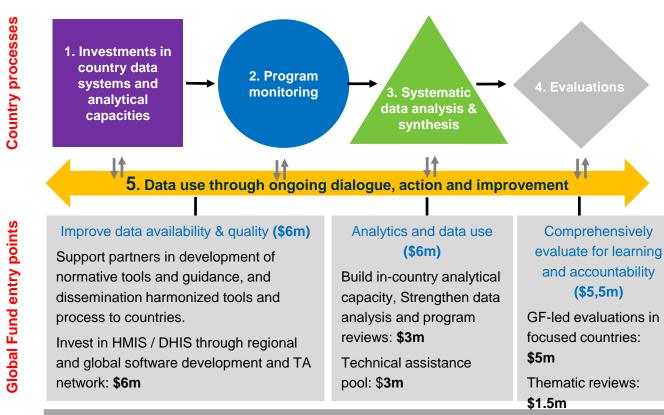
- *Mainstreaming program quality is a key component of the Global Fund strategy*
- The Global Fund is committed to improving program quality to <u>maximize</u> impact at the country level

Identifying and addressing **critical bottlenecks** at different levels of country systems

Identifying and replicating **best practices** leading to better health outcomes

Assurance, monitoring and continuous quality improvement to drive impact

#### 1. Data Use For Action and Improvement (DUFAI) Framework



Coordination with partners

#### Concept on program quality

- Program quality has many dimensions and is complex to measure
- During the grant cycle, program quality assessments are conducted to obtain insight into the quality of GF-supported programs
- A combination of assessment methods as appropriate to the country context are used to monitor program quality and identify issues, define activities for program improvement and follow up on implementation of ongoing quality improvement measures

#### 1. Program Quality

#### The Global Fund supports program monitoring to:

To track program performance To assist in effective management To inform and support timely decision-making through systematic collection of data alongside program implementation Assess adherence to established standards and procedures for program quality Plan for future risk mitigation and assurance activities

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#### 1. Program Monitoring

How does the Global Fund support countries in program monitoring?

- By participation in global efforts to harmonize monitoring frameworks and indicators for unified data collection and reducing reporting burden on countries.
- By the development and application of M&E frameworks for KPs, AGYW, measuring human rights-related barriers to access services and for COEs
- By Strengthening and using existing country reporting platforms to access real time data
- By using a set of tracer indicators to assess program quality.

Each indicator is given a score using a scale of 0 to 4 based on the results, and a composite score is generated resulting in a program quality tracer rating : very high/ high/ moderate / low program quality

#### **Program Quality: HIV**

Components

PLHA know their status: % of estimated people living with HIV knew their positive status

ART coverage: % estimated people living with HIV currently on ART (adults and children)

12 month ART retention: % of people who ever initiated ART are still on ART at 12 months after ART initiation (adult and children)

Viral suppression: % of people who retained in ART for at least 6 months with viral load below 1,000 copies/ml

The sources of the data for these components are either:

(1) Data quality reviews; (2) country evaluations; (3) data quality spot checks; (4) national or disease specific program review; (5) routine programmatic analysis; (6) partner reviews; (7) review of data systems (community/facility); (8) thematic reviews; systematic data quality checks; (9) prospective country evaluations; (10) National program reports (UNAIDS Global AIDS Monitoring, WHO ART Report, Global Cascade Workshop); and (11) national representative; surveys (PEPFAR Population-based HIV Impact Assessment, DHS, AIS); results from PU/DR

#### **Program Quality: Malaria**

Malaria program quality is assessed as a composite of three indicators which are rated according to their performance.

| Component   | Main Data Source / means<br>of implementation | Frequency of<br>Implementation in High<br>Impact and Core countries |
|---|---|---|
| ITN Use: Proportion of population that slept<br>under an insecticide-treated net the previous<br>night                            | MIS or MICS or DHS                            | on average every 2 years  |
| Diagnostic coverage: proportion of suspected malaria cases that receive a parasitological test                                    | HFA, QoC                                      | 1-3 years   |
| IPTp Coverage: % of women who received at least<br>3 doses of IPTp for malaria during ANC visits<br>during their last pregnancy** | HFA, QoC                                      | 1-3 years   |

#### **Program Quality: TB**

| agnosed, new and     |
|----------------------|
| accessfully treated  |
| fully treated (cured |
|                      |

The sources of the data for these components are either:

