



Government of Malawi Ministry of Health

Integrated HIV Program Report April – June 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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HIV Department website: www.hiv.health.gov.mw*

1	EXECUTIVE SUMMARY	2
2	INTEGRATED HIV PROGRAM OVERVIEW	3
3	SUPPORTIVE SITE SUPERVISION	4
3.1	METHODS	4
3.2	SUPERVISION OUTCOMES.....	4
4	INVENTORY OF SITES AND SERVICES	5
4.1	SITES AND SERVICES.....	5
4.2	STAFFING OF HIV SERVICES.....	6
5	HIV TESTING AND COUNSELLING PROGRAM OUTPUTS	7
5.1	HTC DATA.....	8
5.2	HTC ACCESS TYPE.....	8
5.3	AGE AND SEX DISTRIBUTION AMONG HTC CLIENTS	8
5.4	FIRST TIME, REPEAT AND CONFIRMATORY TEST RESULTS.....	8
6	DNA-PCR TESTING FOR EARLY INFANT DIAGNOSIS OF HIV (EID).....	9
7	BLOOD SAFETY	10
8	POST EXPOSURE PROPHYLAXIS (PEP)	11
9	PROVIDER-INITIATED FAMILY PLANNING (PIFP)	11
10	COTRIMOXAZOLE PREVENTIVE THERAPY (CPT)	11
11	TB / HIV INTERVENTIONS	12
11.1	INTENSIFIED CASE FINDING (ICF).....	12
11.2	ISONIAZID PREVENTIVE THERAPY (IPT).....	12
12	HIV-RELATED DISEASES	13
13	HIV-EXPOSED CHILD FOLLOW-UP	13
13.1	METHODS AND DEFINITION OF INDICATORS.....	13
13.2	HIV EXPOSED CHILD REGISTRATION DATA.....	14
13.3	BIRTH COHORT OUTCOMES.....	14
14	PRE-ART	15
14.1	PRE-ART REGISTRATION DATA	15
14.2	CUMULATIVE PRE-ART FOLLOW-UP OUTCOMES.....	15
15	PMTCT / ART	15
15.1	DATA SOURCES AND REPORTING METHODS	15
15.2	ARV COVERAGE AMONG PREGNANT / BREASTFEEDING WOMEN AND EXPOSED INFANTS	17
15.3	HIV SERVICES AT ANC.....	18
15.4	HIV SERVICES AT MATERNITY	18
16	ART ACCESS AND FOLLOW-UP OUTCOMES	19
16.1	NEW ART REGISTRATIONS DURING Q2 2015	19
16.2	CUMULATIVE ART REGISTRATIONS UP JUNE 2015	21
16.3	ART OUTCOMES.....	21
16.4	ART COHORT SURVIVAL ANALYSIS	23
16.5	VIRAL LOAD (VL) MONITORING	27
17	TB / HIV MANAGEMENT.....	29
18	STI TREATMENT.....	29
18.1	ACCESS TO STI TREATMENT AND COVERAGE	30
18.2	CLIENT TYPE AND STI HISTORY.....	30
18.3	HIV STATUS	30
18.4	STI SYNDROMES	30
18.5	REFERRALS	30
19	SUPPLY OF HIV PROGRAM COMMODITIES	30
19.1	QUANTIFICATION AND PROCUREMENT PLANNING.....	30
19.2	BIMONTHLY DISTRIBUTION OF HIV & MALARIA COMMODITIES	31
19.3	QUARTERLY LOGISTICS SUPPORT DURING QUARTER 2 ART/PMTCT SUPERVISION	31
19.4	STOCK STATUS OF HIV COMMODITIES	31
19.5	AVAILABILITY OF STANDARD FIRST LINE ARVS.....	32
20	TRAINING AND MENTORING	34
21	PARTICIPANTS IN Q2 2015 SUPERVISION (SITE VISITS 20 JULY-7 AUGUST 2015)	35
22	APPENDIX (FULL NATIONAL HIV PROGRAM DATA).....	1

1 Executive Summary

A summary of the key achievements between **April and June 2015** is provided below:

- Scale-up of integrated HIV services had reached the following number of sites:
 - **724** static (579 within and 145 outside of health facilities) and 188 outreach HTC sites
 - **711** (static) ART sites
 - **601** PMTCT sites (Option B+, all included in ART sites above)
 - **662** Pre-ART sites
 - **642** sites with HIV-exposed child follow-up
- **493,567** persons were tested and counselled for HIV; **150,728 (31%)** accessed HTC for the first time; **342,839 (69%)** were repeat testers and **9,820 (3%)** of these received confirmatory testing (after having tested positive in the past). This is equivalent to **35%** confirmatory testing coverage among 25,957 patients initiating ART this quarter. **29,285 (6%)** clients received a positive result for the first time.
- **19,596 (95%)** of 20,538 blood units collected were screened for (at least) HIV, hepatitis B and syphilis.
- **141,212 (91%)** of 155,272 women at ANC had their HIV status ascertained; **10,766 (8%)** of these were HIV positive. **120,781 (96%)** of 125,961 women at maternity had their HIV status ascertained **8,967 (7%)** of these were HIV positive.
- **25,957** patients started ART this quarter.
- **568,470** patients were alive and on ART by end of June 2015. This means that **57%** of the estimated 1 million HIV positive population was on ART. ¹ Estimated ART coverage among people in need for treatment¹ was **49%** (49,407 / 101,000) for children (<15 years) and **67%** (519,063 / 779,000) for adults.
- **79%** of adults and **78%** 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 16.4)
- **493,034 (93%)** of 531,106 patients on first line adult ART were on regimen 5A (TDF/3TC/EFV).
- **10,460² (70%)** of an estimated **14,926¹** HIV infected pregnant women in Malawi were on ART this quarter. **5,960 (57%)** of these were already on ART when getting pregnant and **4,500 (43%)** started ART during pregnancy/delivery.
- An additional **1,591²** breastfeeding women started ART due to **Option B+** (in WHO stage 1/2)
- **76%, 72%, 67%** and **69%** of women started under **Option B+** were retained on ART at **6, 12, 24** and **36 months** after initiation, respectively.
- **8,210 (7%)** of infants discharged alive from maternity were known to be HIV exposed, **7,771 (95%)** of these received ARV prophylaxis (nevirapine). **6,563 (80%)** were enrolled in exposed child follow-up before age 2 months.
- A total of **10,396** HIV exposed children and **6,077** 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 HIV-transmission 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 all 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 711 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
- Follow up on action points noted during the previous visit
- Next visit date
- M&E reports from HTC, ANC, maternity, exposed child and pre-ART follow-up, ART and TB
- Physical drug stock-level assessment
- Identification of sites in urgent need of clinical mentoring
- 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

716 public and private sector facilities were visited for **clinical HIV program supervision** between 20th July and 7th August 2015.

The large number of sites was covered by **95** supervisors working in **23** teams that spent a total of **1,991 working hours** at the sites. Each site visit lasted on average **2.8** hours, but up to 2 days were spent at the busiest sites. **372 (52%)** sites were awarded a *Certificate of Excellence* for **excellent performance**. The

number of sites with excellent performance is higher than previous quarter 318. **53 (7%)** 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 Q2

Zone	Total facil. visited*	Supervision hours spent at facilities		Performance (# and % of sites)	
		Total	Average per site	Excellent perform.	Mentoring needed
NZ	124	275	2.2	65 52%	19 15%
CEZ	99	283	2.9	51 52%	5 5%
CWZ	159	404	2.6	91 57%	9 6%
SEZ	165	509	3.1	78 47%	10 6%
SWZ	169	520	3.1	87 51%	10 6%
Malawi	716	1,991	2.8	372 52%	53 7%

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

Table 1 provides a summary of the supervision outcomes by zone. Most facilities were using the standard national M&E tools. **136** sites had cumulatively registered more than 2,000 ART patient and **44** of these had registered more than 5,000. **51 (38%)** of these high burden sites were using electronic data system for ART (EDS). 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 Q2 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 Q2

Zone	Total fac.(1)	Facilities providing HIV services				CD4 count machines (2)			Results
		Exp. child	Pre-ART	PMTCT B+	ART	Installed	Functional	Results	
NZ	134	112 84%	112 84%	95 71%	124 93%	30 22%	28 93%	2,604	
CEZ	99	95 96%	95 96%	88 89%	99 100%	18 18%	15 83%	1,371	
CWZ	162	129 80%	132 81%	127 78%	159 98%	33 20%	25 76%	3,524	
SWZ	170	147 86%	165 97%	141 83%	166 98%	43 25%	35 81%	6,201	
SEZ	164	159 97%	158 96%	150 91%	163 99%	46 28%	39 85%	5,536	
Malawi	729	642 88%	662 91%	601 82%	711 98%	170 23%	142 84%	19,236	

(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.

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

Table 2 shows the distribution of the **729** sites designated to provide clinical HIV services in Q2 2015, by zone. At the national level, there were **711** (static) sites with at least one patient on ART, **601** sites had enrolled women under PMTCT Option B+; **662** sites were providing pre-ART services. **642** had enrolled HIV exposed children for follow-up. ART services were now available at almost all designated sites in the 5 zones.

The CWZ had reached 100% of designated sites with ART services. SEZ had reached 98% of designated sites with Option B+.

CD4 count machines (including 'point of care' machines) were installed at **170** sites, and **142** (84%) of these had produced at least 1 result during Q2 2015. The total number of CD4 results produced decreased from 20,977 in Q1 2015 to **19,236** during Q2 2015. 32% of these outputs were generated by 35 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 will now be started on ART after an initial CD4 test.

4.2 Staffing of HIV Services

4.2.1 HTC 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.

	2014 Q3	2014 Q4	2015 Q1	2015 Q2
Sites visited	720	718	715	716
Sites with any tests done	668 93%	660 92%	664 93%	672 94%
Sites with registered HTC staff	657 91%	659 92%	668 93%	669 93%
Total HTC staff at visited sites	3,583	3,574	3,682	3,820
Staff with any test done	2,276 64%	2,195 61%	2,242 61%	2,487 65%
Staff with 300+ tests done this quarter	322 13%	286 11%	268 11%	326 11%
Logbooks reviewed	2,554 71%	2,526 71%	2,549 69%	2,860 75%
HTC staff participating in PT this quarter	1,380 54%	170 7%	1,649 65%	931 33%
Total tests (HTC register)	550,436	464,292	479,916	493,567
Tests accounted for by individual staff	355,216 65%	331,981 72%	321,539 67%	379,744 77%
Source: logbooks	330,271 93%	303,046 91%	291,206 61%	358,627 94%
Source: HTC register	24,945 7%	28,935 9%	30,333 10%	21,117 6%
Total tests by staff with 300+ tests	154,494 43%	145,300 44%	134,272 42%	166,291 44%

3

669 (93%) of the 716 visited facilities had registered HTC providers and **672** (94%) sites had performed at least one test during Q2 2015. **2,860 (75%)** of **3,820** HTC providers had their logbooks available for review.

According to the 2,860 reviewed logbooks, **931 (33%)** HTC providers had participated in proficiency (panel) testing (PT) this quarter. This is lower than the participation rate from the previous quarters. Only 75% of providers at the visited sites had their logbooks available for review and documentation of PT may have been incomplete. The national HIV reference laboratory is organizing six monthly PT rounds for all practising HTC providers. PT panels are distributed by the district lab supervisors who may not be able to reach all sites in their districts.

³ HTC data in the published report for Q1 2015 included an error that resulted in duplication of outputs from 1 district (Thyolo). This explains the reduction of testing outputs for Q1 2015 shown in the table above.

379,744 (77%) of all 493,567 tests conducted this quarter (according to HTC register reports) were accounted for by individual HTC staff working at the visited sites. **358,627 (94%)** of these tests were documented in the reviewed logbooks and an additional **21,117 (6%)** could be attributed to individual providers from staff codes in the HTC registers. **326 (11%)** of 2,487 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 on 60 working days per quarter). The 326 HTC providers who met or exceeded this target produced **166,291 (44%)** 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 80 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 July 2015, **652** clinicians (physicians, clinical or medical officers); **892** nurses and **891** auxiliary staff (health surveillance assistants, clerks, etc.) were working in ART clinics in Malawi.

	2014 Q3		2014 Q4		2015 Q1		2015 Q2	
Clinicians	594	26%	617	26%	604	27%	652	27%
Nurses	830	37%	855	37%	814	36%	892	36%
Pharmacy staff	11	0%	11	0%	14	1%	12	0%
Auxiliary Staff	831	37%	847	36%	837	37%	891	36%
Total	2,266		2,330		2,269		2,447	

An estimated 2.7 million ART patient visits are currently managed at the 711 ART sites per annum, based on approximately 570,000 patients alive on ART and an average dispensing interval of 2.5 months. With 260 working days per year, an average of 10,500 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 **12** 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 21).

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 HTC data

493,567 people⁴ were tested and counselled for HIV between April and June 2015. **466,265 (94 %)** of these tests were performed at health facilities, **10,190 (2 %)** were done in stand-alone HTC sites and **17,112 (3%)** were done outside of facilities / in the community. Out of a total of **29,285** people newly diagnosed with HIV this quarter, **28,248 (96%)** were tested at health facilities, **594 (2%)** at stand-alone HTC sites and **443 (2%)** in a community-based testing.

5.2 HTC access type

271,719 (55%) of people tested were patients receiving provider-initiated testing and counselling (PITC); **219,989 (45%)** accessed voluntary counselling and testing, door-to-door, community-based testing, etc.; and **1,859 (<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 20,969 FRS issued to index clients this quarter, the successful referral rate for family members was **9%** (1,859 / 20,969). This is lower than in previous quarter (12%). Referral slips have remained under-utilized.

5.3 Age and sex distribution among HTC clients

Out of **493,567** people tested and counselled, **35%** were males and **65 %** were females. **49%** of females were pregnant. The proportion of males (52%) to non-pregnant females (48%) was similar, implying gender balanced access to HTC services. Pregnant women have to be excluded from the comparison of male and female access to HTC because testing among pregnant women is almost entirely provider-initiated and there is no comparable access route targeting males.

119,584 (24%) of all people tested accessed HTC with their partners (as a couple).

52% of all people tested and counselled were 25 years and above, **40 %** were between 15-24 years and **8%** were children below 15 years. **2,277 (<1%)** of rapid tests done were among infants.

5.4 First time, repeat and confirmatory test results

The 2011 and 2014 Malawi Clinical HIV Guidelines stipulate: *All patients 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.*

150,728 (31%) accessed HTC for the first time and **342,839 (69%)** were repeat testers. Based on the cumulative number of people who accessed HTC for the first time, a total of **5,873,428** people have been tested since introduction of the *first time HTC access* indicator in July 2007.

