

Government of Malawi Ministry of Health

Integrated HIV Program Report October – December 2015

- Integrated HIV Program Supervision
- HIV Testing and Counselling / Early Infant Diagnosis
- Blood Safety
- Post Exposure Prophylaxis
- HIV Exposed Child Follow-Up
- Pre-ART
- Prevention of Mother to Child Transmission /
 Antiretroviral Therapy
- TB/HIV
- Sexually Transmitted Infections
- Supply of HIV Program Commodities

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1 Executive Summary

A summary of the key achievements between **October and December 2015** is provided below:

- Scale-up of integrated HIV services had reached the following number of sites:
 - o 724 static (579 within and 145 outside of health facilities) and 188 outreach HTC sites
 - o 716 (static) ART sites
 - o **617** PMTCT sites (Option B+, all included in ART sites above)
 - o 668 Pre-ART sites
 - 652 sites with HIV-exposed child follow-up
- 604,854 persons were tested for HIV and received their results; 172,574 (29%) accessed HTC for the first time; 432,280 (71%) were repeat testers and 15,096 (3%) of these received confirmatory testing (after having tested positive in the past). This is equivalent to 63% confirmatory testing coverage among 24,082 patients initiating ART this quarter. 32,762 (5%) clients received a positive result for the first time.
- 23,692 (>99.9%) of 23,699 blood units collected were screened for (at least) HIV, hepatitis B and syphilis.
- 137,394 (92%) of 149,627 women at ANC had their HIV status ascertained; 10,613 (8%) of these were HIV positive. 122,523 (98%) of 125,204 women at maternity had their HIV status ascertained 9,102 (7%) of these were HIV positive.
- 24,082 patients started ART this quarter.
- **595,186** patients were alive and on ART by end of December 2015. This means that **60%** of the estimated 1 million HIV positive population was on ART. ¹ Estimated ART coverage among people in need for treatment was **51%** (51,105 / 101,000) for children (<15 years) and **69%** (540,071 / 779,000) for adults.
- **76**% of adults and **77**% of children were retained alive on ART at 12 months after initiation. Actual retention rates are thought to be about **10**% higher due to misclassification of 'silent transfers' as defaulters in clinic-based survival/retention analysis. (see section 15.4)
- 517,346 (93%) of 583,674 patients on first line adult ART were on regimen 5A (TDF/3TC/EFV).
- 11,157 ² (79%) of an estimated 14,926 Error! Bookmark not defined. HIV infected pregnant women in Malawi were on ART this quarter. 6,847 (61%) of these were already on ART when getting pregnant and 4,310 (39%) started ART during pregnancy/delivery.
- An additional 1,547 Error! Bookmark not defined. breastfeeding women started ART due to Option B+ (in WHO stage 1/2)
- 76%, 69%, 66% and 65% of women started under *Option B+* were retained on ART at 6, 12, 24 and 36 months after initiation, respectively.
- 8,748 (8%) of infants discharged alive from maternity were known to be HIV exposed, 7,872 (90%) of these received ARV prophylaxis (nevirapine). 7,131 (82%) were enrolled in exposed child follow-up before age 2 months.
- A total of **10,624** HIV exposed children and **5,650** pre-ART patients were enrolled for follow-up in *HIV Care Clinics (HCC)* during this quarter.

¹ 2015 Spectrum estimates based on 2014 definition of eligibility for ART in Malawi (CD4<500, Option B+, UT for U5).

² Adjusted for double counting due to patient transfers / 'failed ART initiations' among women lost to follow-up within 6 months of ART registration.

2 Integrated HIV Program Overview

Malawi implemented a revised HIV Program in all health facilities following the release of the **2011 Malawi Integrated Clinical HIV Guidelines**. The second edition of these guidelines was published in March 2014 and implementation of revised policies commenced in April 2014. Key policies include:

- **PMTCT Option B+:** Universal life-long ART for all HIV infected pregnant and breastfeeding women regardless of clinical or immunological stage.
- Standard **HIV exposed child follow-up** to age 24 months. This program aims to improve early infant diagnosis and ART initiation using DNA-PCR testing for all infants from age 6 weeks; rapid antibody testing is considered diagnostic from age 12 months and repeated at 24 months. HIV exposed child enrolment and follow-up should be integrated with maternal ART follow-up (Option B+) to improve retention and adherence.
- Early ART initiation: universal ART for children under 5 years (confirmed HIV infection, CD4% no longer required), children over 5 years and adults with a CD4 count ≤500, patients with HIV and hepatitis B co-infection.
- Transition to a new first line ART regimens for adults (Malawi regimen 5A: tenofovir / lamivudine / efavirenz) and children (Malawi regimen 2: zidovudine / lamivudine / nevirapine), including provision of paediatric ARV formulations for all children under 25kg. Transition to 5A was completed by end 2013.
- Standardized pre-ART services for all HIV-infected persons not yet eligible for ART. The pre-ART program aims to reduce the incidence of HIV-related diseases and to enable early ART initiation based on the new CD4 cell threshold (500) through scheduled CD4 count monitoring.
- Provider-initiated provision of contraceptives and condoms for all adults in pre-ART and ART clinics
 to reduce the rate of unwanted pregnancies among HIV-infected adults and to reduce HIVtransmission between sexual partners.
- Isoniazid preventive therapy (IPT) for pre-ART patients to reduce the incidence of TB and intensified
 TB case finding (ICF) for all patients in pre-ART and ART follow-up to enable early diagnosis and
 treatment of TB and to reduce TB transmission in HIV clinics.
- Roll-out of scheduled **viral load monitoring** to improve early detection of treatment failure and initiation of second line ART.

Implementation of PMTCT Option B+ requires provision of ART services at <u>all</u> health facilities with Maternal and Child Health services. This required a massive acceleration of the **decentralization of ART services** from 303 (static) sites in June 2011 to currently 716 sites.

3 Supportive Site Supervision

3.1 Methods

The Department for HIV and AIDS has coordinated quarterly supportive supervision visits to all health facilities with ART services since the start of the national treatment program in 2004. Supervision teams are composed of: experienced HIV clinicians; nurses and M&E staff from health facilities in the public and private sector; district and zonal PMTCT and ART coordinators; program officers and technical staff from the Department for HIV and AIDS; technical staff from implementing partners. The TB and HIV programs have fully integrated their respective site supervision exercises since April 2015.

Each quarter, a one-day pre-supervision meeting is organised for all supervisors participating in the upcoming round to share program updates, discuss observations from the previous round, distribute materials and organise logistics, transport and accommodation.

Standard supervision forms are used to guide implementation of the supervision protocol, to update site information and collect M&E reports. Custom forms with previous data for each site are printed from the HIV Department database. The supervision forms include:

- Contact details of HIV service providers at each site
- Quality of service checklist
- o Follow up on action points noted during the previous visit
- Next visit date
- o M&E reports from HTC, ANC, maternity, exposed child and pre-ART follow-up, ART and TB
- o Physical drug stock-level assessment
- o Identification of sites in urgent need of clinical mentoring
- o Semi-structured feedback and performance rating for the supervision teams by facility staff

One copy of the supervision form is returned to the Department for HIV and AIDS, where data are entered in a custom MS Access database to produce national reports and to manage program logistics and the commodity supply chain. A second copy of the supervision form is left at the sites.

The supervision protocol includes a systematic review and verification of primary records (patient cards and registers) at all sites. This effectively provides a quarterly quality audit for M&E records, which has resulted in exceptional accuracy and completeness of HIV Program data in Malawi. At the same time, the systematic chart review helps to identify complex cases or deviations from clinical protocol, allowing the supervision team to provide targeted mentoring and clinical advice. The quarterly supervision exercise also aims to boost staff morale and motivation through *Certificates of Excellence* that are awarded by MOH to sites with an excellent score on the quality of service checklist. A growing number of health workers from sites all over the country participate as supervisors in this quarterly exercise and this has strengthened the national HIV Program identity and has greatly facilitated communication between program staff at the national, zonal, district and facility level.

The HTC Program usually conducts a separate supportive site supervision exercise each quarter, targeting a sample of HTC sites both within and outside of health facilities. Supervision teams consist of district, zonal and national level HTC coordinators, supported by implementing partners.

3.2 Supervision Outcomes

722 public and private sector facilities were visited for **clinical HIV program supervision** between 6th and 19th January 2016.

The large number of sites was covered by **180** supervisors working in **32** teams that spent a total of **2,042** working hours at the sites. Each site visit lasted on average **2.9** hours, but up to 2 days were spent at the busiest sites. **330 (46%)** sites were awarded a *Certificate of Excellence* for **excellent performance.** The

number of sites with excellent performance is similar to previous quarter 330. **73 (9%)** sites had significant weaknesses and were rated to require **intensive mentoring**. The capacity to provide site mentoring will need to be further expanded.

Table 1: Outcomes of integrated HIV services supervision for 2015 Q4

7	Total facil.	Supervision hours	spent at facilities	Performance (#	and % of sites)
Zone	visited*	Total	Average per site	Excellent perform.	Mentoring needed
NZ	127	337	2.7	56 44%	11 9%
CEZ	101	250	2.6	51 50%	16 16%
CWZ	162	441	2.9	66 41%	13 8%
SEZ	164	550	3.4	83 51%	10 6%
SWZ	168	464	2.8	74 44%	23 14%
Malawi	722	2,042	2.9	330 46%	73 10%

^{*} includes facilities that were visited for assessment of readiness, but that may have not (yet) been designated to provide integrated HIV services.

Table 1 summarizes the supervision outcomes by zone. Most facilities were using the standard national M&E tools 138 sites had cumulatively registered more than 2,000 ART patient and 52 of these had registered more than 5,000. 51 (37%) of these high burden sites were using electronic data systems. Some NGO supported sites were using custom tools compatible with the national standard reporting requirements.

4 Inventory of Sites and Services

4.1 Sites and Services

There were 724 static and 188 outreach HTC sites in Q4 2015; 145 of these were outside of health facilities.

Table 2: Facilities with integrated HIV services in the 5 Zones. Availability of services defined by performance (at least 1 patient enrolled) during 2015 Q4

Zana	Total	Facilities providing HIV services			ces	CD4 count machines (2)			
Zone	fac.(1)	Exp. child	Pre-ART	PMTCT B+	ART	Installed	Functional	Results	
NZ	135	115 85%	116 86%	102 76%	126 93%	27 20%	21 78%	2,193	
CEZ	103	99 96%	93 90%	91 88%	101 98%	20 19%	19 95%	1,212	
CWZ	168	131 78%	138 82%	129 77%	159 95%	27 16%	20 74%	2,513	
SWZ	168	148 88%	163 97%	141 84%	167 99%	42 25%	37 88%	6,174	
SEZ	165	159 96%	158 96%	154 93%	163 99%	49 30%	44 90%	4,135	
Malawi	739	652 88%	668 90%	617 83%	716 97%	165 22%	141 85%	16,227	

⁽¹⁾ Total facilities in the public / private sector designated to provide integrated HIV services in this quarter. Individual site selection is reviewed and may change each quarter.

Table 2 shows the distribution of the **739** sites designated to provide clinical HIV services in Q4 2015, by zone. At the national level, there were **716** (static) sites with at least one patient on ART, **617** sites had enrolled women under PMTCT Option B+; **668** sites were providing pre-ART services. **652** had enrolled HIV exposed children for follow-up. ART services were now available at almost all designated sites in the 5 zones.

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⁽²⁾ CD4 machines that have produced at least 1 result during the reporting period are defined as functional.

The SEZ and SWZ had reached 99% of designated sites with ART services. SEZ had reached 93% of designated sites with Option B+.

CD4 count machines (including 'point of care' machines) were installed at **165** sites, and **141** (85%) of these had produced at least 1 result during Q4 2015. The total number of CD4 results produced decreased from 19,513 in Q3 2015 to **16,227** during Q4 2015. 31% of these outputs were generated by 38 machines in the SW zone, implying that many CD4 machines continued to experience down-time or to be running considerably below capacity. The raised CD4 count threshold for ART eligibility may have also resulted in a decrease in the number of pre-ART patients requiring CD4 count monitoring as a large proportion is likely to be started on ART after their first CD4 test.

4.2 Staffing of HIV Services

4.2.1 HIV Testing Services

The Department for HIV and AIDS has maintained a dedicated system for professional registration and performance tracking for HTC Providers since 2011. This separate registration system is needed because HTC providers include lay persons with HTC training who are not registered with any other professional body. All HTC providers are issued with a unique ID and a professional logbook for documentation of duty stations, HTC trainings, sit-in observation and proficiency testing results. Logbook holders are requested to record the total number of tests done at the end of each month. Logbooks are routinely reviewed during quarterly supervision and key performance data for each provider are summarized on the site supervision forms.

	2015 Q1	2015 Q2	2015 Q3	2015 Q4
Sites visited	717	718	727	722
Sites with any tests done	666 93%	674 94%	684 94%	678 94%
Sites with registered HTC staff	670 93%	671 93%	669 92%	679 94%
Total HTC staff at visited sites	3,692	3,830	3,933	3,957
Staff with any test done	2,249 61%	2,495 65%	2,287 58%	2,337 59%
Staff with 300+ tests done this quarter	268 10%	326 11%	474 17%	490 17%
Logbooks reviewed	2,559 69%	2,870 75%	2,856 73%	2,922 74%
HTC staff participating in PT this quart	1,651 65%	931 32%	209 7%	109 4%
Total tests (HTC register)	480,249	494,006	625,803	604,854
Tests accounted for by individual staff	321,858 67%	380,159 77%	443,193 71%	445,638 74%
Source: logbooks	291,525 91%	359,042 94%	420,985 67%	418,870 94%
Source: HTC register	30,333 9%	21,117 6%	22,208 5%	26,768 6%
Total tests by staff with 300+ tests	134,272 42%	166,291 44%	263,234 59%	270,920 61%

679 (94%) of the 722 visited facilities had registered HTC providers and **678**(94%) sites had performed at least one test during Q4 2015. **2,922 (74%)** of **3,957** HTC providers had their logbooks available for review.

According to the 2,922 reviewed logbooks, **109 (4%)** HTC providers had participated in proficiency (panel) testing (PT) this quarter. This is lower than the participation rate from the previous quarters. However, documentation of PT may be incomplete given that not all logbooks were available for review. The national HIV reference laboratory is aiming to organize six monthly PT rounds for all practising HTC providers. However, the distribution of PT panels from the reference lab was delayed this quarter and district lab supervisors did not receive adequate PT supplies for their sites.

445,638 (74%) of all 604,854 tests conducted this quarter (according to HTC register reports) were accounted for by individual HTC staff working at the visited sites. **418,870 (94%)** of these tests were documented in the

reviewed logbooks and an additional **26,768 (6%)** could be attributed to individual providers from staff codes in the HTC registers. **490** (17%) of 2,337 providers with documented activity had tested 300 or more clients this quarter. A dedicated full-time HTC provider is expected serve 300 clients per quarter (average of 5 clients per day for 60 working days per quarter). The **490 staff** who met or exceeded this target provided **270,920 (61%)** of the total number of tests accounted for by individual staff this quarter.

4.2.2 ART/PMTCT

Integrated HIV program supervision has included a staffing census for ART clinics since Q3 2014. This census is implemented during the site visits, indicating all staff members who actually worked at the ART clinic on the most recent clinic day. The census is designed to provide an accurate snapshot of the actual staffing of ART services each quarter. The numbers collected may be slightly lower than longer term averages, because around 100 service delivery staff are themselves participating in the supervision exercise and will not be counted as having worked in their ART clinic during the supervision period. The table below shows that total staffing levels have been fairly consistent over the last 3 quarters.

In January 2016, **684** clinicians (physicians, clinical or medical officers); **1,029** nurses and **961** auxiliary staff (health surveillance assistants, clerks, etc.) were working in ART clinics in Malawi.

	2015 Q1		2015 Q2		2015 Q3		2015 Q4	
Clinicians	606	27%	661	27%	703	26%	684	25%
Nurses	814	36%	895	37%	982	37%	1,029	38%
Pharmacy staff	15	1%	13	1%	16	1%	16	1%
Auxiliary Staff	826	37%	881	36%	956	36%	961	36%
Total	2,261		2,450		2,657		2,690	

An estimated 2.8 million ART patient visits are currently managed at the 716 ART sites per annum, based on approximately 595,000 patients alive on ART and an average dispensing interval of 2.5 months. With 260 working days per year, an average of 10,800 patient visits are therefore managed by the ART sites per working day. At current staffing levels, this translates into an average of 16 ART patient visits per clinician and 10 per nurse per day. This approximate HRH capacity assessment does not take account of site-specific differences in patient burden and staffing levels and there are several medium and high burden sites with sub-optimal staffing. However, the national treatment program is fully decentralized to the health centre level and the program continues to devolve the growing patient burden to peripheral facilities. Since 2011, the steepest increase in ART patient numbers has been recorded at the 300 small peripheral sites that have the largest collective staffing capacity (see Figure 4 on page 20).

5 HIV Testing and Counselling Program Outputs

HTC protocols have been revised in 2013 and a new HTC register was implemented in the course of a national re-training campaign for all HTC providers between May and November 2013. Protocol revisions include:

- Clear recommendations for re-testing based on the client's test result and risk assessment
- Proper documentation of confirmatory testing for clients with a prior positive result (usually performed at enrolment into care).

The full national HTC data are presented in the **Appendix**.

