

HIV MONITORING QUARTERLY REPORT

FOR INTERIOR HEALTH

THIRD 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, BC Vital Statistics, 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.

List of Indicators

Indicator 1. HIV Testing Episodes

Indicator 2. HIV Testing Rate

Indicator 3. New HIV Diagnoses

Indicator 4. Stage of HIV Infection at Diagnosis

Indicator 5. HIV Cascade of Care

Indicator 6. Programmatic Compliance Score (PCS)

Indicator 7. New Antiretroviral Therapy Starts

Indicator 8. CD4 Cell Count at ART Initiation

Indicator 9. Active and Inactive Drug Treatment Program (DTP) Participants

Indicator 10. Antiretroviral Adherence

Indicator 11. Resistance Testing and Results

Indicator 12. AIDS-Defining Illness

Indicator 13. HIV-Related Mortality

Table of Contents

Acknowledgements and Contributions

BC Provincial STOP Program:

A Note on Monitoring and Interpreting HIV Indicators

Indicator 1	Hiv Testing Episodes All HIV Testing Episodes reflect non-prenatal tests. All prenatal tests have been removed.
Figure 1.1	HIV Test Episodes for Interior Health, 2011 Q4–2016 Q3
Figure 1.2	HIV Test Episodes for Interior Health by Gender, 2011 Q4–2016 Q3
Figure 1.3	HIV Test Episodes for Interior Health by Age Category, 2011 Q4–2016 Q3
Figure 1.4	Point-of-Care HIV Tests for Interior Health, 2011 Q4–2016 Q3
Figure 1.5	HIV Test Episodes by HSDA for Interior Health, 2011 Q4–2016 Q3
Figure 1.6	HIV Test Episodes for Non-Prenatal Females in Interior Health by HSDA, 2011 Q4–2016 Q3
Figure 1.7	HIV Test Episodes for Males in Interior Health by HSDA, 2011 Q4–2016 Q3
Indicator 2	HIV Testing Rates All HIV Testing Rates reflect non-prenatal tests. All prenatal tests have been removed.
Figure 2.1	Rate of HIV Testing for Interior Health and HSDA's, 2009–2015
Figure 2.2	Rate of HIV Testing for Interior Health by Gender, 2009–2015
Figure 2.3	Rate of HIV Testing for Interior Health by Age Category, 2009–2015
Indicator 3	New HIV Diagnoses
Figure 3.1	New HIV Diagnoses for Interior Health, 2011 Q4–2016 Q3
Figure 3.2	New HIV Diagnoses for Interior Health by Gender, 2011 Q4–2016 Q3
Figure 3.3	New HIV Diagnoses for Interior Health by Age Category, 2011 Q4–2016 Q3
Figure 3.4	New HIV Diagnoses for Interior Health by Exposure Category, 2011 Q1–2015 Q2
Figure 3.5	New HIV Diagnoses for Interior Health by HSDA, 2011 Q4–2016 Q3
Indicator 4	Stage of HIV Infection at Diagnosis Stage definitions have been altered to remove AIDS diagnosis data. Individuals previously classified as Stage 3 have been re-classified based on CD4 cell count.
Table 1	Staging Classifications of Infection at Time of HIV Diagnosis Based on CDC HIV Surveillance Case Definitions
Figure 4.1	Stage of HIV Infection at Diagnosis for Interior Health, 2011–2015
Figure 4.2	Stage of HIV Infection at Diagnosis for Interior Health by Gender, 2011–2015
Figure 4.3	Stage of HIV Infection at Diagnosis for Interior Health by Age Category, 2011–2015
Figure 4.4	Stage of HIV Infection at Diagnosis for Interior Health by Exposure Category, 2011–2015
Indicator 5	HIV Cascade of Care
Figure 5.1	Estimated Cascade of Care for Interior Health, Year Ending 2016 Q3
Figure 5.2	Estimated Cascade of Care for Interior Health by Gender, Year Ending 2016 Q3
Figure 5.3	Estimated Cascade of Care for Interior Health by Age Category, Year Ending 2016 Q3

