

HIV MONITORING QUARTERLY REPORT

FOR NORTHERN HEALTH

FIRST QUARTER 2016

















Foreword

As part of the BC Centre for Excellence (BC-CFE) in HIV/AIDS's mandate to evaluate the outcomes of STOP HIV/AIDS programming in BC, we have developed quarterly HIV/AIDS monitoring reports. These reports provide up-to-date data on a variety of key HIV-related surveillance and treatment indicators. Selection of these indicators was achieved through a collaborative process with various Health Authority (HA) representatives. There are six reports in total, one for each HA and one for the province of BC as a whole. In addition, there is a technical report which explains how each HIV indicator is calculated. Data used in these reports come from the British Columbia Centre for Disease Control (BCCDC), MSP billings, hospitalization data from the Discharge Abstract Database, the Sunquest Laboratory database at the Provincial Public Health Microbiology and Reference Laboratory, Providence Health Care laboratory and the BC-CFE Drug Treatment Program (DTP) Database.

The objectives of these reports are to:

- 1. Provide timely HA-specific information on key HIV indicators which will guide and inform HIV leaders and innovators in the development of future HIV interventions and programs which will ultimately lead to decreasing the burden of HIV in BC. The indicators will reflect ongoing or past successful public health interventions and highlight areas in the HIV care spectrum which require further attention and support.
- 2. Highlight limitations in our current data due to incomplete or time lagged data and to develop future strategies to improve complete and timely data capture.

These reports are produced for the benefit of individual HA's. As such, we are enthusiastic about your involvement and cooperation regarding the development of these monitoring reports. Please forward your comments and queries to Irene Day, Director of Operations at the BC-CFE at iday@cfenet.ubc.ca.

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Acknowledgements and Contributions



British Columbia Centre for Excellence in HIV/AIDS (BC-CFE): The BC-CFE is responsible for the conception, preparation and ongoing review of this quarterly report. The BC-CFE provides the data and outputs for Indicators 5 (HIV Cascade of Care), 6 (Programmatic Compliance Score), 7 (New Antiretroviral Starts), 8 (CD4 Cell Count at ART Initiation), 9 (Active and Inactive Drug Treatment Program Participants), 10 (Antiretroviral Adherence Level), 11 (Resistance Testing Results by Resistance Category), 12 (AIDS-Defining Illness), and 13 (HIV-Related Mortality). The BC-CFE database provides PVL and CD4 cell count testing data, as well as ART use. All PVL measurements in BC are performed at the St Paul's Hospital virology laboratory, thus PVL data capture is 100%. An estimated 80% of all CD4 count measurements performed in the province are captured in the BC-CFE data holdings. The STOP HIV/AIDS Technical Monitoring Committee—BC-CFE is responsible for oversight of the monitoring report. James Nakagawa wrote, compiled, edited, and published this monitoring report. Paul Sereda, Dr. Viviane Lima and Nada Gataric perform analysis of Indicators 5–13. This report was conceived and guided by Dr. Julio Montaner.



British Columbia Centre for Disease Control (BCCDC): The BCCDC provides the data and outputs for Indicator 1 (HIV Testing Episodes), Indicator 2 (HIV Testing Rate), Indicator 3 (New HIV Diagnoses), Indicator 4 (Stage of HIV at Diagnosis) and Indicator 12 (AIDS-Defining Illness). The BCCDC is the single provincial agency that centralizes all HIV surveillance through the Public Health Microbiology and Reference Laboratory, which does more than 90% of all HIV screening tests in BC and all confirmatory testing. Theodora Consolacion and Dr. Jason Wong are responsible for outputs for Indicators 1–4.

Other Data Sources:

The above databases were supplemented with:

- (I) The BC Vital Statistics database which was used to calculate Indicator 5. The HIV Cascade of Care and Indicator 13. HIV-Related Mortality.
- (II) Linkage and preparation of the de-identified individual-level database used for calculating Indicator 5. The HIV Cascade of Care was facilitated by the British Columbia Ministry of Health.
- (III) The Statistics Canada database: BC and HIV-positive population counts were acquired through the statistics Canada website to calculate HIV-specific mortality rates for Indicator 13. HIV-Related Mortality.

Membership of the STOP HIV/AIDS Technical Monitoring Committee-BC-CfE

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The Seek and Treat for Optimal Prevention (STOP) HIV/AIDS BC Provincial Program: A Note on Monitoring and Interpreting HIV Indicators

The Seek and Treat for Optimal Prevention (STOP) of HIV/AIDS programme is a provincial initiative to improve HIV diagnosis and care delivery in BC through increased HIV-specific funding to all Health Service Delivery Areas (HSDA'S) across BC. The STOP provincial programme is an expansion of a four-year STOP pilot project which was implemented in two Health Service Delivery Areas in March 2010; the Vancouver HSDA which bears the largest burden of the HIV epidemic in the province and the Northern Interior HSDA which bears a high burden of HIV-related mortality. The STOP pilot project demonstrated the urgent need for improved efforts in early diagnosis of HIV and timely initiation of antiretroviral therapy (ART) initiation.

The expansion to a province-wide programme was announced on November 30th 2013 by the BC Ministry of Health with roll out of funding beginning on April 1st, 2013. This funding is intended to be used in the implementation and evaluation of HIV-related diagnosis and care initiatives within individual HA's. Goals of the project include: 1. A reduction in the number of new HIV infections in BC; 2. Improvements in the quality, effectiveness, and reach of HIV prevention services; 3. An increase in early diagnosis of HIV; 4. A reduction in AIDs cases and HIV-related mortality.

The goals of HA-led STOP-funded initiatives are to work toward achieving these goals. To these ends some outcome measures or indicators of progress have been drafted that should be considered in the design and implementation phases of these initiatives.

HIV Testing Episodes and Rates

In this section, the number of HIV test episodes and point of care (POC) HIV tests conducted each quarter in BC is shown. In general terms the goal is to increase the number of tests performed and to maximize testing efficiency. Test episodes are allocated by region according to where the test is performed.

Indicator 1. HIV Testing Episodes

Figure 1.1 HIV Test Episodes for Northern Health

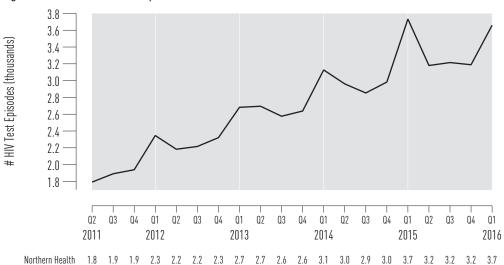


Figure 1.2 HIV Test Episodes by Gender for Northern Health 1,2

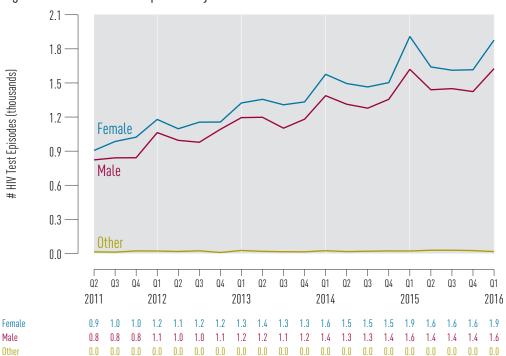


Figure 1.3 HIV Test Episodes by Age Category for Northern Health 1,2

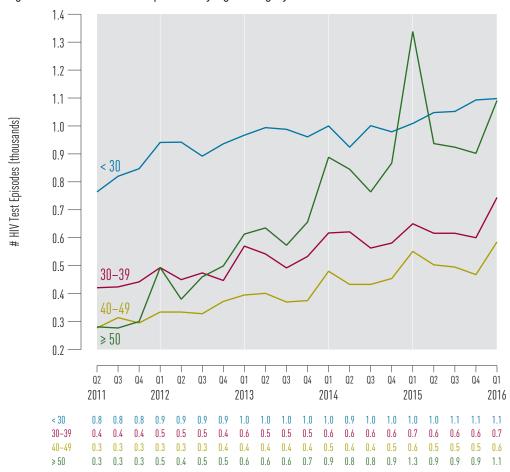
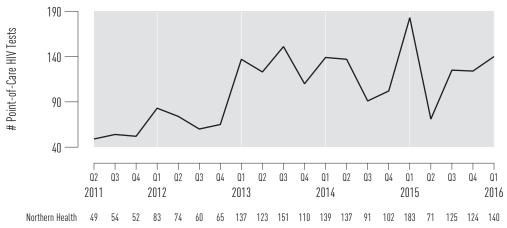


Figure 1.4 Point-of-Care HIV Tests for Northern Health



¹ Data Source: The BC Public Health Microbiology and Reference Laboratory (BCPHMRL) courtesy of the BC Centre for Disease Control (BCCDC).