29,285 (6%) out of all clients received a positive result for the first time. Positive rapid test results among infants (**951**) and inconclusive test results (**1,826**) both accounted for **<1 %** of new results given to clients.

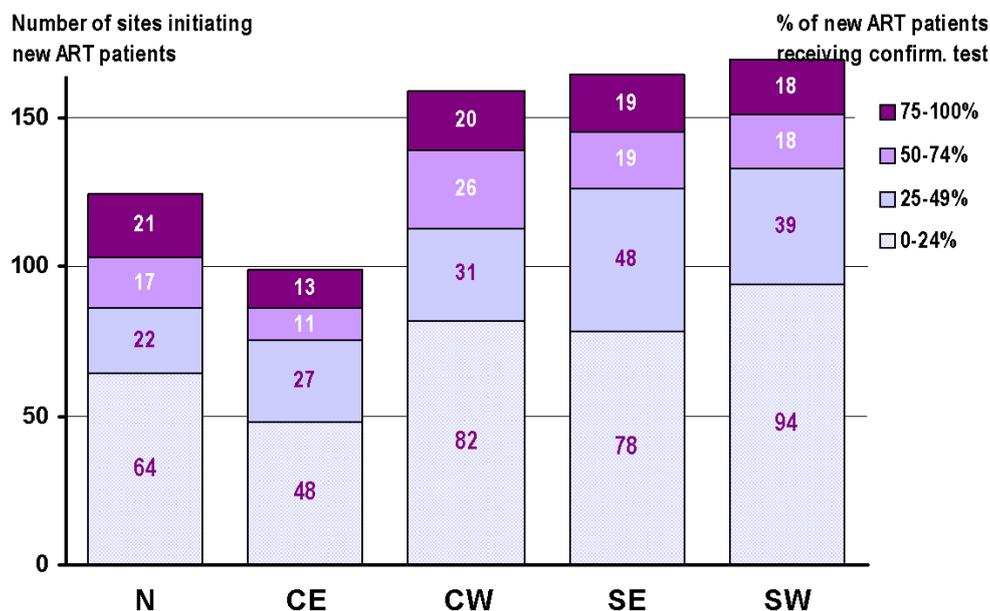
330,885 (97%) of 342,839 repeat testers reported a *last negative* result. **9,820 (3%)** were reported as *previous positives* and all of these should have been classified as receiving a confirmatory test. For most of the **9,820** *previous positives*, testing was probably initiated by a health worker before enrolment into care. *Confirmatory test results* accounted for **9,133 (93%)** of *previous positive* clients. The remainder (687) may have been misclassified as *new positive* or *new inconclusive* because they were among clients who independently sought confirmation of their positive status. **9,133 (96%)** of 9,517 confirmatory tests were

⁴ 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.

concordant positive and **384 (4%)** were classified as *confirmatory inconclusive*. This category includes parallel concordant negative and discordant test outcomes (Determine HIV1/2 and Uni-Gold HIV1/2 are used in parallel for confirmatory testing). This relatively high proportion of clients who did not have a concordant positive confirmation may be 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.

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



The 9,517 confirmatory test results documented this quarter indicate that only **37%** of the 25,957 patients initiating ART this quarter received confirmatory testing and **Figure 1** shows that confirmatory testing was low in all 5 zones. Only **91 (13%)** of facilities throughout the country had performed confirmatory testing for $\geq 75\%$ of patients newly initiated on ART. Implementation of the confirmatory testing policy will be further reinforced over the next quarters.

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.

514 (80%) of 642 sites with HIV exposed children in follow-up had collected and recorded at least 1 DNA-PCR sample during Q2 2015. A total of **7,992** DNA-PCR samples were collected and recorded. By the time the logbooks were reviewed (between 2 and 4 weeks after the end of the quarter), results had been received at the sites for **2,871 (36%)** of these specimens and **1,624 (57%)** of these results had been communicated to the mother. The proportion of results received at the sites was **63%, 36%** and **10%** for samples collected in April, May and June, respectively. A total of **107 (4%)** results received at the sites were positive.

The **8 laboratories** dispatched DNA-PCR test results for **6,013** children in Q2 2015. This is 1,979 (25%) fewer than the number of samples recorded in the DNA-PCR logbooks at health facilities during this quarter. This discrepancy is likely due to the difference in sample collection and result dispatch dates. **3,432 (57%)** of the dispatched results were from samples collected in Q2 2015, while 2,581 (43%) 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 20 and 43 days after sample collection.

3,256 (54%) of all results were from infants under 2 months old at the time of sample collection. 2,151 (36%) were 2-5 months, 410 (7%) were 6-11 months and 49 (1%) were 12 months or older when the sample was collected (date of birth was missing for 147).

Age at sample collection	Tot. Results	Positives	
<2 months	3,256	40	1.3%
2-5 months	2,151	80	3.7%
6-11 months	410	42	10.2%
12 months +	49	7	14.3%

181 (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	537	9%	6	3%
2-5 months	4,672	78%	104	57%
6-11 months	582	10%	50	28%
12 months +	76	1%	10	6%
(missing date)	146	2%	11	6%
Total	6,013	100%	181	100%

Out of 181 positive results dispatched, only 6 (3%) were sent before the child was 2 months old. A total of 110 (60%) positive results were sent before the child was 6 months old and 260 (88%)

were sent before the child was 12 months old. A total of 87 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 **33%** of the positive DNA-PCR results dispatched for children <12 months this quarter.

7 Blood Safety

The Malawi Blood Transfusion Service (MBTS) is striving to provide 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 **20,538** blood units were collected in Malawi during Q2 2015. MBTS collected **11,957 (58%)** of these, **100%** of which were screened comprehensively for the relevant TTIs (HIV, Hepatitis B, Hepatitis C, syphilis, malaria). In addition, **54** hospitals in Malawi collected a total of **8,581** units from replacement donors. **7,639 (89%)** of these units were screened for at least the 3 key TTIs (HIV, HepB and syphilis) and **3,827 (50%)** of these were also screened for HepC and malaria. This means that a total of **19,596 (95%)** of all 20,538 units collected by MBTS and from replacement donors 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, 12 units were screened for HIV and HepB only and 3 were screened only for HIV. 927 were screened with any other combination of tests for TTIs.

A total of **12,962** potential replacement donors were documented in the blood donor registers at the facilities and 8481 (66%) 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, 82% for HepB, 81% for syphilis, 61% for malaria and 35% 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 **1,408** persons received PEP during Q1 2015. This is an 11% increase from the previous quarter (1,271).

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 Q2.

Zone	Pre-ART		ART		Both patient groups	
	Tot. women	On Depo	Tot. women	On Depo	Tot. women	On Depo
NZ	550	146 26%	31,128	9,504 31%	31,677	9,650 30%
CEZ	384	117 30%	26,113	8,152 31%	26,497	8,269 31%
CWZ	3,022	737 24%	66,406	14,862 22%	69,428	15,598 22%
SEZ	2,838	993 35%	102,557	39,888 39%	105,395	40,881 39%
SWZ	4,803	1,122 23%	107,857	31,858 30%	112,660	32,980 29%
Malawi	11,597	3,113 27%	334,061	104,265 31%	345,657	107,378 31%

* 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 **107,378 (31%)** of 345,657 women in care received Depo-Provera from HIV clinics in Q2 2015. The SE Zone had achieved the highest coverage among women in pre-ART and ART. Patient coverage and stock availability has slightly decreased in this quarter. 549 (76%) of ART/PMTCT sites had stocks of Depo-Provera in July 2015 from 82% in April 2015.⁵ The HIV Program is no longer supplementing FP supplies through 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.

Table 4 shows that **634,307 (91%)** of 695,169 all patients in care were on CPT at the end of Q2 2015.

⁵ Many Mission hospitals do not provide family planning.
Malawi Integrated HIV Program Report (April - June 2015)

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 Q2.

Zone	CPT									IPT					
	Exp. child			Pre-ART			ART			All patient groups		Pre-ART			
	Tot. pat.	On CPT	%	Tot. pat.	On CPT	%	Tot. pat.	On CPT	%	Tot. pat.	On CPT	%	Tot. pat.	On IPT	%
NZ	8,060	6,155	76%	2,125	2,038	96%	55,015	53,935	98%	65,200	62,128	95%	2,125	1,912	90%
CEZ	7,861	6,291	80%	1,731	1,696	98%	45,495	44,330	97%	55,087	52,316	95%	1,731	1,610	93%
CWZ	16,035	14,169	88%	9,469	7,368	78%	114,953	106,931	93%	140,457	128,467	91%	9,469	5,889	62%
SEZ	30,417	26,638	88%	11,043	10,205	92%	165,108	152,716	92%	206,568	189,558	92%	11,043	9,700	88%
SWZ	28,163	25,113	89%	15,361	14,477	94%	184,333	162,248	88%	227,857	201,838	89%	15,361	12,086	79%
Malawi	90,536	78,365	87%	39,729	35,783	90%	564,904	520,159	92%	695,169	634,307	91%	39,729	31,197	79%

11 TB / HIV Interventions

11.1 Intensified 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.

552,687 (98%) of all patients retained on ART were screened for TB at their last visit before end of June 2015. As of that visit, **6,160 (1%)** patients were new TB suspects and had presumably been referred for examination by a clinician and for TB investigations. **904 (<1%)** patients had confirmed TB (clinical or lab based). Out of these, **815 (90%)** were confirmed to be on TB treatment and **89 (10%)** 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 not done (Current TB status unknown/ not circ)	12,217	2%
ICF done	552,687	98%
TB not suspected	545,623	99%
TB suspected	6,160	1%
TB confirmed	904	0%
TB confirmed, not on treatment	89	10%
TB confirmed, on TB treatment	815	90%

11.2 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. **31,197 (79%)** of 39,729 patients retained in pre-ART were on IPT by the end of June 2015. Isoniazid was in stock at 621 facilities during the July 2015 supervision visit.

12 HIV-Related Diseases

Table 5 shows the number of patients treated for key HIV-related indicator diseases. **4,287** patients were started on TB treatment this quarter and HIV status was ascertained for **4,073 (95%)**. **2,200 (54%)** of these were HIV positive and **1,513 (69%)** 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 Q2 2015, **459** and **595** patients received Diflucan for acute cryptococcal meningitis and oesophageal candidiasis, respectively. **265** 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).

	TB				KS *	CM *	OC *
	Tot. cases	HIV status asc.	HIV positive	Already on ART	Tot. cases	Tot. cases	Tot. cases
2014 Q3	4,692	4,368 93%	2,149 49%	1,460 68%	310	368	640
2014 Q4	4,204	3,989 95%	2,103 53%	1,546 74%	336	475	663
2015 Q1	4,158	3,765 91%	1,954 52%	1,408 72%	260	865	601
2015 Q2	4,287	4,073 95%	2,200 54%	1,513 69%	265	459	595

13 HIV-Exposed Child Follow-Up

13.1 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.

13.2 HIV Exposed Child Registration Data

10,396 HIV exposed children were newly enrolled into follow-up during Q2 2015; **6,563 (64%)** of these were under the age of 2 months. This represents timely enrolment for **79%** of the 8,210 known HIV exposed children discharged from maternity this quarter. The total number of new enrolments (10,230) exceeds by 2,020 (22%) the total number of known HIV exposed children discharged from maternity (8,210). 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.

13.3 Birth Cohort Outcomes

There were **8,509** infants in the **2 month age cohort**. **2,524 (30%)** had received a DNA-PCR result. **57 (2%)** of these were confirmed HIV infected. An additional **25** infants were diagnosed with *presumed severe HIV disease*, which means that a total of **82** infants were eligible for ART. **47 (57%)** of these had started ART. The proportion of positives starting ART is slightly lower than the previous quarter (60%). Out of the entire 2-month age cohort, **7,806 (93%)** were retained in exposed child follow-up, **47 (1%)** had started ART and **16 (<1%)** were discharged confirmed uninfected⁶. **26 (<1%)** were known to have died and **494 (6%)** had been lost to follow-up.

There were **8,417** children in the **12 month age cohort**. Current HIV infection status was known for **3,594 (43%)** children (DNA-PCR or rapid antibody test) and **165 (5%)** of these were confirmed HIV infected. **24 (<1%)** additional children had been diagnosed with *presumed severe HIV disease*, which means that a total of **189** children were eligible for ART. **160 (85%)** had started ART. Out of the entire age cohort, **6,238 (76%)** were retained in exposed child follow-up, **160 (2%)** had started ART and **61 (<1%)** were discharged confirmed uninfected.⁶ **1,638 (20%)** were lost to follow-up and **71 (<1%)** were known to have died.

There were **8,121** children in the **24 month age cohort**. Current HIV infection status was known for **3,898 (48%)** children (DNA-PCR or rapid antibody test) and **225 (6%)** of these were confirmed HIV infected. **30** additional children had been diagnosed with *presumed severe HIV disease*, which means that a total of **255** children were eligible for ART. **224 (<100%)** of these had started ART. Out of the entire age cohort, **952 (12%)** were retained in exposed child follow-up, **224 (3%)** had started ART and **3,457 (44%)** were discharged confirmed uninfected. **3,061 (39%)** were lost to follow-up and **111 (1%)** were known to have died.

Confirmed HIV-free survival at age 24 months in this quarter remained implausibly low at **44%**. This was related to the fact that only 48% in this cohort had a known HIV status. 4,223 (52%) children were classified as '*current HIV infection status unknown*' and many of these may be among the 3,061 children lost to follow-up and the 111 children who had died. However, 952 (12%) 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.

⁶ 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.

14 Pre-ART

14.1 Pre-ART Registration Data

A total of **6,077** patients were newly registered for pre-ART follow-up in Q2 2015. **472 (4%)** of these were children aged 5-14 years. The number of new pre-ART enrolments continued to decline from the previous quarter (6,223 total, 461 children) due to the introduction of relaxed ART eligibility criteria in April 2014. Several sites had already established pre-ART services before July 2011 and the cumulative number of pre-ART patients ever registered was **193,097**.

14.2 Cumulative Pre-ART Follow-up Outcomes

39,729 (22%) of all patients ever registered were retained in pre-ART follow-up by the end of June 2015; **97,886 (53%)** had started ART; **44,136 (24%)** had been lost to follow-up; **1,549 (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, **3,036** pre-ART patients started ART during Q2 2015. The cumulative number of patients reported as lost to follow-up and died was lower than in the previous quarter, indicating challenges with completeness and accuracy of reporting.

CPT coverage among pre-ART patients was **90%** in Q2 2015 and IPT coverage declined slightly to **79%**. **3,113 (27%)** of 11,597 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.

15 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.