Figure 5.4 Estimated Cascade of Care for Interior Health by Msm Status, Year Ending 2016 Q3 Estimated Cascade of Care for Interior Health by Age Category and Msm Status, Figure 5.5 Year Ending 2016 Q3 Figure 5.6 Estimated Cascade of Care for Interior Health by PWID Status, Year Ending 2016 Q3 Figure 5.7 Estimated Cascade of Care for Interior Health by HSDA, Year Ending 2016 Q3 **Indicator 6 Programmatic Compliance Score (PCS)** Table 2 Probability of Mortality, Immunologic Failure and Virologic Failure Based on the Programmatic Compliance Score Pcs Components for Interior Health, 2014 Q4-2016 Q3 Figure 6.1 Less than 3 CD4 Tests in First Year Less than 3 Viral Load Tests in First Year Not Having Drug Resistance Testing at Baseline Non-Recommended Antiretroviral Therapy Regimen (ART) Baseline CD₄ < 200 cells/ μ L Not Achieving Viral Suppression at 9 Months Figure 6.2 Historical Trends for Pcs Score for Interior Health, 2014 Q4-2016 Q3 **Indicator 7** New Antiretroviral Therapy Starts in Interior Health Figure 7 BC-CfE Drug Treatment Program Enrollment: New Antiretroviral Participants for Interior Health, 2014 Q4-2016 Q3 **Indicator 8 CD4 Cell Count at ART Initiation** Figure 8 CD4 Cell Count at ART Initiation for Interior Health, 2014 Q4–2016 Q3 **Indicator 9** Active and Inactive Drug Treatment Program (DTP) Participants Table 3 Distribution of People on ART in Interior Health, 2016 Q3 Active and Inactive DTP Participants for Interior Health, 2014 Q4-2016 Q3 Figure 9 **Antiretroviral Adherence** Indicator 10 Figure 10 Distribution of Individuals by Adherence Level in 1st Year of Therapy, Based on Pharmacy Refill Compliance for Interior Health, 2014 Q4-2016 Q3 **Resistance Testing and Results** Indicator 11 Figure 11 Cumulative Resistance Testing Results by Resistance Category for Interior Health, 2014 Q4-2016 Q3 **Indicator 12 AIDS-Defining Illness** Figure 12 AIDS Case Rate and Reports for Interior Health, 2008–2015 **Indicator 13 HIV-Related Mortality** Figure 13 HIV-Related Deaths by Year for Interior Health, 2004-2011

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 is responsible for compiling and publishing this report. Lilith Swetland is the editor of this 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. Olga Mazo, 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

Dr. Rolando Barrios, Chair, BC-CFE

Dr. Kate Heath, BC-CFE

Dr. Bohdan Nosyk, BC-CFE

Dr. Viviane Dias Lima, BC-CFE

Irene Day, BC-CFE

Dr. Jean Shoveller, BC-CFE

Dr. Jason Wong, BCCDC

Dr. Mel Krajden, BCCDC

Salman Klar, FHA

Jennifer May-Hadford, іна

Kari Harder, NHA

Dr. Neora Pick, PHSA

Dr. Reka Gustafson, vсна

Dr. Melanie Rusch, vіна

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.

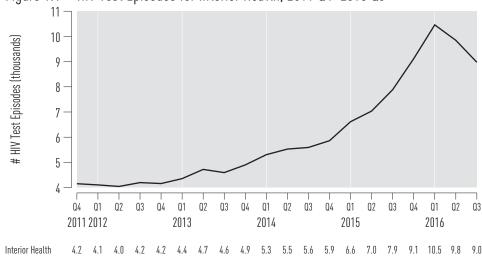
The goals of HA-led STOP-funded initiatives are to work toward achieving these goals. To these ends some outcome measures or indicators of progress have been drafted that should be considered in the design and implementation phases of these initiatives.