Limitation: Repeat tests in individuals who test using various identifiers may not be identified and these individuals may be counted more than once.

2 Testing does not include point of care tests.

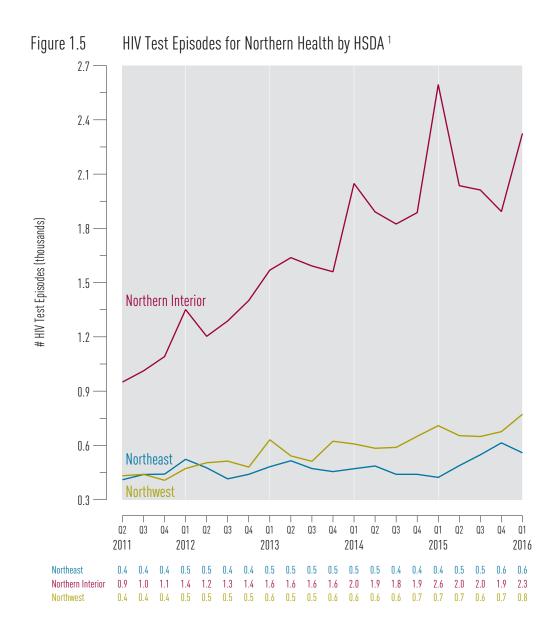
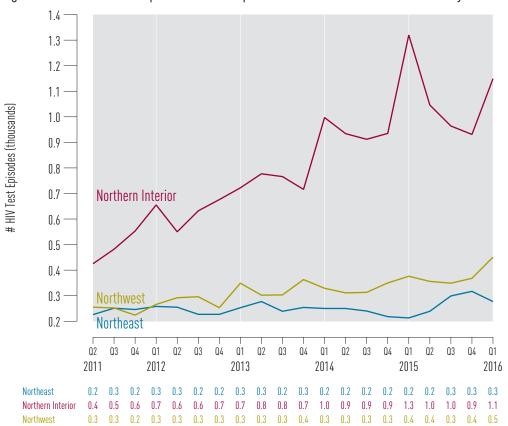
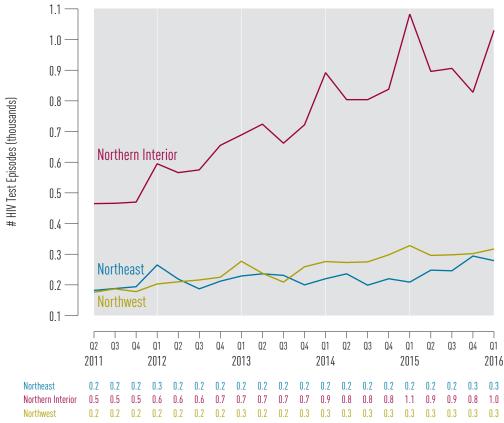


Figure 1.6 HIV Test Episodes for Non-prenatal Females in Northern Health by HSDA 1.2







Indicator 2. HIV Testing Rates

Figure 2.1 Rate of HIV Testing for Northern Health and HSDAs ²

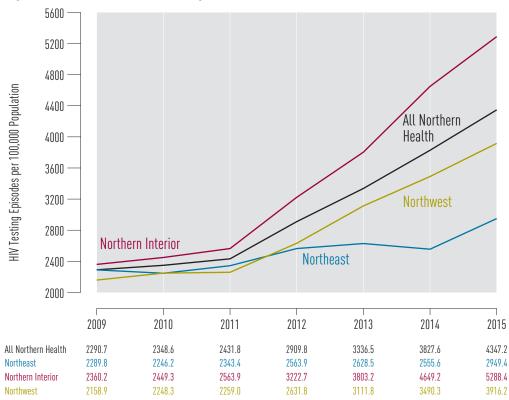
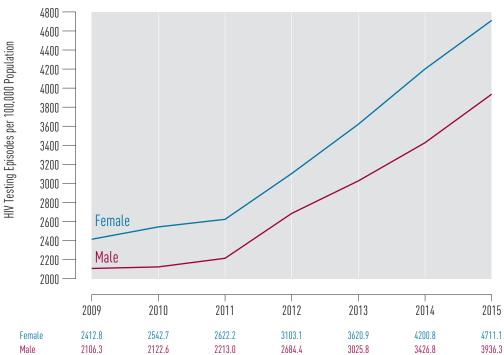


Figure 2.2 Rate of HIV Testing by Gender for Northern Health ²



6000 5400 4800 -30-39 4200 HIV Testing Episodes per 100,000 Population 3600 3000 40-49 2400 < 30 1800 ≥ 50 1200 2009 2010 2011 2012 2013 2014 2015 2400.7 2473.9 2611.1 2971.6 3126.1 3202.5 3492.0 < 30 30-39 4263.7 4336.5 4440.8 4754.5 5253.3 5811.3 6022.0 2670.1 1291.6 3217.5 1975.9 3614.2 2722.4 4326.1 3581.3 2541.9 1243.5 40-49 2448.1 5105.4 4361.9 1198.4

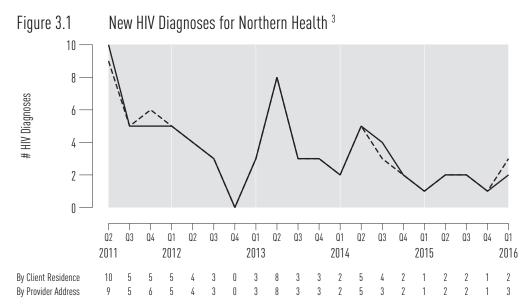
Figure 2.3 Rate of HIV Testing by Age Category for Northern Health ²

² Testing does not include point of care tests.

New HIV Diagnoses

Trends in HIV diagnoses by gender and exposure category are described. Interpreting HIV diagnoses must be done with consideration that trends are influenced by both changes in testing rate as well as changes in transmission rates. It is important to note that new HIV diagnoses cases and rates are not synonymous with HIV incidence as a person may have become infected with HIV long before they tested positive for HIV. However, as there is no reliable method for measuring HIV incidence, we follow trends in HIV diagnoses.

Indicator 3. New HIV Diagnoses



New HIV Diagnoses for Northern Health by Gender ³ Figure 3.2 Male **HIV Diagnoses** Female Q4 Q1 Q2 Q3 Q4 Q1 Q2 Q3 Q4 Q1 Q2 Q3 Q4 Q1 Q3 Q2 Q3 Q4 Q1 2011 2013 2014 2015 2012 2016 Female

Data Source: BCCDC When present, "By Provider Address" is graphed as dashed line in same colour.

Figure 3.3 New HIV Diagnoses for Northern Health by Age Category ³ # HIV Diagnoses < 30 30-39 ≥ 50 Q2 Q3 Q4 Q1 Q2 Q3 Q4 Q2 Q3 Q3 Q4 Q1 Q2 Q3 Q4 Q1 Q1 Q4 Q1 Q2 2011 2012 2013 2014 2015 2016 < 30 0 0 2 30-39 5 0 0 40-49 0 0 0 0 0 ≥ 50 3 0 0

Figure 3.4 New HIV Diagnoses for Northern Health by Exposure Category 3,4 # HIV Diagnoses **PWID** HET **MSM** Other NIR/Unknown Q3 Q3 Q4 Q2 Q4 Q1 Q2 Q3 Q1 Q2 Q1 Q2 Q3 Q4 Q1 Q2 2011 2012 2013 2014 2015 MSM (men who have sex with men) PWID (persons who inject drugs) 3 3 0 0 0 0 0 HET (heterosexual) 3 3 0 0 5 0 0 0 Other (other exposure identified) 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0

NIR/Unknown (no identified exposure)

New HIV Diagnoses for Northern Health by HSDA ³ Figure 3.5 8 # HIV Diagnoses Northern Interior 2 -Northeast Northwest Q2 Q3 Q4 Q2 Q4 Q2 Q3 Q4 Q1 Q3 Q4 Q1 Q2 Q3 Q4 Q2 Q3 Q1 Q1 Q1 2011 2012 2013 2014 2015 2016 Northeast By Provider Address 0 0 0 Northern Interior 3 By Provider Address 0 3 7 0 0 Northwest 0 2 By Provider Address

0 0

³ Data Source: BCCDC. When present, "By Provider Address" is graphed as dashed line in same colour.