15.1 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 starting ANC in the reporting period and the final HIV and ART status of women who had completed 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**) *plus* 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: 2014 Spectrum model for Malawi). There are an estimated 13,317 HIV infected pregnant women in the population per quarter (1/4 of 53,268 in 2014).⁷

⁷ 2015 Spectrum estimates based on current definition of eligibility for ART in Malawi (CD4<500, Option B+, UT for U5).
Malawi Integrated HIV Program Report (April - June 2015)

15.2 ARV Coverage among Pregnant / Breastfeeding Women and Exposed Infants

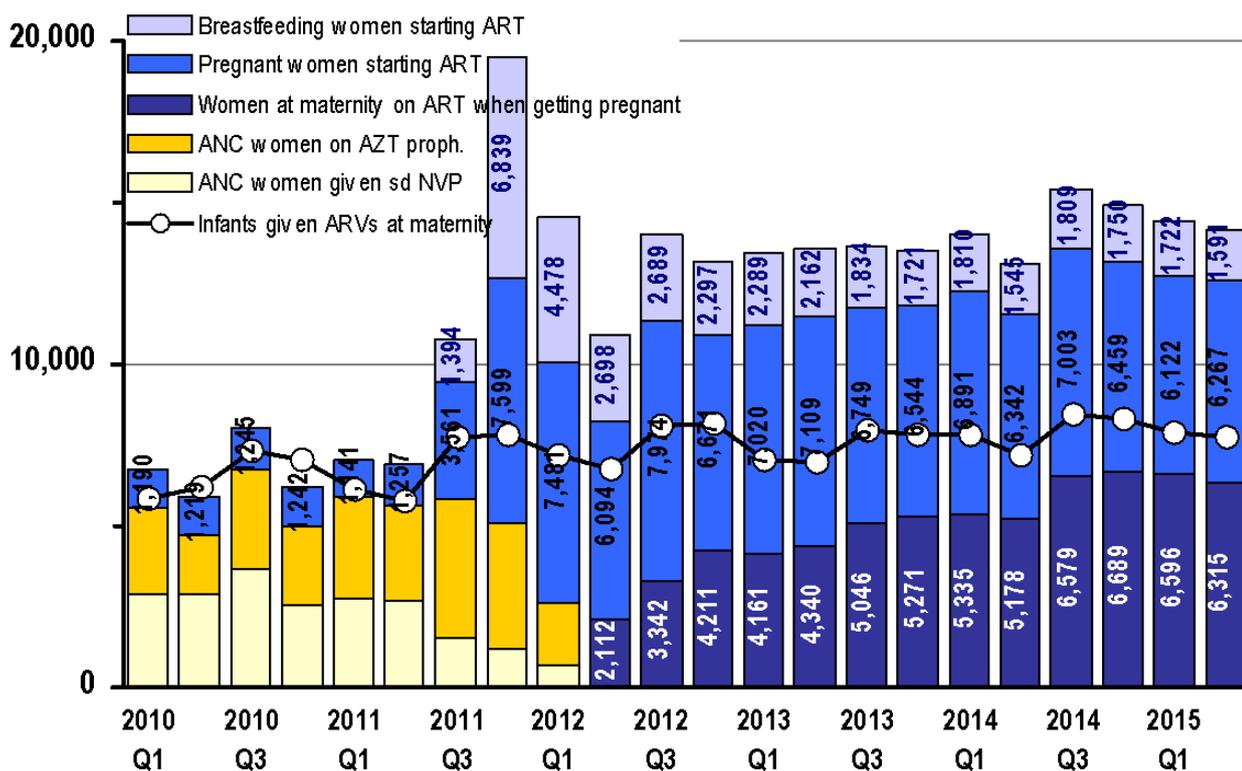
10,460 (70%) of the estimated 14,926 HIV infected pregnant women in Malawi this quarter were on ART. This is based on **5,960**⁸ women at maternity who were already on ART when getting pregnant and **4,500**⁹ women who newly initiated ART in pregnancy. The apparent decline in ART coverage from the previous quarters (85%; 85%) is mainly due to an increase in the estimated number of HIV infected pregnant women in the population in the 2015 Spectrum model.

An additional **1,591**¹⁰ 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 **5,901**. 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. **7,866** 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.



⁸ 6,315 women who started ART before pregnancy admitted at maternity; reduced by 6% to adjust for double-counting of 7,083 referrals among 125,961 total admissions.

⁹ 6,267 women registered at ART clinics who were pregnant at the time of starting ART; a) 8% are discounted to adjust for double-counting of transfers based on 657 of 7,829 women who transferred within 12 months of registration (12 month Option B+ survival analysis); b) 21.6% are discounted to account for presumed failed ART initiations based on 1,788 of 8,275 women lost to follow-up within 6 months of registration (6 month Option B+ survival analysis).

¹⁰ 1,722 women registered at ART clinics who were breastfeeding at the time of starting ART; reduced by 8% to adjust for double-counting of transfers based on 664 out of 8,392 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.

15.3 HIV Services at ANC

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

15.3.1 HIV Ascertainment and ART Coverage

Booking cohort:

144,819 women attended ANC for their first visit between April and June 2015. This is 87 % of the estimated 166,000 pregnant women in the 2015 population during one quarter.¹¹ **123,898 (86%)** of women in the booking cohort had their HIV status ascertained at the first visit. Out of these, **9,879 (8%)** presented with a valid previous test result and **114,019 (92%)** received a new test. A total of **9,879 (8%)** of women were found HIV positive: **5,337 (54%)** of these from a documented previous test and **4,542 (46%)** from a new test. **8,920 (90%)** of all positives were on ART: **4,727 (53%)** of these were already on ART; **3,457 (39%)** started during the 1st or 2nd trimester and **736 (8%)** started in the 3rd trimester of pregnancy.

Outcome cohort:

155,272 women had started ANC between October and December 2014 and their outcomes were reported between April and June 2015. Only **33,938 (22%)** of women in this cohort attended the minimum of 4 focussed ANC visits.

141,212 (91%) of these women had their HIV status ascertained at least once in the course of ANC. This is similar to the previous quarter (91%). In this cohort, **11,914 (8%)** presented with a valid documented previous HIV test result and **129,298 (92%)** received a new HIV test result at ANC. A total of **10,766 (7.6%)** women were found HIV positive. This is slightly lower than the latest Spectrum projections (9.0% HIV prevalence among pregnant women in 2015).⁷

9,740 (90%) of (known) HIV infected women were on ART by the end of ANC. This represents **65%** coverage of the estimated 14,926 HIV positive pregnant women per quarter at the population level. Of the **9,740** ANC women who were known to receive ART, **4,677 (48%)** were already on ART when starting ANC, **4,000 (41%)** initiated before 28 weeks of pregnancy and **1,063 (11%)** initiated during the last trimester of pregnancy. **9,736 (90%)** of HIV infected women at ANC were on Cotrimoxazole Preventive Therapy. **8,908 (83%)** of known HIV infected women attending ANC received the infant dose of ARVs (nevirapine syrup) to take home.

15.3.2 Syphilis Screening

13,259 (9%) of women in the outcome cohort were tested for syphilis and **449 (3%)** were syphilis positive. The low testing rate probably explains the higher (3%) than expected proportion (<1%) of positives as the testing was likely selective of those suspected to be positive.

15.4 HIV Services at Maternity

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

Between April and June 2015, **118,878** women were admitted for delivery to maternity; **7,083** of these were referred to another facility before delivery, resulting in **125,961** total admissions to maternity during Q2 2015. Out of all admissions, **116,472 (96%)** delivered at health facilities, while **4,898 (4%)** had already delivered before reaching a facility. The **116,472** facility deliveries represent **70%** of the estimated 166,000 quarterly deliveries in the population in 2015 which is less than the 83% reported in the Integrated Household Survey Report of 2010-2011.

¹¹ 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

Malawi Integrated HIV Program Report (April - June 2015)

A total of **113,650 (96%)** deliveries were conducted by skilled birth attendants, **575 (<1%)** by paramedical staff and **4,619 (4%)** were not attended by any of the above (probably mainly among women who delivered before reaching maternity). **15,588 (12%)** of women developed obstetric complications. The most common leading complications were obstructed / prolonged labour (**5,270** cases) and post-partum haemorrhage (**1,789** cases). A total of **121,370** babies were born, **116,226 (97%)** were singletons and **4,234 (3%)** were twins/multiples. There were **119,374 (98%)** live births and **1,996 (2%)** stillbirths. **118,182 (99%)** of babies born alive were discharged alive and **1,192 (1%)** died before discharge. **118,750 (>99%)** of women were discharged alive and **94 (<1%)** women died before discharge, which is equivalent to a maternal mortality ratio of **79 per 100,000** live births among women attending maternity.

15.4.1 HIV Ascertainment at Maternity

120,781 (96%) women had their HIV status ascertained at maternity. Out of these, **118,878 (94%)** presented with a valid previous HIV test result and **3,189 (2%)** received a new HIV test result. A total of **8,967 (7%)** women were HIV positive and **111,814 (93%)** were negative. The **120,781** women whose HIV status was ascertained at maternity represent **73%** of the expected 166,000 women delivering in the population.

HIV exposure status was ascertained for **114,093 (97%)** out of 118,182 babies born and discharged alive. **8,210 (7%)** of these were born to a known HIV positive mother.

15.4.2 ARV Coverage at Maternity

A total of **8,779 (98%)** of known HIV infected women admitted to maternity received ART. Out of these, **6,315 (72%)** had started ART before pregnancy, **1,247 (14%)** initiated ART during the 1st or 2nd trimester, **961 (11%)** initiated during the 3rd trimester and **256 (3%)** initiated ART at maternity.

A total of **7,771 (95%)** of 8,210 infants who were known HIV exposed and discharged alive started daily NVP prophylaxis at maternity. This represents **52%** coverage of the estimated 14,926 HIV exposed infants born in the population in this quarter.

16 ART Access and Follow-Up Outcomes

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

16.1 New ART Registrations during Q2 2015

By the end of June 2015, there were **711 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.

The electronic reporting system supported by MSF in Chiradzulu district experienced a prolonged technical fault and accurate ART cohort reports for the 10 affected facilities could not be obtained in time for this report. It was therefore decided to use the previous quarter's results for these 10 sites. This has led to some inconsistencies in the data presented.

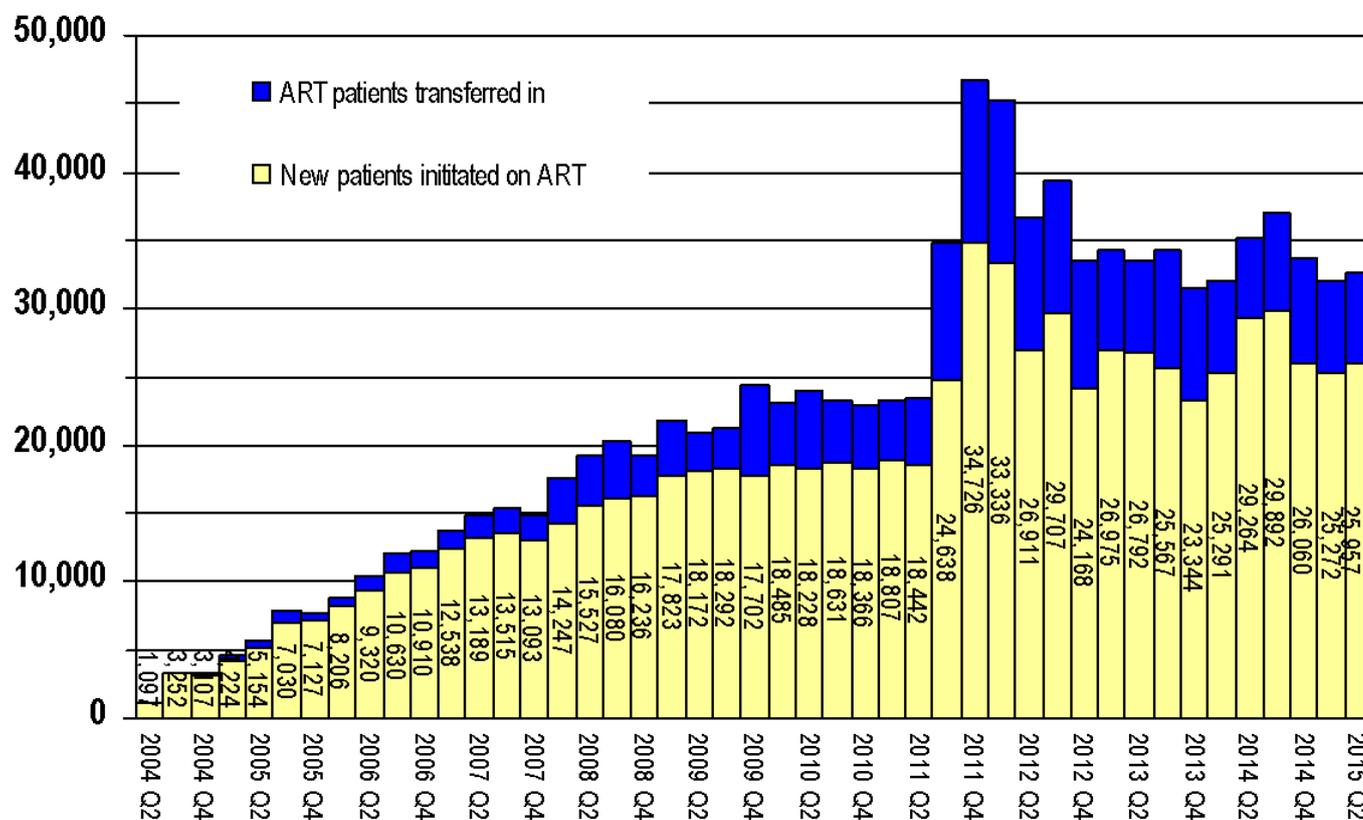
Implementation of the Malawi Integrated Clinical HIV Guidelines started in July 2011, triggering a massive surge in new ART initiations (see **Figure 3**). **25,957** patients initiated ART in Q2 2015 and **6,259** patients were registered as a transfer in (already on treatment; 19% out of all 32,637 clinic registrations). These numbers are similar to previous quarter.

Among all new registrations **36%** were males, **64%** were females. **6,267 (30%)** of females were pregnant. **6,164 (98%)** of pregnant women were started under **Option B+** (in WHO stage 1 or 2 with unknown CD4 or CD4 above 500), while 103 were in more advanced stage of HIV infection. An additional **1,591** women in

WHO stage 1 or 2 were started because of breastfeeding, bringing the total number of women registered as started under **Option B+**¹² to **7,755**.