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 Interior Health, 2011 Q4-2016 Q3



6 5 # HIV Test Episodes (thousands) 3 Female 2 Male 1 -Q1 Q2 Q3 Q4 Q1 Q2 Q3 Q4 Q1 Q2 Q3 Q1 Q2 Q1 Q2

2014

2.6 2.6 2.7

2.4 2.6 2.8

2.2 2.4

2.3 2.3

2015

0.0 0.1

2.8 3.2

2.7 2.9 3.2 3.4 3.8

2016

2013

2.0 2.1 2.3

2.1 2.2

2.0

2011 2012

Female

Male

Other

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

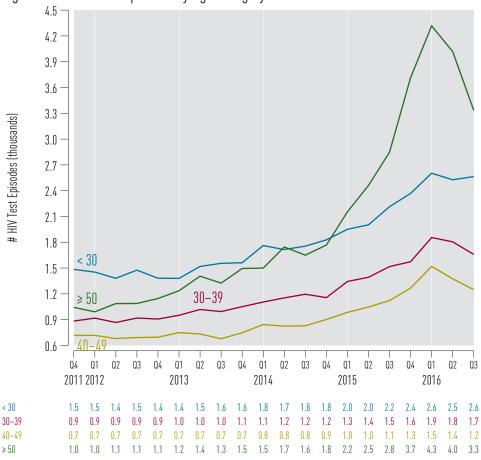
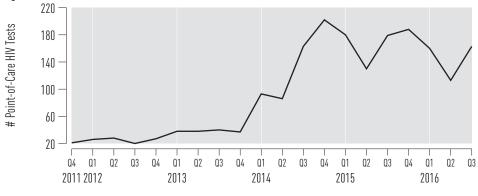


Figure 1.4 Point-of-Care HIV Tests for Interior Health



Interior Health 21.0 26.0 28.0 20.0 27.0 38.0 38.0 40.0 37.0 93.0 86.0 163.0 202.0 180.0 130.0 179.0 188.0 160.0 113.0 163.0

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.

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

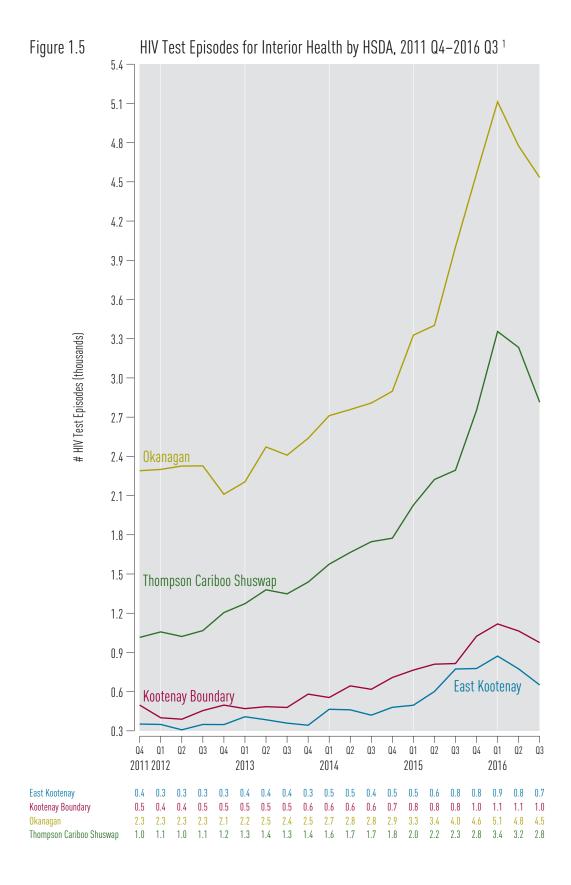
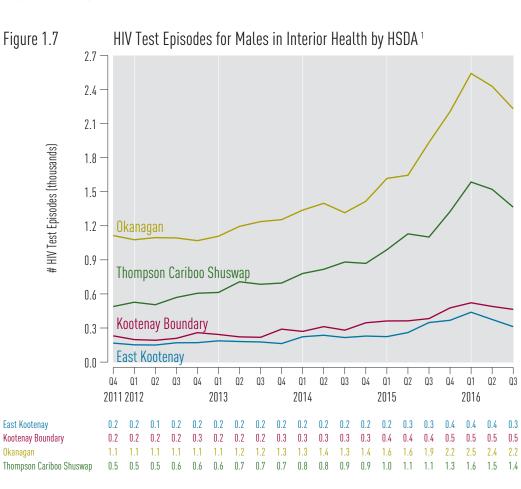