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### 2. Approaches to Program Quality Assessment: Options

| Assurance   | Main service provider   | Assurance   | Service provider   |
|---|---|---|--|
| Review of data systems<br>(community/ facility)                 | Country led (TA as needed);<br>Quality assurance by identified QA service<br>provider/LFA                                     | Partners review   | Partners   |
| Program quality/ data quality spot checks                       | LFA   | Country evaluations   | GF-led with service<br>provider in focused<br>countries                    |
|   |   |   | Country led with TA if needed  |
| Health facility assessment                                      | Country led (TA as  | Thematic reviews  | Secretariat-led with   |
| (national/targeted)   | needed) Targeted HFA<br>by LFA/ identified service provider   |   | service providers)   |
| Data quality reviews  | Country led (TA as needed);   | Prospective Country   | Secretariat (TERG-led)   |
| (national/targeted)   | Quality assurance of national DQR by<br>identified QA service provider<br>Targeted DQR by LFA/ identified service<br>provider | Evaluations   | with service providers   |
| Review of Laboratory systems                                    | Country led (TA as needed);<br>Quality assurance by identified QA service<br>provider/LFA                                     | Population-based surveys<br>(Examples: IBBS, PSE for<br>KP, TBPS, MIS | Country led (TA as<br>needed) or partner<br>contracted service<br>provider |
| Routine programmatic analysis<br>(Examples: Service Cascade-HIV | Country led (TA as needed) or identified service provider in some scenarios   | Community based monitoring  | Country led (TA as needed)   |
| Program reviews   | Country led (TA as needed) with support of identified service provider in some scenarios                                      |   |  |

#### Harmonized Health Facility Assessment Approach

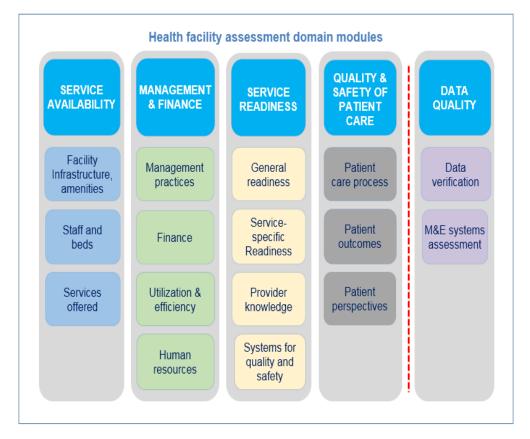
- Joint stakeholder process to harmonize existing health facility assessment implementation, led by WHO and supported by the Global Fund, the World Bank, USAID, GAVI, UNICEF and several other agencies.
- Harmonized indicators, definitions, standard questions (question bank)
- Coordinated implementation at country level and with partners to avoid duplication of efforts and to improve results.
- Improved comparability of results across time, geographical location

# WHO HFA(SARA) Modular approach enables customization of the survey to the specific country context

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#### 2. What is a Health Facility Assessment ?

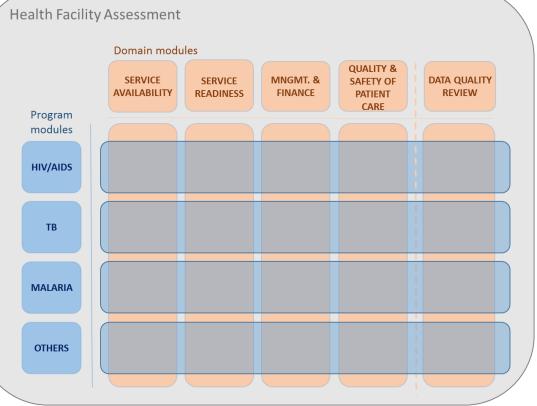
- WHO recommends countries conduct the assessment as a routine aspect of a country's strategic planning cycle.
- Generates evidence to support the planning and managing of a health system.
- Assessment tool designed to assess and monitor the service availability and readiness of the health sector, quality of care, data quality, management
- Address the following key questions(among others) on health services:
  - What is the availability of different health services in a country?
  - To what extent are facilities prepared to provide specific health services?
  - To what extent does the service delivery process follow generally accepted standards of care?



#### HFAs are used by countries for their own strategic planning, but also provide assurances to donors

#### 2. What is a Targeted Health Facility Assessment ?

- While health facility assessments are developed for larger scale implementation, these tools can be used for a more focused assessment with a smaller and targeted sample.
- Country teams can decide to use specific modules
  (e.g. HIV, TB or Malaria) or some specific domains of a module (service availability, service readiness, management and finance or quality of care) depending on the country context.



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#### 2. WHO SARA

Introduction to Service Availability and Readiness Assessment

https://www.who.int/healthinfo/systems/sara\_introduction/en/

Description: The current WHO Service Availability and Readiness HFA survey "plus" additional selected quality of care indicators/questions

• SARA Reference Manual contains the instrument & indicators:

http://www.who.int/healthinfo/systems/sara\_reference\_manual/en/

• SARA Implementation Guide contains guidance on the planning, methodology, sampling, questionnaire adaptation, electronic tools, data collection, supervision, data processing, analysis and reporting: <a href="http://www.who.int/healthinfo/systems/sara\_implementation\_guide/en/">http://www.who.int/healthinfo/systems/sara\_implementation\_guide/en/</a>