5.1 HIV Testing Outputs

606,558 people³ were tested and counselled for HIV between October and December 2015. Testing outputs were 3% lower than the previous quarter. This was most likely due to disruption of services during the Christmas period.

558,890 (92%) of all tests were performed at health facilities, **15,538 (3%)** were done in stand-alone HTC sites and **32,130 (5%)** were done outside of facilities / in the community. Out of a total of **32,762** people newly diagnosed with HIV this quarter; **30,727 (94%)** of these at health facilities; **853 (3%)** at stand-alone HTC sites; and **1,182 (4%)** in a community-based testing. The 'yield' for new diagnoses was **5.6%** at health facilities, **5.7%** at stand-alone HTC sites and **3.8%** in community settings (excluding clients with a previous positive result from the denominator for all 3 settings).

5.2 HIV testing access type

357,020 (59%) of people tested were patients receiving provider-initiated testing and counselling (PITC); **246,061 (41%)** accessed voluntary testing and counselling, door-to-door, community-based testing, etc.; and **3,477 (1%)** came for testing with a *Family HTC Referral Slip* (FRS) that was issued to a family member at a prior HTC encounter. Based on a total of 23,867 FRS issued to index clients this quarter, the successful referral rate for family members was **15%** (3,477 /23,867). This is lower than in previous quarter (17%). Referral slips have remained under-utilized.

5.3 Age and sex distribution among HIV testing clients

Out of **606,558** people tested and counselled, **34%** were males and **66%** were females. **40%** of females were pregnant. The proportion of males (46%) to non-pregnant females (54%) was similar, implying gender balanced access to HTC services. Pregnant women have to be excluded from this comparison because testing among pregnant women is almost entirely provider-initiated and there is no comparable access route targeting males.

133,196 (22%) of all people tested accessed HTC with their partners (as a couple).

50% of all people tested and counselled were 25 years and above, **40%** were between 15-24 years and **11%** were children below 15 years. **2,641 (<1%)** of rapid tests done were among infants.

5.4 First time, repeat and confirmatory test results

All HIV positive patients enrolled in care need a confirmatory HIV test to rule out any possibility of mix-up of test results or fraudulent access to ART: either at enrolment into pre-ART follow-up, or before starting ART if the test to confirm was not done in pre-ART. Children under 12 months starting ART with a positive DNA-PCR do not need another confirmatory test before starting ART, but all need a confirmatory rapid antibody test at age 12 and 24 months.

173,165 (29%) accessed HTC for the first time and **433,393 (71%)** were repeat testers. Based on the cumulative number of people who accessed HTC for the first time, a total of **6,236,644** people have been tested since introduction of the *first time HTC access* indicator in July 2007.

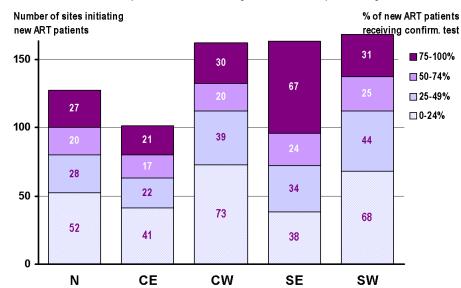
32,766 (5.5%) out of all clients received a positive result for the first time. Positive rapid test results among infants (1,229) and inconclusive test results (1,446) both accounted for <1 % of new results given to clients.

³ Reports from the HTC register are based on client encounters. It is not possible to de-duplicate people who access HTC multiple times in the reporting period. However, very few individuals come for repeat testing in less than 3 months and the number of HTC encounters in one quarter is therefore assumed to represent individuals.

415,718 (96%) of 433,393 repeat testers reported a *last negative* result. **14,666 (3%)** were reported as *previous positives* and all of these should have been classified as receiving a confirmatory test. For most of these previous *positives*, testing was probably initiated by a health worker before enrolment into care. *Confirmatory test results* exceeded by **472** the number of *previous positive* clients, indicating misclassification or data errors. **14,581 (96%)** of 15,138 confirmatory test results were concordant positive and **557 (4%)** were classified as *confirmatory inconclusive*. This category includes parallel concordant

Figure 1: Confirmatory HIV testing coverage at ART sites in the 5 zones

Num.: total confirmatory HIV tests documented in HTC registers. Denom.: total new patients initiating ART at the site



and discordant outcomes (Determine HIV1/2 and Uni-Gold HIV1/2 are used in parallel for confirmatory testing). relatively high proportion of clients who did not have a concordant confirmation positive may explained by selective confirmatory testing among clients with doubts about their previous positive status, but it underscores the importance of both routine confirmatory testing before ART initiation as well as the need to continue strengthening HTC quality assurance processes.

The 14,581 confirmatory test results

documented this quarter indicate that **60**% of the 24,082 patients initiating ART this quarter received confirmatory testing. **Figure 1** shows that the number of sites that implemented routine confirmatory testing remained low in all 5 zones: Only **176 (21%)** of facilities throughout the country had performed confirmatory testing for ≥75% of patients newly initiated on ART. However, the number and proportion of patients who received confirmatory test results has improved from 51% in the previous quarter and further increases are expected as the new HDA cadre is getting fully established.

6 DNA-PCR testing for Early Infant Diagnosis of HIV (EID)

DNA-PCR testing is performed at 8 labs (Mzuzu Central Hospital, Mzimba District Hospital, Kamuzu Central Hospital, Queen Elizabeth Central Hospital, DREAM Blantyre, DREAM Balaka, Zomba Central Hospital and Partners in Hope, Lilongwe). EID counsellors collect infant blood samples as dried blood spots on filter paper. Health facilities are requested to maintain a standard EID DNA-PCR logbook to document EID samples and to track results. The logbook includes the dates of collection, dispatch, receipt of result from the lab and communication of the result to the mother. Supervision teams were asked to collect basic data from these logbooks.

511 (78%) of 652 sites with HIV exposed children in follow-up had collected and recorded at least 1 DNA-PCR sample during Q4 2015. A total of **7,474** DNA-PCR samples were collected and recorded. By the time the logbooks were reviewed (between 1 and 3 weeks after the end of the quarter), results had been received at the sites for **2,724** (36%) of these specimens and **1,406** (52%) of these results had been communicated to the mother. The proportion of results received at the sites was **53**%, **38**% and **11**% for samples collected in October, November and December, respectively. A total of **102** (4%) results received at the sites were positive.

The **8 laboratories** registered the **receipt** of **4,339** DNA-PCR samples that were collected during Q4 2015. This represents 58% of the 7,474 samples recorded in the logbooks at the sites. 3,299 (76%) of the 4,339 registered samples arrived in the same quarter.

A total of **5,223** valid DNA-PCR results were dispatched from the labs in Q4 2015. **3,300 (63%)** of the dispatched results were from samples collected in Q4 2015, while 1,923 (37%) were from samples collected in the previous quarters. The median time between sample collection and dispatch of the result was **29 days**; 75% of results were dispatched between 22 and 41 days after sample collection.

2,736 (52%) of all results were from infants under 2 months old at the time of sample collection. 1,956 (38%) were 2-5 months, 442 (9%) were 6-11 months and 15 (<1%) were 12-17 months. 15 results were from older children or adults, presumably from samples sent to the lab as 'tie-breaker' for inconclusive rapid test results. The date of birth was missing for 74 samples.

Age at sample collection	Tot. Results	Positives	
<2 months	2736	34	1.2%
2-5 months	1956	70	3.6%
6-11 months	442	46	10.4%
12 months +	15	2	13.3%
(missing)	74	5	6.8%

157 (3.0%) of all results dispatched were positive. The age-specific number (%) of positive results is shown on the left. Receipt of the DNA-PCR result at the health facility is a prerequisite to updating of patient records and for appropriate clinical management. Considering the delays between

sample collection and dispatch of the test result from the lab, the child's age at the time of dispatch of the result from the lab is a useful indicator for the process of early infant diagnosis and early initiation of ART for confirmed infected infants. The table below shows the distribution of ages when results were dispatched from the lab.

Age when result disp. from lab	Tot. Results	(Col %)	Positives	(Col %)
<2 months	522	10%	5	3%
2-5 months	3,777	75%	82	53%
6-11 months	660	13%	60	38%
12 months +	28	<1%	4	3%
(missing)	79	<2%	6	3%
Total	5,066	100%	157	100%

Out of 157 positive results dispatched, only 5 (3%) were sent before the child was 2 months old. A total of 87 (56%) positive results were sent before the child was 6 months old and 147 (94%) were

sent before the child was 12 months old. A total of 93 infants were started on ART in WHO stage 1 or 2 on the basis of confirmed HIV infection (see ART section below). This is equivalent to **63%** of the number of positive DNA-PCR results dispatched for children <12 months this quarter.

7 Blood Safety

The Malawi Blood Transfusion Service (MBTS) is striving to provide safe blood products for the entire country using voluntary non-remunerated donors and quality assured screening for transfusion transmissible infections (TTIs). For the last years, MBTS has not been able to meet the national demand and several hospitals continue to supplement or rely entirely on blood units collected from replacement donors. Complete reports from MBTS have been available throughout, but blood safety reports from health facilities have not been consistently available and it has been challenging to compile national reports relying on the data passively submitted by the sites. Therefore, the PMTCT/ART supervision teams were tasked with active collection of blood donor and cross-matching data from all visited health facilities. Some of the visited laboratories were not using the standard MOH registers and the aggregation of data for reporting may have been affected by incomplete documentation at some sites.

A total of **23,699** blood units were collected in Malawi during Q4 2015. MBTS collected **17,109** (**72%**) of these, **100**% of which were screened comprehensively for the relevant TTIs (HIV, Hepatitis B, Hepatitis C, syphilis, malaria). In addition, **58** hospitals in Malawi collected a total of **6,890** units from replacement donors. **6,583** (**96%**) of these units were screened for at least the 3 key TTIs (HIV, HepB and syphilis) and **4,606** (**70%**) of these were also screened for HepC and malaria. This means that a total of **23,692** (**99%**) of all units collected this quarter were screened at least for HIV, HepB and syphilis. Based on the blood donor registers at the sites that collected blood from replacement donors, **17** units were screened for HIV and HepB only and **2** were screened only for HIV. **1,109** were screened with any other combination of tests for TTIs.

A total of **9,567** potential replacement donors were documented in the blood donor registers at the facilities and **6,890** (72%) of these ended up donating. Facilities may have used different screening algorithms and potential donors may have been excluded on the basis of different criteria, including TTIs, blood group, haemoglobin concentration and/or clinical conditions. Testing for less prevalent TTs may have only been carried out for donors who passed the screening for more common conditions. In total, 82% of potential donors were tested for HIV, 81% for HepB, 79% for syphilis, 69% for malaria and 41% for HepC. Detailed data on outcomes of individual tests among all potential blood donors are presented in the Appendix.

8 Post Exposure Prophylaxis (PEP)

A total of 1,608 persons received PEP during Q4 2015. This is very similar to the previous quarter (1,593).

9 Provider-Initiated Family Planning (PIFP)

The Integrated Clinical HIV Guidelines encourage health workers to routinely provide condoms to all adults in pre-ART and ART clinics. Women should also be offered at least the standard injectable contraceptive (Depo-Provera) during any pre-ART or ART visit. This policy aims to address the significant unmet need for family planning that had been observed among HIV patients in Malawi and to reduce the number of unwanted pregnancies among HIV-infected women (*PMTCT Prong 2*). HIV program reporting on PIFP is limited to women who received an injection of Depo-Provera in pre-ART and ART clinics during the last quarter. The report does not account for family planning need nor does it include women who accessed family planning services outside of HIV clinics.

Table 3: Number and % of women retained in HIV care * who were on injectable contraceptives (Depo) by the end of 2015 Q4.

	Pre-	ART	Α	RT	Both patient groups		
Zone	Tot. women	On Depo	Tot. women	On Depo	Tot. women	On Depo	
NZ	537	188 <i>35%</i>	33,638	12,346 37%	34,175	12,534 37%	
CEZ	358	37 10%	27,364	3,658 13%	27,722	3,695 13%	
CWZ	3,345	782 23%	69,405	19,350 28%	72,750	20,132 28%	
SEZ	2,779	559 20%	107,195	24,459 23%	109,974	25,018 23%	
SWZ	5,412	824 15%	112,052	20,685 18%	117,464	21,509 18%	
Malawi	12,431	2,390 19%	349,653	80,498 23%	362,084	82,888 23%	

^{*} estimated from the total number of patients retained in pre-ART and ART, multiplied by the proportions of females and adults registered

Table 3 shows that 82,888 (23%) of 362,084 women in care received Depo-Provera from HIV clinics in Q4 2015. The CW Zone had achieved the highest coverage among women in pre-ART and ART. Patient coverage has slightly decreased in this quarter. 579 (81%) of ART/PMTCT sites had stocks of Depo-Provera in January 2015 compared with 82% in October 2015.4 The HIV Program is no longer supplementing FP through supplies procurement and distribution of additional Depo-Provera to sites.

10 Cotrimoxazole Preventive Therapy (CPT)

All patients in HIV care are universally eligible for CPT in order to reduce the frequency and severity of several HIV-related diseases. Patients with confirmed HIV infection are provided lifelong CPT in pre-ART and ART clinics. CPT is also given to HIV exposed children until exposure to breast milk has stopped and HIV infection has been ruled out (usually around age 24 months). Fewer than 5% of patients are expected to require stopping of CPT due to toxicity.

⁴ Many Mission hospitals do not provide family planning.

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Table 4: Number and % of patients retained in HIV care who were on cotrimoxazole and isoniazid preventive therapy (CPT, IPT) by the end of 2015 Q4.

		СРТ						
	Ex	p. child	Pr	e-ART	ART		All patient groups	Pre-ART
Zone	Tot. pat.	On CPT	Tot. pat.	On CPT	Tot. pat.	On CPT	Tot. pat. On CPT	Tot. pat. On IPT
NZ	8,782	7,113 <i>81%</i>	2,235	2,188 98%	59,437	58,133 98%	70,454 67,435 96%	2,235 2,031 91%
CEZ	8,096	6,674 82%	1,689	1,677 99%	47,676	41,029 86%	57,461 49,379 86%	1,689 1,556 92%
CWZ	16,475	13,865 <i>84%</i>	10,231	7,029 69%	120,177	107,901 90%	146,883 128,795 <i>88%</i>	10,231 6,081 59%
SEZ	31,367	27,388 87%	11,318	11,164 99%	172,961	170,128 98%	215,646 208,681 97%	11,318 10,666 9 <i>4%</i>
SWZ	29,201	23,681 81%	17,660	12,978 73%	191,674	174,719 91%	238,535 211,378 <i>89%</i>	17,660 11,385 64%
Malawi	93,921	78,721 <i>84%</i>	43,133	35,036 81%	591,925	551,911 93%	728,979 665,668 91%	43,133 31,720 74%

Table 4 shows that **665,668 (91%)** of 728,979 all patients in care were on CPT at the end of Q4 2015.

10.1 Intensified TB Case Finding (ICF)

TB is one of the most important HIV-related diseases in Malawi and a considerable proportion of (mainly early) deaths on ART are attributed to undiagnosed TB. ICF is carried out using a standard symptom checklist at every HIV patient visit. ICF outcomes are documented on HIV exposed child, pre-ART and ART patient cards, but routine M&E reporting is currently limited to ART patients in order to reduce the burden of reporting secondary cohort outcomes. It is assumed that implementation of ICF is similar in pre-ART and exposed child follow-up.

578,825 (98%) of all patients retained on ART were screened for TB at their last visit before end of December 2015. As of that visit, **3,686 (1%)** patients were new TB suspects and had presumably been referred for examination by a clinician and for TB investigations. **853 (<1%)** patients had confirmed TB (clinical or lab based). Out of these, **672 (79%)** were confirmed to be on TB treatment and **181 (21%)** had not yet started or had interrupted TB treatment. An excerpt from the data in the **Annex** (*Cumulative ART outcomes*) is shown below.

Current TB status among ART patients (ICF)

ICF n	ot done (Current TB status unknown/ not circ)	13,100	2%
ICF d	one	578,825	98%
	TB not suspected	574,286	99%
	TB suspected	3,686	1%
	TB confirmed	853	0%
	TB confirmed, not on treatment	181	21%
	TB confirmed, on TB treatment	672	79%

10.2 Isoniazid Preventive Therapy (IPT)

All pre-ART patients with a negative screening outcome for TB symptoms are eligible for IPT. The first (large scale) distribution of isoniazid and pyridoxine for the HIV programs reached the sites during July 2012. **31,720** (74%) of 43,133 patients retained in pre-ART were on IPT by the end of December 2015. Isoniazid was in stock at 583 facilities during the January 2015 supervision visit.

11 HIV-Related Diseases

Table 5 shows the number of patients treated for key HIV-related indicator diseases. **3,924** patients were started on TB treatment this quarter and HIV status was ascertained for **3,744** (95%). **2,032** (54%) of these were HIV positive and **1,660** (82%) of all HIV positives were already on ART when starting TB treatment. The Diflucan (fluconazole) donation program has received renewed attention and significant stocks of fluconazole were distributed to all sites in November 2012. In Q4 2015, **972** and **1,233** patients received Diflucan for acute cryptococcal meningitis and oesophageal candidiasis, respectively. **296** patients with Kaposi sarcoma were registered for ART in this quarter.