Figure 1.6 HIV Test Episodes for Non-prenatal Females in Interior Health by HSDA ¹ 2.6 2.4 2.2 2.0 # HIV Test Episodes (thousands) 1.8 1.6 1.4 Okanagan 1.2 1.0 8.0 Thompson Cariboo Shuswap 0.6 0.4 Kootenay Boundary 0.2 East Kootenay 0.0 Q1 Q2 Q3 Q4 Q1 Q2 Q3 Q4 Q1 02 Q3 Q4 Q1 Q2 Q3 Q4 Q1 Q2 2011 2012 2013 2014 2015 2016 0.2 0.2 0.2 0.2 0.2 0.2 East Kootenay 0.2 0.2 0.2 0.2 0.2 0.2 0.3 0.3 0.4 0.4 0.4 0.2 0.2 0.2 0.2 0.2 0.3 0.3 0.3 0.3 0.3 0.3 0.3 0.5 Kootenay Boundary 1.1 1.2 1.2 1.2 1.0 1.0 1.2 1.1 1.2 1.3 1.4 1.4 1.6 1.7 1.9 2.2 2.4 Okanagan Thompson Cariboo Shuswap 0.5 0.5 0.5 0.5 0.6 0.7 0.7 0.7 0.7 0.8 0.8 0.8 0.8 1.0 1.0 1.1 1.4 1.7



Indicator 2. HIV Testing Rates

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

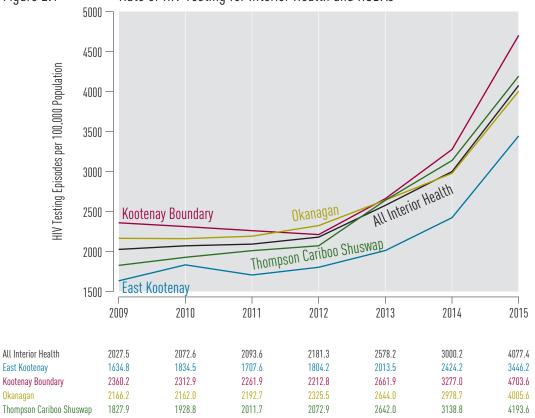
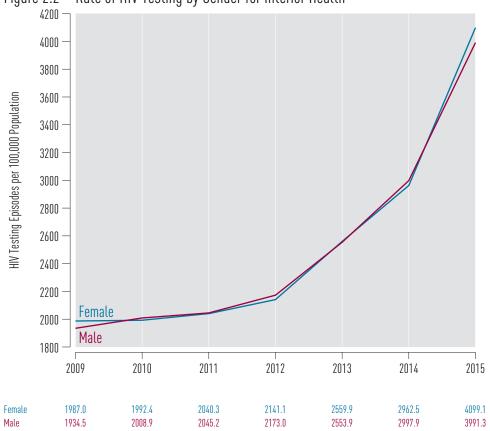
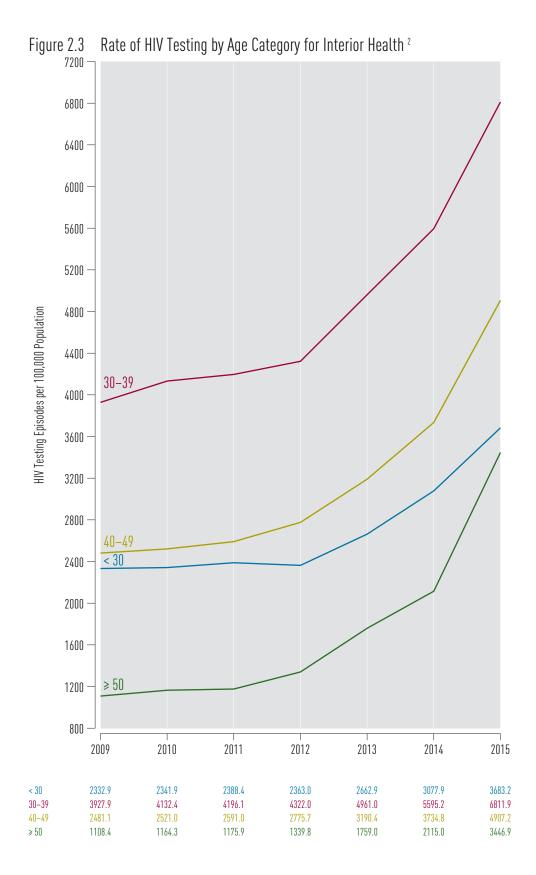