#### 2. Comparison of a nationally representative HFA and the Targeted HFA

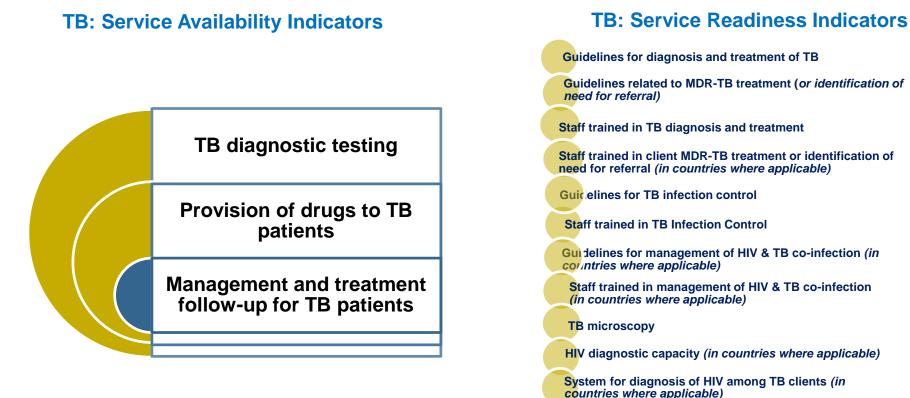
|                         | Nationally representative HFA   | Targeted HFA  |
|-------------------------|---|---|
| ΤοοΙ                    | Harmonized Health Facility Assessment tool<br>-or other HFA tool that includes core indicators and<br>quality of care (e.g. SARA plus, SPA, SDI)  | Harmonized Health Facility Assessment tool<br>-or other HFA tool that includes core indicators and<br>quality of care (e.g. WHO SARAplus) |
| Components              | Entire Health Facility Assessment survey tool<br>-Comprehensive assessment includes general and<br>specific services (e.g. RMNCH, HIV, TB, malaria, non-<br>communicable diseases, etc.)<br>-Include all facility management, supervision, HR,<br>finance, lab, pharmacy, etc. sections | Only the HIV, TB and/or malaria components of the Health Facility Assessment survey tool  |
| Sample Size             | Nationally representative (census or representative sampling)   | Will vary, but generally 20 - 40 sites  |
| Inference of<br>results | Nationally representative   | Will vary based on sampling used if inference is possible beyond the sites visited and to what extent                                     |

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### 3. Comparison of Planning a National and Targeted HFA

| Nationally HFA   | Targeted HFA  |
|--|---|
| Global Fund with or without Partners; Partners   | Global Fund Country Team  |
| Coordinating group of country stakeholders   | LFA M&E team in collaboration with the ,PR, MoH and in-country stakeholders, i.e, WHO   |
| Country-led with TA as needed  | LFA M&E team  |
| Census or representative sampling, requires Master Facility<br>List  | Will vary, but generally 20 - 40 sites  |
| <ul> <li>Select modules based on objectives;</li> <li>Guidance</li> <li>May Adapt questionnaires based on objectives</li> <li>Data Entry CSPRO or similar</li> <li>Data Analysis Excel Workbook</li> <li>Develop a data analysis plan</li> </ul> | <ul> <li>Focus on HIV, TB &amp; Malaria services &amp; Quality of Care ;</li> <li>Guidance</li> <li>May adapt Questionnaires based on objectives</li> <li>Data Entry -Excel sheet</li> <li>Data Analysis-Excel sheet</li> <li>Develop a data analysis plan</li> </ul>   |
| <ul> <li>✓ Training</li> <li>✓ Pilot &amp; Modification as needed</li> <li>✓ Conduct Actual Survey</li> <li>✓ Data Management</li> <li>Yes</li> </ul>  | <ul> <li>✓ Conduct Actual Survey</li> <li>✓ Data management</li> <li>Will vary, but generally 20 - 40 sites</li> </ul>  |
| Nationally representative  | Will vary based on sampling used if inference is possible beyond the sites visited and to what extent   |
|  | Global Fund with or without Partners; Partners         Coordinating group of country stakeholders         Country-led with TA as needed         Census or representative sampling, requires Master Facility<br>List         ✓       Select modules based on objectives;         •       Guidance         •       May Adapt questionnaires based on objectives         •       Data Entry CSPRO or similar         •       Data Analysis Excel Workbook         •       Develop a data analysis plan         ✓       Training         ✓       Pilot & Modification as needed         ✓       Conduct Actual Survey         ✓       Data Management |

#### 4. Outputs - Priority SARA Indicators: TB



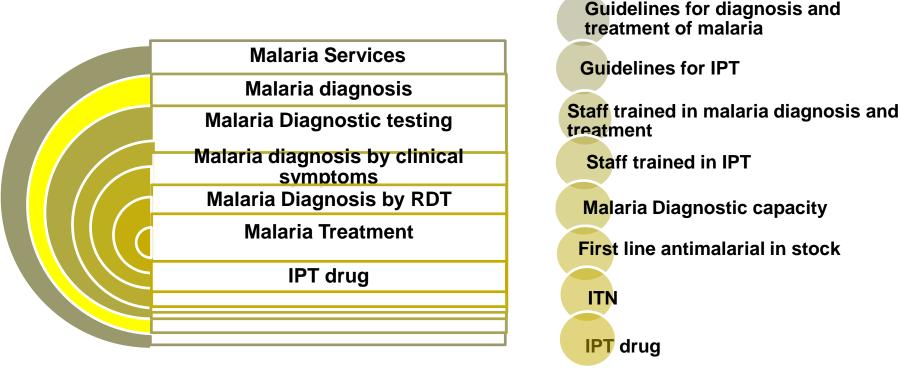
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First-line TB medications in stock