⁴ MSM=men who have sex with men; PWID=people who inject drugs; HET=heterosexual. NIR=No identified risk/exposure.

Stage of HIV Infection at Diagnosis

Classification of stage of HIV infection, in the absence of information regarding recent testing history, is reliant on clinical information available at the time of diagnosis, including first CD4+ cell count and laboratory results suggestive of acute HIV infection (Table 1). The benefits of Treatment as Prevention (TasP) are maximized when antiretroviral therapy (ART) is initiated at high CD4 cell counts. Accordingly, it is preferable that individuals newly diagnosed with HIV be in the early stages of HIV infection (stage 0 or 1) to allow for early ART initiation.

N.B. Interpretation of Stage of HIV Infection at Diagnosis should proceed with caution. Early increases in diagnosis at late stage (i.e., low CD4 counts) may represent a "catching up" of previously missed long term infected individuals rather than a trend toward diagnosis at later stage of infection.

Indicator 4. Stage of HIV Infection at Diagnosis

Table 1 Staging Classifications of Infection at Time of HIV Diagnosis Based on CDC HIV Surveillance Case Definitions

Stage	Criteria									
0	Laboratory criteria met for acute HIV infection previous negative or indeterminate HIV test wi 180 days of first confirmed positive HIV test.									
1			CD4 ≥500							
2a			CD4 350-499							
2b	Stage 0 not met	and	CD4 200-349							
3	HOLHIEL		CD4 <200							
Unknown			No available CD4							
Updated 2016 Q1: AIDS diagnosis date is no longer used in this indicator.										

Figure 4.1 Stage of HIV Infection at Diagnosis for Northern Health, 2011–2015 5

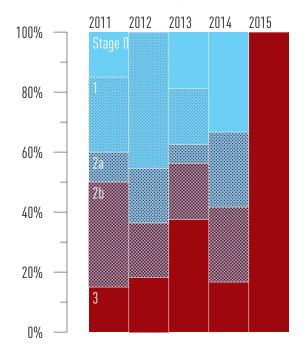
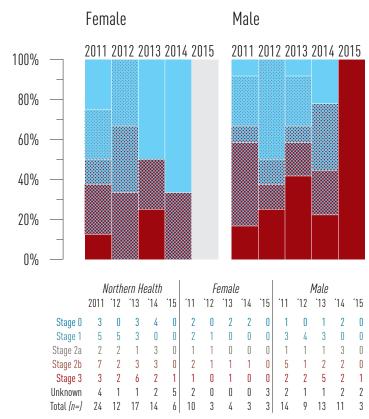


Figure 4.2 Stage of HIV Infection at Diagnosis by Gender for Northern Health, 2011–2015 ⁵



Data Source: BCCDC

Figure 4.3 Stage of HIV Infection at Diagnosis by Age Category for Northern Health, 2011–2015 ⁵

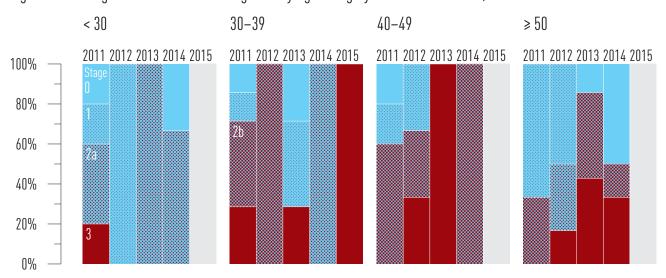
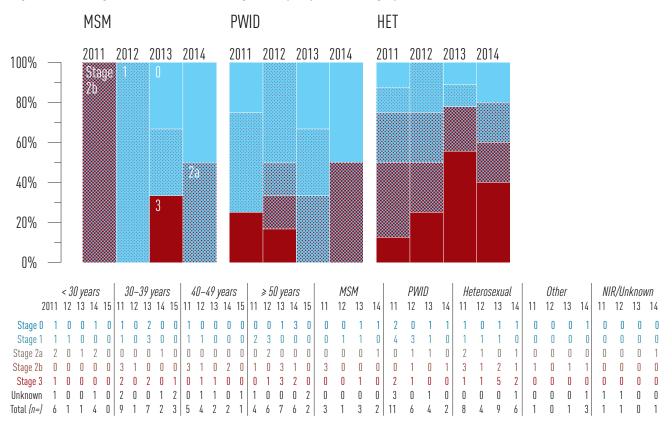


Figure 4.4 Stage of HIV Infection at Diagnosis by Exposure Category for Northern Health, 2011–2014 5.6



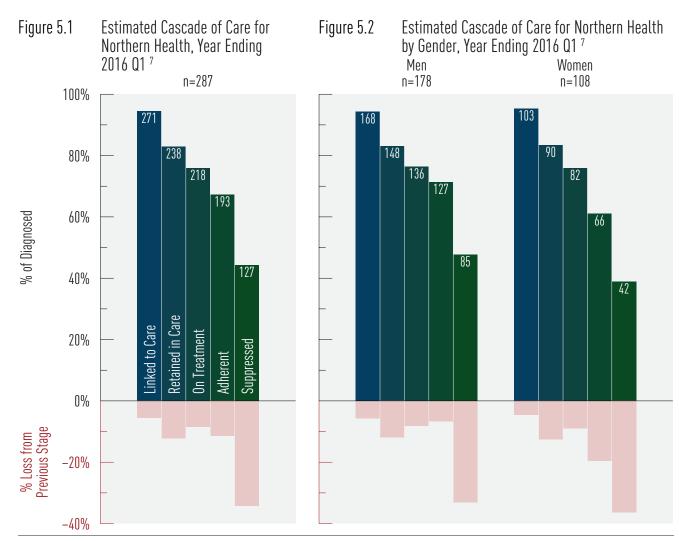
⁵ Data Source: BCCDC

⁶ MSM=men who have sex with men; PWID=people who inject drugs; HET=heterosexual. NIR=No identified risk/exposure.

HIV Cascade of Care

Indicator 5. HIV Cascade of Care

The success of seek, test, treat and retain (STTR) strategies like STOP is reliant on early diagnosis of HIV, linking newly diagnosed HIV-positive persons with ongoing care, retaining persons in HIV-care; initiating ART based on best evidenced practices and maintaining optimal ART adherence to ensure a suppressed viral load. These stages of HIV-care can be summarized as: 1. HIV diagnosis, 2. Linked to HIV care, 3. Retained in HIV care, 4. On ART, 5. Adherent to ART and 6. Achieving a suppressed VL; collectively, they are referred to as the cascade of care. Attrition between any of these stages of HIV-care means a reduction in the potential of ART as a benefit to the HIV-positive individual and as an HIV transmission prevention method on a population level. Thus, when interpreting trends in the cascade of care, we strive to see increases along each step of the cascade of care (i.e. reduced attrition) with the ultimate goal being 100% within each stage of the cascade. Monitoring the Cascade of Care provides a picture as to where deficiencies lie in the delivery and uptake of HIV-care. In this section we present the cascade of care for the period 2015 Q2–2016 Q1 in Northern Health and stratified by sex and age.

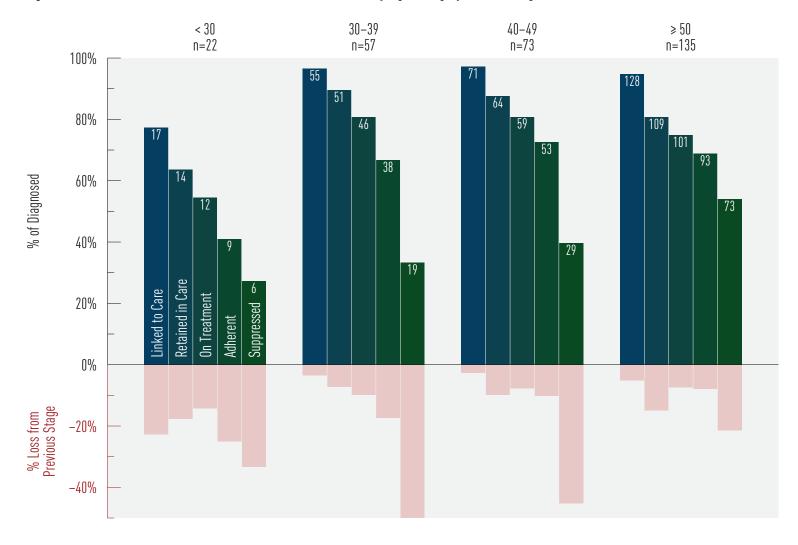


- 7 Data is for the period 2015 Q2-2016 Q1. Data Sources:
 - i British Columbia Centre for Excellence Drug Treatment Program (DTP) Database (ARV use, VL and CD4 count).
 - ii Administrative data (ex. MSP billings; hospitalization data from the Discharge Abstract Database (DAD)).