Figure 2.2 Rate of HIV Testing by Gender for Interior Health ²



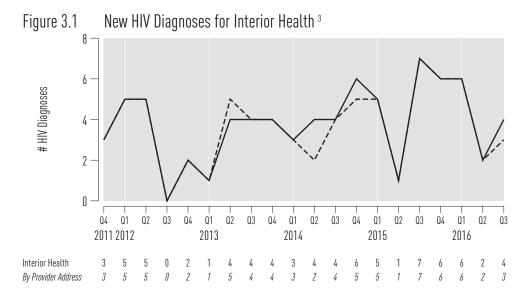


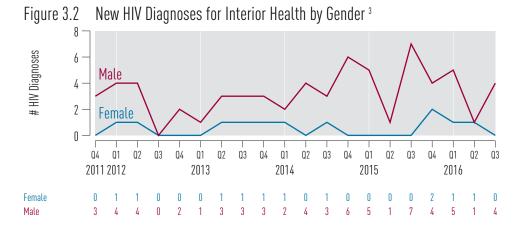
² Testing does not include point of care tests.

New HIV Diagnoses

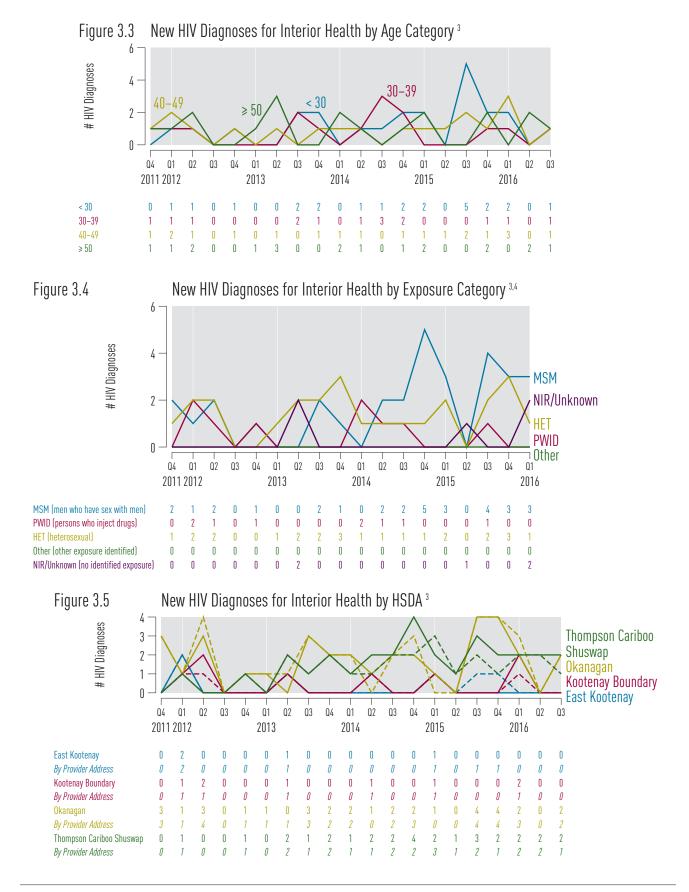
Trends in HIV diagnoses by gender and exposure category are described. Interpreting HIV diagnoses must be done with consideration that trends are influenced by both changes in testing rate as well as changes in transmission rates. It is important to note that new HIV diagnoses cases and rates are not 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