4. Outputs- Priority SARA Indicators: Malaria

#### Malaria: Service Availability Indicators

**Malaria: Service Readiness Indicators** 



#### 4. Outputs- Priority SARA Indicators: HIV

| HIV: Service Availability Indicators                    | HIV:Service Readiness Indicators (1)<br>for discussion with Disease Advisors | HIV:Service Readiness Indicators (2)<br>For discussion with disease advisors |
|---|--|--|
|   | Guidelines on HIV Counseling & Testing                                       | Staff trained in ART prescription and management                             |
|   | Guidelines for clinical management of HIV                                    | CD4 or Viral load  |
| HIV Counselling & Testing                               | Staff trained in HIV Counselling and testing                                 | Three first-line Anti-Retrovirals  |
| HIV/AIDS care and support services                      |  |  |
| Treatment of opportunistic infections                   | Staff trained in clinical management of HIV                                  | Dried blood spot (DBS) filter paper for                                      |
| Intravenous treatment of fungal infections              |  | diagnosing HIV in newborns   |
| Care for Paediatric HIV/AIDS patients                   | HIV Diagnostic capacity  | Nevirapine (NVP) syrup   |
| ARV prescription or ARV treatment follow<br>up services |  |  |
| Treatment follow-up services for persons<br>on ART      | System for diagnosis of TB among HIV+ client                                 | System for diagnosis of TB among HIV+<br>clients                             |
|   | IV treatment fungal infections   | Maternal ARV prophylaxis   |
|   | Guide/nes for Antiretroviral therapy,  | Zidovudine (AZT) syrup   |

Guidelines for Antiretroviral therapy,

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#### **Existing LFA Programmatic Spot Checks**

| HIV   | ТВ  | Malaria   | AGYW  | PWID   | DHIS   | M&E Plan &<br>Implementati<br>on                    | Supervision  | Training   | HPM related                                    |
|---|---|---|---|--|--|---|--|--|--|
| 1. Community<br>based HIV<br>testing among<br>key populations | 2. Community<br>based TB<br>activities<br>(including MDR-<br>TB where<br>appropriate) | 3.Implementation<br>of malaria vector<br>control<br>interventions<br>(LLINS<br>distribution and<br>IRS) | <b>4</b> . Adolescent<br>Girls and<br>Young Women<br>programs | 5.Assessment of<br>implementation<br>of Opioid<br>Substitution<br>Therapy<br>program | 6. DHIS<br>implementation<br>and hospital<br>data recording/<br>coding system                                    | 7. Assessment<br>of M&E plan and<br>implémentations | 8.<br>Assessment<br>of supervision<br>effectiveness<br>in a given<br>health<br>program | <b>9.</b> Training activities and related expenses | <b>10</b> . HPM Supply<br>Management<br>Review |
| 11. Prevention<br>services among<br>key populations           | 12. HPM<br>Procurement<br>Review  | 13. Community<br>based malaria<br>case<br>management<br>activities                                      | 14.<br>M&E Ass<br>essment                                     | <b>15.</b> Assessment<br>of training<br>activities and<br>related<br>expenses        | <b>16</b> . HPM Quality<br>Monitoring for<br>Pharmaceuticals<br>HPM Quality<br>Monitoring for<br>Pharmaceuticals | 17. HPM LMIS Im<br>Review                           | plementation   | <b>18</b> . HPM ACT<br>co-payment mechar<br>Buyers | isms First Line                                |

# **Planning Programmatic Spot Checks**

Country Team request LFA to implement Spot checks based on risk, country context and assurance needs

## 2

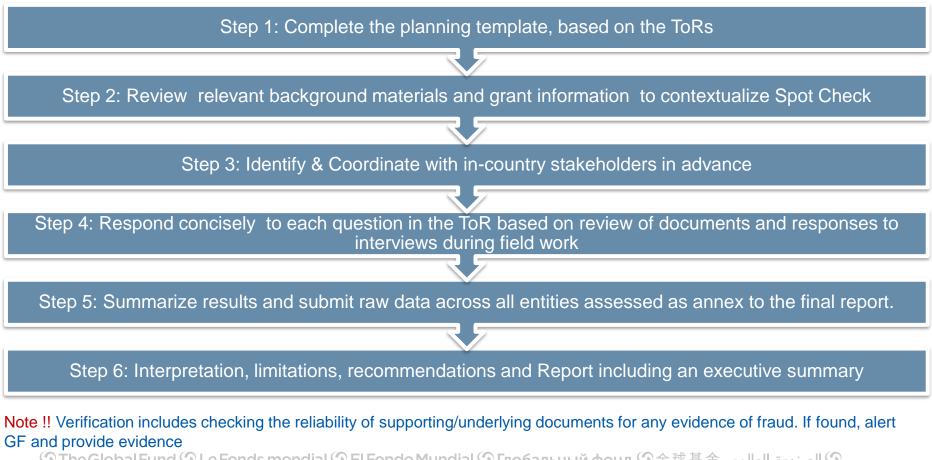
Based on knowledge of the local context, LFA can also recommend a need for a Spot Check

3

Standard Terms of Reference for Spot Checks developed; may be customized based on objectives and context LFA & Country team to agree on the Scope of Work, Sampling approach, Experts and LoEs before Spot check starts

4

#### Implementing a Spot Check



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LFA is expected to analyze data across all facilities and across all key informants interviewed. This applies to both qualitative and quantitative data.