Figure 3: Patients newly initiated 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 **20,876 (64%)** of all patients registered started in WHO stage 1 or 2 and **12,391 (59%)** of these started due a low CD4 count. **9,895 (30%)** of patients registered started in WHO stage 3 and **1,438 (4%)** started in stage 4.

2,773 children were registered at ART sites in Q2 2015. **635** 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. **113** children started ART with presumed severe HIV disease, is similar to the previous quarter (109). **87** infants in WHO stage 1 or 2 who started due to confirmed HIV infection through DNA-PCR, which is lower than the previous quarter (116). 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 8,210 HIV exposed infants were identified at maternity and assuming a 2% transmission rate among the 98% of HIV positive mothers at maternity who received ART (and 20% transmission in the 2% who did not receive ART)¹³, only about 213 of these known HIV exposed infants may have been infected perinatally during Q2 2015. However, considering the projected 1,560 new infant HIV infections in the 2015 population per quarter⁷, early infant treatment coverage remains low at an estimated **6% (87 / 1,560)**. The most significant bottleneck for early infant treatment remains the identification of HIV infected pregnant / breastfeeding women.

931 (3%) out of all ART clinic registrations were patients with TB: **589 (2%)** had a current and **342 (1%)** a recent history of TB. **265 (1%)** of patients registered had Kaposi's sarcoma.

¹² Universal ART for all HIV infected pregnant and breastfeeding women in WHO stage 1 or 2, independent of CD4 count

¹³ UNAIDS Reference Group on Estimates Modelling and Projections (2011). Working paper on mother-to-child-transmission rates for use in Spectrum. Geneva, UNAIDS.

16.2 Cumulative ART Registrations up June 2015

By the end of June 2015, there were a cumulative total of **1,025,754** clinic registrations, representing **820,367 (80%)** patients who newly initiated ART and **193,867 (19%)** patients who transferred between clinics. **11,520 (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 **30,901 (3.0%)** of total patient registrations.

16.3 ART Outcomes

568,470 patients were alive on ART by the end of June 2015. This is equivalent to **57% ART coverage** among the estimated 1 million HIV positive population in Malawi in 2015. The number of patients on ART includes an estimated 3,566 patients in transit between sites (50% of the 7,122 patients newly registered as transferred out at sites across the country).

Out of the **820,367** patients ever initiated on ART, **568,470 (69%)** were retained alive on ART, **77,342 (9%)** were known to have died, **186,074 (23%)** were lost to follow-up and **3,366 (<1%)** were known to have stopped ART. An estimated **519,063** adults and **49,407** children (<15 years) were alive on ART by the end of June 2015.

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

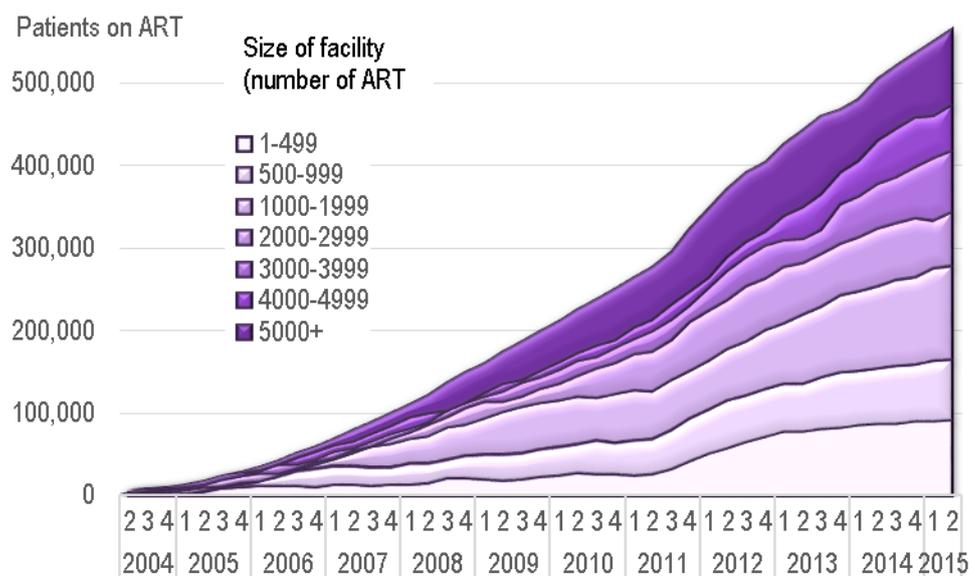


Figure 4 shows the increase of patients alive on ART by the end of each quarter. The number of patients alive on ART **increased by 16,904** in Q2 of 2015. **Figure 4** also illustrates the ongoing decentralization of Malawi's ART program. From Q3 2011, the greatest increase in ART patient numbers was seen at sites with fewer than 500 patients alive on ART. By the end of June 2015, **49%** 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)

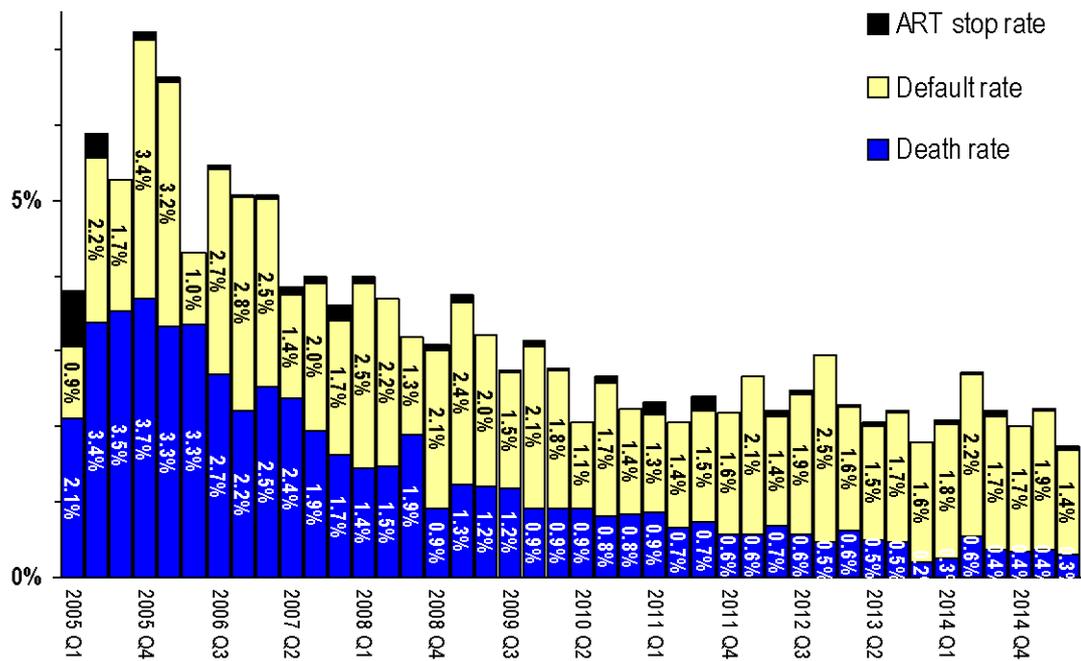


Figure 5 shows the considerable decrease of ART drop-out rates since the start of the national program. There were **1,806** new deaths, **8,002** new defaulters and **257** new stops in Q2 2015. This translates into a quarterly death rate of **0.3%** and a defaulter rate of **1.4%** among the patients 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 June 2015, a cumulative **77,342 (9%)** patients were known to have died **186,074 (22%)** were lost to follow-up and **3,366 (<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)

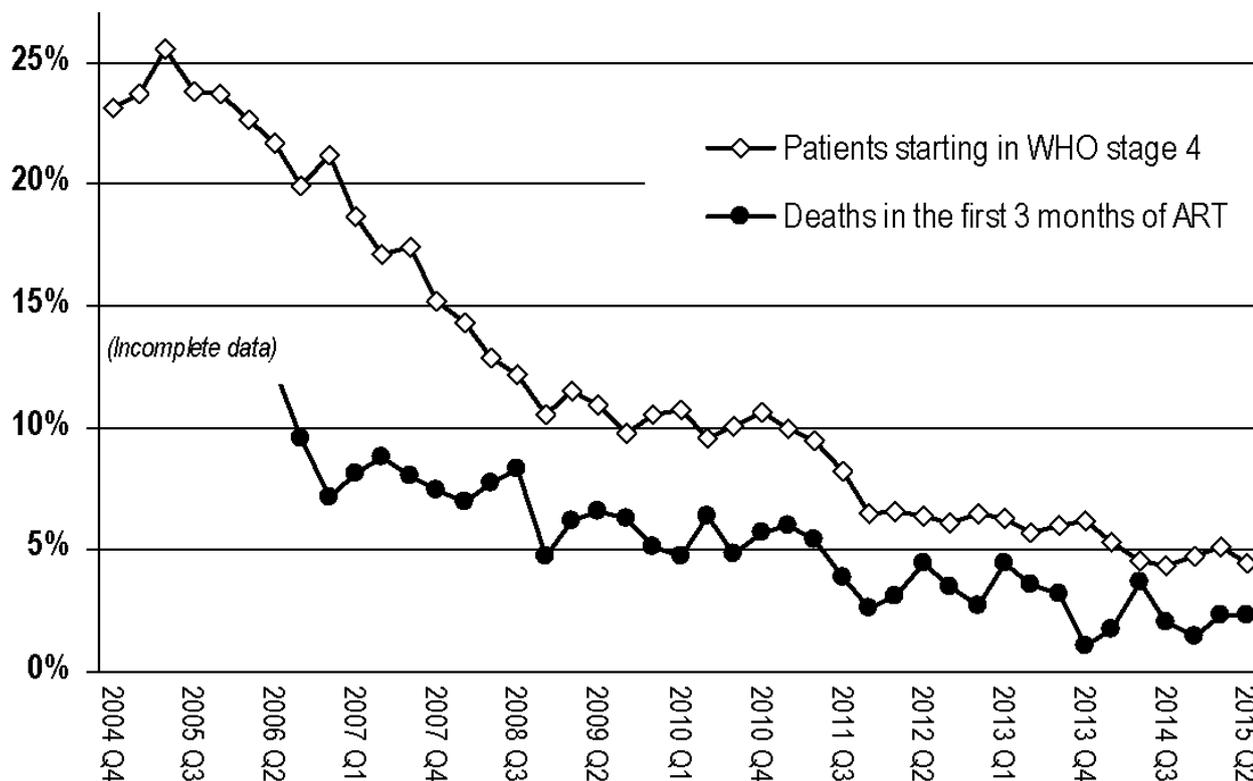


Figure 6 shows the considerable decline in **early mortality** since the start of the program. In Q2 of 2006 11% of new patients died within the first 3 months after ART initiation. There has been a remarkable decline in early mortality and the lowest point thus far has been reached in Q4 2013. The decrease 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 4% in Q2 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).

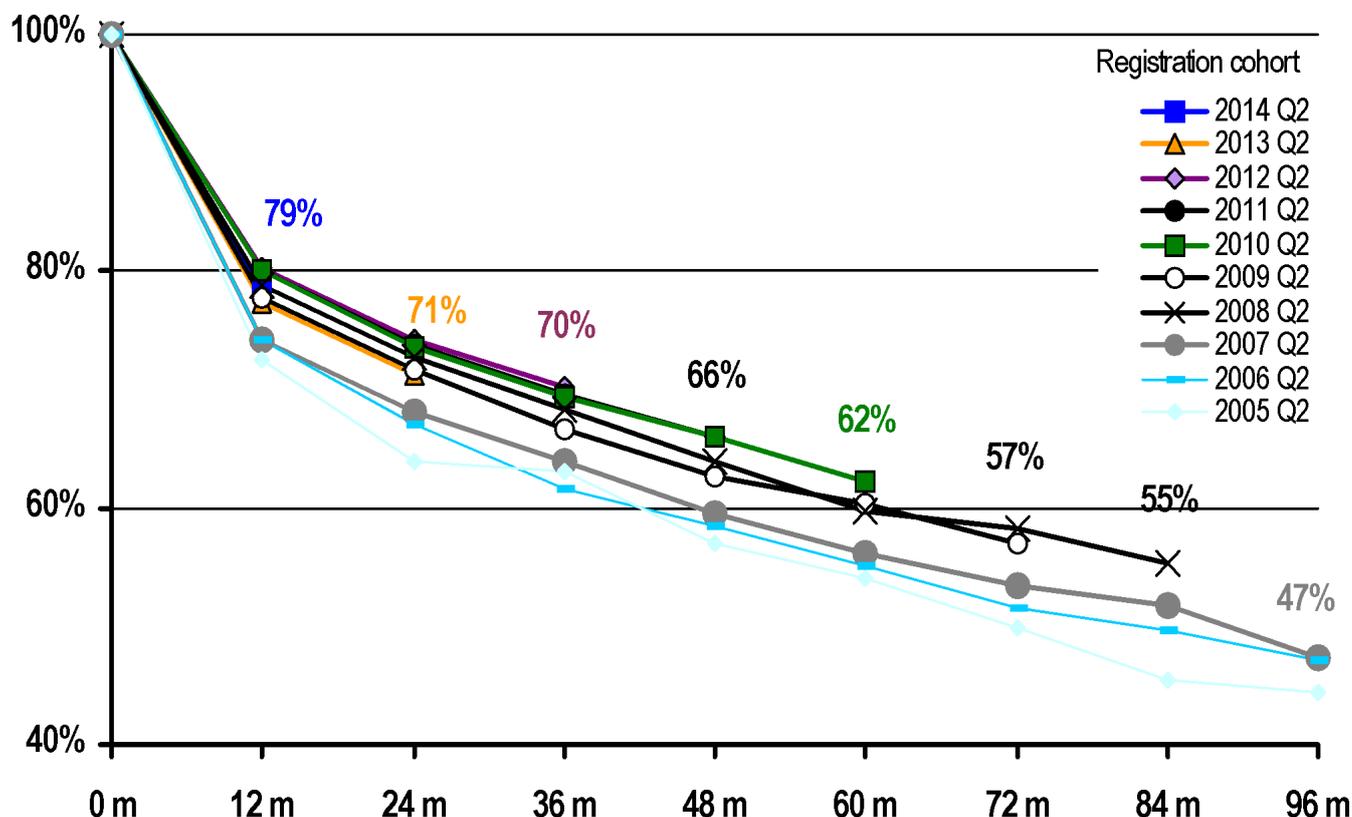
16.4 ART Cohort Survival Analysis

A 12, 24, 36, 48, 60, 72, 84, 96 and 108-month **'cohort outcome survival analysis'** was conducted for patients registered in Q2 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 Q2 2014. For the 12th time, a further subgroup analysis was done for women who started ART under **Option B+** during Q2 2013, Q1 2015 and Q4 2014. **79% of adults and 78% 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. However, actual retention rates are thought to be about **10%** higher due to 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.¹⁴

Figure 7 shows the continuous improvement of long-term treatment outcomes over time. **58%** and **52%** 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 **8,275 (99%)** of the 8,362 women registered as having started ART under *Option B+* in Q4 2014.¹⁵ This number represents **588 (7%)** women who transferred out and are therefore double counted and **7,687 (93%)** patients not transferred. **5,845 (76%)** of these were retained at 6 months after registration. **1,788 (97%)** of those not retained were lost to follow-up, **27 (<2%)** were known to have stopped ART and **27 (<2%)** were known to have died.