Table 5: Number new cases of key HIV-related diseases registered per quarter (KS = Kaposi Sarcoma, CM = cryptococcal meningitis, OC = oesophageal candidiasis).

		T	В	KS *	CM *	OC *	
	Tot. cases	HIV status asc.	HIV positive	Already on ART	Tot. cases	Tot. cases	Tot. cases
2015 Q1	4,158	3,765 91%	1,954 52%	1,408 72%	260	865	610
2015 Q2	4,288	4,074 95%	2,200 <i>54%</i>	1,513 69%	265	459	599
2015 Q3	4,346	3,973 91%	2,230 56%	1,573 71%	323	525	808
2015 Q4	3,924	3,744 95%	2,032 <i>54%</i>	1,660 82%	296	972	1,233

12 HIV-Exposed Child Follow-Up

12.1 Methods and Definition of Indicators

There are multiple entry points into HIV exposed child follow up: children of HIV infected mothers may be enrolled at birth at maternity / postnatal ward; they may be found at Under 1 or Under 5 Clinics through active screening for HIV exposure; they may be identified when presenting sick to OPD; or they may be seen with their mothers in ART follow-up. Although the targeted enrolment age is below 2 months, children may theoretically be enrolled up to 23 months of age (when HIV infection can be ruled out by rapid antibody test and breast milk exposure is likely to have stopped).

Initial registration data and details for every visit are recorded on an *Exposed Child Patient Card* and a subset of the registration data is copied in the *HIV Care Clinic (HCC) register* (one record per patient). Registration data are reported from the HCC register on a quarterly basis. Follow-up outcomes are reported monthly, selecting children who were **2**, **12 and 24 months** old in the respective reporting month. Outcomes are determined from the latest visit details recorded on each card. HIV infection status is evaluated as *known negative* if a negative DNA-PCR or rapid test result was available at the last visit; HIV infection status is evaluated as *known positive* if a positive DNA-PCR result was available at any age or a positive rapid antibody test was available from age 12 months; HIV infection status is counted as *unknown* if HIV infection has not been confirmed and/or a negative test result pre-dated the last visit (assuming on-going HIV exposure through breast milk). All children under 24 months with confirmed HIV infection and those under 12 months with confirmed HIV infection through DNA-PCR or HIV antibody and symptoms of *presumed severe HIV disease* are *eligible for ART*.

The main outcome indicator for the HIV exposed child follow-up program is *HIV-free survival at 24 months of age*. This is defined as the proportion of children who were discharged as confirmed HIV uninfected by the age of 24 months.

12.2 HIV Exposed Child Registration Data

10,624 HIV exposed children were newly enrolled into follow-up during Q4 2015; **7,131 (68%)** of these were under the age of 2 months. This represents timely enrolment for **82%** of the 8,748 known HIV exposed children discharged from maternity this quarter. The total number of new enrolments (10,624) exceeds by 1,876 (21%) the total number of known HIV exposed children discharged from maternity (8,748). This apparent discrepancy may be explained by delayed enrolment of infants born in previous quarters; by double-counting of infants who transferred between sites; or by identification and enrolment of additional HIV exposed infants after birth. Overall, enrolment into follow-up for known HIV exposed infants appears to be almost complete.

The documentation of follow-up outcomes, particularly the updating of DNA-PCR results on patient cards, remained incomplete at several sites. This has led to an underreporting of ascertainment of HIV status among the 2-month old cohort.

12.3 Birth Cohort Outcomes

There were **8,774** infants in the **2-month age cohort**. **2,916 (33%)** had received a DNA-PCR result. **40 (1%)** of these were confirmed HIV infected. An additional **9** infants were diagnosed with *presumed severe HIV disease*, which means that a total of **49** infants were eligible for ART. **37 (76%)** of these had started ART. The proportion of positives starting ART is higher than the previous quarter (52%). Out of the entire 2-month age cohort, **7,929 (92%)** were retained in exposed child follow-up, **37 (<1%)** had started ART and **19 (<1%)** were discharged confirmed uninfected⁵. **30 (<1%)** were known to have died and **577 (7%)** had been lost to follow-up.

There were **9,767** children in the **12-month age cohort**. Current HIV infection status was known for **4,589 (47%)** children (DNA-PCR or rapid antibody test) and **190 (4%)** of these were confirmed HIV infected. **17 (<1%)** additional children had been diagnosed with *presumed severe HIV disease*, which means that a total of **207** children were eligible for ART. **169 (82%)** had started ART. Out of the entire age cohort, **7,028 (74%)** were retained in exposed child follow-up, **169 (2%)** had started ART and **78 (<1%)** were discharged confirmed uninfected. **5 2,117 (22%)** were lost to follow-up and **68 (<1%)** were known to have died.

There were **8,832** children in the **24 month age cohort**. Current HIV infection status was known for **4,429** (50%) children (DNA-PCR or rapid antibody test) and **225** (6%) of these were confirmed HIV infected. **8** additional children had been diagnosed with *presumed severe HIV disease*, which means that a total of **233** children were eligible for ART. **215** (92%) of these had started ART. Out of the entire age cohort, **767** (9%) were retained in exposed child follow-up, **215** (3%) had started ART and **4,094** (48%) were discharged confirmed uninfected. **3,376** (39%) were lost to follow-up and **109** (1%) were known to have died.

Confirmed HIV-free survival at age 24 months in this quarter remained implausibly low at 48%. This was related to the fact that only 50% in this cohort had a known HIV status. 4,403 (50%) children were classified as 'current HIV infection status unknown' and many of these may be among the 3,376 children lost to follow-up and the 109 children who had died. However, 767 (9%) were retained in follow-up beyond age 24 months and a final rapid test was not available for these children, possibly due to continued breast feeding. There are still problems with scheduled HIV testing (and documentation of test results) at 6 weeks, 12 and 24 months of age.

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⁵ A small number of children may be rightfully discharged as 'confirmed uninfected' by 2 or 12 months of age, provided that HIV exposure through breast milk has definitely stopped (e.g. maternal death) and a negative HIV test was obtained at least 6 weeks thereafter.

13 Pre-ART

13.1 Pre-ART Registration Data

A total of **5,650** patients were newly registered for pre-ART follow-up in Q4 2015. **441 (8%)** of these were children aged 5-14 years. The number of new pre-ART enrolments slightly decreased from the previous quarter (6,929 total, 587 children). Several sites had already established pre-ART services before July 2011 and the cumulative number of pre-ART patients ever registered was **206,528**.

13.2 Cumulative Pre-ART Follow-up Outcomes

43,133 (22%) of all patients ever registered were retained in pre-ART follow-up by the end of December 2015; **104,983 (53%)** had started ART; **48,965 (25%)** had been lost to follow-up; **1,897 (1%)** were known to have died. The proportion of patients starting ART is bound to increase in the cumulative pre-ART cohort analysis over time. Based on a subtraction of cumulative outcomes from the previous quarter, **4,411** pre-ART patients started ART during Q4 2015; **2,241** were lost to follow-up and **29** died. The quarterly number of died is lower than in the previous quarter, indicating challenges with completeness and accuracy of reporting.

CPT coverage among pre-ART patients was **81**% in Q4 2015 and IPT coverage declined slightly to **74**%. **2,390** (**19%**) of 12,431 women had received Depo-Provera from their pre-ART clinic. Further details on CPT, Depo-Provera and IPT are presented in **Tables 3** and **4** in the sections above.

14 PMTCT / ART

The implementation of **PMTCT Option B+** has effectively integrated PMTCT and ART services. The program aims to initiate lifelong ART for all HIV infected women as early as possible in pregnancy. ART may be started and continued at ANC, labour and delivery, and at ART clinics. All infants born to HIV-infected women are supposed to start daily nevirapine prophylaxis for the first 6 weeks of life. Nevirapine syrup is given to women at ANC at the earliest opportunity to take home with instructions how to give it to the new-born.

14.1 Data Sources and Reporting Methods

New standard M&E tools for ANC and maternity were implemented in January 2010 and revised in Q2 2012 to reflect the Option B+ policy. ANC and maternity clinic registers and reporting forms include patient management information and all relevant data elements for the maternal and child health and HIV programs. The ANC register was specifically designed to avoid data duplication that previously affected PMTCT reports from ANC due to the inability to account for individual women's outcomes in the course of multiple visits. The cohort reporting system is designed to aggregate women's outcome data after they have completed their ANC visits. The outcome report is completed for women who started ANC 6 months before the reporting period.

From **Q2 2015**, the PMTCT data elements (HIV ascertainment and ART status) were also added to the first section of ANC reporting form that captures women's status at their first (booking) visit. The ANC report now includes the HIV and ART status at the first visit for women <u>starting</u> ANC in the reporting period and the final HIV and ART status of women who had <u>completed</u> ANC by the end of the reporting period. This addition aims to monitor PMTCT service implementation more closely in time, allowing for corrective action in the course of subsequent visits.

Data from ANC and maternity are collated and presented separately because records do not allow identification of individual women and hence are subject to double counting if not separated.

All patients starting ART are recorded using standard program monitoring tools (ART patient treatment cards and ART clinic registers). **ART baseline data** for all patients registered are reported each quarter from ART

clinic registers. **ART outcomes** of all patients ever registered are reported after reviewing the cards of all new patients and of those who were on ART at the end of the previous quarter, updating the status of patients who have subsequently died, stopped or been lost to follow-up. Secondary outcomes such as current regimen, CPT status, side effects, adherence and TB status are reported for all patients retained on ART.

ART scale-up has resulted in a growing proportion of HIV-infected women who are already on ART when getting pregnant. Implementation of *Option B+* will further increase ART coverage in this group. **Maternal ART coverage** is estimated from the number of pregnant women who were already on ART when getting pregnant (**maternity reports**) <u>plus</u> those who newly started ART when pregnant (**ART reports**).

Maternity reports capture ART status at the time of delivery (up to the time of discharge from the postnatal ward). The timing of ART initiation is categorized into: (any time) before pregnancy; during 1st / 2nd trimester; during 3rd trimester; during labour. About 97% of pregnant women in Malawi attend ANC, but only 83% of women in the general population deliver at a health facility in Malawi. Maternity reports therefore have the potential for undercounting the number of mothers and infants receiving ARVs. However, there is evidence from ANC and maternity reports that almost all of the known HIV infected women deliver at health facilities. ARV coverage among known positives is therefore reliably calculated from maternity reports. Women admitted at maternity who are referred to another facility before / after delivery are double-counted in aggregated maternity data. Assuming the probability of referral is independent of ART status, the number of women already on ART when getting pregnant is therefore adjusted by the overall proportion of referrals among women admitted to maternity.

ART program reports capture pregnancy (and breastfeeding) status at the time of *ART initiation*, providing information on the number of new women starting ART while pregnant (or while breastfeeding). ART reports do not capture women who become pregnant after starting ART. For the estimation of maternal ART coverage, the number of women starting ART in pregnancy is **adjusted for:**

- a) Double-counting of women starting ART in pregnancy and subsequently transferring to another site. These women are counted multiple times as 'pregnant at the time of starting ART' in the quarterly ART cohort reports because the disaggregation of age, sex and reason for starting ART applies to all patients newly registered in the quarter, including transfers in. Separate *ART 'survival' analyses* are collected each quarter for women started under Option B+. The proportion of women transferred within 12 months of registration is used to adjust the quarterly number of pregnant women starting ART for transfers.
- b) Failed ART initiation is thought to be the main underlying reason for early loss to follow-up among the Option B+ cohort. Patients are recorded on patient cards and in clinic registers when the first supply of ARVs is dispensed and all new entrants are counted as ART initiations in the quarterly ART cohort report. Recent operational studies indicate that most pregnant women lost to follow-up within the first 6 months never return after this first dispensing visit and many of these may have never actually started taking ART. The proportion of women lost to follow-up in the 6-month survival analysis is therefore used to adjust the number of pregnant women starting ART in the quarterly ART cohort reports for failed initiations.

Infant PMTCT coverage is estimated from maternity reports, based on the number of infants born to known HIV-infected women and discharged alive who started nevirapine prophylaxis.

Coverage is calculated by dividing the number of patients served by population denominators. The denominators are derived from expected pregnancies based on population projections and HIV prevalence from epidemiological surveillance (source: 2015 Spectrum model for Malawi). There are an estimated 14,926 HIV infected pregnant women in the population per quarter (1/4 of 59,704 in 2015).⁶

⁶ 2015 Spectrum estimates based on current definition of eligibility for ART in Malawi (CD4<500, Option B+, UT for U5).

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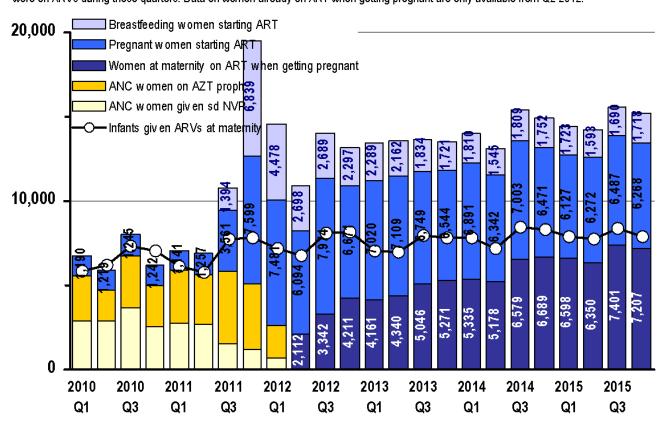
14.2 ARV Coverage among Pregnant / Breastfeeding Women and Exposed Infants

11,157 (75%) of the estimated 14,926 HIV infected pregnant women in Malawi this quarter were on ART. This is based on **6,847** ⁷ women at maternity who were already on ART when getting pregnant and **4,310** ⁸ women who newly initiated ART in pregnancy. This is an increase in ART coverage from 70% in the previous quarter.

An additional **1,547** ⁹ breastfeeding women started ART due to *Option B+* (in WHO clinical stage 1 or 2), bringing the total number newly started on ART under *Option B+* to **5,857**. Most women starting ART while breastfeeding were probably identified late in maternity or early in the postnatal period, but this group may also include some women who re-initiated after interrupting ART in pregnancy. **7,872** infants were confirmed to have started NVP prophylaxis at maternity.

Figure 2 shows the transition from prophylactic ARV regimens for HIV infected mothers to universal ART under *Option B+* (registration data; not adjusted as above). The (less effective) single dose NVP regimen and AZT combination prophylaxis had been phased out by April 2012. The average number of pregnant women registered for ART each quarter **increased almost 6-fold** from **1,221** in the 12-month period before introduction of Option B+ to an average of around **6,500** since Q4 2011.

Figure 2: Transition from prophylactic ARV regimens for PMTCT to Option B+ in Malawi
Women who moved to Option B+ from sdNVP / AZT were double counted between Q3 2011 - Q1 2012. It is likely that <12,000 total women were on ARVs during these quarters. Data on women already on ART when getting pregnant are only available from Q2 2012.



⁷ 7,207 women who started ART before pregnancy admitted at maternity; reduced by 5% to adjust for double-counting of 6,361 referrals among 125,743 total admissions.

⁸ 6,268 women registered at ART clinics who were pregnant at the time of starting ART; a) 10% are discounted to adjust for double-counting of transfers based on 824 of 8,556 women who transferred within 12 months of registration (12 month Option B+ survival analysis); b) 23.6% are discounted to account for presumed failed ART initiations based on 1,704 of 7,227 women lost to follow-up within 6 months of registration (6 month Option B+ survival analysis).

⁹ 1,718 women registered at ART clinics who were breastfeeding at the time of starting ART; reduced by 10% to adjust for double-counting of transfers based on 824 out of 8,556 women who transferred within 12 months of registration (12 month Option B+ survival analysis). Failed ART initiations are thought to be less common among this group, so no further adjustment is made.

14.3 HIV Services at ANC

The full national data from ANC are presented in the **Appendix**.

14.3.1 HIV Ascertainment and ART Coverage

Booking cohort:

139,322 women attended ANC for their first visit between October and December 2015. This is 84% of the estimated 166,000 pregnant women in the 2015 population during one quarter.¹⁰ It is also lower than the number of women who attended ANC for the first time between July and September 2015 (150,456). 125,149 (90%) of women in the booking cohort had their HIV status ascertained at the first visit. Out of these, 10,087 (8%) presented with a valid previous test result and 115,056 (92%) received a new test. A total of 9,623 (8%) of women were found HIV positive: 5, 079 (53%) of these from a documented previous test and 4,544 (47%) from a new test. 8,687 (90%) of all positives were on ART: 4,498 (52%) of these were already on ART when starting ANC and 4,189 (48%) newly started ART at their first ANC visit. Out of these, 3,532 (84%) were in their 1st or 2nd trimester and 657 (16%) were in the 3rd trimester of pregnancy.

Outcome cohort:

150,212 women had started ANC between April and June 2015 and their outcomes were reported between October and December 2015. Only **36,606 (25%)** of women in this cohort attended the recommended minimum of 4 focussed ANC visits.