#### For qualitative data:

(i) conduct a <u>thematic analysis</u> and present synthesized findings across the individuals interviewed or facilities assessed

(ii) focus on examining data and recording emerging patterns or themes within the data to explain key issues and to serve as basis for recommendations.

#### **For quantitative data:**

(i) for categoric responses[ yes/no/partially]: Use frequency tables & charts

(ii) where results are presented in %, provide the counts (n) : # of sites/ people interviewed

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Step 6: Classify findings into Major and Minor issues and list them in order priority

| Major issue   | Minor Issues:   |
|---|---|
| Significant gaps that pose <u>major risks</u> to the successful implementation of the assessed activity | The identified gaps pose <u>minor risks</u> that can be managed and/or strengthening measures can be implemented within a short timeframe |

The LFA's recommendations should be **specific**, **relevant** and **realistically achievable** in the implementation context. The LFA should provide an **appropriate timeframe** for the implementation of each recommendation

#### 3. Linking SMART recommendations to Data Use for Programme Improvements Specific, Measurable, Achievable, Result-oriented, Time-bound

| 1. PCU has no supervision plan  | PCU to develop a supervision plan to<br>enhance the supervision of Disease<br>Programmes and Implementation<br>Partners.                         | By 30 March,<br>2020 | PCU  |
|---|--|----------------------|--|
| 2. LFA found HIV tests' data discrepancy<br>between Laboratory register and<br>Monthly Integrated Activity Report | LFA recommends that health facility in-<br>charge should make sure the correct<br>figures are transferred from Lab<br>register to Monthly Report | By June, 2020        | Health Facility in-charge<br>and BPHS/EPHS<br>implementers |

3. Developing comprehensive supervision policies and plans to guide the support supervision activities for Community TB interventions, covering both the community sputum collection points and Community mobilization and demand. The documents should articulate the different types of SS visits and objectives of each visit i.e. programmatic, M&E, financial etc.

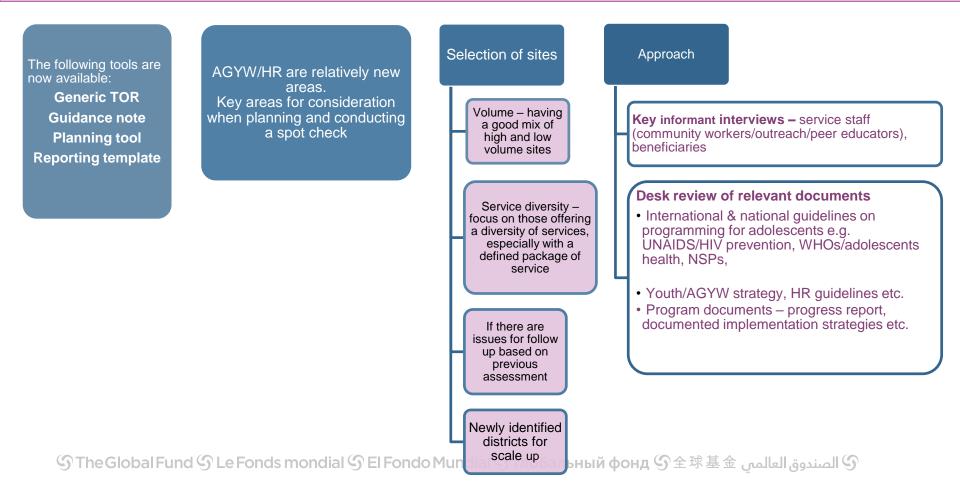
4. The integrity of data at the ABCD site is questionable, with 100% of the documents reviewed having at least one data inconsistency/ error. There should be remedial training and close monitoring by the PR of CPAI PEs on adherence to good data recording practices

Programmatic Spot Checks Spotlight:

# 1) Adolescent Girls & Young Women 2) Human Rights

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#### Key recommendations: Planning and Implementation



#### Key areas for consideration at implementation

1. Mapping of available service providers targeting AGYW/Youth

- Accessibility & services provided
- Coordination and how that is supporting referrals/linkage.

2. Targeting and enrollment of AGYW into the program – vulnerability assessment where applicable, risk assessment etc

• Is there a structured process of continually identifying needs of AGYW, and whether the program is addressing those needs in a timely manner.

3. Defined package of service provided and linkage to national guidance where this exists.

- Link to reporting YP 2 indicator how is this computed? When is a beneficiary counted as "reached" and reported in YP 2 indicator?
- What is the definition of "reached"?