12-month group cohort survival outcomes were known for **7,829 (<100%)** of the 7,855 women registered as having started ART under *Option B+* in Q2 2014.¹⁵ This number represents **657 (8%)** women who transferred out and are therefore double counted and **7,172 (92%)** patients not transferred. **5,168 (72%)** of these were retained at 12 months after registration. **1,925 (96%)** of those not retained were lost to follow-up, **26 (1%)** were known to have stopped ART and **53 (4%)** were known to have died.

24-month group cohort survival outcomes were known for **8,600 (95%)** out of the 9,072 women registered as having started ART under *Option B+* in Q2 2013.¹⁵ This number represents **845 (10%)** women who transferred out and are therefore double counted and **7,755 (90%)** patients not transferred. **5,225 (67%)** of

¹⁴ 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.0000000000000252

¹⁵ 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. *Malawi Integrated HIV Program Report (April - June 2015)*

these were retained at 24 months after registration. **2,411 (95%)** of those not retained were lost to follow-up, **37 (>1%)** were known to have stopped ART and **82 (3%)** were known to have died.

2,162 (24%) of the women in the 24 month Option B+ survival cohort had initiated ART in the breastfeeding period and **2,129 (23%)** 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 **67% and 69% 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 the same as in 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 clinic registrations	8,275	100%
Transfers out (double counted)	588	7%
Total not transferred out (patients in cohort)	7,687	93%
Total alive on ART	5,845	76%
Total not retained	1,842	24%
Defaulted	1,788	97%
Stopped ART	27	1%
Died	27	1%

12 month survival OptionB+

Survival and retention in ART program

*

ART cohort registration group outcomes

Total ART clinic registrations	7,829	100%
Transfers out (double counted)	657	8%
Total not transferred out (patients in cohort)	7,172	92%
Total alive on ART	5,168	72%
Total not retained	2,004	28%
Defaulted	1,925	96%
Stopped ART	26	1%
Died	53	3%

24 month survival OptionB+

Survival and retention in ART program

*

ART cohort registration group outcomes

Total ART clinic registrations	8,600	100%
Transfers out (double counted)	845	10%
Total not transferred out (patients in cohort)	7,755	90%
Total alive on ART	5,225	67%
Total not retained	2,530	33%
Defaulted	2,411	95%
Stopped ART	37	1%
Died	82	3%

36 month survival OptionB+

Survival and retention in ART program

*

ART cohort registration group outcomes

Total ART clinic registrations	9,269	100%
Transfers out (double counted)	1,334	14%
Total not transferred out (patients in cohort)	7,935	86%
Total alive on ART	5,462	69%
Total not retained	2,473	31%
Defaulted	2,292	93%
Stopped ART	40	2%
Died	141	6%

16.4.1 Secondary outcomes of patients retained on ART

Secondary outcomes are known for the **564,904** patients alive on ART who remained at their sites at end of the quarter.

ART Regimens

557,673 (99%) of patients were on first line and **6,758 (1%)** were on second line regimens; **473 (<1%)** 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, **26,567 (5%)** were on paediatric formulations and **25,401 (96%)** of these were on the new standard first line for children (regimen 2P: AZT/3TC/NVP). By the end of June 2015, **493,034 (93%)** of patients on adult first line were receiving regimen **5A** (tenofovir / lamivudine / efavirenz). **28,298 (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 **1,158 (<1%)** were on regimen 1A (stavudine / lamivudine / nevirapine).

Adherence to ART

Pill counts and the number of missed doses were documented for **560,794 (99%)** out of all patients retained on ART and **510,901 (91%)** of these were classified as >95% adherent in Q2 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

410,760 (73%) patients on ART had information on drug side effects documented at their last clinic visit before end of June 2015. This decrease from the previous quarter (93%) was due to a software update for the national EMR that interfered with the side effect recording. **9,666 (2%)** 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.

16.5 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 Q2 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 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 **16,267** VL results were produced between April and June 2015. **4,810 (30%)** of samples processed were plasma and **11,455 (70%)** were DBS.

Lab	Samples Processed				Turn-around Time (Days) [§]
	Plasma	DBS	Oth/unk	Total	
DREAM Blantyre	1,248	866	0	2,114	48
DREAM Balaka	1,023	0	0	1,023	56
Kamuzu CH	1,223	513	0	1,736	31
Mzimba DH	0	97	0	97	7
Mzuzu CH	0	2,113	0	2,113	39
Partners in Hope	871	3,431	0	4,302	32
QUECH	8	829	0	837	92
Thyolo DH	437	2,367	1	2,805	13
Zomba CH	0	1,239	1	1,240	52
Total	4,810	11,455	2	16,267	36

§ Median days between sample collection and printing of results in the lab

Partners in Hope lab (Lilongwe) achieved the highest outputs, contributing 24% of all results this quarter. The median interval between sample collection and printing of results was **36 days** at the national level, ranging from **7 days** at Mzimba DH to **92 days** at Kamuzu CH. The most significant delays occurred between sample receipt and processing in the lab (median 18 days), while on average only 7 days elapsed between sample collection and receipt in the lab.

Reason	0-999		1000-4999		5000+		Total
Routine	13,560	85%	670	4%	1,578	10%	15,808
Targeted	151	65%	13	6%	68	29%	232
Other/unk	174	77%	8	4%	45	20%	227
Total	13,885	85%	691	4%	1,691	10%	16,267

15,808 (97%) of all VL samples were classified as *routine scheduled*. This is equivalent to **23%** of the estimated 70,000 ART patients passing a VL monitoring milestone this quarter. **232 (1%)** of samples were classified as *targeted (suspected treatment failure / repeat)* and for **227 (1%)** the reason for the sample was other or not specified. **13,885 (85%)** of all results were undetectable / below 1,000 copies/ml. The proportion of results with 5,000+ copies was higher among targeted samples (29%) than among *routine* samples (10%).

The time on ART was entered for only **3,780 (24%)** of 15,808 routine samples registered on the LIMS and only **675 (18%)** 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 **88%, 88%, 84%, 88%, 82%** and **80%** 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 (**83%**) or those with unknown timing (**84%**).

Patient age was recorded for 14,461 (91%) of routine monitoring samples. Among these, 6%, 7%, 12%, 34% and 41% 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: **76%**) and adolescents (10-19 yrs: **78%**) compared with adults (**84%, 84%** and **85%** for the age groups 20-29, 30-39, 40+ years, respectively).

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

Given the relatively low access to VL monitoring (estimated 23% of all patients on ART), the measured **85% 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 over-represent patients with poor adherence and/or treatment failure. Conservatively, the national viral suppression rate can be estimated as **483,199 (85%)** of 568,470 patients on ART, which is equivalent to **48%** of the total 1 million HIV infected population.

17 TB / HIV Management

Approximately **82%** of HIV infected TB patients were receiving ART in Q2 2015. This estimate is based on the following triangulation of TB and ART program data:

TB Program Data: A total of **4,288** TB patients were registered during Q2 2015. Assuming an average HIV prevalence of 60% among TB patients, **2,573** TB patients were HIV positive and therefore in need of ART. Given that **1,513** TB patients registered were already on ART at the time of starting TB treatment, $2,573 - 1,513 = 1,060$ TB patients needed to initiate ART.

ART Program Data: An estimated **589** patients¹⁶ started ART with a current or recent episode of TB in Q2 2015. This is **56%** (589 of 1060) of the TB patients who needed to start ART. This means that a total of $1,513 + 589 = 2,102$ (**82%**) of the estimated 2,573 HIV infected TB patients were receiving ART in Q2 2015.

TB program report

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TB clinic registrations

Total TB patients registered	4,288	100%
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HIV status ascertainment

HIV status not ascertained	214	5%
HIV status ascertained	4,074	95%
HIV negative	1,874	46%
HIV positive	2,200	54%
Already on ART	1,513	69%
Not on ART when starting TB treatment	687	31%

TB / ART program triangulation

*

HIV-burden among TB patients (estimated)

HIV negative (est. 40%)	1,715	40%
HIV positive (est. 60%) in need of ART	2,573	60%
Not on ART	319	12%
Total on ART (coverage)	2,253	88%
Already on ART (TB prog)	1,513	67%
Started ART within 24m of TB diagnosis (ART prog)	740	33%
ART initiations with current TB (ART prog)	468	63%
ART initiations after recent TB (ART prog)	272	37%

18 STI Treatment

STI reports were actively collected during the Integrated HIV Program Supervision exercise for the 9th 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 640 out of 928 facilities offering STI management according to the *2013-14 Service Provision Assessment*¹⁷ in Malawi. The site-level reports included here may therefore only represent 69% 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.

¹⁶ 21% of the 1,012 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.

¹⁷ 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>
Malawi Integrated HIV Program Report (April - June 2015)

18.1 Access to STI treatment and coverage

Based on the data collected at the facilities, a total of **54,937** STI cases were treated in Q2 2015. Considering the 69% site-level completeness of reporting, this number is estimated to represent a total of **79,619** STI cases treated. This is equivalent to **81% STI treatment coverage** of the expected 98,600 STI cases in the population.

Out of **54,937** documented clients treated, **22,377** (41%) were male and **32,560** (59%) were female. **3,980** (12%) of female STI clients were pregnant. **36,776** clients (67%) were 25 years and above, **13,151** (24%) were 20-24 years and **5,010** (9%) were under 20 years old.

18.2 Client Type and STI History

48,855 (89%) of clients were symptomatic and **6,082** (11%) were asymptomatic (treated as partners). Among symptomatic clients, **44,700** (91%) of were index cases and **4,155** (9%) were partners. A total of **18,380** partner notification slips were issued, equivalent to an average of 0.41 slips per index case. Considering the 18,380 partner notification slips issued, **56%** (10,237) of those notified presented to the clinic. **40,891** (74%) of clients presented with their first lifetime episode of STI, **9,656** (18%) clients reported to have had an STI more than 3 months ago and **4,390** (8%) 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.

18.3 HIV Status

HIV status was ascertained for **28,563** (52%) clients and **6,953** (24%) of these were HIV positive. **1,547** (22%) of positives were identified through a new test initiated at the STI clinic, while **5,406** (78%) presented with a documented previous positive HIV test result. **4,260** (79%) of clients with a previous positive HIV test result were on ART.

The rate of HIV status ascertainment at STI clinics remained low. This is likely due to poor implementation of provider initiated testing and counselling, combined with weak back-referral systems which may lead to incomplete documentation of new HIV test results at the STI clinics. 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.

18.4 STI Syndromes

The most common syndrome was abnormal vaginal discharge (AVD) with **17,272** (30%) cases, followed by urethral discharge (UD, **13,256** cases), genital ulcers (GUD, **10,406** cases) and lower abdominal pain (LAP, **8,564** cases). Similar to previous reports, balanitis, bubo, warts and neonatal conjunctivitis each accounted for 1 – 2% of cases.

18.5 Referrals

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. **19,767** (41%) of the 47,984 STI clients with unknown or new negative test result were referred for repeat HTC. **1,052** (68%) of 1,547 clients who were newly tested HIV positive were referred for ART eligibility assessment.

19 Supply of HIV Program Commodities

19.1 Quantification and procurement planning

The program updated the quarterly quantification and procurement planning for all HIV commodities in July 2015 to inform procurement planning and budgeting for HIV commodities for the period ending December

2017. The Global Fund made available transition funding for the procurement of essential HIV Program commodities in Q2 2015 to facilitate a smooth transition into the New Funding Model Grant which is expected to start in January 2016.

During Q2 2015, antiretroviral drugs (ARVs) and medicines for opportunistic infections (OIs) worth \$33 million were received by the Bollore Africa Logistics-managed warehouses dedicated for Department of HIV and AIDS and National Malaria Control Program commodities. This comprised of Tenofovir/Lamivudine/Efavirenz 300/300/600mg (Regimen 5A; 85% of the value of adult ARVs) and other ARV formulations (altogether 97% of the value for all medicines received during the period). To maintain adequate stocks and ensure uninterrupted supply for subsequent orders, the Ministry has continued processing HIV commodity orders for ARVs, OIs, rapid diagnostic tests (RDTs) and other related commodities through Partnership for Supply Chain Management (ARVs and RDTs) and IDA Foundation (laboratory commodities and medicines for OIs).

19.2 Bimonthly distribution of HIV & Malaria Commodities

Two scheduled bimonthly rounds of distribution for HIV and Malaria commodities (Distribution Round 22 and 23) took place between April and June 2015. A total of 64 different commodities (anti-malarials, ARVs, OI medicines, STI medicines and laboratory commodities) were distributed to 726 health facilities. These were the third and fourth successful consolidated distributions for HIV and malaria commodities.

The DHA Logistics Team developed distribution lists for all HIV commodities for 726 sites for both rounds, covering the consumption period up to end September 2015. The team also coordinated over **1,500** individual commodity transactions between sites to avert stock outs and/or expiries. All transactions were managed using the toll-free HIV Department Supply Chain Hot Line that was set up to facilitate communication between the health facilities and the department. Health workers are able to communicate in a timely manner supply chain issues that need to be resolved by the technical team at the department.

19.3 Logistics support during integrated site supervision

Supply Chain and Logistics Officers provided stock management support at 292 sites during the Q2 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. Health care providers have continued to use RDT daily activity registers and relocation books for registration of redistributed commodities to health facilities. Authorization codes must be obtained for all such transactions as a measure for tracking, audit and to foster accountability at all sites.

19.4 Stock Status of HIV Commodities

Physical stock counts for ARVs and medicines for OIs were performed at all sites during the supervision in July 2015. **Table 6** shows the total medicine stocks found at the sites and the estimated consumption rates.

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.

493,431 patients were on regimen 5A, which was 19,267 (**3.8%**) less than projected in the previous forecast for the end of this quarter (**512,698**). The national ART program forecast and quantification was updated in July 2015 to inform procurement planning and budgeting for HIV commodities for the period ending December 2017.