137,958 (92%) of the outcome cohort had their HIV status ascertained at least once in the course of ANC. This is similar to the previous quarter (91%). **10,343 (7%)** presented with a valid documented previous HIV test result and **127,615 (93%)** received a new HIV test result at ANC. A total of **10,667 (7.7%)** women were found HIV positive. This is slightly lower than the latest Spectrum projections (9.0% HIV prevalence among pregnant women in 2015).⁶

10,199 (96%) of (known) HIV infected women were on ART by the end of ANC. This represents 68% coverage of the estimated 14,926 HIV positive pregnant women per quarter at the population level. Of the 10,199 ANC women who were known to receive ART, 5,266 (52%) were already on ART when starting ANC, 3,962 (39%) initiated before 28 weeks of pregnancy and 971 (10%) initiated during the last trimester of pregnancy. 10,144 (95%) of HIV infected women at ANC were on Cotrimoxazole Preventive Therapy. 9,370 (88%) of known HIV infected women attending ANC received the infant dose of ARVs (nevirapine syrup) to take home.

14.3.2 Syphilis Screening

36,839 (25%) of women in the outcome cohort were tested for syphilis and **607 (2%)** were syphilis positive. The low testing rate probably explains the higher (2%) than expected proportion (<1%) of positives as the testing was likely selective of those suspected to be positive.

14.4 HIV Services at Maternity

The full national data from maternity are presented in the **Appendix**.

Between July and September 2015, **119,382** women were admitted for delivery to maternity; **6,361** of these were referred to another facility before delivery, resulting in **125,743** total admissions to maternity during Q4 2015. Out of all admissions, **116,55** (96%) delivered at health facilities, while **5,360** (4%) had already delivered before reaching a facility. The **116,555** facility deliveries represent **70**% of the estimated 166,000

¹⁰ Estimated as ¼ of 664,000 births projected for 2015 (Demographic Proj Spectrum 2015).

Malawi National Statistics Office. (2008). Malawi Population Projections 2008-2030.

Retrieved from http://www.nsomalawi.mw/images/stories/data_on_line/demography/census_2008/MainReport/ThematicReports/Population Projections Malawi.pdf

quarterly deliveries in the population in 2015 which is less than the 83% reported in the Integrated Household Survey Report of 2010-2011.

A total of 113,890 (95%) deliveries were conducted by skilled birth attendants, 640 (<1%) by paramedical staff and 5,143 (4%) were not attended by any of the above (probably mainly among women who delivered before reaching maternity). 14,536 (12%) of women developed obstetric complications. The most common leading complications were obstructed / prolonged labour (4,810 cases) and post-partum haemorrhage (1,574 cases). A total of 121,915 babies were born, 117,815 (97%) were singletons and 4,100 (3%) were twins/multiples. There were 119,729 (98%) live births and 2,186 (2%) stillbirths. 118,586 (99%) of babies born alive were discharged alive and 1,143 (1%) died before discharge. 119,567 (>99%) of women were discharged alive and 106 (<1%) women died before discharge, which is equivalent to a maternal mortality ratio of 89 per 100,000 live births among women attending maternity.

14.4.1 HIV Ascertainment at Maternity

123,039 (98%) women had their HIV status ascertained at maternity. Out of these, **119,597** (97%) presented with a valid previous HIV test result and **3,442** (3%) received a new HIV test result. A total of **9,148** (7%) women were HIV positive and **113,891** (93%) were negative. The **123,039** women whose HIV status was ascertained at maternity represent **74**% of the expected 166,000 women delivering in the population.

HIV exposure status was ascertained for **116,290 (98%)** out of 118,586 babies born and discharged alive. **8,748 (8%)** of these were born to a known HIV positive mother.

14.4.2 ARV Coverage at Maternity

A total of **9,073 (99%)** of known HIV infected women admitted to maternity received ART. Out of these, **7,207 (79%)** had started ART before pregnancy, **1,090 (12%)** initiated ART during the 1st or 2nd trimester, **657 (7%)** initiated during the 3rd trimester and **119 (1%)** initiated ART at maternity.

A total of **7,872 (90%)** of 8,748 infants who were known HIV exposed and discharged alive started daily NVP prophylaxis at maternity. This represents **53%** coverage of the estimated 14,926 HIV exposed infants born in the population in this quarter.

15 ART Access and Follow-Up Outcomes

The full national data from the ART Program are shown in the **Appendix**.

15.1 New ART Registrations during Q4 2015

By the end of December 2015, there were **716 static ART sites** in Malawi, managed by government, mission, NGOs and the private sector. Out of these, **91** were ART facilities in the private sector, charging a nominal MK500 per monthly prescription of drugs per patient.

Implementation of the Malawi Integrated Clinical HIV Guidelines started in July 2011, triggering a massive surge in new ART initiations (see **Figure 3**). **24,082** patients initiated ART in Q4 2015 and **7,843** patients were registered as a transfer in (already on treatment; 24% out of all 32,402 clinic registrations). These numbers are similar to previous quarter.

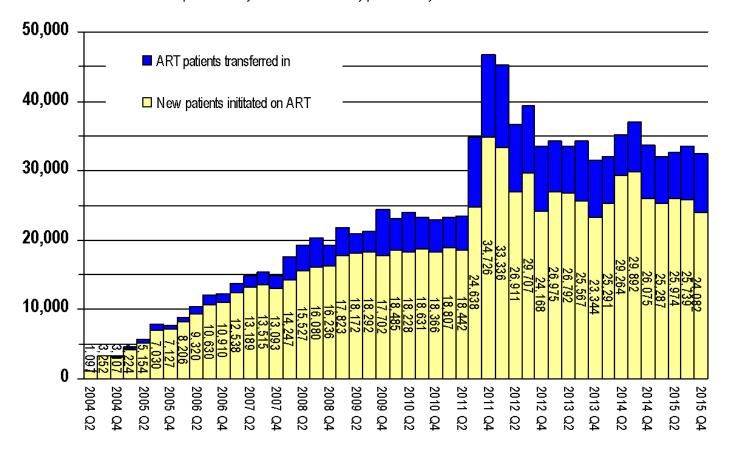
Among all new registrations **36%** were males, **64%** were females. **6,268 (31%)** of females were pregnant. **6,264** (99.9%) of pregnant women were started under **Option B+** (in WHO stage 1 or 2 with unknown CD4 or CD4 above 500), while 4 were in more advanced stage of HIV infection. An additional **1,718** women in

WHO stage 1 or 2 were started because of breastfeeding, bringing the total number of women registered as started under *Option B+* 11 to **7,982**.

Figure 3: Patients newly inititated on ART and total ART clinic registrations per quarter

Total ART clinic registrations include patients who transferred between sites. This results in double counting of patients at the

national level. For 'patients newly initiated on ART' every patient is only counted once.



A total of **20,231 (63%)** of all patients registered started in WHO stage 1 or 2 and **11,544 (57%)** of these started due a low CD4 count. **10,237 (32%)** of patients registered started in WHO stage 3 and **1,456 (5%)** started in stage 4.

2,605 children were registered at ART sites in Q4 2015. **608 (26%)** of these were registered under the expanded policy of universal ART for children aged 12-59 months in WHO stage 1 or 2, independent of CD4 count. **149 (6%)** of children started ART with presumed severe HIV disease. This is higher than the previous quarter (104). **93** infants in WHO stage 1 or 2 started due to confirmed HIV infection through DNA-PCR, which is lower than the previous quarter (103). Early paediatric ART access has remained below expectations, but the relatively low number is consistent with reduced transmission rates due to Option B+: considering that 8,748 HIV exposed infants were identified at maternity and assuming a 2% transmission rate among the 90% of HIV positive mothers at maternity who received ART (and 20% transmission in the 10% who did not receive ART)¹², only about 333 of these known HIV exposed infants may have been infected perinatally during Q4 2015. However, considering the projected 1,560 new infant HIV infections in the 2015 population per quarter⁶, early infant treatment coverage remains low at an estimated **6%** (93 / 1,560). The most significant

¹¹ Universal ART for all HIV infected pregnant and breastfeeding women in WHO stage 1 or 2, independent of CD4 count

¹² UNAIDS Reference Group on Estimates Modelling and Projections (2011). Working paper on mother-to-child-transmission rates for use in Spectrum. Geneva, UNAIDS.

bottleneck for early infant treatment remains the identification of HIV infected pregnant / breastfeeding women.

862 (3%) out of all ART clinic registrations were patients with TB: **490 (2%)** had a current and **372 (1%)** a recent history of TB. **296 (1%)** of patients registered had Kaposi's sarcoma.

15.2 Cumulative ART Registrations up to December 2015

By the end of December 2015, there were a cumulative total of **1,091,656** clinic registrations, representing **872,567 (80%)** patients who newly initiated ART and **206,755 (19%)** patients who transferred between clinics. **12,334 (1%)** out of all clinic registrations were patients who re-initiated ART after treatment interruption. Out of all registrations, **36**% were males and **64**% were females, **91**% were adults and **9**% were children (<15 years). Private sector clinics accounted for **32,597** (3.0%) of total patient registrations.

15.3 ART Outcomes

595,186 patients were alive on ART by the end of December 2015. This is equivalent to **60% ART coverage** among the estimated 1 million HIV positive population in Malawi in 2015. The number of patients on ART includes an estimated 3,261 patients in transit between sites (50% of the 6,521 patients newly registered as transferred out at sites across the country).

Out of the **872,567** patients ever initiated on ART, **595,186** (**68%**) were retained alive on ART, **81,471** (**9%**) were known to have died, **207,968** (**24%**) were lost to follow-up and **4,293** (<1%) were known to have stopped ART. An estimated **540,071** adults and **51,105** children (<15 years) were alive on ART by the end of December 2015.

Figure 4 Patients alive on ART at the end of each quarter, stratified by size of facility (number of patients alive on ART)

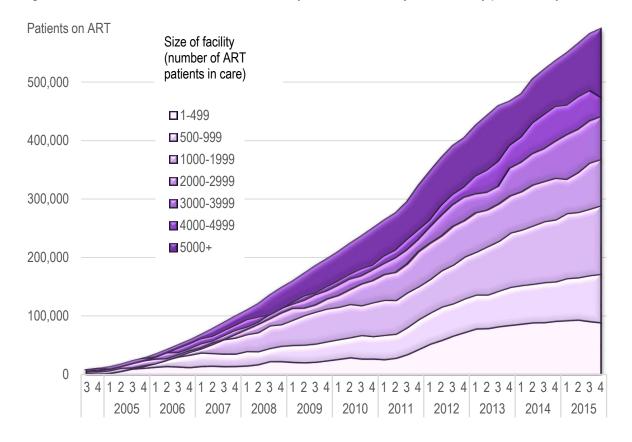


Figure 4 shows the increase of patients alive on ART by the end of each quarter. The number of patients alive on ART **increased by 9,526** in Q4 of 2015. **Figure 4** also illustrates the ongoing decentralization of Malawi's ART program. From Q3 2011, the greatest increase in ART patient numbers was seen at sites with fewer than

500 patients alive on ART. By the end of September 2015, **48**% of the national ART patient cohort was in care at sites with fewer than 2,000 patients.

Figure 5: Quarterly rates of ART drop out (ART stop, defaulters and deaths)

Numerator: new ART stops, new defaulters and new deaths in the respective quarter

Denominator: total patients retained alive at the end of the previous quarter plus new patients registered in the respective quarter)

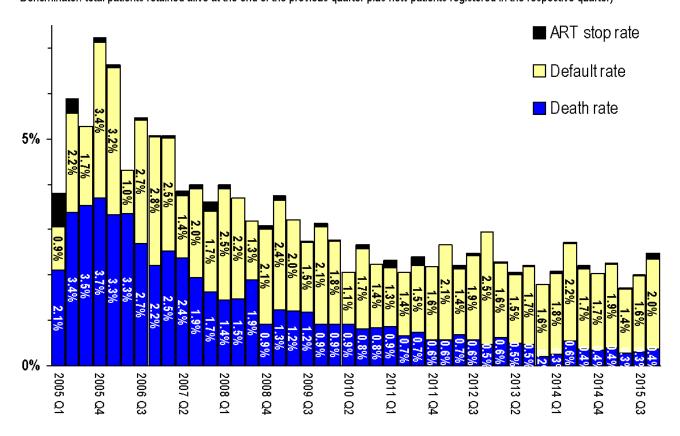


Figure 5 shows the considerable decrease of ART drop-out rates since the start of the national program. There were 2,191 new deaths, 11,986 new defaulters and 845 new stops in Q4 2015. This translates into a quarterly death rate of 0.4% and a defaulter rate of 2.0% among the patients alive and on treatment in this quarter. These rates were similar to the previous quarter. Based on previous operational studies, about half of the patients in stage 3 and 4 who are classified as lost to follow-up are thought to have died. There is also an indication that 10-15% of pregnant women who were registered as 'initiated on ART' under Option B+ may have never actually started taking ARVs due to inadequate preparation at ANC. Importantly, the ascertainment of loss to follow-up requires updating of patient treatment cards after analysis of the most recent dispensing visit. Any lack in rigour in this process will lead to a misclassification of patients who have been lost to follow-up as 'retained alive on ART'.

By end of December 2015, a cumulative **81,471 (9%)** patients were known to have died **207,968 (24%)** were lost to follow-up and **4,293 (<1%)** were known to have **stopped ART.**

Figure 6: Patients starting ART in WHO stage 4 and deaths in the first 3 months after ART initiation. (Shown as proportions among new patients registered each quarter)

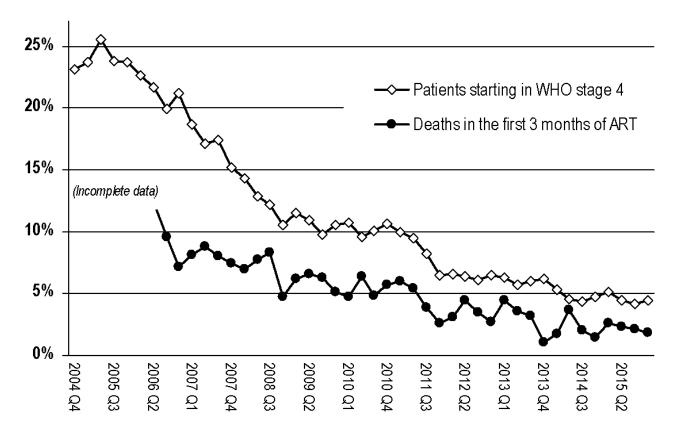


Figure 6 shows the considerable decline in **early mortality** since the start of the program. In Q2 of 2006 11% of new patients died within the first 3 months of ART initiation. There has been a remarkable decline in early mortality and the lowest point thus far has been reached in Q4 2013. The decrease in early mortality is probably mainly due to earlier ART initiation (patients in WHO stage 2 with a CD4 count below the threshold or in stage 3). It correlates well with the decline in the proportion of patients starting ART in WHO clinical stage 4 from 25% in 2005 Q2 to **5%** in Q4 2015. Slight fluctuations in the calculated early mortality rates are mainly due to inconsistent classification of month of death at the sites with electronic patient record systems. The 2014 guidelines have led to further reduction in early mortality, as more patients are started in WHO stage 1 and 2 (CD4 threshold for eligibility <500; universal ART for HIV infected pregnant and breastfeeding women and children under 5 years).

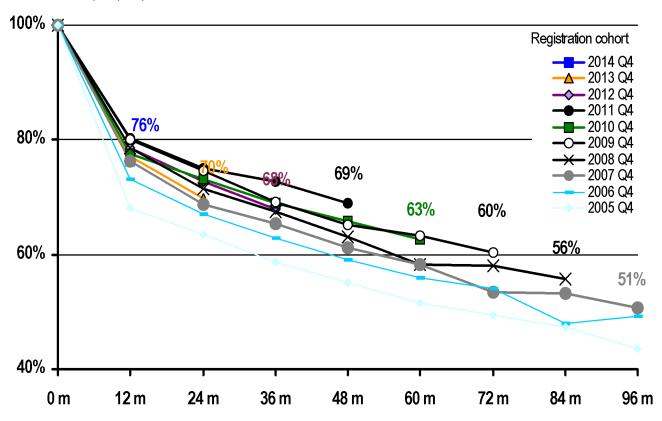
15.4 ART Cohort Survival Analysis

A 12, 24, 36, 48, 60, 72, 84, 96 and 108-month 'cohort outcome survival analysis' was conducted for patients registered in Q4 of 2006, 2007, 2008, 2009, 2010, 2011, 2012, 2013 and 2014, respectively. A separate 12-month cohort outcome analysis was conducted for children who were under 15 years at the time of ART initiation and who registered for ART in Q4 2014. A further subgroup analysis was done for women who started ART under *Option B+* during Q4 2012, Q4 2013, Q4 2014 and Q2 2015. **76% of adults** and **77% of children** were retained alive on ART after 12 months on treatment. This is similar to the previous quarter and remains below the WHO target of 85%. The majority of patients classified as lost to follow-up are likely to have stopped/ interrupted ART, but others will have transferred to another facility without notifying the previous site. Actual retention rates are thought to be about **10%** higher due to this misclassification of 'silent transfers' as 'defaulters' in clinic-based survival/retention analysis. A population-based study in Karonga

district with individual linkage showed that **92**% of patients started in 2011-2012 were retained after 12 months on ART while routine monitoring data showed **79**% retention rates for the same period.¹³

Figure 7 shows the continuous improvement of long-term treatment outcomes over time. **61%** and **54%** of patients registered 5 and 7 years ago had been retained alive on ART.