4. Are services tailored to specific age groups based on country led guidance?

• Community strengthening - are there activities/services provided to families, households & communities.

5. Compliance/adherence to interventions – availability and use of SOPs/guideline for delivering interventions

6. Relevance and appropriateness of service/s provided and if aligned to expectations and needs.

- Individual beneficiaries
- Households, families and communities SEI Fondo Mundial 分 Глобальный фонд 分全球基金 الصندوق العالمي の

#### Key areas for consideration at implementation level

## Adequacy of M&E system with regards to the following:

Availability of tools that support identification & enrollment of vulnerable AGYW into the program, track services provided & reporting, changes occurring to the beneficiaries, clarity & understanding of tools by providers

- > When individuals are counted as being "reached" depending on the intervention provided
- Reporting of unique individuals who receive a defined package of service, is there a system to address potential double reporting within the program?
- Design of data collection and reporting tools among different service providers and alignment to a particular system to avoid double counting; support the counting of AGYW and youth having received multiple services
- Monitoring of referrals and linkages made to services not offered by the PR/SR
- Mechanisms that ensure data quality at all levels
- > Data use is there evidence that data is analyzed and use beyond reporting to the next level?

#### Human Rights(HR) areas for consideration at implementation level

| 1. Availability of systems and structures that<br>support implementation of HR programs;<br>dedicated staff, capacity building for the staff etc. | 6. Implementation modality – is this effective in reaching, identifying and addressing HR issues |
|---|--|
| 2. Specific interventions/services delivered  | 7. Availability and use of tools to track services/activities                                    |
| 3. Which populations/individuals are targeted?  | 8. Clarity for use by providers  |
| 4. How are beneficiaries identified?  | 9. Effectiveness in tracking progress  |
| 5. Identify areas for improvement   |  |

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# Recommendations

- Quality Recommendations are a fundamental part of program monitoring process
- Taking sufficient time in developing them is essential for many reasons:

i) Without recommendations, a report has reduced chance of being used

ii) Recommendations are the result of the LFA Spot Check & analysis

- iii) They define and prioritize actions for program improvements
- iv) Recommendations should make a constructive contribution to the correcting the identified issue

v) Recommendations should form the basis for follow-up

#### SMART RECOMMENDATIONS

Specific: Address one specific issue and one or more specific related actions

Measurable: For the future, to assess the extent to which a recommendation has been implemented. Add baseline information and an indicator in the report for future comparison

Achievable: Highlight what must be done within reason; should possible in practical terms

Results-oriented: Actions suggested should be designed to lead to a concrete result Time-bound: Important for prioritization, create pressure for action, enhances accountability

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#### Group Work: Critique of Programmatic Spot Check Report

#### Step 1

 Quick review of objectives, ToR, Background, Methodology, Findings & recommendations of your respective area

#### Answer these questions

- Based on your experience, what do you think about the quality of the Spot Check?
- How well does the report address the objectives?
- Are the recommendations linked to the Findings?
- Are the recommendations SMART?

#### Groups

- Group 1: Community TB
- Group 2: Female Sex Workers(FSW)
- Group 3 Adolescent Girls and Young Women(AGYW)



October 2019

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## Experience with assessing QOC using RR during HFA

| COUNTRY         | SAMPLE <sup>2</sup><br>(all @ 5<br>records/observa<br>tions per<br>facility) | TOOLS USED   |                    |              |             |         |     |     |                          |  |
|-----------------|--|--|--------------------|--------------|-------------|---------|-----|-----|--------------------------|--|
|                 |  | ANC  | PMTC<br>T w<br>ANC | PMTC<br>T PP | Malari<br>a | ТВ      | HTS | ART | Child<br>obser<br>vation |  |
| ZAMBIA          | 8 facilities (60-<br>70)   | ~  | ~                  | ~            | ✓           | ✓       | ✓   | ✓   | ✓                        |  |
| Liberia         | (44+ facilities)<br>(250-400 patients<br>per topic)                          |  |                    | √            | √<br>child  | Partial |     | √   |                          |  |
| Sierra<br>Leone | 138 facilities/690<br>patients   | <ul> <li>✓</li> <li>Added<br/>newborn<br/>and<br/>delivery<br/>care</li> </ul> |                    |              | V           | V       | V   |     |                          |  |
| Burkina<br>Faso | @80 FACILITIES<br>300 PATIENTS   |  | ✓                  |              | √           | √       |     | √   |                          |  |
| Nigeria         | 3300 facilities<br>1-4 records   | ✓  | ✓                  |              | ✓           | ✓       |     | ✓   | ✓                        |  |

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### **TOOLS FOR RR QOC**

- Record review of documented care process for individual patient
  - HIV/ART/HTS;
  - Malaria/IPT;
  - TB;
  - MNH (PMTCT for ANC);
  - PMTCT for HIV positive women and exposed infants
  - partograph review for delivery care,
  - Outpatient care for sick child (focus pneumonia/diarrhea/malaria);
  - VMMC;
  - Outpatient care for chronic conditions (hypertension, diabetes)[not yet tested]

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## **QOC INDICATORS AND METHODS FOR HFA (1)**