19.5 Availability of standard first line ARVs

493,431 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. As of July 2015, the total stock of this regimen was equivalent to **5.5 and 5.9 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 July 2015 confirmed that 699 (**98.4%**) of all 710 ART sites with patients on this regimen had available stocks. This translates into a 'stock-out' rate of only 1.6% 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.

Table 6: Total stocks of HIV program commodities at all sites visited during the 2015 Q2 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 30/06/2015

Inventory unit	Item	Sites with any Stock	Total Physical Stock		Consumption/ Month	Months of Stock *	
			At Sites	In Warehouse		At Sites	Wareh.
tins	ABC / 3TC 60 / 30mg tins (60 tabs)	166	13,414	34,878	4,590	2.9	7.6
	ABC / 3TC 600 / 300mg tins (30 tabs)	39	1,356	2,558	663	2.0	3.9
	ATV / r 300 / 100mg tins (30 tabs)	193	13,011	43,290	5,896	2.2	7.3
	AZT / 3TC / NVP 300 / 150 / 200mg tins (60 tabs)	625	119,180	375,783	28,298	4.2	13.3
	AZT / 3TC / NVP 60 / 30 / 50mg tins (60 tabs)	640	592,565	531,650	63,503	9.3	8.4
	AZT / 3TC 300 / 150mg tins (60 tabs)	491	30,399	962	2,669	11.4	0.4
	AZT / 3TC 60 / 30mg tins (60 tabs)	580	28,146	14,288	2,464	11.4	5.8
	d4T / 3TC / NVP 30 / 150 / 200mg tins (60 tabs)	159	23,482	2,144	1,158	20.3	1.9
	d4T / 3TC 30 / 150mg tins (60 tabs)	219	12,566		123	102.2	
	EFV 200mg tins (90 tabs)	157	2,070	2,164	332	6.2	6.5
	EFV 600mg tins (30 tabs)	220	11,492	6,687	893	12.9	7.5
	LPV / r 100 / 25mg tins (60 tabs)	72	4,377	12,238	2,586	1.7	4.7
	LPV / r 200 / 50mg tins (120 tabs)	62	1,100	2,341	447	2.5	5.2
	NVP 200mg tins (60 tabs)	450	41,637	75,419	7,533	5.5	10.0
	NVP 50mg tins (60 tabs)	144	10,628	15,641	1,670	6.4	9.4
	TDF / 3TC / EFV 300 / 300 / 600mg tins (30 tabs)	704	2,689,119	2,911,454	493,034	5.5	5.9
	TDF / 3TC 300 / 300mg tins (30 tabs)	635	69,183	80,544	12,468	5.5	6.5
bottles	Fluconazole (Diflucan) 50mg / 5ml bottles (35 ml)	19	2,272		71	32.1	
	NVP 10mg/ml bottles (25 ml)	584	166,232	4,675	15,188	10.9	0.3
vials	Benzathine Penicillin 1.44g vials (50 each)	578	362,835		39,289	9.2	
	Bleomycine 15,000IU vials (1 each)	25	6,894				
	Ceftriaxone 1g vials (50 each)	362	91,025		106,048	0.9	
	Depo-Provera 150mg/1ml vials (25 each)	549	858,671	601,650	269,248	3.2	2.2
	Gentamicin 80mg / 2ml vials (50 each)	514	208,433	250,000	99,796	2.1	2.5
	Streptomycin 1 gm vials (50 each)	47	32,575				
Vincristine 1mg / 1ml vials (1 each)	85	33,449	29,950	3,180	10.5	9.4	
tabs	Aciclovir 200mg blister packs (500 tabs)	638	4,560,350	1,392,000	639,256	7.1	2.2
	Azithromycin 500mg blister packs (3 tabs)	243	203,854	18,000	10,550	19.3	1.7
	Ciprofloxacin 500mg blister packs (100 tabs)	307	1,258,469	874,400	302,378	4.2	2.9
	Clotrimazole 500mg boxes (1 each)	281	40,964	45,031	38,868	1.1	1.2
	Codeine 30mg tins (100 tabs)	69	347,890		50,045	7.0	
	Cotrimoxazole 100 / 20mg blister packs (1000 tabs)	621	35,082,486	14,477,000	6,951,399	5.0	2.1
	Cotrimoxazole 400 / 80mg tins (1000 tabs)	605	31,571,311	15,087,000	16,764,374	1.9	0.9
	Cotrimoxazole 960mg blister packs (1000 tabs)	664	29,521,726	10,721,000	17,776,210	1.7	0.6
	Doxycycline 100mg tins (1000 tabs)	521	5,602,000	1,505,000	4,480,304	1.3	0.3
	E thambutol (E) 100 mg blister packs (100 tabs)	44	78,811				
	E thambutol (E) 400 mg blister packs (672 tabs)	11	27,368				
	Erythromycin 250mg tins (1000 tabs)	523	3,370,588	1,240,000	4,008,086	0.8	0.3
	Fluconazole (Diflucan) 200mg tins (28 tabs)	341	799,425	134,092	39,399	20.3	3.4
	Ibuprofen 200mg tins (100 tabs)	238	919,592		856,938	1.1	
	Isoniazid (H) 100mg blister packs (100 tabs)	156	670,195		144,931	4.6	
	Isoniazid (H) 300mg blister packs (672 tabs)	63	478,216		1,059,811	0.5	
	Isoniazid (H) 300mg tins (1000 tabs)	621	14,859,065	10,059,000	1,060,764	14.0	9.5
	Metronidazole 200mg tins (1000 tabs)	576	13,614,618		4,867,064	2.8	
	Morphine 10mg blister packs (60 tabs)	113	21,397,704		218,379	98.0	
Pyridoxine 50mg tins (1000 tabs)	573	13,233,414	1,000,000	1,132,277	11.7	0.9	
sheets	ART pat. card adult (yellow) bundles (100 sheets)	657	260,634		9,955	26.2	
	ART pat. card paed. (blue) bundles (100 sheets)	579	151,350		924	163.7	
	Exposed child card (pink) bundles (50 sheets)	581	56,534		3,465	16.3	
	Polythene sleeve bundles (100 sheets)	582	163,293		16,370	10.0	
	Pre-ART pat. card (green) bundles (100 sheets)	566	134,173		2,026	66.2	
tests	DBS kit (filter paper, lancet, etc.) boxes (50 each)	533	59,220	41,750	32,404	1.8	1.3
	Determine HIV1/2 boxes (100 each)	663	1,138,297	617,400	108,725	10.5	5.7
	Determine syphilis boxes (100 each)	341	104,302	265,900			
	Uni-Gold HIV1/2 boxes (20 each)	576	85,850	50,420	13,641	6.3	3.7
pieces	Condoms female boxes (1000 each)	419	725,621		187,660	3.9	
	Condoms male boxes (144 each)	516	9,084,166	1,465,200	4,462,760	2.0	0.3

* '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.

20 Training and Mentoring

ART/PMTCT trainings

19 were newly trained in the 2014 national clinical guidelines. 11 were nurse and 8 clinicians.

21 Participants in Q2 2015 Supervision (Site visits 20 July-7 August 2015)

Absalom Kaunda (CO, MOH, Mzimba DHO)
 Afred Kamoto (Logistics Fellow, MOH)
 Agnes Kalitsiro (Nurse, Mlambe Mission Hospital)
 Alefa Fikira (CMT, MOH)
 Alexander Malunguza (, NTP)
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 Andrew Dimba (, NTP)
 Andrew Gompho (Clinician, MOH)
 Andrew Mganga (M&E Fellow, Dept for HIV and AIDS)
 Annie Biza (Nurse, MDF)
 Austin Nkute (CLINICAL CORDINATOR, indep NGO)
 Austins Namondwe (CO, CHAM)
 Batoni Upindi (TB Zonal Supervisor, MOH)
 Beatrice Malonje (Nurse, MOH)
 Benardetta Chunda (Nurse, Lighthouse)
 Catherine Kassam (, MOH)
 Cecelia Tenesi (Nurse, MOH)
 Cecilia Manyawa (Nurse, MOH)
 Cecilia Sambakunsi (Logistics Fellow, HIV Dept)
 Chancy Kamba (, NTP)
 Charles F Sekani (CO, .)
 Chifundo Makuluni (Nurse, MOH)
 Chikayiko Majamanda (Nurse, MOH)
 Chikumbutso Pendame (MA, MOH)
 Chimwemwe Francis Mkandawire (IT Fellow, Dept for HIV and AIDS)
 Chisomo Ngwalo (, COM)
 Chisomo Thondolo (Nurse, EGPAF)
 Chrissy Gondwe (, Dignitas)
 Chrissy Lizengo (, MOH)
 Christopher Mkwezalamba (CO, MOH)
 Clement Chiphota (CO, MoH)
 Cornelias Kang'ombe (, NTP)
 Cornelius Kang'ombe (, NTP)
 Dalitso Midiani (PMTCT Officer, MOH)
 Dan Chimbayo (CO, NGO)
 Daniel Nyirenda (Clerk, MOH)
 Davie Nkosi (, MOH)
 Deliwe Msiska (, JHPIEGO)
 Diana Chipande (, MOH)
 Dorica Chirwa (Logistic officer, MOH)
 Dr Simon Chiumia (, Private)
 Edgar Lungu (, UN AIDS)
 Edith Thaulo (Nurse, MOH)
 Elizabeth Chatsika (CO, CHAM)
 Elton Masina (CO, EGPAF)
 Enipher Kalengamaliro (, MOH)
 Erick Mtemang'ombe (CO, CHAM)
 Erik Mittochi (CO ART coord), MOH)
 Evans Kagwira (TB Zonal Supervisor, MOH)
 Everista Mkandawire (Nurse, MOH)
 Everson Mwandira C (, MOH)
 Ezra Majoni (Nurse, MOH)
 Fainala Muyila (Nurse, MOH)
 Fatsireni Mapulanga (, MOH)
 Felix Magwira (Clinical Cordinator, indep NGO)
 Frazer Mkawa (Nurse, MOH)
 Geoffrey Makhalaria (, NTP)
 Gerald Zomba (Program Officer, Dept for HIV and AIDS)
 Grant Gondwe (, NTP)
 Hanna Tenthani (Nurse, MOH)
 Hannock Matupi (ARV clinician, MOH, Rumphidh)
 Harrison Tembo (CO, MOH)
 Harry Tsapa (CO, MOH)
 Henry Kanyerere (TB/HIV Program Officer, MOH)
 Isaac Makhonya (, PIH)
 Isaiah Dambe (, NTP)
 Issa Sulemani (, MOH)
 Janet Chikonda (Nurse, MOH)
 Jean Kayamba (Nurse, MOH)
 Jesse Lobeni (Nurse, MOH)
 John Kabichi (CO, MOH)
 Johnbosco Mwafilaso (Clerk, MOH)
 Juliana Soko (ARV nurse, MOH, Livingstonia MH)
 Julie Kazima (Nurse, MSH)
 Juliet Nyirenda (Nurse, MOH)
 Justice Kaphiri (, NTP)
 Kelvin Makina (Logistics, Kasungu)
 Kingsley Makwale (MA, MOH)
 Kingsley Mbewa (CO, MOH)
 Knox Banda (TB Zonal Supervisor, MOH)
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 Laphoid Chisuwu (, NTP)
 Leonard Kadongola (, MOH)
 Lilian Kachali (Nurse, MOH)
 Lincy Chalunda (CO, MOH)
 Linda Dziweni (Nurse, Baylor)
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 Mary Gosten (MA, 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.

3rd November 2015

22 Appendix (Full National HIV Program Data)

HTC site report

Malawi (national)

2015 Q2 (1st month of quarter, 2nd month of quarter, 3rd month of quarter)

Clients at health facility (static)

HTC client details

*

Total HTC clients served

Total HIV tested	466,265	100%
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Sex

Males tested	160,000	34%
Females tested	306,265	66%
Females non-pregnant	150,738	49%
Females pregnant	155,527	51%

Age

Children 0-14 yrs	34,270	7%
Children below 12 mths (Age group A)	2,264	7%
Children 12 mths - 14 yrs (Age group B)	32,006	93%
Adults 15+ years	431,995	93%
Young adults 15-24 years (Age group C)	188,229	44%
Older adults 25+ yrs (Age group D)	243,766	56%

HTC access type

PITC	263,084	56%
Family Referral Slip (FRS)	1,821	0%
Other (VCT, etc.) HTC access	201,360	43%

HTC first time / repeat

Never tested before	140,983	30%
Previously accessed HTC	325,282	70%
Last negative	313,551	96%
Last positive	9,608	3%
Last exposed infant	1,702	1%
Last inconclusive	421	0%

Counseling session type / Partner present

Counseled with partner / partner present	116,796	25%
Counseled alone / Partner not present	349,469	75%

Outcome summary (HIV test)

Single test negative	426,700	92%
Single test positive	431	0%
Test 1&2 negative	377	0%
Test 1&2 positive	37,416	8%
Test 1&2 discordant	1,341	0%

Final result given to client

Results among clients never tested / last negative	456,853	98%
New negative	426,360	93%
New positive	28,248	6%
New exposed infants	941	0%
New inconclusive	1,304	0%
Confirmatory results (previous positive clients)	9,412	2%
Confirmatory positive	9,040	96%
Confirmatory inconclusive	372	4%

HTC site report

Malawi (national)

2015 Q2 (1st month of quarter, 2nd month of quarter, 3rd month of quarter)

HTC client details

*

Partner / Family HTC referral slips

Sum of slips given	20,454	100%
Total clients presenting with referral slip	1,821	9%
Total failed referrals (slips not returned)	18,633	91%

Clients tested in the community

HTC client details

*

Total HTC clients served

Total HIV tested	17,112	100%
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Sex

Males tested	8,565	50%
Females tested	8,547	50%
Females non-pregnant	6,875	80%
Females pregnant	1,672	20%

Age

Children 0-14 yrs	2,831	17%
Children below 12 mths (Age group A)	7	0%
Children 12 mths - 14 yrs (Age group B)	2,824	100%
Adults 15+ years	14,281	83%
Young adults 15-24 years (Age group C)	7,261	51%
Older adults 25+ yrs (Age group D)	7,020	49%

HTC access type

PITC	3,885	23%
Family Referral Slip (FRS)	12	0%
Other (VCT, etc.) HTC access	13,215	77%

HTC first time / repeat

Never tested before	6,998	41%
Previously accessed HTC	10,114	59%
Last negative	10,029	99%
Last positive	77	1%
Last exposed infant	2	0%
Last inconclusive	6	0%

Counseling session type / Partner present

Counseled with partner / partner present	1,579	9%
Counseled alone / Partner not present	15,533	91%

Outcome summary (HIV test)

Single test negative	15,321	90%
Single test positive	2	0%
Test 1&2 negative	3	0%
Test 1&2 positive	518	3%
Test 1&2 discordant	1,268	7%