Figure 7: Group cohort survival analysis: Proportion of patients retained alive on ART 12, 24, 36, 48, 60, 72, 84 and 96 months after ART initiation



6-month group cohort survival outcomes were known for **7,716 (98%)** of the 7,862 women registered as having started ART under *Option B+* in Q2 2015. ¹⁴ The 7,716 represents 489 (6%) women who transferred out and are therefore double counted and **7,227 (94%)** patients not transferred. **5,482 (76%)** of these were retained at 6 months after registration. **1,704 (98%)** of those not retained were lost to follow-up, **19 (<1%)** were known to have stopped ART and **22 (<1%)** were known to have died.

12-month group cohort survival outcomes were known for **8,556** women registered as having started ART under *Option B+* in Q4 2014. ¹⁴ This number is 178 (2%) higher than the number of women that registered as having started ART of 8,378. This discrepancy is likely due to data abstraction inaccuracies. The 8,556 represents **824 (10%)** women who transferred out and are therefore double counted and **7,732 (90%)** patients not transferred. **5,319 (69%)** of these were retained at 12 months after registration. **2,318 (96%)** of those not retained were lost to follow-up, **31 (1%)** were known to have stopped ART and **64 (2%)** were known to have died.

24-month group cohort survival outcomes were known for **8,171** women registered as having started ART under *Option B+* in Q4 2013. ¹⁴ This number is 265 (3%) higher than the number of women that registered as having started ART of 7,906. Similar to the 12 months cohorts, the discrepancy is likely due to data abstraction inaccuracies. The **8,171** number represents **955** (12%) women who transferred out and are

¹³ Koole, O., Houben, R. M. G. J., Mzembe, T., Van Boeckel, T. P., Kayange, M., Jahn, A., Crampin, A. C. (2014). Improved retention of patients starting antiretroviral treatment in Karonga District, northern Malawi, 2005-2012. Journal of Acquired Immune Deficiency Syndromes (2014), 67(1), e27–33. doi:10.1097/QAI.000000000000252

¹⁴ Group cohort survival analyses were not available from some sites with electronic data systems. 'Reason for starting' may be reclassified for some patients, leading to minor inconsistencies in patients included in group cohort survival analyses.

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therefore double counted and **7,216 (88%)** patients not transferred. **4,790 (66%)** of these were retained at 24 months after registration. **2,319 (96%)** of those not retained were lost to follow-up, **31 (<1%)** were known to have stopped ART and **76 (>1%)** were known to have died.

1,721 (22%) of the women in the 24-month Option B+ survival cohort had initiated ART in the breastfeeding period and **1,869 (24%)** started in the third trimester / in labour; considering the 24 month median breastfeeding period in Malawi (2010 MDHS), more than half of the women in this cohort can be assumed to have stopped breastfeeding. The **66% and 65% retention rate at 24 and 36 months** after ART initiation confirms that a high proportion of women started under Option B+ **remain on ART beyond the cessation of breastfeeding**.

The 6-month retention rate is slightly higher than the previous quarter. These are satisfactory results. Most of the women lost to follow-up failed to return after their first visit and many of these may have never actually started ART due to inadequate counselling and preparation in the initial phase of implementation. The program is examining the different modes of delivery closely to further improve uptake and retention on *Option B+*.

6 month survival OptionB+

Survival and retention in ART program

ART cohort registration group outcomes

Total ART din	Total ART clinic registrations			
Trans	Transfers out (double counted)			
Total r	Total not transferred out (patients in cohort)			
	Total alive on ART			
	Total not retained	1,745	24%	
	Defaulted	1,704	98%	
	Stopped ART	19	1%	
	Died	22	1%	

12 month survival OptionB+

Survival and retention in ART program

ART cohort registration group outcomes

Total ART clinic registrations		100%
Transfers out (double counted)		10%
Total not transferred out (patients in cohort)		90%
Total alive on ART		69%
Total not retained		31%
Defaulted	2,318	96%
Stopped ART	31	1%
Died	64	3%

24 month survival OptionB+

Survival and retention in ART program

ART cohort registration group outcomes

Total A	Total ART clinic registrations		8,171	100%
	Transfers out (double counted)		955	12%
	Total not transferred out (patients in cohort)		7,216	88%
	Total alive on ART		4,790	66%
	Total not retained		2,426	34%
		Defaulted	2,319	96%
		Stopped ART	31	1%
		Died	76	3%

36 month survival OptionB+

Survival and retention in ART program

ART cohort registration group outcomes

Total .	Total ART clinic registrations		9,625	100%
	Transfers out (double counted)		1,310	14%
	Total not transferred out (patients in cohort)		8,315	86%
	Total alive on ART		5,420	65%
	Total not retained		2,895	35%
		Defaulted	2,689	93%
		Stopped ART	46	2%
		Died	160	6%

15.4.1 Secondary outcomes of patients retained on ART

Secondary outcomes are known for the **591,925** patients alive on ART who remained at their sites at end of the quarter.

ART Regimens

583,674 (99%) of patients were on first line regimens. The number of patients on second line regimens increased by **435** from 7,334 in Q3 to **7,769** this quarter. **482 (<1%)** patients were on non-standard regimens. Non-standard regimens are not necessarily substandard regimens and include patients continuing an ART regimen that was started outside Malawi, patients in research programmes and patients in specialist care.

Among patients on first line regimens, **26,064 (4%)** were on paediatric formulations and **25,001 (96%)** of these were on the new standard first line for children (regimen 2P: AZT/3TC/NVP). By the end of December 2015, **517,346 (93%)** of patients on adult first line were receiving regimen **5A** (tenofovir / lamivudine / efavirenz). **28,686 (5%)** were on regimen 2A (zidovudine / lamivudine / nevirapine), which was the main alternative regimen for patients with stavudine side-effects before transition to regimen 5A and **985 (<1%)** were on regimen 1A (stavudine / lamivudine / nevirapine).

Adherence to ART

Pill counts and the number of missed doses were documented for **586,453 (99%)** out of all patients retained on ART and **530,724 (90%)** of these were classified as >95% adherent in Q4 2015. Manual estimation of adherence from pill counts is practically difficult and classification can be misleading. The ART program has switched to a direct evaluation of doses missed in 2010 to improve on accuracy of adherence assessment and plausible adherence levels are recorded with this method. However, there have also been persistent challenges with the analysis of adherence levels at sites with Electronic Data Systems (EDS) and adherence data from several of these sites could not be included in this report.

ART Side Effects

551,464 (93%) patients on ART had information on drug side effects documented at their last clinic visit before end of December 2015. This is an increase from the previous quarter (83%). This was due to a software update for the national EMR that interfered with the side effect recording. **7,621 (1%)** of patients with information had documented side-effects. The prevalence of side effects seems to have stabilized at very low levels following the full transition to regimen 5A (tenofovir / lamivudine / efavirenz) that started in July 2013.

15.5 Viral Load (VL) Monitoring

The National Treatment Program has started rolling out routine VL monitoring for patients on ART to facilitate early detection of treatment failure and timely switching to second line ART. Routine VL monitoring is scheduled at 6 months after ART initiation, at 2 years and every 24 months thereafter. Additional targeted VL testing may be carried out for patients with clinically suspected treatment failure. During Q4 2015, 9 laboratories in the national program provided VL testing for patients enrolled at the respective facilities and associated sites. All labs used the MOH lab information management system (LIMS) for registration of samples and storage of results. The following results are based on an analysis of exported LIMS data.

A total of **20,509** VL results were produced between October and December 2015. **3,868 (19%)** of samples processed were plasma and **16,587 (11%)** were DBS. For 54 results, the specimen type was not specified.

Lab		Turn-around				
	Plasma	DBS	Oth/unk	Total	Time (Days)§	
DREAM Blantyre	1,011	136	0	1,147	12	
DREAM Balaka	176	1,020	5	1,197	38	
Kamuzu CH	1,076	1,093	1	2,170	52	
Mzimba DH	0	233	0	233	31	
Mzuzu CH	0	1,493	0	1,493	54	
Partners in Hope	731	4,035	1	4,766	30	
QUECH	0	2,794	0	2,794	50	
Thyolo DH	874	4,048	50	4,972	30	
Zomba CH	0	1,735	2	1,737	55	
Total	3,868	16,587	54	20,509	37	
§ Median days between sample collection and printing of results in the lab						

Thyolo DH lab achieved the highest outputs, contributing 24% of all results this quarter. The median interval between sample collection and printing of results was **37 days** at the national level, ranging from **12 days** at DREAM Blantyre to **54 days** at Mzuzu CH. The most significant delays occurred between sample receipt and processing in the lab (median 25 days), while on average only 7 days elapsed between sample collection and receipt in the lab.

Reason	0-9	99	1000	4999	50	000+	Total
Routine	17,317	87%	909	5%	1,775	9%	20,001
Targeted	156	58%	20	7%	91	34%	267
Other/unk	133	55%	33	14%	75	31%	241
Total	17,606	86%	962	5%	1,941	9%	20,509

20,001 (98%) of all VL samples were classified as *routine scheduled*. This is equivalent to **27%** of the estimated 75,000 ART patients passing a VL monitoring milestone this quarter. **267 (1%)** of samples were classified as *targeted (suspected treatment failure / repeat)* and for **241 (1%)** the reason for the sample was 'other' or not specified. **17,606 (85%)** of all results were below 1,000 copies/ml. The proportion of results with 5,000+ copies was higher among the *targeted* samples (34%) and those with *unspecified* reason (31%), compared with 9% among *routine* samples.

The time on ART was entered for only **6,766 (25%)** of 27,162 routine samples registered on the LIMS and only **2,105 (31%)** of these were drawn on schedule (from 1 month before to 3 months after a VL milestone). The proportion of patients with VL < 1000 was **87%**, **85%**, **85%**, **90%**, **84%** and **88%** at 6, 24, 48, 72, 96 and 120 months on ART respectively. Viral suppression rates of samples drawn on schedule were similar to those of 'catch-up' (extra-schedular) samples (**82%**) or those with unknown timing (**85%**).

Patient age was recorded for all routine monitoring samples. Among these, 5%, 6%, 11%, 31% and 48% were from the age groups 0-9, 10-19, 20-29, 30-39 and 40+ years. Viral suppression rates (VL<1000/ml) were significantly lower among children (0-9 yrs: **61%**) and adolescents (10-19 yrs: **60%**) compared with adults (**81%**, **85%** and **88%** for the age groups 20-29, 30-39, 40+ years, respectively).

VL monitoring outputs are expected to increase further over the next quarters.

Given the relatively low access to VL monitoring (estimated 39% of all ART patients due for VL monitoring this quarter), the measured **86% viral suppression rate** may not be representative for the entire national ART cohort. With generally limited access to testing, the VL samples analyzed this quarter may overrepresent patients with poor adherence and/or treatment failure. Conservatively, the national viral suppression rate can be estimated as **503,668** (86%) of 585,660 patients on ART, which is equivalent to **50%** of the total 1 million HIV infected population.

16 TB / HIV Management

Approximately **89%** of HIV infected TB patients were receiving ART in Q4 2015. This estimate is based on the following triangulation of TB and ART program data:

TB Program Data: A total of **3,924** TB patients were registered during Q4 2015. Assuming an average HIV prevalence of 60% among TB patients, **2,354** TB patients were HIV positive and therefore in need of ART. Given that **1,660** TB patients registered were already on ART at the time of starting TB treatment, 2,354 – 1,660 = **694** TB patients needed to initiate ART.

ART Program Data: An estimated **655** patients¹⁵ started ART with a current or recent episode of TB in Q42015. This is **94%** (655 of 694) of the TB patients who needed to start ART. This means that a total of 1,660 + 655 = **2,315** (**98%**) of the estimated 2,354 HIV infected TB patients were receiving ART in Q4 2015.

TB program repor	t		*
TB clinic registration	s		
Total TB patients regist	ered	3,924	100%
HIV status ascertainr	nent		
HIV status not ascertain	ned	180	5%
HIV status ascertained		3,744	95%
HIV negative		1,712	46%
HIV positive		2,032	54%
Already	on ART	1,660	82%
Not on	ART when starting TB treatment	372	18%
TB / ART program	triangulation		*
. •	B patients (estimated)		
HIV negative (est. 40%)	1,570	40%
HIV positive (est. 60%) in need of ART	2,354	60%
Not on ART	Not on ART		2%
Total on ART (coverage)		2,301	98%
Already	on ART (TB prog)	1,660	72%
Started	ART within 24m of TB diagnosis (ART prog)	641	28%
	ART initiations with current TB (ART prog)	364	57%
	ART initiations after recent TB (ART prog)	276	43%

17 STI Treatment

STI reports were actively collected during the Integrated HIV Program Supervision exercise for the 10th time this quarter. This decision was taken due to the persistent challenges with 'passive' reporting based on data aggregated by the district STI coordinators. This quarter, supervision teams collected STI data from 653 out of 928 facilities offering STI management according to the *2013-14 Service Provision Assessment*¹⁶ in Malawi. The site-level reports included here may therefore only represent 70% of all STI services in Malawi. The supervision teams re-emphasized the importance of complete and accurate documentation at the sites and the data quality is expected to improve with resumption of regular site supervision for the STI program. The complete set of STI program data collected is included in the Appendix.

¹⁵ 24% of the 862 ART patients who were registered this quarter with a recent or current episode of TB at the time of ART initiation were assumed to be transfers and were subtracted to adjust for double-counting.

¹⁶ Ministry of Health, & ICF International. (2015). Malawi Service Provision Assessment (SPA) 2013-14. Lilongwe, Malawi and Rockville, Maryland, USA. Retrieved from http://dhsprogram.com/pubs/pdf/SPA20/SPA20.pdf

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17.1 Access to STI treatment and coverage

Based on the data collected at the facilities, a total of **55,938** STI cases were treated in Q4 2015. Considering the 70% site-level completeness of reporting, this number is estimated to represent a total of **79,911** STI cases treated. This is equivalent to **81% STI treatment coverage** of the expected 98,600 STI cases in the population.

Out of **55,938** documented clients treated, **22,324** (40%) were male and **33,614** (60%) were female. **4,036** (12%) of female STI clients were pregnant. **37,054** clients (66%) were 25 years and above, **13,584** (24%) were 20-24 years and **5,300** (9%) were under 20 years old.

17.2 Client Type and STI History

49,986 (89%) of clients were symptomatic and **5,952** (11%) were asymptomatic (treated as partners). Among symptomatic clients, **45,685** (91%) of were index cases and **4,301** (9%) were partners. A total of **14,919** partner notification slips were issued, equivalent to an average of 0.33 slips per index case. Considering the 14,919 partner notification slips issued, **69%** (10,253) of those notified presented to the clinic. **41,091** (73%) of clients presented with their first lifetime episode of STI, **10,569** (71%) clients reported to have had an STI more than 3 months ago and **4,278** (29%) of clients reported having had an STI within the last three months. Re-occurrence of an STI after a recent episode may be due to re-infection or treatment failure.

17.3 HIV Status

HIV status was ascertained for **34,879** (62%) clients and **8,693** (25%) of these were HIV positive. **2,297** (26%) of positives were identified through a new test initiated at the STI clinic, while **6,393** (74%) presented with a documented previous positive HIV test result. **5,137** (80%) of clients with a previous positive HIV test result were on ART.

The rate of HIV status ascertainment at STI clinics has gradually improved over time. This is likely due to increased numbers of dedicated testing staff available at the sites (HDAs). Weak back-referral systems which may lead to incomplete documentation of new HIV test results at the STI clinics, so the actual HIV ascertainment rates are likely to be higher. It is worth noting that a substantial proportion of clients who are aware of their HIV infection present with a new episode of an STI. This may suggest poor translation of positive living strategies promoted during counselling, but could also be due to the increased risk of recurrence of HSV-2 and balanitis among HIV-infected clients.

17.4 STI Syndromes and Referrals

The most common syndrome was abnormal vaginal discharge (AVD) with **19,631** (31%) cases, followed by urethral discharge (UD, **14,664** cases), genital ulcers (GUD, **9,705** cases) and lower abdominal pain (LAP, **8,841** cases). Similar to previous reports, balanitis, bubo, warts and neonatal conjunctivitis each accounted for 1-2% of cases.

Given the high risk of recent HIV infection among STI clients, all clients with unknown status and those with a new negative test result should be referred for (repeat) HIV testing and counselling. **18,107 (80%)** of the 21,059 STI clients with unknown or new negative test result were referred for repeat HTC. **1,350 (59%)** of 2,297 clients who were newly tested HIV positive were referred for ART eligibility assessment.

18 Supply of HIV Program Commodities

18.1 Quantification, procurement planning and distribution

The program conducted the quarterly quantification, procurement plan budget update for antiretroviral and opportunistic infection medicines to be procured under New Funding model (NFM). This covers the consumption period ending December 2017 plus 9 months buffer.

During Q4 2015, ARVs, medicines for opportunistic infections and laboratory health products worth \$21 million were received by the Bollore Africa Logistics managed warehouses dedicated for Department of HIV and National Malaria Control Program commodities. This comprised of Tenofovir/Lamivudine/Efavirenz 300/300/600mg (Regimen 5A; 86% of the value of adult ARVs). To maintain adequate stocks in the pipeline and hence ensure uninterrupted supply for subsequent orders, the Ministry has continued processing HIV commodity orders for ARVs, OI, RDTs and other related commodities through Partnership for Supply Chain Management (ARVs and RDTs) and IDA Foundation (laboratory commodities and medicines for opportunistic infections).

During the implementation period, there were delayed deliveries for Co-trimoxazole 960mg with 3 months of stock available at central level for distribution. IDA Foundation was requested to fast track all subsequent Co-trimoxazole shipments.