### <u>RECORDED</u> EVIDENCE THE PATIENT CARE PROCESS IS IN ADHERENCE TO STANDARDS

- What examinations, tests, diagnosis, treatment, and counseling are recorded for individual patients
  - Measured by reviewing *recorded* information for individual patients

### Strengths

- Can select sample based on the types of cases desired
- Promotes importance of recording patient information—critical for continuity of care, sharing information between providers
- Can assess care over time for chronic/long-term treatment conditions (e.g., TB, HIV)

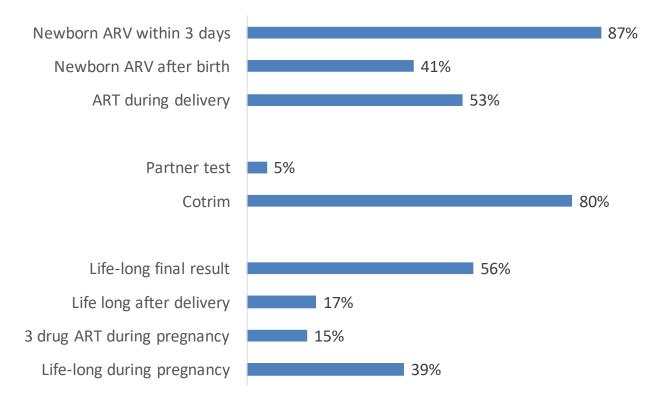
### Limitations

- Depends on availability of records and completeness of recording
- Must assume what is recorded actually occurred

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## **EXAMPLES OF FINDINGS (1)**

#### Liberia 2017 PMTCT positive women (n=176)



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## **EXAMPLES OF FINDINGS (2)**

#### Table A7.b1 Number of observed patients where the indicated suspect malaria eligibility criteria was met

| Total<br>observed<br>sick |          | Patients<br>meeting<br>criteria for | Among suspect malaria patients                    |                                      | Among<br>patients with<br>malaria test                       | Among patients            | All<br>treated<br>correctl  |                         |  |     |
|---------------------------|----------|-------------------------------------|---|--------------------------------------|--|---------------------------|---|-------------------------|--|-----|
|                           | children | suspect<br>malaria <sup>1</sup>     | Observation<br>identifies<br>patient had<br>fever | Any<br>malaria<br>test<br>prescribed | prescribed,<br>test result<br>received prior<br>to departure | Positive test<br>recorded | Among<br>positive test<br>results,<br>those<br>treated<br>correctly | Negative<br>test result | Amon<br>g<br>negati<br>ves,<br>those<br>treated<br>correct<br>ly | У   |
|                           | (a)      | (b)                                 | (c)   | (d)                                  | (e)  | (f)                       | (g)   | (h)                     | (i)  | (j) |
|                           | Ν        | N (%)                               |   | N (%)                                | N (%)  | N (%)                     | N (%)   |                         |  |     |
| Zambia                    | 62       | 7F<br>50 O                          | 94%   | 55%                                  | 73%  | 0%                        | na  | 100%                    | 100%   | 45% |
| Nigeria                   | 3132 O   | 1146 F<br>3027 O                    | 88%   | 63%                                  | 63%  | 51%                       | 52%   | 49%                     | 43%  | 46% |

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# **EXAMPLES OF FINDINGS (3)**

| Table A5.1 Antiretroviral therapy  |                           |                            |                      |      |        |                       |                          |                          |                                   |                           |  |
|--|---------------------------|----------------------------|----------------------|------|--------|-----------------------|--------------------------|--------------------------|-----------------------------------|---------------------------|--|
|  | of of                     | Number<br>of<br>facilities | Months patients have |      |        | Among all patients    |                          |                          |                                   |                           |  |
|  |                           |                            | been on ART          |      |        | Confirmator<br>y test |                          | CD4 within past 6 months | Viral load<br>at least            | Most<br>recent            |  |
|  |                           |                            | 0-6                  | 7-12 | >12m   | ,                     | 1 <sup>st</sup> 2 months |                          | once                              | VL non-<br>detectab<br>le |  |
|  | (a)                       | (b)                        | (c)                  | (d)  | (e)    | (f)                   | (g)                      | (h)                      | (i)                               | (j)                       |  |
| Sierra<br>Leone  |                           |                            |                      |      | INFORM | IATION NOT COLLECTED  |                          |                          |                                   |                           |  |
| Zambia   | 70                        | 7                          | 0                    | 100% | 0      | 79%                   | 79%                      | 44%                      | 7%                                | 4%                        |  |
| Liberia  | 187                       | 44                         | 79%                  | 18%  | 4%     | 96%                   | 11%                      | 71%                      | 10%                               | 2%                        |  |
| Burkina<br>Faso  | 178                       | 62                         | 1%                   | 7%   | 92%    | 98%                   | 73%                      | 27% <sup>1</sup>         | 7%<br>VL6m:<br>6%<br>VL12m:<br>6% | 6%                        |  |
| Nigeria  | INFORMATION NOT COLLECTED |                            |                      |      |        |                       |                          |                          |                                   |                           |  |
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### **ISSUES FOR RECORD REVIEW: DRAWING SAMPLE**