HTC site report

Malawi (national)

2015 Q2 (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	17,033	100%
New negative	16,546	97%
New positive	443	3%
New exposed infants	10	0%
New inconclusive	34	0%
Confirmatory results (previous positive clients)	79	0%
Confirmatory positive	72	91%
Confirmatory inconclusive	7	9%

Partner / Family HTC referral slips

Sum of slips given	161	100%
Total clients presenting with referral slip	12	7%
Total failed referrals (slips not returned)	149	93%

Clients at stand-alone HTC sites

HTC client details

*

Total HTC clients served

Total HIV tested	10,190	100%
------------------	--------	------

Sex

Males tested	5,097	50%
Females tested	5,093	50%
Females non-pregnant	4,374	86%
Females pregnant	719	14%

Age

Children 0-14 yrs	170	2%
Children below 12 mths (Age group A)	6	4%
Children 12 mths - 14 yrs (Age group B)	164	96%
Adults 15+ years	10,020	98%
Young adults 15-24 years (Age group C)	3,443	34%
Older adults 25+ yrs (Age group D)	6,577	66%

HTC access type

PITC	4,750	47%
Family Referral Slip (FRS)	26	0%
Other (VCT, etc.) HTC access	5,414	53%

HTC first time / repeat

Never tested before	2,747	27%
Previously accessed HTC	7,443	73%
Last negative	7,305	98%
Last positive	135	2%
Last exposed infant	0	0%
Last inconclusive	3	0%

Counseling session type / Partner present

Counseled with partner / partner present	1,209	12%
Counseled alone / Partner not present	8,981	88%

HTC site report

Malawi (national)

2015 Q2 (1st month of quarter, 2nd month of quarter, 3rd month of quarter)

HTC client details

*

Outcome summary (HIV test)

Single test negative	9,454	93%
Single test positive	3	0%
Test 1&2 negative	4	0%
Test 1&2 positive	701	7%
Test 1&2 discordant	28	0%

Final result given to client

Results among clients never tested / last negative	10,164	100%
New negative	9,466	93%
New positive	594	6%
New exposed infants	0	0%
New inconclusive	104	1%
Confirmatory results (previous positive clients)	26	0%
Confirmatory positive	21	81%
Confirmatory inconclusive	5	19%

Partner / Family HTC referral slips

Sum of slips given	354	100%
Total clients presenting with referral slip	26	7%
Total failed referrals (slips not returned)	328	93%

Blood safety

Malawi (national)

2015 Q2 (1st month of quarter, 2nd month of quarter, 3rd month of quarter)

Infect. disease screening among potential donors

*

HIV screening

HIV testing not done	2,373	18%
Tested for HIV	10,648	82%
HIV negative	10,055	94%
HIV positive	593	6%

Hepatitis B screening

HepB testing not done	2,382	18%
Tested for Hepatitis B	10,639	82%
HepB Negative	10,149	95%
HepB Positive	490	5%

Hepatitis C screening

HepC testing not done	8,373	64%
Tested for Hepatitis C	4,648	36%
HepC Negative	4,578	98%
HepC Positive	70	2%

Syphilis screening

Syphilis testing not done	2,431	19%
Tested for Syphilis	10,590	81%
Syphilis Negative	10,277	97%
Syphilis Positive	313	3%

Malaria screening

Malaria testing not done	5,011	38%
Tested for malaria	8,010	62%
Malaria Negative	7,146	89%
Malaria Positive	864	11%

Summary screening outcome

Not donated	4,402	34%
Donated	8,619	66%
Screened for at least HIV, HepB and syphilis	7,675	89%
Screened for HIV, HepB, HepC, Syphilis, Malaria	3,859	50%
Screened for HIV, HepB, Syphilis	3,816	50%
Screened for HIV, HepB	14	0%
Screened for HIV only	3	0%
Screened with any other combination of tests	927	11%

Cross-matching report

*

Blood group typing (for units and patients)

Total blood group typing done	25,827	100%
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Blood units cross-matched (by source)

Total blood units cross-matched	16,424	100%
Total units from MBTS (estimated)	7,805	48%
Total units from replacement donors	8,619	52%

Blood units cross-matched by patient group

Units cross-matched for maternity	1,973	12%
Units cross-matched for paediatrics	8,209	50%
Units cross-matched for other ward	6,242	38%

Blood safety

Malawi (national)

2015 Q2 (1st month of quarter, 2nd month of quarter, 3rd month of quarter)

Cross-matching report

*

Transfusion reactions

Units transfused without adverse events	16,412	100%
Units with suspected transfusion reactions	9	0%
Units with confirmed transfusion reactions	3	0%

HIV exposed child follow-up

Malawi (national)

2015 Q2 (1st month of quarter, 2nd month of quarter, 3rd month of quarter)

Age 2 months

Age cohort outcomes

*

Total children in birth cohort

Total children registered	8,509	100%
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CPT status

On CPT	7,723	91%
Not on CPT	786	9%

HIV status

Current HIV infection status unknown	5,985	70%
HIV infection not confirmed, not ART eligible	5,960	100%
HIV infection not confirmed, ART eligible (PSHD)	25	0%
Current HIV infection status known	2,524	30%
Confirmed not infected	2,467	98%
Confirmed infected (ART eligible)	57	2%

ART eligibility summary

Not eligible for ART	8,427	99%
ART eligible	82	1%
ART not initiated	35	43%
Initiated ART	47	57%

Primary follow-up outcome

Discharged uninfected	16	0%
Continue follow-up	7,806	93%
Started ART	47	1%
Defaulted	494	6%
Died	26	0%

Transfers between sites

Total not transferred out	8,389	99%
Transferred out	120	1%

Age 12 months

Age cohort outcomes

*

Total children in birth cohort

Total children registered	8,417	100%
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CPT status

On CPT	6,278	75%
Not on CPT	2,139	25%

HIV status

Current HIV infection status unknown	4,823	57%
HIV infection not confirmed, not ART eligible	4,799	100%
HIV infection not confirmed, ART eligible (PSHD)	24	0%
Current HIV infection status known	3,594	43%
Confirmed not infected	3,429	95%
Confirmed infected (ART eligible)	165	5%

HIV exposed child follow-up

Malawi (national)

2015 Q2 (1st month of quarter, 2nd month of quarter, 3rd month of quarter)

Age cohort outcomes

*

ART eligibility summary

Not eligible for ART	8,228	98%
ART eligible	189	2%
ART not initiated	29	15%
Initiated ART	160	85%

Primary follow-up outcome

Discharged uninfected	61	1%
Continue follow-up	6,238	76%
Started ART	160	2%
Defaulted	1,638	20%
Died	71	1%

Transfers between sites

Total not transferred out	8,168	97%
Transferred out	249	3%

Age 24 months

Age cohort outcomes

*

Total children in birth cohort

Total children registered	8,121	100%
---------------------------	-------	------

CPT status

On CPT	1,192	15%
Not on CPT	6,929	85%

HIV status

Current HIV infection status unknown	4,223	52%
HIV infection not confirmed, not ART eligible	4,193	99%
HIV infection not confirmed, ART eligible (PSHD)	30	1%
Current HIV infection status known	3,898	48%
Confirmed not infected	3,673	94%
Confirmed infected (ART eligible)	225	6%

ART eligibility summary

Not eligible for ART	7,866	97%
ART eligible	255	3%
ART not initiated	31	12%
Initiated ART	224	88%

Primary follow-up outcome

Discharged uninfected	3,457	44%
Continue follow-up	952	12%
Started ART	224	3%
Defaulted	3,061	39%
Died	111	1%

Transfers between sites

Total not transferred out	7,805	96%
Transferred out	316	4%

2015 Q2 (Quarter)

Registration details

*

HCC clinic registrations

Total HCC registrations	16,473	100%
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Registration type

Patients enrolled first time	15,676	95%
Patients re-enrolled	46	0%
Patients transferred in	751	5%

Sex

Males (all ages)	7,598	46%
Females (all ages)	8,875	54%
Non-pregnant	8,856	100%
Pregnant	19	0%

Age at registration

Adults 15+ yrs	5,676	34%
Children 0-14 yrs	10,797	66%
Children 24 months - 14 years	472	4%
Children below 24 months (exposed children)	10,325	96%
Children 2 - below 24 months	3,762	36%
Infants below 2 months	6,563	64%

Reason for HCC registration

Exposed infants	10,396	63%
Confirmed infected patients (pre-ART)	6,077	37%

2015 Q2 (Cumulative)

Registration details

*

HCC clinic registrations

Total HCC registrations	354,234	100%
-------------------------	---------	------

Registration type

Patients enrolled first time	341,821	96%
Patients re-enrolled	1,131	0%
Patients transferred in	11,282	3%

Sex

Males (all ages)	153,115	43%
Females (all ages)	201,119	57%
Non-pregnant	200,125	100%
Pregnant	994	0%

Age at registration

Adults 15+ yrs	176,773	50%
Children 0-14 yrs	177,461	50%
Children 24 months - 14 years	16,254	9%
Children below 24 months (exposed children)	161,207	91%
Children 2 - below 24 months	76,528	47%
Infants below 2 months	84,679	53%

Reason for HCC registration

Exposed infants	160,999	45%
Confirmed infected patients (pre-ART)	193,235	55%

Pre-ART follow-up outcome

*

Primary follow-up outcomes

Total retained in pre-ART	39,791	22%
Started ART	97,923	53%
Defaulted	44,168	24%
Died	1,550	1%

Transfers between sites

Total not transferred out	183,826	95%
Transferred out	9,409	5%

Antenatal Care

Malawi (national)

2015 Q2 (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	144,819	100%
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ANC cohort analysis

*

Trimester of first visit

Started ANC 0-12 wks	14,665	10%
Started ANC 13+ wks	130,154	90%

HIV status ascertainment

HIV status not ascertained	20,921	14%
HIV status ascertained	123,898	86%
Valid previous test result	9,879	8%
Previous negative	4,542	46%
Previous positive	5,337	54%
New test at ANC	114,019	92%
New negative	109,477	96%
New positive	4,542	4%

HIV status summary

Total women HIV negative	114,019	92%
Total women HIV positive	9,879	8%

PMTCT regimen mother

No ARVs	959	10%
Any ARVs	8,920	90%
ART (by time of initiation)	8,920	100%
Already on ART when starting ANC	4,727	53%
Started ART at 0-27 weeks of pregnancy	3,457	39%
Started ART at 28+ weeks of preg.	736	8%

ANC women after 6 months

ANC cohort analysis

*

Total women completing ANC in the reporting period

Total women in booking cohort	155,272	100%
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Visits per woman

Women with 1 visit	34,772	22%
Women with 2 visits	41,294	27%
Women with 3 visits	45,268	29%
Women with 4 visits	27,508	18%
Women with 5+ visits	6,430	4%

Pre-eclampsia

No pre-eclampsia	152,944	99%
Pre-eclampsia	2,328	1%

TTV doses

0-1 TTV doses	78,058	50%
2+ TTV doses	77,214	50%

SP tablets

0 SP doses	17,773	11%
1 SP dose (1 x 3 tabs)	41,753	27%
6+ SP tablets (2 x 3 tabs)	95,746	62%

Antenatal Care

Malawi (national)

2015 Q2 (1st month of quarter, 2nd month of quarter, 3rd month of quarter)

ANC cohort analysis

*

FeFo tablets

0-119 FeFo tablets	118,171	76%
120+ FeFo tablets	37,101	24%

Albendazole (Deworming)

0 Albend. doses	30,621	20%
1 Albend. dose	125,375	80%

ITN (bednets)

No ITN	34,635	22%
ITN received	120,810	78%

Syphilis status

Not tested for syphilis	142,013	91%
Tested for syphilis	13,259	9%
Syphilis negative	12,810	97%
Syphilis positive	449	3%

HIV status ascertainment

HIV status not ascertained	14,060	9%
HIV status ascertained	141,212	91%
Valid previous test result	11,914	8%
Previous negative	6,484	54%
Previous positive	5,430	46%
New test at ANC	129,298	92%
New negative	123,962	96%
New positive	5,336	4%

HIV status summary

Total women HIV negative	130,446	92%
Total women HIV positive	10,766	8%

CPT status (among HIV pos)

Not on CPT	1,030	10%
On CPT	9,736	90%

PMTCT regimen mother

No ARVs	1,026	10%
Any ARVs	9,740	90%
ART (by time of initiation)	9,740	100%
Already on ART when starting ANC	4,677	48%
Started ART at 0-27 weeks of pregnancy	4,000	41%
Started ART at 28+ weeks of preg.	1,063	11%

Baby's ARVs dispensed

No ARVs dispensed for infant	1,858	17%
ARVs dispensed for infant	8,908	83%

Maternity

Malawi (national)

2015 Q2 (1st month of quarter, 2nd month of quarter, 3rd month of quarter)

Maternal details

*

Admissions in the reporting period

Total admissions (referrals double-counted)	125,961	100%
Not referred to other site (total women)	118,878	94%
Referred out before delivery (multiple admissions)	7,083	6%

HIV status ascertainment

HIV status not ascertained	5,146	4%
HIV status ascertained	120,781	96%
Valid previous test result	117,592	97%
Previous negative	108,871	93%
Previous positive	8,721	7%
New test at maternity	3,189	3%
New negative	2,943	92%
New positive	246	8%

HIV status summary

Total women HIV negative	111,814	93%
Total women HIV positive	8,967	7%

ARVs during pregnancy (among HIV pos)

No ARV in pregnancy	188	2%
Any ARVs	8,779	98%
ART (by time of initiation)	8,779	100%
ART initiated before pregnancy	6,315	72%
ART initiated in 1st / 2nd trimester	1,247	14%
ART initiated in 3rd trimester	961	11%
ART initiated during labour	256	3%

Obstetric complications

No obstetric complications	110,339	88%
Any obstetric complications	15,588	12%
Haemorrhage	2,547	16%
Haemorrhage ante-partum	758	30%
Haemorrhage post-partum	1,789	70%
Obstr / prol labour	5,270	34%
(pre-) Eclampsia	906	6%
Maternal sepsis	136	1%
Ruptured uterus	109	1%
Other obstetric complications	6,620	42%

Emergency obstetric care

Oxytocin	116,468	94%
Anticonvulsive	656	1%
Antibiotics	6,277	5%
Blood transfusion	321	0%
Manual removal of placenta	470	0%

Vitamin A

Vit A not given	36,871	29%
Vit A given	89,056	71%

Maternity

Malawi (national)