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



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	previous	ńegativ	ria met for acute HIV infection, or ve or indeterminate HIV test within confirmed positive HIV test.
1	,		CD4 ≥500
2a		and	CD4 350-499
2b	Stage 0		CD4 200-349
3	not met		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 Interior Health, 2011–2015 ⁵

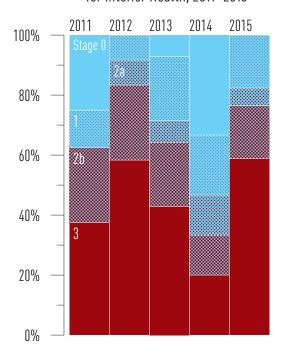
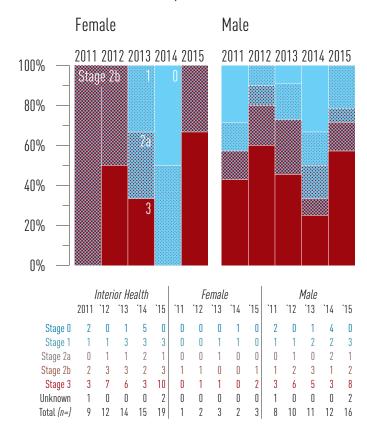
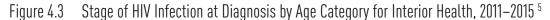


Figure 4.2 Stage of HIV Infection at Diagnosis by Gender for Interior Health, 2011–2015 ⁵





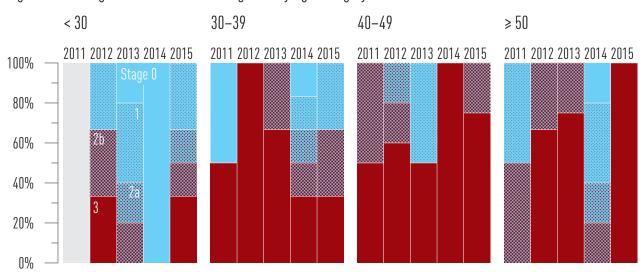
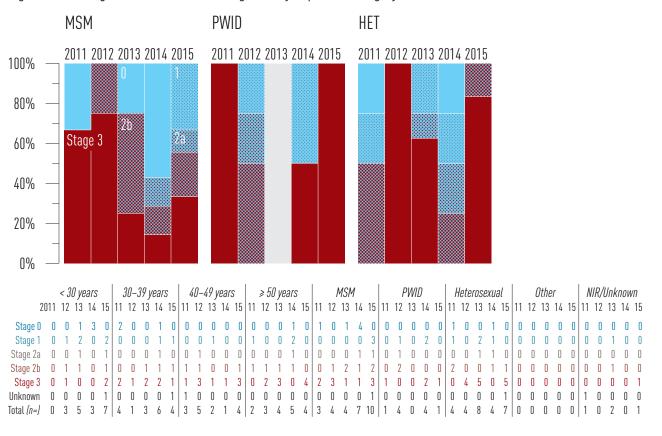


Figure 4.4 Stage of HIV Infection at Diagnosis by Exposure Category for Interior Health, 2011–2015 5,6



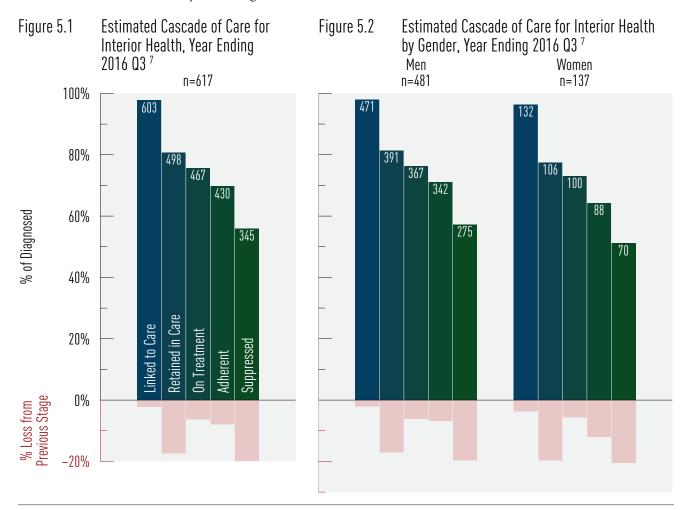
⁵ Data Source: BCCDC

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 Q4–2016 Q3 in Interior Health and stratified by sex and age.