18.2 Logistics support during integrated site supervision

District and central level Supply Chain and Logistics Officers provided stock management support at over 200 sites during the Q4 2015 integrated ART/PMTCT site supervision. This included a physical inventory at all sites and ad-hoc mentoring in stock management at health facilities with poor performance. There was an overall improvement in the logistics management of ARVs and medicines for OI medicines. Some health facilities visited had storage constraints hence providers had to conduct physical inventory at multiple locations.

Health care providers have continued to use RDT daily activity registers and relocation books for registration of redistributed commodities to health facilities. However at selected health facilities, it was noted that RDT daily activity registers are not updated in real time.

18.3 Stock Status of HIV Commodities

Physical stock counts for ARVs and other medicines for HIV-related diseases were performed at all sites during the supervision visits in January 2016. **Table 6** shows the total medicine stocks found at the sites and the estimated consumption patterns.

Minimum stocks of TDF/3TC 300/300mg and AZT/3TC 300/150mg are maintained at all sites for post-exposure prophylaxis (PEP) and the total stocks at the sites therefore far exceeds the actual consumption from patients using this regimen in alternative ART regimens or as PEP. There are some residual quantities of stavudine- and efavirenz-containing regimens at the warehouse and site level following the transition to a tenofovir-based regimens. The program continues to monitor the trend of patients on such regimens to inform future procurements for alternative regimens such as Abacavir/Lamivudine formulations given that stavudine containing regimens will be phased out completely in 2016.

517,731 patients were on regimen 5A, which was **23,249 (4.2%)** less than projected in the previous forecast for the end of this quarter **(540,980)**. The national ART program forecast and quantification was updated in March 2016 to inform procurement planning and budgeting for HIV commodities for the period ending December 2017.

18.4 Availability of standard first line ARVs

517,731 of all ART patients were on the standard first line regimen (5A; tenofovir / lamivudine / efavirenz). This is equivalent to 87% of patients overall or 93% of patients on first line adult regimens. By end January/beginning February 2016, the total stock of this regimen was equivalent to 5.7 and 4.5 months of consumption at the warehouse and site-level, respectively. Total national stocks may conceal stock imbalances at the facility level and a key supply management indicators is therefore the availability of this regimen at each site. The physical stock count carried out during supportive supervision in January 2016 confirmed that 714 (99.7%) of all 716 ART sites with patients on this regimen had available stocks. This

translates into a 'stock-out' rate of only 0.3% of sites. Such stock-out events typically affect small peripheral sites and are usually short due to the bi-monthly scheduled distribution cycle and the ad-hoc stock relocation facility coordinated through the toll-free supply hotline. This healthy supply chain has enabled the program to consistently implement three monthly drug dispensations for patients.

18.5 Bimonthly distribution of HIV & Malaria Commodities

One scheduled bimonthly distribution of HIV & Malaria commodities (Distribution Round 26) took place from December 2015 to January 2016. A total of 148 different commodities (anti-malarial, ARVs, OI medicines, STI medicines and laboratory commodities) were distributed to 726 health facilities. This was the seventh successful consolidated distribution for HIV and malaria commodities.

12 selected health facilities in Mulanje, Phalombe, Zomba, Balaka, Ntcheu and Lilongwe were visited to verify items received in distribution round 23 and 24. The supply chain trail exercise is routinely conducted at systematically selected health facilities to validate signed delivery notes provided by the third party logistics provider and to review adherence to stock management procedures. There were no deviations from the signed proofs of delivery.

During Q4 2015, the logistics team at the Department of HIV and AIDS also coordinated a total of 1,372 individual commodity transactions at 362 ART sites to manage stock imbalances. All transactions are managed and authorized using the toll-free HIV Department Supply Chain Hot Line that was set up to facilitate communication between the health facilities and the central office. Health workers are able to consult on supply chain and other HIV commodities related issues that need to be resolved by the technical team at the department in a timely manner.

Table 6: Total stocks of HIV program commodities at all sites visited during the 2015 Q4 supportive site supervision. Stock positions are from the date of the visit (between 1-4 weeks after the end of the quarter). Warehouse stock positions are from 10/02/2016

Inventory	Item	Sites with Total Phys		sical Stock	Consump-	- Months of Stock	
unit		any Stock	At Sites	In Warehouse	tion/ Month	At Sites	
tins	ABC / 3TC 60 / 30mg tins (60 tabs)	201	27,781	65,188	5,013	5.5	13.0
	ABC / 3TC 600 / 300mg tins (30 tabs)	85	3,885	19,318	837	4.6	23.1
	ATV / r 300 / 100mg tins (30 tabs)	235	30,595	21,295	6,703	4.6	3.2
	AZT / 3TC / NVP 300 / 150 / 200mg tins (60 tabs)	645	139,362	177,306	28,691	4.9	6.2
	AZT / 3TC / NVP 60 / 30 / 50mg tins (60 tabs)	651	386,761	864,016	62,458	6.2	13.8
	AZT / 3TC 300 / 150mg tins (60 tabs)	403	13,320	75	3,294	4.0	0.0
	AZT / 3TC 60 / 30mg tins (60 tabs)	509	21,654	25,481	2,284	9.5	11.2
	d4T / 3TC / NVP 30 / 150 / 200mg tins (60 tabs)	165	24,795	27	985	25.2	0.0
	d4T / 3TC 30 / 150mg tins (60 tabs)	203	11,790	21	103	114.5	0.0
	EFV 200mg tins (90 tabs)	172	2,784	6,081	305	9.1	19.9
	EFV 600mg tins (30 tabs)	243	13,264	5	820	16.2	0.0
	LPV / r 100 / 25mg tins (60 tabs)	95	12,357	13,053	3,198	3.9	4.1
	LPV / r 200 / 50mg tins (00 tabs)	70	980	3,157	466	2.1	6.8
	NVP 200mg tins (60 tabs)	489	41,826	73,772	8,820	4.7	8.4
	NVP 50mg tins (60 tabs)	157	11,664	16,561	1,513	7.7	10.9
	TDF / 3TC / EFV 300 / 300 / 600mg tins (30 tabs)	716	2,335,950	2,945,694	517,731	4.5	5.7
	TDF / 3TC 300 / 300mg tins (30 tabs)	654	60,715	121,826	14,016	4.3	8.7
1 441				121,020			0.1
bottles	Fluconazole (Diflucan) 50mg / 5ml bottles (35 ml)	36	3,869	70 774	117	33.0	44.0
	NVP 10mg/ml bottles (100 ml)	267	39,277	76,774	6,603	5.9	11.6
	NVP 10mg/ml bottles (25 ml)	561	103,656	55	16,440	6.3	0.0
vials	Benzathine Penicillin 1.44g vials (50 each)	525	296,483		41,153	7.2	
	Bleomycine 15,000IU vials (1 each)	23	6,581	1,562			
	Ceftriaxone 1g vials (50 each)	615	181,730		111,078	1.6	
	Depo-Provera 150mg/1ml vials (25 each)	579	650,530	264,750	268,563	2.4	1.0
	Gentamicin 80mg / 2ml vials (50 each)	556	171,501		104,530	1.6	
	Streptomycin 1 gm vials (50 each)	61	51,556				
	Vincristine 1mg / 1ml vials (1 each)	79	25,513		3,528	7.2	
tabs	Aciclovir 200mg blist packs (500 tabs)	400	1,972,373		669,578	2.9	
	Azithromycin 500mg blist packs (3 tabs)	454	74,593		11,050	6.8	
	Ciprofloxacin 500mg blist packs (100 tabs)	417	481,039	1,961,500	316,720	1.5	6.2
	Clotrimazole 500mg boxes (1 each)	467	176,920	13,791	40,711	4.3	0.3
	Codeine 30mg tins (100 tabs)	420	777,856		52,419	14.8	
	Cotrimoxazole 100 / 20mg blist packs (1000 tabs)	545	24,036,929	58,104,000	7,859,593	3.1	7.4
	Cotrimoxazole 400 / 80mg tins (1000 tabs)	642	41,423,804	11,969,000	17,559,556	2.4	0.7
	Cotrimoxazole 960mg blist packs (1000 tabs)	571	27,693,104	61,318,000	18,665,178	1.5	3.3
	Doxycycline 100mg tins (1000 tabs)	588	4,671,827		4,692,818	1.0	
	E thambutol (E) 100 mg blist packs (100 tabs)	47	106,992				
	E thambutol (E) 400 mg blist packs (672 tabs)	10	34,868				
	Erythromycin 250mg tins (1000 tabs)	567	6,482,362	1,001,000	4,198,201	1.5	0.2
	Fluconazole (Diflucan) 200mg tins (28 tabs)	190	759,400	23,436	73,767	10.3	0.3
	lbuprofen 200mg tins (100 tabs)	118	1,079,566		897,585	1.2	
	Isoniazid (H) 100mg blist packs (100 tabs)	85	439,484		157,488	2.8	
	Isoniazid (H) 300mg blist packs (672 tabs)	99	1,478,054	4,800,096	1,151,630	1.3	4.2
	Isoniazid (H) 300mg tins (1000 tabs)	584	14,710,852	5,162,000	1,152,666	12.8	4.5
	Metronidazole 200mg tins (1000 tabs)	631	20,346,770	7,426,000	5,097,923	4.0	1.5
	Morphine 10mg blist packs (60 tabs)	85	950,575		228,737	4.2	
	Pyridoxine 50mg tins (1000 tabs)	546	10,060,554	3,997,000	1,230,374	8.2	3.2
	RH 150 / 75 mg blist packs (672 tabs)	130	463,819				

Inventory	ltem	Sites with	Total Phys	sical Stock	Consump-	Months of Stock *	
unit		any Stock	At Sites	In Warehouse	tion/ Month	At Sites	Wareh.
	RH 60 / 60 mg blist packs (84 tabs)	9	18,490				
	RHE 150 / 75/ 275 mg blist packs (1000 tabs)	53	211,556				
	RHZ 60 / 30/ 150 mg blist packs (84 tabs)	34	47,336				
	RHZE 150/75/400/275mg blist packs (672 tabs)	193	483,178				
sheets	ART pat. card adult (yellow) bundles (100 sheets)	641	652,919	7,050	9,929	65.8	0.7
	ART pat. card paed. (blue) bundles (100 sheets)	526	71,621		868	82.5	
	Exposed child card (pink) bundles (50 sheets)	637	75,335	6,350	3,545	21.2	1.8
	Family HTC Referral Slip bundles (100 sheets)	219	35,195				
	Polythene sleeve bundles (100 sheets)	498	97,313		16,227	6.0	
	Pre-ART pat. card (green) bundles (100 sheets)	543	127,677		1,885	67.7	
	STI Partner Referral Slip bundles (100 sheets)	318	30,985				
tests	DBS kit (filter paper, lancet, etc.) boxes (50 each)	574	108,412	141,750	33,839	3.2	4.2
	Determine HIV1/2 boxes (100 each)	625	907,281	379,300	198,778	4.6	1.9
	Determine syphilis boxes (100 each)	345	164,370	286,900	50,029	3.3	5.7
	Uni-Gold HIV1/2 boxes (20 each)	641	158,582	48,240	21,400	7.4	2.3
pieces	Condoms female boxes (1000 each)	380	747,550		196,562	3.8	
	Condoms male boxes (144 each)	563	11,922,117	4,004,928	5,409,990	2.2	0.7

^{* &#}x27;Consumption per month' and 'Months of stock' for ARVs, CPT, INH and HIV test kits are based on the respective patient-regimen groups in the standard service reports. Estimates are based on the number of patients on the respective regimen at the end of the quarter evaluated and do not account for potential (positive or negative) growth. Facility stock positions for OI and STI drugs include HIV Program and other supply sources. Total national consumption and MoS estimates are used for these commodity groups. 'Months of stock' is calculated from the day of the physical stock count, which is on average 1 month after the end of the quarter.

19 Training and Mentoring

19.1 ART/PMTCT trainings

13 clinicians and 20 nurses were newly trained and certified as ART/PMTCT providers this quarter. Trainings were based on the 2014 guidelines.

64 ART / PMTCT program coordinators and other members of the district health management teams from **9** districts from the central region participated in quarterly review meeting. **64** members attended. District health management team (DHMT), ART coordinator and PMTCT coordinators consisted participants from each district. The objective of the review meeting is to strengthen data use.

160 nurses, clinicians, HTC counsellors and ART clerks participated in **4** district workshops aiming to improve data use.

240 health workers from the 3 districts participated EID mentorship meetings.

Health workers from 9 districts in the north and central region participated in E-MTCT review meetings that focused on the use dashboard in tracking indicators.

19.2 HIV Testing Services

21 HIV Diagnostic Assistants and 20 other HTS providers were newly trained and certified.

HIV testing master trainers and officers from the HIV Department monitored and supervised the intensive skills trainings in all the districts.

20 Participants in Q4 2015 Supervision (Site visits 6 - 19 January 2015)

Abgail Nyasulu (Nurse, MOH)

Absalom Kaunda (CO, MOH, Mzimba DHO)

Adamson Munthali (, BAYLOR)

Agnes Kalitsiro (Nurse, Mlambe Mission Hospital)

Alefa Fikira (CMT, MOH) Alexander Malunguza (, NTP) Andraida Mtoseni (Nurse, MOH) Andrew Dimba (, NTP)

Andrew Gompho (Clinician, MOH)

Andrew Mganga (M&E Officer, Dept for HIV & AIDS)

Andrew Ntenge (, JHPIEGO) Annex Kwekwesa (, moh) Annie Biza (Nurse, MDF) Annie Kamoto (Nurse, Private) Austins Namondwe (CO, CHAM) Batoni Upindi (TB Zonal Supervisor, MOH) Beatrice Malonje (Nurse, MOH) Benjamin Mazalo (CO, SUCOMA Clinic) Brighton Zumazuma (CO, NGO) Catherine Kassam (, MOH) Cecelia Tenesi (Nurse, MOH) Cecilia Manyawa (Nurse, MOH)

Cecilia Sambakunsi (Logistics Fellow, HIV Dept)

Chifundo Chomanika (MA, MHO) Chifundo Makuluni (Nurse, MOH) Chikayiko Majamanda (Nurse, MOH) Chikumbutso Pendame (MA, MOH)

Chimwemwe Francis Mkandawire (IT Fellow, Dept

for HIV and AIDS)

Chimwemwe Mlenga (, MOH) Chipulumutso Kambanje (, Dignitas) Chisomo Thondolo (Nurse, EGPAF) Chiukepo Longwe (CO, Private) Chrissy Lizengo (, MOH) Christopher Mkwezalamba (CO, MOH)

Clement Chiphota (CO, MoH) Cornelias Kang'ombe (, NTP) Dalitso Midiani (PMTCT Officer, MOH) Damison Msiska (CO, Dwangwa) Dan Nyirenda (, MOH)

Davie Maseko (CO, SOS) Davie Nkosi (, MOH) Deliwe Msiska (, JHPIEGO) Diana Chipande (, MOH)

Dorica Chirwa (Logistic officer, MOH) Dorica Sambo (Nurse, MOH) Dr Simon Chiumia (, Private) Edith Thaulo (Nurse, MOH) Egnatius Mtambalika (, DTO) Elton Masina (CO, EGPAF) Envance Njaidi (MA, MOH) Erik Mittochi (CO (ART coord), MOH)

Eustice Mhango (ART officer, MOH, Department of

HIV and AIDS)

Evans Kagwira (TB Zonal Supervisor, MOH) Everista Mkandawire (Nurse, MOH) Ezra Majoni (Nurse, MOH) Fainala Muyila (Nurse, MOH)

Fatsireni Mapulanga (, MOH) Francis Munthali (, COM) Frank Pondani (, Lighthouse)

Gabriel Lavout Kachere (clinician, MOH) Gerald Zomba (Program Officer, Dept for HIV and

AIDS)

Gift Pelani (, Baylor)

Golden Kachimanga (MA, MOH)

Grant Gondwe (, NTP)

Hannock Matupi (ARV clinician, MOH, Rumphi DH) Harold Malinda (CO, MOH, QECH)

Harry Tsapa (CO, MOH) Harvey Mafuta (, Machinga) Henry Banda (CO, MOH)

Henry Kanyerere (TB/HIV Program Officer, MOH)

Innocent Mainjeni (Logistics, MOH) Ireen Magongwa (, MSH) Isaiah Dambe (, NTP) Ivy Chibwana (, Dignitas) James Chilinda (, MOH) Janet Chikonda (Nurse, MOH) Jean Kayamba (Nurse, MOH) Jesse Lobeni (Nurse, MOH) John Kabichi (CO, MOH) Jotham Nyasulu (, MOH)

Judith Ntopa (Nurse, Cobbe Barracks)

Juliana Soko (ARV nurse, MOH, Livingstonia MH)

Juliet Nyirenda (Nurse, MOH) Justice Kaphiri (, NTP) Kelvin Makina (Logistics, Kasungu) Khumbo Ng'ona (, HIV Dept) Kingsley Makwale (MA, MOH) Kingsley Mbewa (CO, MOH)
Knox Banda (TB Zonal Supervisor, MOH)

Kondwani Chikoti (CO, MOH) Kondwani Kautsa (, MOH) Kumbukani Mataka (, JHPIEGO) Kuzani Mbendera (, NTP) Lameck Mlauzi (, NTP(MOH)) Lameck Mzava (, NTP) Laphoid Chisuwo (, NTP) Lawrence Sakali (, CHAM) Laywel Nyirenda (, EGPAF) Leonard Kadongola (, MOH) Licy Kadziweni (NMT, MOH) Lilian Kachali (Nurse, MOH) Limbani Kumambala (CO, Private) Limbani Mbetewa (, DTO) Lincy Chalunda (CO, MOH) Lioyd Wella (CO, MOH) Little Banda (, MOH)

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Lucy Banda (, MOH)

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Rodrick Kaulere (CO, CHAM (Sister Tereza))

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Tekia Banda (Nurse, MOH) Thoko Kalua (, HIV DEPT) Timothy Mwenyedini (MA, MOH) Vera Kajawa (Nurse, MOH)

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We thank all facility staff for their sincere welcome and co-operation with the HIV Department and its partners during these supportive visits. We congratulate all staff for their excellent work.