Limitations: HA assignment is based on the most recent HA of residence of the patient, if not available of the HIV-care provider. If the most recent HA of residence is not updated then the designated HA may be incorrect.

NB: Transgender have been assigned to their biological sex.





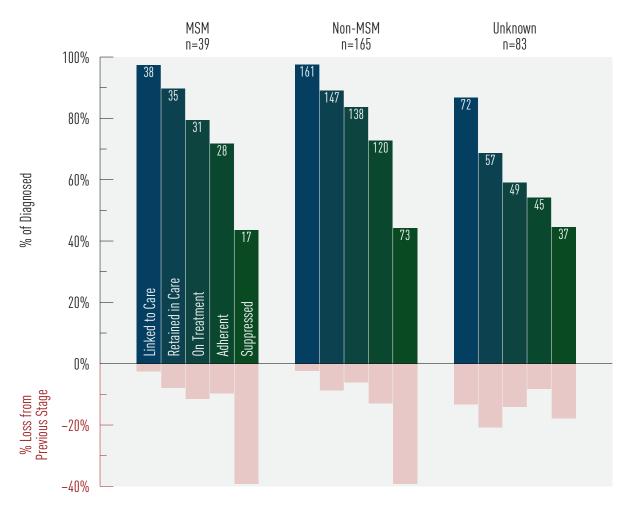
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ii Administrative data (ex. MSP billings; hospitalization data from the Discharge Abstract Database (DAD)).





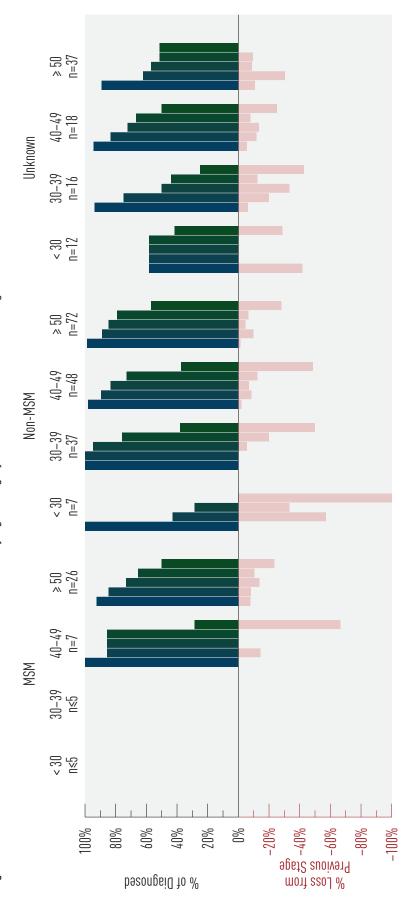
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ii Administrative data (ex. MSP billings; hospitalization data from the Discharge Abstract Database (DAD)).

Figure 5.5 Estimated Cascade of Care for Northern Health by Age Category and MSM Status, Year Ending 2016 Q1 9



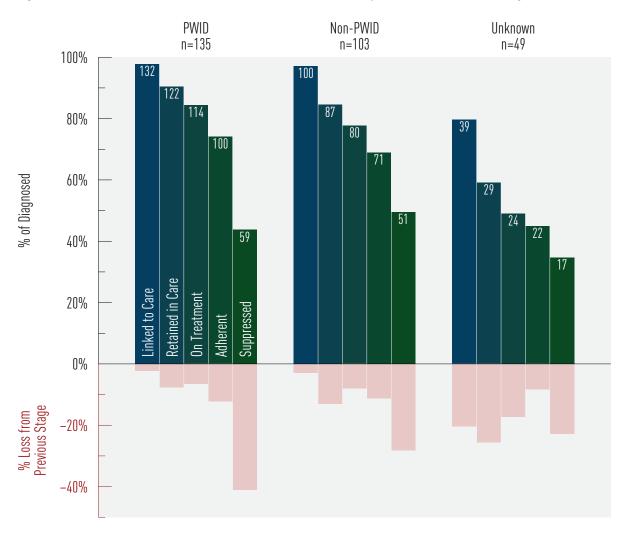
Data is for the period 2015 Q2-2016 Q1. Data Sources:

British Columbia Centre for Excellence Drug Treatment Program (DTP) Database (ARV use, VL and CD4 count).

Limitations: HA assignment is based on the most recent HA of residence of the patient, if not available of the HIV-care provider. If the most recent HA of residence is not updated then the designated HA may be incorrect.

ii Administrative data (ex. MSP billings; hospitalization data from the Discharge Abstract Database (DAD)).





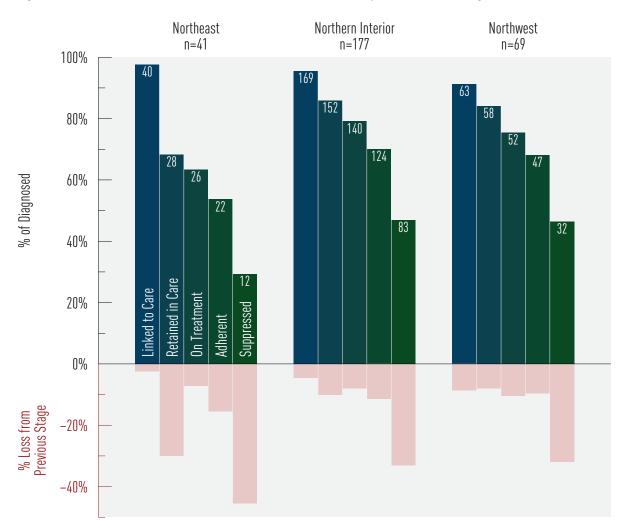
Limitations: HA assignment is based on the most recent HA of residence of the patient, if not available of the HIV-care provider. If the most recent HA of residence is not updated then the designated HA may be incorrect.

⁹ Data is for the period 2015 Q2-2016 Q1. Data Sources:

i British Columbia Centre for Excellence Drug Treatment Program (DTP) Database (ARV use, VL and CD4 count).

ii Administrative data (ex. MSP billings; hospitalization data from the Discharge Abstract Database (DAD)).





Limitations: HA assignment is based on the most recent HA of residence of the patient, if not available of the HIV-care provider. If the most recent HA of residence is not updated then the designated HA may be incorrect.

⁹ Data is for the period 2015 Q2-2016 Q1. Data Sources:

i British Columbia Centre for Excellence Drug Treatment Program (DTP) Database (ARV use, VL and CD4 count).

ii Administrative data (ex. MSP billings; hospitalization data from the Discharge Abstract Database (DAD)).

Programmatic Compliance Score

Indicator 6. Programmatic Compliance Score (PCS)

The Programmatic Compliance Score (PCS) is a summary measure of risk of future death, immunologic failure and virologic failure from all causes for people who are starting ART for the first time. It is composed of patient- and physician-driven effects. PCs scores range from o−6 with higher scores indicative of poorer health outcomes and greater risk of death. Table 1 provides mortality, immunologic failure and virologic failure probabilities for given PCs scores. We interpret an individual with a PCs≥4 as being 22 times more likely to die, almost 10 times more likely to have immunologic failure and nearly 4 times as likely to demonstrate virologic failure compared to those individuals with a PCs score of o. A detailed description of how the PCs score is calculated and its validation can be found in the technical report. In short, PCs scores are calculated by summing the results (yes=1, no=0) of six un-weighted non-performance indicators based on IAS−USA treatment guidelines:

- having <3 CD4 cell count tests in the first year after starting antiretroviral therapy (ART);
- 2. having <3 plasma viral load (VL) tests in the first year after starting ART;
- 3. not having drug resistance testing done prior to starting ART;
- 4. starting on a non-recommended ART regimen;
- 5. starting therapy with CD4<200 cells/μL; and
- 6. not achieving viral suppression within 9 months since ART initiation.

In this section we provide PCS scores and their components over time for the province of BC. A decline to 0%, (i.e., all individuals having a score of o) is the eventual goal.