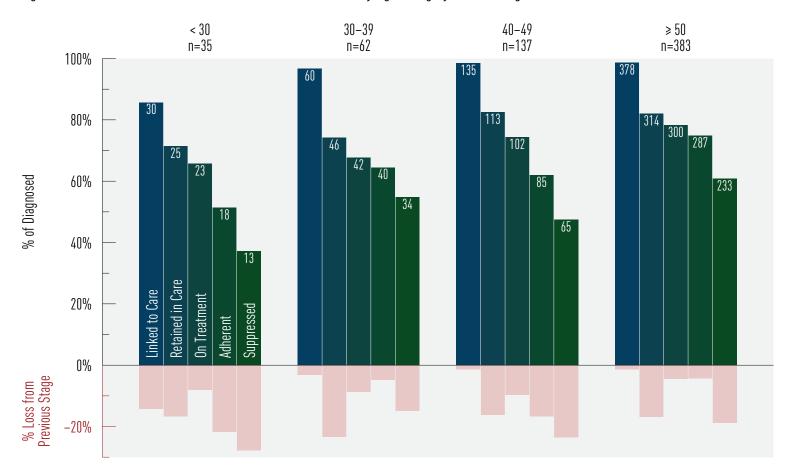


- 7 Data is for the period 2015 Q4-2016 Q3. Data Sources:
 - i British Columbia Centre for Excellence Drug Treatment Program (DTP) Database (ARV use, VL and CD4 count).
 - ii Administrative data (ex. MSP billings; hospitalization data from the Discharge Abstract Database (DAD)).

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.

Figure 5.3 Estimated Cascade of Care for Interior Health by Age Category, Year Ending 2016 Q3 8



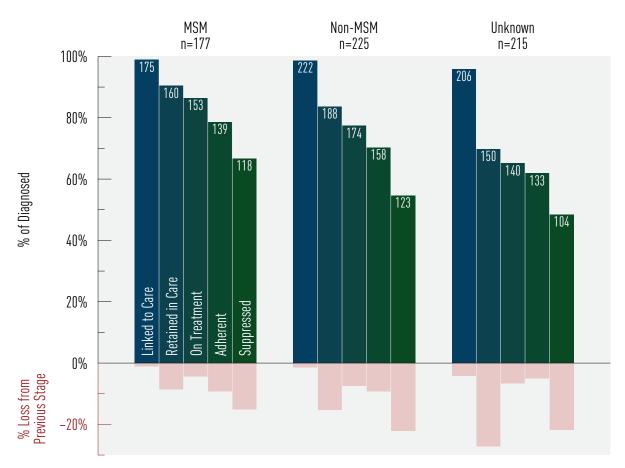
Limitations: HA assignment is based on the most recent HA of residence of the patient, if not available of the HIV-care provider. If the most recent HA of residence is not updated then the designated HA may be incorrect.

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





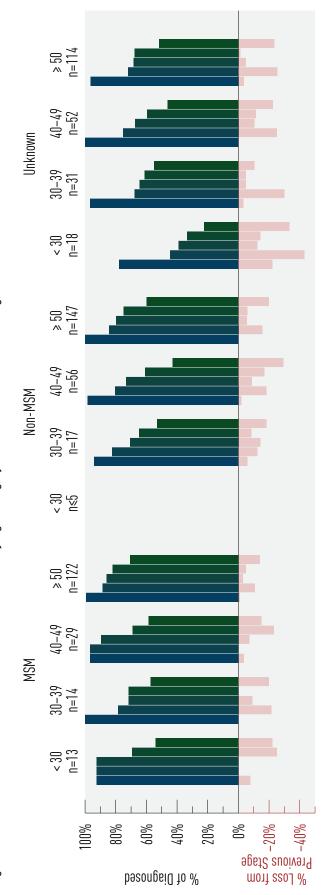
Limitations: HA assignment is based on the most recent HA of residence of the patient, if not available of the HIV-care provider. If the most recent HA of residence is not updated then the designated HA may be incorrect.