26 May 2016

21 Appendix (Full National HIV Program Data)

2015 Q4 (1st month of quarter, 2nd month of quarter, 3rd month of quarter)

Clients at health facility (static)

HTC client details

Total	HT	C	clie	nts	sen	hev
ı Otai		v	UIIC	IILO	361	/CU

Total HIV tested	558,890	100%
Sex	·	
Males tested	186,481	33%
Females tested	372,409	67%
Females non-pregnant	216,987	58%
Females pregnant	155,422	42%
Age		
Children 0-14 yrs	54,441	10%
Children below 12 mths (Age group A)	2,615	5%
Children 12 mths - 14 yrs (Age group B)	51,826	95%
Adults 15+ years	504,449	90%
Young adults 15-24 years (Age group C)	224,094	44%
Older adults 25+ yrs (Age group D)	280,355	56%
HTC access type		
PITC	334,086	60%
Family Referral Slip (FRS)	3,250	1%
Other (VCT, etc.) HTC access	221,554	40%
HTC first time / repeat		
Never tested before	158,396	28%
Previously accessed HTC	400,494	72%
Last negative	384,120	96%
Last positive	13,403	3%
Last exposed infant	2,386	1%
Last inconclusive	585	0%
Counseling session type / Partner present		
Counseled with partner / partner present	128,316	23%
Counseled alone / Partner not present	430,574	77%
Outcome summary (HIV test)		
Single test negative	511,776	92%
Single test positive	635	0%
Test 1&2 negative	670	0%
Test 1&2 positive	43,854	8%
Test 1&2 discordant	1,955	0%
Final result given to client		
Results among clients never tested / last negative	545,002	98%
New negative	511,669	94%
New positive	30,731	6%
New exposed infants	1,213	0%
New inconclusive	1,389	0%
Confirmatory results (previous positive clients)	13,888	2%
Confirmatory positive	13,349	96%
Confirmatory inconclusive	539	4%

2015 Q4 (1st month of quarter, 2nd month of quarter, 3rd month of quarter)

HTC client details

Partner / Family	HTC referral slips
------------------	--------------------

S	um of slips given 18,993	100%
	Total clients presenting with referral slip 3,250	17%
	Total failed referrals (slips not returned) 15,743	83%

Clients tested in the community

HTC client details

Total HIV tested

Total HTC clients served

Sex		
Males tested	12,769	40%
Females tested	19,361	60%
Females non-pregnant	16,311	84%
Females pregnant	3,050	16%

Age

Children 0-14 yrs	9,179	29%
Children below 12 mths (Age group A)	19	0%
Children 12 mths - 14 yrs (Age group B)	9,160	100%
Adults 15+ years	22,951	71%
Young adults 15-24 years (Age group C)	11,010	48%
Older adults 25+ yrs (Age group D)	11,941	52%

HTC access type

ĺ	PITC	10,773	34%
	Family Referral Slip (FRS)	27	0%
	Other (VCT, etc.) HTC access	21,330	66%

HTC first time / repeat

I	Never tested before	11,146	35%
ı	Previously accessed HTC	20,984	65%
	Last negative	20,235	96%
	Last positive	722	3%
	Last exposed infant	10	0%
	Last inconclusive	17	0%

Counseling session type / Partner present

Counseled with partner / partner present	2,164	7%
Counseled alone / Partner not present	29,966	93%

Outcome summary (HIV test)

Single test negative	30,175	94%
Single test positive	0	0%
Test 1&2 negative	13	0%
Test 1&2 positive	1,897	6%
Test 1&2 discordant	45	0%

32,130

100%

2015 Q4 (1st month of quarter, 2nd month of quarter, 3rd month of quarter)

HTC client details

Final result given to client

Results among clients never tested / last negative	31,401	98%
New negative	30,170	96%
New positive	1,182	4%
New exposed infants	9	0%
New inconclusive	40	0%
Confirmatory results (previous positive clients)	729	2%
Confirmatory positive	717	98%
Confirmatory inconclusive	12	2%
Partner / Family HTC referral cline	_	

Partner / Family HTC referral slips

Sum of slips given	4,346	100%
Total clients presenting with referral slip	27	1%
Total failed referrals (slips not returned)	4,319	99%

Clients at stand-alone HTC sites

HTC client details

Total HIV tested

Total HTC clients served

	•	
Sex		
Males tested	5,802	37%
Females tested	9,736	63%
Females non-pregnant	7,806	80%
Females pregnant	1,930	20%

Age

Children 0-14 yrs		901	6%
(Children below 12 mths (Age group A)	7	1%
(Children 12 mths - 14 yrs (Age group B)	894	99%
Adults 1	5+ years	14,637	94%
)	Young adults 15-24 years (Age group C)	5,651	39%
(Older adults 25+ yrs (Age group D)	8,986	61%

HTC access type

PITC	12,161	78%
Family Referral Slip (FRS)	200	1%
Other (VCT, etc.) HTC access	3,177	20%

HTC first time / repeat

١	Never tested before 3,623	23%
F	Previously accessed HTC 11,915	77%
	Last negative 11,363	95%
	Last positive 541	5%
	Last exposed infant 4	0%
	Last inconclusive 7	0%

Counseling session type / Partner present

Counseled with partner / partner present	2,716	17%
Counseled alone / Partner not present	12,822	83%

15,538

100%

2015 Q4 (1st month of quarter, 2nd month of quarter, 3rd month of quarter)

HTC client details

Outcome summary (HIV test)

Single test negative	14,102	91%
Single test positive	6	0%
Test 1&2 negative	2	0%
Test 1&2 positive	1,401	9%
Test 1&2 discordant	27	0%

Final result given to client

Results amo	ong clients never tested / last negative	5,017	97%
New	negative 1	4,140	94%
New	positive	853	6%
New	exposed infants	7	0%
New	rinconclusive	17	0%
Confirmator	y results (previous positive clients)	521	3%
Conf	firmatory positive	515	99%
Conf	firmatory inconclusive	6	1%

Partner / Family HTC referral slips

3	Sum of slips given	528	100%
	Total clients presenting with referral slip	200	38%
	Total failed referrals (slips not returned)	328	62%

Blood safety Malawi (national)

2015 Q4 (1st month of quarter, 2nd month of quarter, 3rd month of quarter)

Infect. disease screening among potential donors

HIV	scree	חחוחמ
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HIV testing not done	1,713	18%
Tested for HIV	7,854	82%
HIV negative	7,322	93%
HIV positive	532	7%

Hepatitis B screening

HepB testing not done	1,809	19%
Tested for Hepatitis B	7,758	81%
HepB Negative	7,308	94%
HepB Positive	450	6%

Hepatitis C screening

HepC testing not done	5,598	59%
Tested for Hepatitis C	3,969	41%
HepC Negative	3,877	98%
HepC Positive	92	2%

Syphilis screening

Syphilis testing not done	2,004	21%
Tested for Syphilis	7,563	79%
Syphilis Negative	7,379	98%
Syphilis Positive	184	2%

Malaria screening

Malaria testing not done	3,003	31%
Tested for malaria	6,564	69%
Malaria Negative	6,102	93%
Malaria Positive	462	7%

Summary screening outcome

Not d	onated	2,677	28%
Dona	ted	6,890	72%
	Screened for at least HIV, HepB and syphilis		
	Screened for HIV, HepB, HepC, Syphilis, Malaria	4,606	70%
	Screened for HIV, HepB, Syphilis	1,977	30%
	Screened for HIV, HepB	2	0%
Screened for HIV only			0%
Screened with any other combination of tests			4%

Cross-matching report

Blood group typing (for units and patients)

Total blood group typing done	24,969	100%
Blood units cross-matched (by source)		

Total blood units cross-matched	17,305	100%
Total units from MBTS (estimated)	10,415	60%
Total units from replacement donors	6,890	40%

Blood units cross-matched by patient group

Units cross-matched for maternity	3,509	20%
Units cross-matched for paediatrics	4,949	29%
Units cross-matched for other ward	8,847	51%

Page 1 of 2

Blood safety Malawi (national)

2015 Q4 (1st month of quarter, 2nd month of quarter, 3rd month of quarter)

Cross-matching report

Transfusion reactions

Units transfused without adverse events	17,285	100%
Units with suspected transfusion reactions	16	0%
Units with confirmed transfusion reactions	4	0%

2015 Q4 (1st month of quarter, 2nd month of quarter, 3rd month of quarter)

Age 2 months

Age	col	hort	outc	omes
Ayu	UU	IVI L	Juli	

Total children in birth cohort		
Total children registered	8,774	100%
CPT status		
On CPT	7,863	90%
Not on CPT	911	10%
HIV status		
Current HIV infection status unknown	5,858	67%
HIV infection not confirmed, not ART eligible	5,849	100%
HIV infection not confirmed, ART eligible (PSHD)	9	0%
Current HIV infection status known	2,916	33%
Confirmed not infected	2,876	99%
Confirmed infected (ART eligible)	40	1%
ART eligibility summary		
Not eligible for ART	8,725	99%
ART eligible	49	1%
ART not initiated	12	24%
Initiated ART	37	76%
Primary follow-up outcome		
Discharged uninfected	19	0%
Continue follow-up	7,929	92%
Started ART	37	0%
Defaulted	577	7%
Died	30	0%
Transfers between sites		
Total not transferred out	8,592	98%
Transferred out	182	2%
Age 12 months		
Age cohort outcomes		
Total children in birth cohort		*
Total children registered	9,767	100%
CPT status	,	
On CPT	7,051	72%
Not on CPT	2,716	28%
HIV status	, , ,	
Current HIV infection status unknown	5,178	53%
HIV infection not confirmed, not ART eligible	5,161	100%
HIV infection not confirmed, ART eligible (PSHD)	17	0%
Current HIV infection status known	4,589	47%
	.,,,,	

Confirmed not infected

Confirmed infected (ART eligible)

4,399

190

96%

4%

2015 Q4 (1st month of quarter, 2nd month of quarter, 3rd month of quarter)

Age cohort outcomes

ADT 11 11 11 11		
ART eligibility summary		
Not eligible for ART	9,560	98%
ART eligible	207	2%
ART not initiated	38	18%
Initiated ART	169	82%
Primary follow-up outcome		
Discharged uninfected	78	1%
Continue follow-up	7,028	74%
Started ART	169	2%
Defaulted	2,117	22%
Died	68	1%
Transfers between sites		
Total not transferred out	9,460	97%
Transferred out	307	3%
Total children registered	8,832	100%
CPT status		
On CPT	1,015	11%
Not on CPT	7,817	89%
HIV status		
Current HIV infection status unknown	4,403	50%
HIV infection not confirmed, not ART eligible	4,395	100%
HIV infection not confirmed, ART eligible (PSHD)	8	0%
Current HIV infection status known	4,429	50%
Confirmed not infected	4,226	95%
Confirmed infected (ART eligible)	203	5%
ART eligibility summary		
Not eligible for ART	8,621	98%
ART eligible	211	2%
ART not initiated	-4	-2%

١	Not eligible for ART 8,621	98%
Α	ART eligible 211	2%
	ART not initiated -4	-2%
	Initiated ART 215	102%

Primary follow-up outcome

Discharged uninfected 4	1,094	48%
Continue follow-up	767	9%
Started ART	215	3%
Defaulted 3	3,376	39%
Died	109	1%

Transfers between sites

Total not transferred out	8,561	97%
Transferred out	271	3%

Malawi (national) **Antenatal Care**

2015 Q4 (1st month of quarter, 2nd month of quarter, 3rd month of quarter)

New ANC registrations in reporting period

Women with first visit in reporting period		
New women registered	139,322	100%
ANC cohort analysis		*
Trimester of first visit		
Started ANC 0-12 wks	16,173	12%
Started ANC 13+ wks	123,149	88%
HIV status ascertainment		
HIV status not ascertained	14,179	10%
HIV status ascertained	125,143	90%
Valid previous test result	10,087	8%
Previous negative	5,008	50%
Previous positive	5,079	50%
New test at ANC	115,056	92%
New negative	110,512	96%
New positive	4,544	4%
HIV status summary		
Total women HIV negative	115,520	92%
Total women HIV positive	9,623	8%
PMTCT regimen mother		
No ARVs	936	10%
Any ARVs	8,687	90%
ART (by time of initiation)	8,687	100%
Already on ART when starting ANC	4,498	52%
Started ART at 0-27 weeks of pregnancy	3,532	41%
Started ART at 28+ weeks of preg.	657	8%
ANC women after 6 months		
ANC cohort analysis		*
Total women completing ANC in the reporting period		^
Total women in booking cohort	150,212	100%
Visits per woman	,	
Women with 1 visit	31,149	21%
Women with 2 visits	37,278	25%
Women with 3 visits	45,179	30%
Women with 4 visits	29,459	20%
Women with 5+ visits	7,147	5%
Pre-eclampsia Pre-eclampsia	,	
No pre-eclampsia	147,247	98%
Pre-eclampsia	2,965	2%
TTV doses	_,,,,,	
0-1 TTV doses	72,485	48%
2+ TTV doses	72,465 77,727	40% 52%
	11,121	JL /0
SP tablets	10.454	440/
0 SP doses	16,154	11%
1 SP dose (1 x 3 tabs)	33,956	23% 67%
6+ SP tablets (2 x 3 tabs)	100,102	67%

Antenatal Care Malawi (national)

2015 Q4 (1st month of quarter, 2nd month of quarter, 3rd month of quarter)

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ANC conort analysis		*
FeFo tablets		
0-119 FeFo tablets	117,796	78%
120+ FeFo tablets	32,416	22%
Albendazole (Deworming)		
0 Albend. doses	24,711	17%
1 Albend. dose	124,682	83%
ITN (bednets)		
No ITN	17,978	12%
ITN received	134,137	88%
Syphilis status		
Not tested for syphilis	113,373	75%
Tested for syphilis	36,839	25%
Syphilis negative	36,232	98%
Syphilis positive	607	2%
HIV status ascertainment		
HIV status not ascertained	12,254	8%
HIV status ascertained	137,958	92%
Valid previous test result	10,343	7%
Previous negative	4,671	45%
Previous positive	5,672	55%
New test at ANC	127,615	93%
New negative	122,620	96%
New positive	4,995	4%
HIV status summary		
Total women HIV negative	127,291	92%
Total women HIV positive	10,667	8%
CPT status (among HIV pos)		
Not on CPT	523	5%
On CPT	10,144	95%
PMTCT regimen mother		
No ARVs	468	4%
Any ARVs	10,199	96%
ART (by time of initiation)	10,199	100%
Already on ART when starting ANC	5,266	52%
Started ART at 0-27 weeks of pregnancy	3,962	39%
Started ART at 28+ weeks of preg.	971	10%
Baby's ARVs dispensed		
No ARVs dispensed for infant	1,297	12%
ARVs dispensed for infant	9,370	88%

Malawi (national)

2015 Q4 (1st month of quarter, 2nd month of quarter, 3rd month of quarter)

Maternal details *

Admissions	in	the	reporting	period
,	•••			P00

Total admissions (referrals double-counted)	125,743	100%
Not referred to other site (total women)	119,382	95%
Referred out before delivery (multiple admissions)	6,361	5%

HIV status ascertainment

HIV status not ascertained	2,995	2%
HIV status ascertained	123,039	98%
Valid previous test result	119,597	97%
Previous negative	110,750	93%
Previous positive	8,847	7%
New test at maternity	3,442	3%
New negative	3,141	91%
New positive	301	9%

HIV status summary

	Total women HIV negative	113,891	93%
•	Total women HIV positive	9,148	7%

ARVs during pregnancy (among HIV pos)

No ARV in pr	No ARV in pregnancy		1%
Any ARVs	Any ARVs		99%
ART (by time of initiation)	9,073	100%
	ART initiated before pregnancy	7,207	79%
	ART initiated in 1st / 2nd trimester	1,090	12%
	ART initiated in 3rd trimester	657	7%
	ART initiated during labour	119	1%

Obstetric complications

·		
No obstetric complications 111,498		88%
Any obstetric complications 14,5		12%
Haemorrhage	2,284	16%
Haemorrhage ante-partum	710	31%
Haemorrhage post-partum	1,574	69%
Obstr / prol labour	4,810	33%
(pre-) Eclampsia	1,087	7%
Maternal sepsis	339	2%
Ruptured uterus	194	1%
Other obstetric complications	5,822	40%

Emergency obstetric care

Oxytocin 112,874	95%
Anticonvulsive 524	0%
Antibiotics 5,193	4%
Blood transfusion 318	0%
Manual removal of placenta 494	0%