- □ ANC, delivery, PMTCT, HIV, TB all have registers where eligible cases are easy to identify
  - Problem if ANC records are not maintained at facility—then must take

opportunistic sample of patients present day of survey

- Difficulties for outpatient curative care (malaria and sick child)
  - □ Some facilities do not maintain patient records at facility
  - U Where records are maintained:
    - Information in register varies—some have presenting symptoms,

some final diagnosis, some neither

Patients (particularly sick children) may have multiple diagnoses

(see following slide)—this must be taken into account during analysis

## **ELIGIBILITY DIAGNOSIS AND ADDITIONAL DIAGNOSES FOR CHILDREN**

| Table 1.2 Comparison of records selected from outpatient register, that met the criteria for the indicated diagnosis and the diagnoses noted in the patient record |                       |  |            |                      |                    |           |                 |                         |  |  |
|--|-----------------------|--|------------|----------------------|--------------------|-----------|-----------------|-------------------------|--|--|
| selected for reco  | Total patient records | Diagnoses noted in the reviewed patient record |            |                      |                    |           |                 |                         |  |  |
|  | selected              | Malaria  | Pneumonia  | Other<br>respiratory | Watery<br>diarrhea | Dysentery | Other diagnosis | noted in patient record |  |  |
|  | а                     | b  | С          | d                    | е                  | f         | g               | Н                       |  |  |
| Suspect<br>malaria   | 58                    | 33 (56.9%)                                     | 7 (12.1%)  | 13 (22.4%)           | 6 (10.3%)          | 1 (1.7%)  | 22 (37.9%)      | 4 (6.7%)                |  |  |
| Respiratory<br>illness   | 59                    | 4 (6.8%)                                       | 13 (22%)   | 38 (64.4%)           | 2 (3.4%)           | 1 (1.7%)  | 7 (11.9%)       | 5 (8.5%)                |  |  |
| Diarrheal<br>illness   | 59                    | 0 (0%)   | 3 (5.2%)   | 10 (17.2%)           | 50 (86.2%)         | 3 (5.2%)  | 5 (8.6%)        | 4 (6.9%)                |  |  |
| Total  | 175                   | 37 (21.1%)                                     | 23 (13.1%) | 61 (34.9%)           | 58 (33.1%)         | 5 (2.9%)  | 34 (19.4%)      | 13 (7.4%)               |  |  |
| <sup>1</sup> More than one diagnosis might have been recorded  |                       |  |            |                      |                    |           |                 |                         |  |  |

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### **ISSUES: ELIGIBILITY CRITERIA**

□ PMTCT, HIV, and TB

- To capture process of care for TB, HIV, HIV + women and exposed infants: the patient must have been under treatment for a specific amount of time (or be a number of weeks postpartum)
- Eligibility information must be captured for each record. Do not assume sample records all met the official eligibility criteria
- If management of positive cases (ANC + for HIV or + for malaria) are desired, the sample should specify positive cases otherwise sample may be too small to provide a good picture of care
- □ If different management of children and adults is a concern, the sample should specify children or adults, otherwise sample may be too small to provide a good picture of care

## **ISSUES WITH COLLECTING INFORMATION**

- Reading charts—abbreviations and illegible handwriting
  - Facility nurses are very helpful in reading charts—they know individual doctor's writing and abbreviations
- Time required for finding sample records: varies by organization of facility.
  - □ In general this was not a problem
  - Data collectors must identify replacements when records are not found

## **ISSUES WITH ANALYZING INFORMATION**

□ Want to <u>use facility as unit of measure</u> (average percentage/measure for the facility and then calculate the average across district/regional/national facilities)

### Rational:

- Eliminates bias if sample sizes vary—equal representation of patient care in large volume and small volume facilities
- Eliminates bias if distribution of positive and negative test results (malaria and HIV) are skewed by facility—picture of care of positives may primarily be from a few facilities.
- Except for outpatient curative care—facility practices are similar across patients and providers

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- Found many problems with presenting results using different denominators (subsets of subsets)
- Either need clear tables that show how the denominator changed or the same denominator for items in graph
  - One country had a graph showing high percentages of HIV exposed infants being tested and results available; e.g., (illustrative)
    - ✓ 20 HIV + women records reviewed
    - ✓ 10 infants came to PMTCT clinic postpartum
    - $\checkmark$  Blood for PCR test drawn on 5 infants
    - ✓ Results returned for 4
    - ✓ Presented result: 40% of exposed infants had blood drawn and results returned: actual is 20%
    - $\checkmark$  No explanation for the 10 infants never brought to clinic

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Encourage local supervision to use the tools as a guide to assess care

Reinforces to providers and supervisors what the standards are

Education to managers/supervisors how to use information

Comparing QOC results with readiness results to help identify source of problem with QOC –lack of guidelines? Lack of tests or drugs? Non-trained staff?

Develop job aids to reinforce the standards being assessed poster where patient and provider can see key points;

# Back up slides

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