Table 2. Probability of Mortality, Immunologic Failure and Virologic Failure based on the Programmatic Compliance Score

Programmatic Compliance Score	Mortality Risk Ratio (95% Confidence Interval)	Immunologic Failure Risk Ratio (95% CI)	Virologic Failure Risk Ratio (95% CI)
·			
O (Best score)	1 (-)	1 (-)	1 (-)
1	3.81 (1.73-8.42)	1.39 (1.04–1.85)	1.32 (1.05–1.67)
2	7.97 (3.70–17.18)	2.17 (1.54–3.04)	1.86 (1.46–2.38)
3	11.51 (5.28–25.08)	2.93 (1.89-4.54)	2.98 (2.16–4.11)
4 or more (Worst score)	22.37 (10.46–47.84)	9.71 (5.72–16.47)	3.80 (2.52–5.73)

Reference: Lima VD, Le A, Nosyk B, Barrios R, Yip B, et al. (2012) Development and Validation of a Composite Programmatic Assessment Tool for HIV Therapy. PLoS ONE 7(11): e47859. doi:10.1371/journal.pone.0047859

Figure 6.1 PCS Components for Northern Health, 2014 Q2-2016 Q1 10

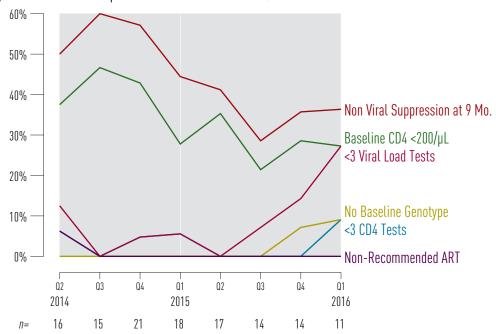
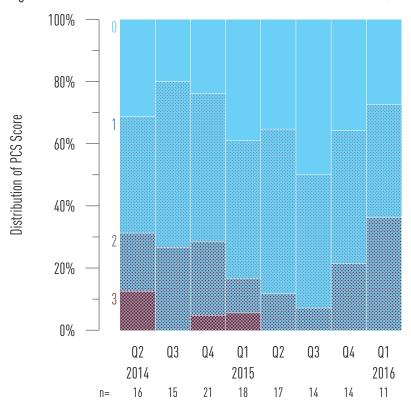


Figure 6.2 Historical Trends for PCS Score for Northern Health, 2014 Q2-2016 Q1 10,11



Data Source: British Columbia Centre for Excellence Drug Treatment Program (DTP) Database. Limitations: CD4 cell count capture is approximately 80%.

Each quarter's data is calculated as the sum of the 4 quarters leading up to it. e.g. 2013 Q1 is calculated from 2012 Q2 – 2013 Q1. NB: A score of 0 is the best score and a score of 4 or more is the worst score.

Antiretroviral Uptake

In this section we present trends in ART uptake, the number and proportion of new HIV treatment initiations and the number of active and inactive DTP participants. Trends in ART uptake should be interpreted under the consideration of changing BC HIV treatment guidelines. BC HIV treatment guidelines are updated regularly by the BC-CFE Therapeutic Guidelines Committee and reflect those of the International AIDS Society. Most recent changes were made in 2012 and HIV treatment is now recommended for all HIV-positive adults regardless of CD4 cell count; as evidence demonstrates that early initiation of HIV treatment maximizes both the individual's health outcomes as well as the potential of ART as a form of HIV transmission prevention at a population level. As such, trends in the number and proportion of persons on ART and new ART starts (in both naïve and experienced persons) are expected to increase over time at higher CD4 cell counts.

Indicator 7. New Antiretroviral Therapy Starts in Northern Health

Figure 7 BC-CfE Drug Treatment Program Enrollment: New ART Participants in Northern Health, 2014 Q2-2016 Q1 12



Indicator 8. CD4 Cell Count at ART Initiation

Figure 8 CD4 Cell Count at ART Initiation of ART-Naïve DTP Participants in Northern Health, 2014 Q2-2016 Q1 13

The majority of cells in this figure have $n \le 5$, which is considered statistically insignificant as well as a possible risk to patient privacy. For this reason, this figure has been omitted. Authorized parties may contact the British Columbia Centre for Excellence in HIV/AIDS to obtain this information.

Data Source: Drug Treatment Program Database

Limitation: DTP participants are designated to an HA based on most current residence provided by the participant.

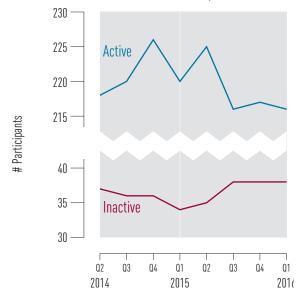
¹³ Data Source: Drug Treatment Program Database Limitations: CD4 cell count data is approximately 80% complete.

Indicator 9. Active and Inactive DTP Participants

Table 3. Distribution of People on ART for Northern Health, 2016 Q1 14

Age	< 30	10
	30-39	45
	40-49	64
	≥ 50	97
Gender	Male	136
	Female	80
Exposure	MSM	31
	PWID	113
Total		216

Figure 9 Active and Inactive DTP Participants for Northern Health, 2014 Q2-2016 Q1 ¹⁵



14 Data Source: Drug Treatment Program Database
Limitation: DTP participants are designated to an HA based
on most current residence provided by the participant.

Definition:

'On antiretroviral therapy' defined as being on treatment in the current quarter

15 Active DTP participants: An individual who has had medication prescribed at least once in the preceding quarter. Inactive DTP participants: Persons no longer prescribed drugs through the HIV/AIDS Drug Treatment Program in the last quarter.

Antiretroviral Adherence Level

In this section we present trends in prescription refill adherence levels for individuals in their first year of treatment. Given that the benefits of ART are compromised in the presence of imperfect ART adherence, we expect to see the proportion of persons on ART achieving near perfect adherence (ie. $\geq 95\%$) to increase with time. Furthermore, it is important that trends in the proportion of ART users achieving prescription refill adherence of $\geq 95\%$ keep pace with new ART starts and increase among those continuing on ART.

Indicator 10. Antiretroviral Adherence

Figure 10 Distribution of Individuals by Adherence Level in 1st Year of Therapy, Based on Pharmacy Refill Compliance for Northern Health, 2014 Q2–2016 Q1 ¹⁶

The majority of cells in this figure have $n \le 5$, which is considered statistically insignificant as well as a possible risk to patient privacy. For this reason, this figure has been omitted. Authorized parties may contact the British Columbia Centre for Excellence in Hiv/Aids to obtain this information.

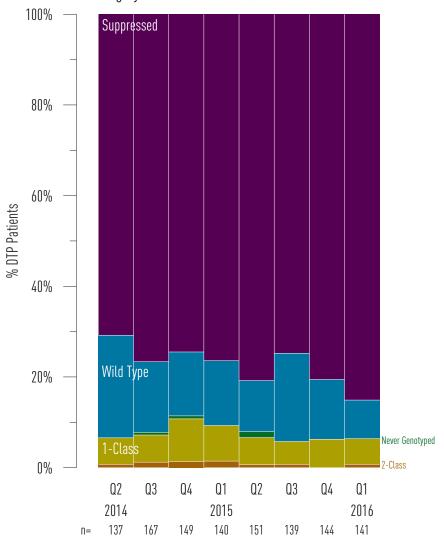
¹⁶ Data Source: Drug Treatment Program Database Limitation: Prescription refill adherence is used as a proxy for patient adherence.

Resistance Testing and Results

Indicator 11. Resistance Testing and Results

In this section, we present trends in cumulative resistance testing by resistance category: Suppressed (where a DTP participant's viral load is too low to be genotyped); Wild Type (where no HIV treatment resistances were discovered), Never Genotyped, and Resistances to one, two, three, or four HIV treatment classes. Resistance testing prior to ART initiation is recommended in the BC HIV treatment primary care guidelines. Thus, it is expected that trends over time should find all persons enrolled in the DTP to have been genotyped. Trends over time should also show an increase in the proportion of DTP participants achieving a suppressed status and an increase in resistance testing should not lead to an increase in the number of ART resistances occurring.