2015 Q2 (1st month of quarter, 2nd month of quarter, 3rd month of quarter)

Maternal details

*

Staff conducting delivery

Category A: MO, CO, nurse/midwife, MA	113,650	96%
Category B: PA, WA, HSA	575	0%
Category C: Other	4,619	4%

Mother survival

Mother alive	118,750	100%
Mother died	94	0%

Infant details

*

Single babies / multiple deliveries

Total babies delivered	121,370	100%
Single babies	117,136	97%
Twin / multiple babies	4,234	3%

Delivery place

Total deliveries at a health facility	116,472	96%
This facility	116,226	100%
Other facility	246	0%
Total deliveries before reaching the facility	4,898	4%
In transit	3,257	66%
Home / TBA	1,641	34%

Delivery mode

Spontaneous vaginal	109,105	90%
Vacuum extraction	1,557	1%
Breech	2,146	2%
Caesarean section	8,562	7%

Infant complications

No infant complications	105,255	87%
Total infants with complications	16,115	13%
Prematurity	3,930	24%
Weight less 2500g	5,021	31%
Asphyxia	4,952	31%
Sepsis	567	4%
Other newborn complication	1,645	10%

Infant survival

Total live births	119,374	98%
Discharged alive	118,182	99%
Neonatal deaths	1,192	1%
Stillbirths	1,996	2%
Stillbirth, fresh	1,012	51%
Stillbirth, macerated	984	49%

Maternity

Malawi (national)

2015 Q2 (1st month of quarter, 2nd month of quarter, 3rd month of quarter)

Infant details

*

HIV exposure / ARV proph. (among discharged alive)

Infants with unknown HIV exposure status	4,089	3%
Infants with known HIV exposure status	114,093	97%
Not HIV exposed	105,883	93%
HIV exposed	8,210	7%
Received no ARVs	439	5%
Received ARVs	7,771	95%
Nevirapine	7,771	100%

Breastfeeding initiated

BF not started within 60min	10,334	9%
BF started within 60min	111,036	91%

Tetracycline eye ointment given

TO not given	16,562	14%
TO given	104,808	86%

ART cohort analysis

Malawi (national)

2015 Q2 (Quarter)

Registration details

*

ART clinic registrations

Total ART clinic registrations	32,637	100%
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Registration type

First time ART initiations (total patients)	25,957	80%
ART re-initiations	421	1%
ART transfers in	6,259	19%

Sex

Males	11,901	36%
Females	20,736	64%
Non-pregnant	14,469	70%
Pregnant	6,267	30%

Age at ART initiation

Adults 15+ yrs	29,864	92%
Children 0-14 yrs	2,773	8%
Children 2-14 yrs	2,132	77%
Children below 24 mths	641	23%

Reason for starting ART

Presumed severe HIV Disease	113	0%
Confirmed HIV infection	32,524	100%
WHO stage 1 or 2	20,876	64%
Total lymphocytes <threshold	8	0%
CD4 below threshold	12,391	59%
CD4 unknown or >threshold	8,477	41%
PCR infants	87	1%
Children 12-59 mths	635	7%
Pregnant women	6,164	73%
Breastfeeding mothers	1,591	19%
WHO stage 3	9,895	30%
WHO stage 4	1,438	4%
Unknown / reason outside of guidelines	315	1%

TB at ART initiation

Never TB / TB > 24 months ago	31,706	97%
TB within the last 24 months	342	1%
Current episode of TB	589	2%

Kaposi's sarcoma at ART initiation

No KS	32,372	99%
Patients with KS	265	1%

ART cohort analysis

Malawi (national)

2015 Q2 (Cumulative)

Registration details

*

ART clinic registrations

Total ART clinic registrations	1,025,754	100%
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Registration type

First time ART initiations (total patients)	820,367	80%
ART re-initiations	11,520	1%
ART transfers in	193,867	19%

Sex

Males	369,284	36%
Females	656,470	64%
Non-pregnant	536,145	82%
Pregnant	120,325	18%

Age at ART initiation

Adults 15+ yrs	936,603	91%
Children 0-14 yrs	89,151	9%
Children 2-14 yrs	68,735	77%
Children below 24 mths	20,416	23%

Reason for starting ART

Presumed severe HIV Disease	3,520	0%
Confirmed HIV infection	1,022,234	100%
WHO stage 1 or 2	447,066	44%
Total lymphocytes <threshold	267	0%
CD4 below threshold	297,890	67%
CD4 unknown or >threshold	148,909	33%
PCR infants	2,861	2%
Children 12-59 mths	6,911	5%
Pregnant women	101,636	68%
Breastfeeding mothers	37,501	25%
WHO stage 3	467,590	46%
WHO stage 4	100,379	10%
Unknown / reason outside of guidelines	7,199	1%

TB at ART initiation

Never TB / TB > 24 months ago	953,839	93%
TB within the last 24 months	36,587	4%
Current episode of TB	35,328	3%

Kaposi's sarcoma at ART initiation

No KS	1,006,268	98%
Patients with KS	19,486	2%

ART cohort analysis

Malawi (national)

2015 Q2 (Cumulative)

ART outcomes

*

Primary follow-up outcomes

Total alive on ART	565,105	68%
Alive on ART at site of last registration	564,904	100%
ART patients in transit between sites	201	0%
Defaulted	186,074	22%
Stopped ART	3,366	0%
Total died	77,342	9%
Died month 1	19,087	25%
Died month 2	11,998	16%
Died month 3	7,075	9%
Died month 4+	39,182	51%

Transfers between sites

Total not transferred out	831,686	81%
Transferred out	194,068	19%

ART regimens

First line regimens	557,673	99%
Adult formulation	531,106	95%
Regimen 0A	190	0%
Regimen 1A	1,158	0%
Regimen 2A	28,298	5%
Regimen 3A	123	0%
Regimen 4A	770	0%
Regimen 5A	493,034	93%
Regimen 6A	7,533	1%
Paed. formulation	26,567	5%
Regimen 0P	443	2%
Regimen 1P	225	1%
Regimen 2P	25,401	96%
Regimen 3P	54	0%
Regimen 4P	444	2%
Second line regimens	6,758	1%
Adult formulation	5,896	87%
Regimen 7A	4,466	76%
Regimen 8A	1,430	24%
Paed. Formulation	862	13%
Regimen 9P	862	100%
Other regimen (adult / paed)	473	0%

Adherence

Adherence unknown (not recorded)	4,110	1%
Adherence recorded	560,794	99%
0-3 doses missed	510,901	91%
4+ doses missed	49,893	9%

ART side effects

Side effects unknown (not recorded)	154,144	27%
Side effects recorded	410,760	73%
No side effects	401,094	98%
Any side effects	9,666	2%

ART cohort analysis

Malawi (national)

2015 Q2 (Cumulative)

ART outcomes

*

Current TB status among ART patients (ICF)

ICF not done (Current TB status unknown/ not circ)	12,217	2%
ICF done	552,687	98%
TB not suspected	545,623	99%
TB suspected	6,160	1%
TB confirmed	904	0%
TB confirmed, not on treatment	89	10%
TB confirmed, on TB treatment	815	90%

2015 Q2 (Quarter)

12 month survival children**Survival and retention in ART program**

*

ART cohort registration group outcomes

Total ART clinic registrations	2,925	100%
Transfers out (double counted)	273	9%
Total not transferred out (patients in cohort)	2,652	91%
Total alive on ART	2,067	78%
Total not retained	585	22%
Defaulted	495	85%
Stopped ART	7	1%
Died	83	14%

12 month survival all ages**Survival and retention in ART program**

*

ART cohort registration group outcomes

Total ART clinic registrations	33,757	100%
Transfers out (double counted)	2,895	9%
Total not transferred out (patients in cohort)	30,862	91%
Total alive on ART	24,274	79%
Total not retained	6,588	21%
Defaulted	5,561	84%
Stopped ART	51	1%
Died	976	15%

24 month survival all ages**Survival and retention in ART program**

*

ART cohort registration group outcomes

Total ART clinic registrations	31,231	100%
Transfers out (double counted)	3,967	13%
Total not transferred out (patients in cohort)	27,264	87%
Total alive on ART	19,405	71%
Total not retained	7,859	29%
Defaulted	6,407	82%
Stopped ART	81	1%
Died	1,371	17%

36 month survival all ages**Survival and retention in ART program**

*

ART cohort registration group outcomes

Total ART clinic registrations	35,012	100%
Transfers out (double counted)	5,554	16%
Total not transferred out (patients in cohort)	29,458	84%
Total alive on ART	20,666	70%
Total not retained	8,792	30%
Defaulted	6,766	77%
Stopped ART	122	1%
Died	1,904	22%

2015 Q2 (Quarter)

48 month survival all ages**Survival and retention in ART program**

*

ART cohort registration group outcomes

Total ART clinic registrations	22,708	100%
Transfers out (double counted)	5,779	25%
Total not transferred out (patients in cohort)	16,929	75%
Total alive on ART	11,166	66%
Total not retained	5,763	34%
Defaulted	3,970	69%
Stopped ART	85	1%
Died	1,708	30%

60 month survival all ages**Survival and retention in ART program**

*

ART cohort registration group outcomes

Total ART clinic registrations	22,590	100%
Transfers out (double counted)	5,799	26%
Total not transferred out (patients in cohort)	16,791	74%
Total alive on ART	10,441	62%
Total not retained	6,350	38%
Defaulted	4,201	66%
Stopped ART	68	1%
Died	2,081	33%

72 month survival all ages**Survival and retention in ART program**

*

ART cohort registration group outcomes

Total ART clinic registrations	19,824	100%
Transfers out (double counted)	5,822	29%
Total not transferred out (patients in cohort)	14,002	71%
Total alive on ART	7,984	57%
Total not retained	6,018	43%
Defaulted	3,963	66%
Stopped ART	83	1%
Died	1,972	33%

84 month survival all ages**Survival and retention in ART program**

*

ART cohort registration group outcomes

Total ART clinic registrations	18,306	100%
Transfers out (double counted)	5,454	30%
Total not transferred out (patients in cohort)	12,852	70%
Total alive on ART	7,111	55%
Total not retained	5,741	45%
Defaulted	3,421	60%
Stopped ART	85	1%
Died	2,235	39%

ART survival analysis

Malawi (national)

2015 Q2 (Quarter)

96 month survival all ages

Survival and retention in ART program

*

ART cohort registration group outcomes

Total ART clinic registrations	13,486	100%
Transfers out (double counted)	4,140	31%
Total not transferred out (patients in cohort)	9,346	69%
Total alive on ART	4,431	47%
Total not retained	4,915	53%
Defaulted	2,947	60%
Stopped ART	51	1%
Died	1,917	39%

108 month survival all ages

Survival and retention in ART program

*

ART cohort registration group outcomes

Total ART clinic registrations	10,423	100%
Transfers out (double counted)	3,351	32%
Total not transferred out (patients in cohort)	7,072	68%
Total alive on ART	3,130	44%
Total not retained	3,942	56%
Defaulted	1,981	50%
Stopped ART	39	1%
Died	1,922	49%

120 month survival all ages

Survival and retention in ART program

*

ART cohort registration group outcomes

Total ART clinic registrations	4,867	100%
Transfers out (double counted)	1,639	34%
Total not transferred out (patients in cohort)	3,228	66%
Total alive on ART	1,286	40%
Total not retained	1,942	60%
Defaulted	874	45%
Stopped ART	25	1%
Died	1,043	54%

6 month survival OptionB+

Survival and retention in ART program

*

ART cohort registration group outcomes

Total ART clinic registrations	8,275	100%
Transfers out (double counted)	588	7%
Total not transferred out (patients in cohort)	7,687	93%
Total alive on ART	5,845	76%
Total not retained	1,842	24%
Defaulted	1,788	97%
Stopped ART	27	1%
Died	27	1%

2015 Q2 (Quarter)

12 month survival OptionB+**Survival and retention in ART program**

*

ART cohort registration group outcomes

Total ART clinic registrations	7,829	100%
Transfers out (double counted)	657	8%
Total not transferred out (patients in cohort)	7,172	92%
Total alive on ART	5,168	72%
Total not retained	2,004	28%
Defaulted	1,925	96%
Stopped ART	26	1%
Died	53	3%

24 month survival OptionB+**Survival and retention in ART program**

*

ART cohort registration group outcomes

Total ART clinic registrations	8,600	100%
Transfers out (double counted)	845	10%
Total not transferred out (patients in cohort)	7,755	90%
Total alive on ART	5,225	67%
Total not retained	2,530	33%
Defaulted	2,411	95%
Stopped ART	37	1%
Died	82	3%

36 month survival OptionB+**Survival and retention in ART program**

*

ART cohort registration group outcomes

Total ART clinic registrations	9,269	100%
Transfers out (double counted)	1,334	14%
Total not transferred out (patients in cohort)	7,935	86%
Total alive on ART	5,462	69%
Total not retained	2,473	31%
Defaulted	2,292	93%
Stopped ART	40	2%
Died	141	6%

STI site report

Malawi (national)

2015 Q2 (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 treated	54,937	100%
Index patients treated (symptomatic)	44,700	81%
Partners treated	10,237	19%

Sex

Males	22,377	41%
Females	32,560	59%
Non-pregnant	28,580	88%
Pregnant	3,980	12%

Age group

Age group A (0-19 years)	5,010	9%
Age group B (20-24 years)	13,151	24%
Age group C (25+ years)	36,776	67%

Client type

Symptomatic cases	48,855	89%
Index cases	44,700	91%
Partners symptomatic	4,155	9%
Partners asymptomatic	6,082	11%

STI treatment history

Never treated for STI	40,891	74%
Previously treated for STI	14,046	26%
Old >3 months ago	9,656	69%
Recent ≤3 months ago	4,390	31%

STI syndromic diagnosis

GUD	10,406	18%
UD	13,256	23%
AVD	17,272	30%
Low risk	6,560	38%
High risk	10,712	62%
LAP	8,564	15%
SS	954	2%
BU	661	1%
BA	1,233	2%
NC	237	0%
Genital Warts	538	1%
Syphilis RPR VDRL	1,483	3%
Other STI	3,600	6%

STI partner notification

Total partner notification slips issued	18,380	100%
Total partners returned	10,237	56%
Total partners not seen	8,143	44%

STI site report

Malawi (national)

2015 Q2 (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	26,374	48%
HIV status ascertained	28,563	52%
HIV negative (new test)	21,610	76%
HIV positive	6,953	24%
New positive	1,547	22%
Previous positive	5,406	78%
Not on ART	1,146	21%
On ART	4,260	79%

STI clients referred for services

Lab	787	3%
Gynae review	480	2%
Surgical review	260	1%
Repeat HTC	19,767	83%
ART (for assessment)	1,052	4%
PMTCT	189	1%
Other (service referrals)	1,357	6%