Vitamin A

Vit A not given	34,697	28%
Vit A given	91,337	72%

Malawi (national)

2015 Q4 (1st month of quarter, 2nd month of quarter, 3rd month of quarter)

Maternal det	tails		*

Staff conducting delivery

Category A: MO, CO, nurse/midwife, MA	113,890	95%
Category B: PA, WA, HSA	640	1%
Category C: Other	5,143	4%

Mother survival

Mother alive	119,567	100%
Mother died	106	0%

Infant details *

Single babies / multiple deliveries

1	Total babies delivered	121,915	100%
	Single babies	117,815	97%
	Twin / multiple babies	4,100	3%

Delivery place

Total	deliveries at a health facility	116,555	96%
	This facility	116,086	100%
	Other facility	469	0%
Total deliveries before reaching the facility		5,360	4%
	In transit	3,607	67%
	Home / TBA	1,753	33%

Delivery mode

Spontaneous vaginal	109,863	90%
Vacuum extraction	1,718	1%
Breech	2,191	2%
Caesarean section	8,143	7%

Infant complications

No in	No infant complications 104,548		86%
Total	infants with complications	17,367	14%
	Prematurity	3,663	21%
	Weight less 2500g	5,211	30%
	Asphyxia	5,387	31%
	Sepsis	1,292	7%
	Other newborn complication	1,814	10%

Infant survival

Total live births		119,729	98%
	Discharged alive	118,586	99%
	Neonatal deaths	1,143	1%
Stillbir	ths	2,186	2%
	Stillbirth, fresh	1,226	56%
	Stillbirth, macerated	960	44%

Malawi (national)

2015 Q4 (1st month of quarter, 2nd month of quarter, 3rd month of quarter)

Infant details *

HIV exposure / ARV proph. (among discharged alive)

III CAPO	saire / Arty propii. (among discharged anve)		
Infants with unknown HIV exposure status		2,296	2%
Infants w	Infants with known HIV exposure status	116,290	98%
N	ot HIV exposed	107,542	92%
H	IV exposed	8,748	8%
	Received no ARVs	876	10%
	Received ARVs	7,872	90%
	Nevirapine	7,872	100%
Breastfe	eding initiated		
BF not st	arted within 60min	11,232	9%
BF starte	d within 60min	110,683	91%
Tetracyc	line eye ointment given		

TO not given	17,949	15%
TO given	103,966	85%

Registration details

HCC clinic registrations

HCC clinic registrations		
Total HCC registrations	16,274	100%
Registration type		
Patients enrolled first time	15,185	93%
Patients re-enrolled		0%
Patients transferred in	1,048	6%
Sex		
Males (all ages)	7,605	47%
Females (all ages)	8,669	53%
Non-pregnant	8,648	100%
Pregnant	21	0%
Age at registration		
Adults 15+ yrs	5,295	33%
Children 0-14 yrs	10,979	67%
Children 24 months - 14 years	441	4%
Children below 24 months (exposed children)	10,538	96%
Children 2 - below 24 months	3,407	32%
Infants below 2 months	7,131	68%
Reason for HCC registration		
Exposed infants	10,624	65%
Confirmed infected patients (pre-ART)	5,650	35%
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HCC clinic registrations

HCC clinic registrations		
Total HCC registrations	388,424	100%
Registration type		
Patients enrolled first time	373,866	96%
Patients re-enrolled	1,205	0%
Patients transferred in	13,353	3%
Sex		
Males (all ages)	169,516	44%
Females (all ages)	218,908	56%
Non-pregnant	218,034	100%
Pregnant	874	0%
Age at registration		
Adults 15+ yrs	188,099	48%
Children 0-14 yrs	200,325	52%
Children 24 months - 14 years	17,259	9%
Children below 24 months (exposed children)	183,066	91%
Children 2 - below 24 months	83,599	46%
Infants below 2 months	99,467	54%
Reason for HCC registration		
Exposed infants	181,896	47%
Confirmed infected patients (pre-ART)	206,528	53%
Pre-ART follow-up outcome		*
Primary follow-up outcomes		
Total retained in pre-ART	43,133	22%
Started ART	104,983	53%
Defaulted	48,965	25%
Died	1,897	1%
Transfers between sites		
Total not transferred out	199,048	96%
Transferred out	7,480	4%

Registration details

ART clinic registrations

ART clinic registrations		
Total ART clinic registrations	1,091,656	100%
Registration type		
First time ART initiations (total patients)	872,567	80%
ART re-initiations	12,334	1%
ART transfers in	206,755	19%
Sex		
Males	393,498	36%
Females	698,158	64%
Non-pregnant	564,661	81%
Pregnant	133,497	19%
Age at ART initiation		
Adults 15+ yrs	997,286	91%
Children 0-14 yrs	94,370	9%
Children 2-14 yrs	72,623	77%
Children below 24 mths	21,747	23%
Reason for starting ART		
Presumed severe HIV Disease	3,707	0%
Confirmed HIV infection		100%
WHO stage 1 or 2	488,873	45%
Total lymphocytes <threshold< td=""><td>250</td><td>0%</td></threshold<>	250	0%
CD4 below threshold	323,010	66%
CD4 unknown or >threshold	165,613	34%
PCR infants	2,988	2%
Children 12-59 mths	8,097	5%
Pregnant women	113,931	69%
Breastfeeding mothers	40,597	25%
WHO stage 3	487,560	45%
WHO stage 4	103,674	10%
Unknown / reason outside of guidelines	7,842	1%
TB at ART initiation		
Never TB / TB > 24 months ago	1,017,057	93%
TB within the last 24 months	38,357	4%
Current episode of TB	36,242	3%
Kaposi's sarcoma at ART initiation		
No KS	1,071,606	98%
Patients with KS	20,050	2%

ART outcomes *

Primary f	follow-up	outcomes
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Total	Total alive on ART 591,176		67%
	Alive on ART at site of last registration	591,925	100%
	ART patients in transit between sites	-749	0%
Defau	Defaulted 207,968		24%
Stopp	Stopped ART 4,293		0%
Total	Total died 81,471		9%
	Died month 1	19,542	24%
	Died month 2	12,275	15%
	Died month 3	7,393	9%
	Died month 4+	42,261	52%

Transfers between sites

Total not transferred out	885,650	81%
Transferred out	206,006	19%

ART regimens

First line regimens 583,6	74 99%
Adult formulation 557,6	10 96%
Regimen 0A	55 0%
Regimen 1A	85 0%
Regimen 2A 28,6	86 5%
Regimen 3A	03 0%
Regimen 4A 7	16 0%
Regimen 5A 517,3	46 93%
Regimen 6A 9,4	19 2%
Paed. formulation 26,0	64 4%
Regimen 0P 5	08 2%
Regimen 1P	97 0%
Regimen 2P 25,0	01 96%
Regimen 3P	36 0%
Regimen 4P	22 2%
Second line regimens 7,7	69 1%
Adult formulation 6,7	03 86%
Regimen 7A 4,6	60 70%
Regimen 8A 2,0	43 30%
Paed. Formulation 1,0	66 14%
Regimen 9P 1,0	66 100%
Other regimen (adult / paed)	82 0%

Adherence

ĺ	Adherence unknown (not recorded) 5,472		1%
	Adherence recorded	586,453	99%
	0-3 doses missed	530,724	90%
	4+ doses missed	55,729	10%

ART side effects

Side effects unknown (not recorded) 40,461		7%
Side effects recorded	551,464	93%
No side effects	543,843	99%
Any side effects	7,621	1%

ART outcomes *

Current TB status among ART patients (ICF)

ICF not done (Current TB status unknown/ not circ) 13,100		2%
ICF done 578,825		98%
TB not suspec	ed 574,286	99%
TB suspected		1%
TB confirmed		0%
TB con	firmed, not on treatment 181	21%
TB con	firmed, on TB treatment 672	79%

Registration details

ART clinic re	gistrations		
Total ART clin	ic registrations	32,402	100%
Registration	уре		
First time ART	initiations (total patients)	24,082	74%
ART re-initiation	ons	477	1%
ART transfers	in	7,843	24%
Sex			
Males		11,753	36%
Females		20,649	64%
Non-pr	egnant	14,381	70%
Pregna	int	6,268	30%
Age at ART in	nitiation		
Adults 15+ yrs		29,797	92%
Children 0-14	yrs	2,605	8%
Childre	n 2-14 yrs	1,973	76%
Childre	n below 24 mths	632	24%
Reason for st	arting ART		
Presumed sev	ere HIV Disease	149	0%
Confirmed HIV infection 32,25		32,253	100%
WHO stage 1 or 2		20,231	63%
	Total lymphocytes <threshold< td=""><td>4</td><td>0%</td></threshold<>	4	0%
	CD4 below threshold	11,544	57%
	CD4 unknown or >threshold	8,683	43%
	PCR infants	93	1%
	Children 12-59 mths	608	7%
	Pregnant women	6,264	72%
14//10	Breastfeeding mothers	1,718	20%
WHO s		10,237	32%
WHO s	· ·	1,456	5%
	wn / reason outside of guidelines	329	1%
TB at ART ini		04 540	070/
	> 24 months ago	31,540	97%
	ast 24 months	372	1%
Current episod		490	2%
•	coma at ART initiation	00.400	000/
No KS	40	32,106	99%
Patients with I	(S	296	1%

12 month survival children

Survival and retention in ART program

ART cohort registration group outcomes

Total A	ART clin	ic regis	trations	2,633	100%
	Transfers out (double counted)		285	11%	
	Total n	ot trans	eferred out (patients in cohort)	2,348	89%
	Total alive on ART 1,80		1,806	77%	
	Total not retained 54		542	23%	
			Defaulted	449	83%
			Stopped ART	12	2%
	Died 81		15%		

12 month survival all ages

Survival and retention in ART program

ART cohort registration group outcomes

Total AR	Total ART clinic registrations 32,421		100%	
Т	Transfers out (double counted)		3,116	10%
Т	Γotal not trar	nsferred out (patients in cohort)	29,305	90%
	Total alive on ART		22,332	76%
	Total	not retained	6,973	24%
		Defaulted	5,929	85%
		Stopped ART	100	1%
		Died	944	14%

24 month survival all ages

Survival and retention in ART program

ART cohort registration group outcomes

Total ART cl	Total ART clinic registrations 30,011		100%
Trans	Transfers out (double counted)		13%
Total	Total not transferred out (patients in cohort)		87%
	Total alive on ART		70%
	Total not retained	7,884	30%
	Defaulted	6,549	83%
	Stopped ART	133	2%
	Died	1,202	15%

36 month survival all ages

Survival and retention in ART program

Total /	Total ART clinic registrations 32,17		32,177	100%	
	Transfers out (double counted)		4,894	15%	
	Total n	ot trans	sferred out (patients in cohort)	27,283	85%
	Total alive on ART		18,532	68%	
		Total r	not retained	8,751	32%
			Defaulted	6,926	79%
			Stopped ART	154	2%
			Died	1,671	19%

48 month survival all ages

Survival and retention in ART program

ART cohort registration group outcomes

Total A	Total ART clinic registrations 44,10		44,109	100%	
	Transfers out (double counted)		7,443	17%	
	Total n	ot trans	ferred out (patients in cohort)	36,666	83%
	Total alive on ART		25,295	69%	
		Total r	not retained	11,371	31%
			Defaulted	8,938	79%
			Stopped ART	179	2%
			Died	2,254	20%

60 month survival all ages

Survival and retention in ART program

ART cohort registration group outcomes

Total ART	Total ART clinic registrations 22,309		100%	
Tra	Transfers out (double counted)		5,447	24%
То	otal not trans	ferred out (patients in cohort)	16,862	76%
	Total alive on ART		10,555	63%
	Total r	not retained	6,307	37%
		Defaulted	4,349	69%
		Stopped ART	136	2%
		Died	1,822	29%

72 month survival all ages

Survival and retention in ART program

ART cohort registration group outcomes

Total ART c	Total ART clinic registrations		100%
Trans	Transfers out (double counted)		26%
Total	Total not transferred out (patients in cohort)		74%
	Total alive on ART		60%
	Total not retained		40%
	Defaulted	4,543	68%
	Stopped ART	98	1%
	Died	2,080	31%

84 month survival all ages

Survival and retention in ART program

Total ART clir	Total ART clinic registrations 18,942		100%
Transf	Transfers out (double counted)		28%
Total r	Total not transferred out (patients in cohort)		72%
	Total alive on ART		56%
	Total not retained	6,000	44%
	Defaulted	3,789	63%
	Stopped ART	115	2%
	Died	2,096	35%

96 month survival all ages

Survival and retention in ART program

ART cohort registration group outcomes

Total A	Total ART clinic registrations 14,581		100%		
	Transfers out (double counted)		4,453	31%	
	Total n	ot trans	eferred out (patients in cohort)	10,128	69%
	Total alive on ART		5,137	51%	
		Total r	not retained	4,991	49%
			Defaulted	3,159	63%
			Stopped ART	59	1%
			Died	1,773	36%

108 month survival all ages

Survival and retention in ART program

ART cohort registration group outcomes

Total AF	Total ART clinic registrations 12		12,107	100%
7	Transfers out (double counted)		3,551	29%
1	Total not tra	insferred out (patients in cohort)	8,556	71%
	Total alive on ART		4,034	47%
	Tota	al not retained	4,522	53%
		Defaulted	2,440	54%
		Stopped ART	65	1%
		Died	2,017	45%

120 month survival all ages

Survival and retention in ART program

ART cohort registration group outcomes

Total Al	Total ART clinic registrations		7,337	100%
	Transfers out (double counted)		2,238	31%
	Total not transferred out (patients in cohort)		5,099	69%
	Total alive on ART		2,076	41%
	Total not retained		3,023	59%
		Defaulted	1,485	49%
		Stopped ART	39	1%
		Died	1,499	50%

6 month survival OptionB+

Survival and retention in ART program

Total /	Total ART clinic registrations		7,716	100%	
	Transfers out (double counted)		489	6%	
	Total not transferred out (patients in cohort)		7,227	94%	
	Total alive on ART		5,482	76%	
		Total n	ot retained	1,745	24%
			Defaulted	1,704	98%
			Stopped ART	19	1%
			Died	22	1%

12 month survival OptionB+

Survival and retention in ART program

ART cohort registration group outcomes

Total A	Total ART clinic registrations 8,		8,556	100%
	Transfers out (double counted)		824	10%
	Total not transferred out (patients in cohort)		7,732	90%
	Total alive on ART		5,319	69%
		Total not retained	2,413	31%
		Defaulted	2,318	96%
		Stopped ART	31	1%
		Died	64	3%

24 month survival OptionB+

Survival and retention in ART program

ART cohort registration group outcomes

Total ART clinic registrations		8,171	100%
Transfers out (double counted)		955	12%
Total not transferred out (patients in cohort)		7,216	88%
Т	otal alive on ART	4,790	66%
Т	Total not retained		34%
	Defaulted	2,319	96%
	Stopped ART	31	1%
	Died	76	3%

36 month survival OptionB+

Survival and retention in ART program

Total ART clinic registrations		9,625	100%
Transfers out (double counted)		1,310	14%
Total not transferred out (patients in cohort)		8,315	86%
Total	Total alive on ART		65%
Total not retained		2,895	35%
	Defaulted	2,689	93%
	Stopped ART	46	2%
	Died	160	6%

2015 Q4 (1st month of quarter, 2nd month of quarter, 3rd month of quarter)

STI clients treated in the reporting period

Total STI clients

Total STI clients		
Total STI clients treated	55,938	100%
Index patients treated (symptomatic)	45,685	82%
Partners treated	10,253	18%
Sex		
Males	22,324	40%
Females	33,614	60%
Non-pregnant	29,578	88%
Pregnant	4,036	12%
Age group		
Age group A (0-19 years)	5,300	9%
Age group B (20-24 years)	13,584	24%
Age group C (25+ years)	37,054	66%
Client type		
Symptomatic cases	49,986	89%
Index cases	45,685	91%
Partners symptomatic	4,301	9%
Partners asymptomatic	5,952	11%
STI treatment history		
Never treated for STI	41,091	73%
Previously treated for STI	14,847	27%
Old >3 months ago	10,569	71%
Recent ≤3 months ago	4,278	29%
STI syndromic diagnosis		
GUD	9,705	15%
UD	14,664	23%
AVD	19,631	31%
Low risk	7,515	38%
High risk	12,116	62%
LAP	8,841	14%
SS	816	1%
BU	806	1%
BA	1,031	2%
NC	402 586	1%
Genital Warts		1%
Syphilis RPR VDRL 1,649		3% 8%
Other STI 5,036		
STI partner notification		
Total partner notification slips issued	14,919	100%
Total partners returned	10,253	69%
Total partners not seen	4,666	31%

2015 Q4 (1st month of quarter, 2nd month of quarter, 3rd month of quarter)

STI clients treated in the reporting period

HIV test / ART status

HIV status not ascertained 21,		21,059	38%		
HIV status ascertained		34,879	62%		
HIV negative (new test)		26,186	75%		
	HIV positive		8,693	25%	
		New po	sitive	2,297	26%
		Previous	s positive	6,396	74%
			Not on ART	1,259	20%
			On ART	5,137	80%

STI clients referred for services

Lab	477	2%
Gynae review	394	2%
Surgical review	259	1%
Repeat HTC	18,107	80%
ART (for assessment)	1,350	6%
PMTCT	133	1%
Other (service referrals)	1,845	8%