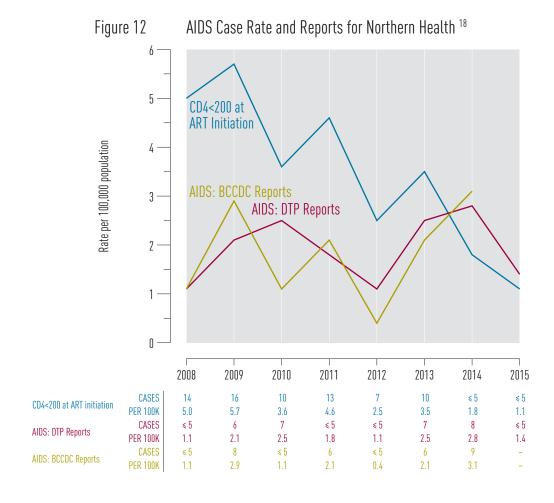
¹⁷ Data Source: Drug Treatment Program Database

Limitation: DTP participants are designated to a HA based on most current residence provided by the participant.

AIDS-Defining Illness

Indicator 12. AIDS-Defining Illness

Improvements in ART and the expansion of ART province-wide has led to very low numbers of recorded AIDS cases across BC. However, interpreting trends in AIDS cases is challenging as AIDS reporting is passive in BC and it is likely that they are under-reported across all Health Authorities. In addition to under-reporting, methods of reporting AIDS cases are inconsistent across HA's and do not truly reflect the current reality of new AIDS diagnoses. Efforts will need to be made to improve under- and inconsistent reporting of AIDS cases across all HA's. The table below shows AIDS cases using three definitions. First, AIDS cases were defined as the number of physician-reported AIDS defining illness (ADI) in a given year. AIDS case reporting is a passive process and physicians can voluntarily report AIDS cases to the BCCDC or DTP. As such, we have plotted both BCCDC reports and DTP reported AIDS cases. We also show the proportion of persons initiating ART with a CD4<200 cells/µL.



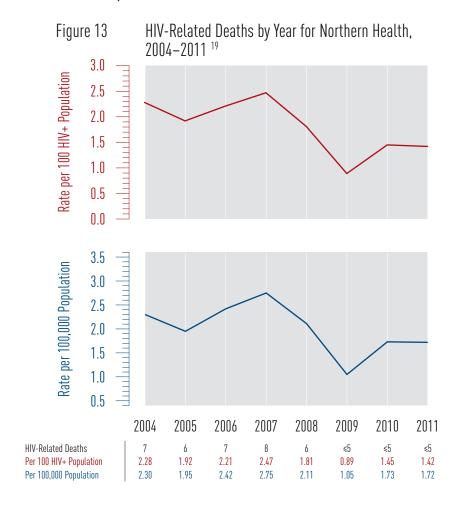
Data Source: DTP AIDS cases are obtained from the Drug Treatment Program Database; BCCDC AIDS cases are obtained from the BCCDC; CD4<200 at ART initiation data came from the DTP database.

Limitation: AIDS case reporting was investigated using 3 definitions: First, using AIDS cases reported in AIDS case report forms from the DTP; Second, using AIDS cases reported via the BCCDC and third, using a CD4 cell count of <200 cells/µL at time of ART initiation using DTP data. AIDS case reporting is passive in BC, thus; AIDS case reporting is not well captured. The DTP sends out AIDS reporting forms to physicians annually. The BCCDC uses DTP AIDS case reports as well as physician AIDS case reports made directly to the BCCDC. Interpreting AIDS case reports should be done with these limitations in mind. AIDS data is updated annually as very few AIDS cases reports are reported in general and trends would be difficult to notice if reported quarterly.

HIV-Related Mortality

Indicator 13. HIV-Related Mortality

Evidence indicates that individuals who initiate treatment with recommended ART in a timely fashion may live near normal lifespans. Excess mortality among HIV positive persons is, therefore, an important measure of HIV care with a goal of minimizing HIV-related mortality in British Columbia.



Limitation:

¹⁹ Data Source: BC Vital Statistics

^{1.} DTP participants are designated to an HA based on most current residence provided by the participant.

^{2.} Mortality data is updated annually.

^{3.} The most recent available data was used.

Appendices

Indicator 1 Episodes (: Test thousands)	<u> </u>	Q3	Q4	2012 Q1	Q2	Q3		2013 Q1	Q2	Q3	Q4		Q2	Q3	Q4	2015 Q1	Q2			Q4	2016 Q1
Northern H	Iealth	1.8	1.9	1.9	2.3	2.2	2.2	2.3	2.7	2.7	2.6	2.6	3.1	3.0	2.9	3.0	3.7	3.2	3.	.2	3.2	3.7
Gender	Female	0.9	1.0	1.0	1.2	1.1	1.2	1.2	1.3	1.4	1.3	1.3	1.6	1.5	1.5	1.5	1.9	1.6	1.	.6	1.6	1.9
	Male	0.8	0.8	0.8	1.1	1.0	1.0	1.1	1.2	1.2	1.1	1.2	1.4	1.3	1.3	1.4	1.6	1.4	1.	.4	1.4	1.6
	Other	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.	.0	0.0	0.0
Age	< 30	0.8	0.8	0.8	0.9	0.9	0.9	0.9	1.0	1.0	1.0	1.0	1.0	0.9	1.0	1.0	1.0	1.0	1.	.1	1.1	1.1
	30-39	0.4	0.4	0.4	0.5	0.5	0.5	0.4	0.6	0.5	0.5	0.5	0.6	0.6	0.6	0.6	0.7	0.6	0.	.6	0.6	0.7
	40-49	0.3	0.3	0.3	0.3	0.3	0.3	0.4	0.4	0.4	0.4	0.4	0.5	0.4	0.4	0.5	0.6	0.5	0.	.5	0.5	0.6
	≥ 50	0.3	0.3	0.3	0.5	0.4	0.5	0.5	0.6	0.6	0.6	0.7	0.9	0.8	0.8	0.9	1.3	0.9	0.	.9	0.9	1.1
POC HIV 7		49	54	52	83	74	60	65	137	123	151	110	139	137	91	102	183	71	12	25 1	124	140
Northeast		0.4	0.4	0.4	0.5	0.5	0.4	0.4	0.5	0.5	0.5	0.5	0.5	0.5	0.4	0.4	0.4	0.5	0	.5	0.6	0.6
Female		0.2	0.3	0.2	0.3	0.3	0.2	0.2	0.3	0.3	0.2	0.3	0.2	0.2	0.2	0.2	0.2	0.2	0	.3	0.3	0.3
Male		0.2	0.2	0.2	0.3	0.2	0.2	0.2	0.2	0.2	0.2	0.2	0.2	0.2	0.2	0.2	0.2	0.2	0	.2	0.3	0.3
Northern I	nterior	0.9	1.0	1.1	1.4	1.2	1.3		1.6	1.6	1.6	1.6	2.0	1.9	1.8	1.9	2.6	2.0			1.9	2.3
Female		0.4	0.5	0.6	0.7	0.6	0.6		0.7	0.8	0.8	0.7	1.0	0.9	0.9	0.9	1.3	1.0			0.9	1.1
Male		0.5	0.5	0.5	0.6	0.6	0.6		0.7	0.7	0.7	0.7	0.9	0.8	0.8	0.8	1.1	0.9			0.8	1.0
Northwest		0.4	0.4	0.4		0.5	0.5		0.6	0.5	0.5	0.6		0.6	0.6	0.7	0.7	0.7			0.7	0.8
Female		0.3	0.3	0.2	0.3	0.3	0.3		0.3	0.3	0.3	0.4		0.3	0.3	0.3	0.4	0.4			0.4	0.5
Male		0.3	0.3	0.2		0.3	0.3		0.3	0.3	0.3	0.4	0.3	0.3	0.3	0.3	0.4	0.3	0.		0.4	0.3
						0.2	0.2	0.2	0.3	0.2	0.2	0.3	0.3	0.3	0.3	0.3	0.3	0.3	0.	.5	0.3	0.5
Indicator 2	: Rate of H	IV Testing	per 10 200	-		10		2011		201	2	20	013	2	2014		2015	5				
All Norther	n Health		2290	.7	234	8.6	2	431.8		2909.	8	333	6.5	38	27.6		4347.2	2				
Northeast			2289	.8	224	6.2	2	343.4		2563.	9	262	8.5	25	55.6		2949.4	1				
Northern Ir	nterior		2360.2			2449.3 2563.9				3222.7 3803.			803.2 4649.2			5288.4						
Northwest			2158.9			2248.3 2259.0					2631.8			3111.8 3490.3			3916.2					
Gender	Female		2412.8			2.7	2622.2		3103.1		3620.9		4200.8			4711.1	l					
	Male		2106.3			2122.6 2213.0				2684.4			3025.8		3426.8		3936.3					
Age	< 30		2400	.7	247	3.9	2	611.1		2971.	6	312	6.1	32	02.5		3492.0)				
0	30-39		4263	.7	433	6.5	4	440.8		4754.	5	525	3.3	58	11.3		6022.0)				
	40-49		2448		4550.5 2541.9			670.1		3217.5		3614.2		4326.1		5105.4						
	≥ 50		1198					291.6 1975.9				272		3581.3			4361.9					
	= 30		1170		2011	J.J)12			2013	2,2	.2.1	201)15				201
Indicator 3	: New HIV	Diagnoses			Q2	Q3 (2 Q3			Q2_(Q3 Q		Q2	Q3	Q4 (Q2	Q3		
Northern H	Iealth	By Client	Resid	ence	10	5	5	5	4 3	0	3	8	3	3 2	5	4	2	1	2	2	1	2
		By Provid	er Ada	dress	9	5	6	5	4 3	0	3	8	3	3 2	5	3	2	1	2	2	1	3
Gender		Female			5	1	3	1	2 0	0	0	3	1	0 1	0	1	1	0	0	2	1	2
		Male			5	4	2	4	2 3	0	3	5	2	3 1	5	3	1	1	2	0	0	(
Age		< 30			4	1	2	1	0 0	0	0	2	0	0 0	1	2	1	0	0	0	0	2
		30-39			4	1	2		1 0	0	0	5		1 1		0	1	0	2	1	0	
		40-49			1	2	0		2 2		1	0		1 1		0	0	1	0	0	0	
		≥ 50			1	1	1		1 1		2	1		1 0		2	0	0	0	1	1	(
Exposure		MSM			1	1	1		0 1	0	1	1		1 0		1	0	0	0	0	_	
-I - 2 - 2 - 2		PWID			5	1	3		2 1	0	0	2		2 1		0	1	0	0	0	_	
		HET			3	3	1		2 0		2	5		0 1		1	1	0	1	0	_	
		Other			0	0	0		0 0		0	0		0 0		1	0	0	0	0		
			nour																		_	
		NIR/Unk	nown		1	0	0	0	0 1	0	0	0	U	0 0	0	1	0	1	1	2	_	_

Indicator 3	: Nev	w Hl							Q3	Q4					21 Q		Q4)15)1 Q			
Northeast				•			idenc <i>ddres</i>				0 0	0	0 0	0		$\begin{array}{ccc} 0 & 0 \\ 0 & 0 \end{array}$		0 0	0	0 0	1 1	0		$egin{array}{ccc} 1 & 0 \\ 1 & 0 \end{array}$	0
Northern Ir	nterio	or		-			idenc				2	4	2	0		6 2		1	2	3	1	1		1 1	2
				-			ddres		4	4	2	4	2	0		7 2		1	3	3	1	1		1 1	3
Northwest				•			idenc <i>ddres</i>				3	0 0	1 1	0		2 1 1 1	0	1 1	3 2	1 0	0	0		$\begin{array}{ccc} 0 & 0 \\ 0 & 0 \end{array}$	0
Indicator 4:	Stag	ge of		_																					
	No	orthe	rn H	Iealt	h		Fen	nale			Ma		.			years				years				years	
Stage 0	3	12	13 3	$\frac{14}{4}$	15	2	0	$\frac{3 \ 14}{2 \ 2}$		11 `		3 `14 1 2		11	0	13 '14 0 1	15	11 1	$\frac{12}{0}$	$\frac{13 \ 14}{2}$		11	$\frac{12}{0}$	$\frac{.3}{0}$ $\frac{.14}{0}$	
Stage 1	5	5	3	0	0	2		0 0		3		3 0		1	1	0 0		1	0	3 0		1	1	0 0	
Stage 2a	2	2	1	3	0	1	1	0 0	0	1	1	1 3	0	2	0	1 2	0	0	0	0 1	. 0	0	0	0 0	0
Stage 2b	7	2	3	3	0	2	1	1 1	0	5	1	2 2	0	0	0	0 0	0	3	1	0 0	0	3	1	0 2	0
Stage 3	3	2	6	2	1	1	0	1 0	0	2	2	5 2	1	1	0	0 0	0	2	0	2 0) 1	0	1	1 0	0
Unknown	4	1	1	2	5	2	0	0 0	3	2	1	1 2	2	1	0	0 1	0	2	0	0 1	. 2	0	1	1 0	1
Total	24	12	17	14	6	10	3	4 3	3	14	9 1	3 11	3	6	1	1 4	0	9	1	7 2	2 3	5	4	2 2	1
	'11		0 yea '13		'1 E	' 11	MS '12	SM '13	' 14	' 11	PW.	ID '13	' 14	11 11	Hetero	sexua '13	l '14	Oth '11	er E '12	xposu '13	re '14	N] '11	R/U1 12	nknow '13	vn '14
Stage 0	0	0	1	3	0	0	0	13	1	2	0	1	1	1	0	13	1	0	0	0	1	0	0	0	0
Stage 1	2	3	0	0	0	0	1	1	0	4	3	1	0	1	1	1	0	0	0	0	0	0	0	0	0
Stage 2a	0	2	0	0	0	0	0	0	1	0	1	1	0	2	1	0	1	0	0	0	0	0	0	0	1
Stage 2b	1	0	3	1	0	3	0	0	0	0	1	0	1	3	1	2	1	1	0	1	1	0	0	0	0
Stage 3	0	1	3	2	0	0	0	1	0	2	1	0	0	1	1	5	2	0	0	0	0	0	0	0	0
Unknown	1	0	0	0	2	0	0	0	0	3	0	1	0	0	0	0	1	0	0	0	1	1	1	0	0
Total	4	6	7	6	2	3	1	3	2	11	6	4	2	8	4	9	6	1	0	1	3	1	1	0	1
Indicator 5			scad	le of	Ca	re		DIAG			:	LINKI		R	ETAIN			ON A		AI	OHER		SU	PPRES	
Northern F									28				71		2	238		2	218			193			127
Age Catego	ory	< 30								22			17			14			12			9			6
		30-3								57			55			51			46			38			19
		$40-4$ ≥ 50							13	73			71 28			64 109		1	59 .01			53 93			29 73
Aga Cataga		≥ SU MSI				30				5			28 5			109 ≤ 5			.01 ≤ 5			93 ≤ 5			/3 ≤ 5
Age Catego and MSM	и у	10131	VI			. 30 0–39				5			5			≤ 5 ≤ 5			≤ 5 ≤ 5			≤ 5 ≤ 5			≤ 5 ≤ 5
Status						0-39 0-49			_	7		_	7			≥ <i>5</i>		-	≤ <i>5</i>			≥ <i>3</i>			≥ 3 2
						: 50			2	26			24			22			19			17			13
		Non	-MS	M		30				7			7			3			2			0			0
						0-39			3	37		1	37			37			35			28			14
					4	0-49			4	18		4	17			43			40			35			18
																						57			41
					≥	50				72			71			64			61			3/			41
		Unk	now	n					7	72						64 7			61 7			7			5
		Unk	now	n	<	50			7 1			7	71												
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				n	3	50 30 0–39			7 1 1 1 3	12 16 18 37			71 7 15 17 33			7 12 15 23			7 8 13 21			7 7 12 19			5 4 9 19
Gender		Mal	e	n	3	50 30 0–39 0–49			7 1 1 1 3 17	12 16 18 37		10	71 7 15 17 33]	7 12 15 23 148		1	7 8 13 21 36			7 7 12 19 127			5 4 9 19 85
		Male Fem	e ale	n	3	50 30 0–39 0–49			7 1 1 3 17 10	12 16 18 37 78		10	71 7 15 17 33 68			7 12 15 23 148 90		1	7 8 13 21 36 82			7 7 12 19 127 66			5 4 9 19 85 42
Injection		Male Fem	e ale ID		3	50 30 0–39 0–49			77 11 13 33 177 100 13	12 16 18 37 78 08		10 11	71 7 15 17 33 68 03			7 12 15 23 148 90		1	7 8 13 21 36 82 14			7 7 12 19 127 66 100			5 4 9 19 85 42 59
		Male Fem PW	e ale ID -PW	'ID	3	50 30 0–39 0–49			77 11 13 17 10 13 10	12 16 18 37 78 08 35		10 10 10 10	71 7 15 17 33 68 03 32			7 12 15 23 148 90 122 87		1	7 8 13 21 36 82 14 80			7 7 12 19 127 66 100 71			5 4 9 19 85 42 59 51
Injection Drug Use		Male Fem PWI Non Unk	e ale ID -PW now	'ID	3	50 30 0–39 0–49			77 11 13 177 100 133 100 44	12 16 18 37 78 08 35 03		10 10 13 10	71 7 15 17 33 68 03 32 00			7 12 15 23 148 90 122 87 29		1	7 8 13 21 36 82 14 80 24			7 7 12 19 127 66 100 71 22			5 4 9 19 85 42 59 51 17
Injection	.s	Male Fem PW! Non Unk MS!	e ale ID -PW now M	TID n	3	50 30 0–39 0–49			77 11 13 17 10 13 10 4	12 16 18 37 78 08 35 03 19		10 10 13 10	71 7 15 17 33 58 03 32 00 39		1	7 12 15 23 148 90 122 87 29 35		1	7 8 13 21 36 82 14 80 24 31			7 7 12 19 127 66 100 71 22 28			5 4 9 19 85 42 59 51 17
Injection Drug Use	.s	Male Fem PWI Non Unk MSN Non	e ale ID -PW now M	TID n	3	50 30 0–39 0–49			17 10 13 10 13 10 4	12 16 18 37 78 08 35 03 19 39		10 10 13 10 3	71 7 15 17 33 68 03 32 00 39 38		1	7 12 15 23 148 90 122 87 29 35		1 1	7 8 13 21 36 82 14 80 24 31 38			7 7 12 19 127 66 100 71 22 28 120			5 4 9 19 85 42 59 51 17 17 73
Injection Drug Use MSM Statu	.s	Male Fem PWI Non Unk MSN Non Unk	e ale ID -PW now M -MS now	TID n M n	3	50 30 0–39 0–49			10 13 17 10 13 10 4 3 16	12 16 18 37 78 98 95 93 94 95 95 93 93 93 93 93 93 93 93 93 93 93 94 95 95 95 95 95 95 95 95 95 95 95 95 95		10 10 10 10 3 10	71 7 7 115 117 115 117 117 117 117 117 1		1	7 12 15 23 148 90 122 87 29 35 147 57		1 1	7 8 13 21 36 82 14 80 24 31 38 49			7 7 12 19 127 66 100 71 22 28 120 45			5 4 9 19 85 42 59 51 17 73 37
Injection Drug Use MSM Statu Health	s	Male Fem PWI Non Unk MSI Non Unk	e ale ID -PW now M -MS now	TID n M n	< 33 44 ≥ ≥	50 30 0–39 0–49 50			77 11 11 12 17 10 13 11 10 44 43 16 88 84	12 16 18 37 78 08 35 03 19 39 55 33		10 10 13 10 10 10 10 10 10 10 10 10 10 10 10 10	71 7 7 115 115 117 115 117 117 117 117 1		1	7 12 15 23 148 90 122 87 29 35 147 57 28		1 1	7 8 13 21 36 82 14 80 24 31 38 49 26			7 7 12 19 127 66 100 71 22 28 120 45 22			5 4 9 19 85 42 59 51 17 73 37 12
Injection Drug Use MSM Statu	S	Mald Fem PW! Non Unk MSN Non Unk Nor	e ale ID -PW now M -MS now	rIID n M n st	< 33 44 ≥ ≥	50 30 0–39 0–49 50			77 11 13 13 17 10 13 11 10 4 4 4 4 17	12 16 18 37 78 08 35 03 19 39 55 33		10 10 10 10 10 10 10 10 10 10 10 10 10 1	71 7 7 115 117 115 117 117 117 117 117 1		1	7 12 15 23 148 90 122 87 29 35 147 57		1 1 1	7 8 13 21 36 82 14 80 24 31 38 49			7 7 12 19 127 66 100 71 22 28 120 45			5 4 9 19 85 42 59 51 17 73 37

Indicator 6: Program	matic Com	pliance Sco	re (PCS)								
		2014 Q2	Q3		Q4	2015 Q1	Q2	(Q3	Q4	2016 Q1
< 3 CD4 Tests		6.2%	0.0%		4.8%	5.6%	0.0%	0.0		0.0%	9.1%
< 3 Viral Load Tests		12.5%	0.0%		4.8%	5.6%	0.0%	7.1		14.3%	27.3%
No Baseline Genotyp		0.0%	0.0%		0.0%	0.0%	0.0%	0.0		7.1%	9.1%
Baseline CD4 < 200 c		37.5%	46.7%		2.9%	27.8%	35.3%	21.4		28.6%	27.3%
Non-Recommended		6.2%	0.0%		0.0%	0.0%	0.0%	0.0		0.0%	0.0%
Non Viral suppressio	n at 9 Mo.	50.0%	60.0%	5	57.1%	44.4%	41.2%	28.6		35.7%	36.4%
PCS Score: 0		5	3		5	7	6		7	5	3
PCS Score: 1		6	8		10	8	9		6	6	4
PCS Score: 2		3	4		5	2	2		1	3	4
PCS Score: 3		2	0		1	1	0		0	0	0
PCS Score: 4 or more		0	0		0	0	0		0	0	0
Total (n=)		16	15		21	18	17		14	14	11
Indicator 7: New DT	P ARV Parti										
First Starts		2	4		5	3	4		3	4	1
Experienced Starts		8	6		2	3	5		4	4	6
Indicator 8: CD4 Cel	l Count at A	RT Initiati	on for ARV	-Naïve	DTP Parti	icipants					
CD4 ≥ 500		-	-		-	-	_		_	_	-
CD4 350-499		_	_		_	_	_		_	_	_
CD4 200-349		_	_		_	_	_		_	_	_
CD4 50-199		_	_		_	_	_		_	_	_
CD4 < 50		_	_		_	_	_		_	_	_
CD4 Median (cells/µL	(,)	_	_		-	-	_		_	-	_
Total (n=)		≤ 5	≤ 5		≤ 5	≤ 5	≤ 5	<u>≤</u>	5	≤ 5	≤ 5
Indicator 9: Active ar	nd Inactive I	OTP Partici	pants								
Active DTP Participa		218	220		226	220	225	2.	16	217	216
Inactive DTP Particip		37	36		36	34	35		38	38	38
mactive D11 Turtien	Juito	3,	30		30	31	33	•	<i>.</i>	30	30
Indicator 10: Antiret	roviral Adh	erence									
≥ 95%		_	5		5	_	-		-	_	_
80% to < 95%		-	2		2	_	-		-	-	-
40% to < 80%		_	0		0	_	-		-	_	_
< 40%		-	0		0	-	-		-	-	-
Total (n=)		≤ 5	7		7	≤ 5	≤ 5	\leq	5	≤ 5	≤ 5
Indicator 11: Resistar	nce Testing a	and Results									
Suppressed		97	128		111	107	122	10	04	116	120
Wild Type		31	26		21	20	17	2	27	19	12
Never Genotyped		0	1		1	0	2		0	0	0
1-Class		8	10		14	11	9		7	9	8
2-Class		1	2		2	2	1		1	0	1
3-Class		0	0		0	0	0		0	0	0
Total (n=)		137	167		149	140	151	13	39	144	141
Indicator 12: AIDS-I	Defining Illn	ess	2007	2008	2009	2010	2011	2012	2013	2014	2015
CD4 < 200 at	Cases		15	14	16	10	13	7	10	≤ 5	≤ 5
ART initiation	Rate per 1	00,000	5.4	5.0	5.7	3.6	4.6	2.5	3.5	1.8	1.1
AIDS Cases	Cases	,	6	≤ 5	6	7	≤ 5	≤ 5	7	8	≤ 5
(DTP Reports)	Rate per 1	00,000	2.1	1.1	2.1	2.5	1.8	1.1	2.5	2.8	1.4
AIDS Cases	Cases	,	≤ 5	≤ 5	8	≤ 5	6	≤ 5	6	9	-
(BCCDC Reports)	Rate per 1	00,000	1.8	1.1	2.9	1.1	2.1	0.4	2.1	3.1	_
Indicator 13: HIV-Re	elated Morto	lity	2004	2005	2006	2007	2008	2009	2010	2011	
Northern Health	171UI ld		7	6	7	8	6	<u>2009</u> ≤5	<u>2010</u> ≤5	<u>2011</u> ≤5	
Per 100 HIV+ Popula	ntion		2.28	1.92	2.21	2.47	1.81	0.89	1.45	1.42	
Per 100,000 Population			2.30	1.95	2.42	2.75	2.11	1.05	1.73	1.72	
1 er 100,000 i opuialio	J11		2.50	1.75	2.72	4.73	2.11	1.03	1./3	1./ 4	