⁹ Data is for the period 2015 Q4–2016 Q3. 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)).

Figure 5.5 Estimated Cascade of Care for Interior Health by Age Category and MSM Status, Year Ending 2016 Q3 9



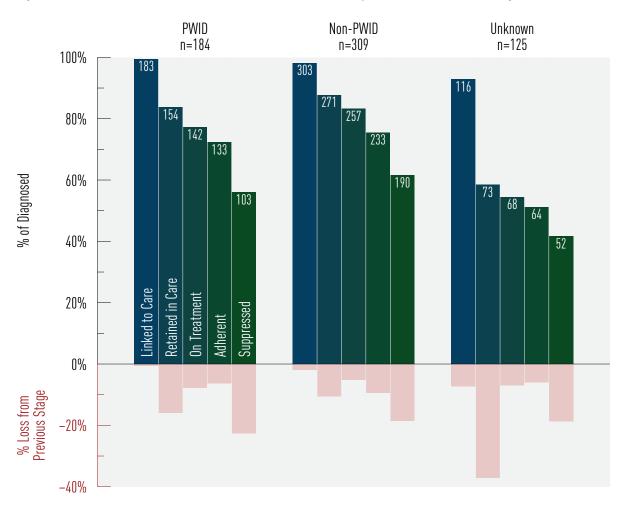
Data is for the period 2015 Q4-2016 Q3. Data Sources:

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.

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

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





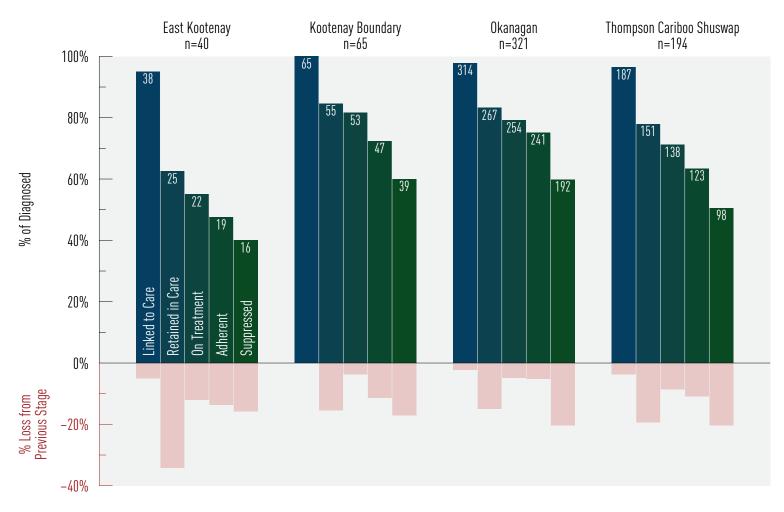
Limitations: HA assignment is based on the most recent HA of residence of the patient, if not available of the HIV-care provider. If the most recent HA of residence is not updated then the designated HA may be incorrect.

⁹ Data is for the period 2015 Q4–2016 Q3. Data Sources:

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

 $ii \quad \ \, Administrative \ data \ (ex. \ {\tt MSP} \ billings; hospitalization \ data \ from \ the \ Discharge \ Abstract \ Database \ ({\tt DAD})).$

Figure 5.7 Estimated Cascade of Care for Interior Health by HSDA, Year Ending 2016 Q3 9



Data Sources:

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 Q4-2016 Q3.

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

ii Administrative data (ex. мsp 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 2 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 Interior Health, 2014 Q4-2016 Q3 10

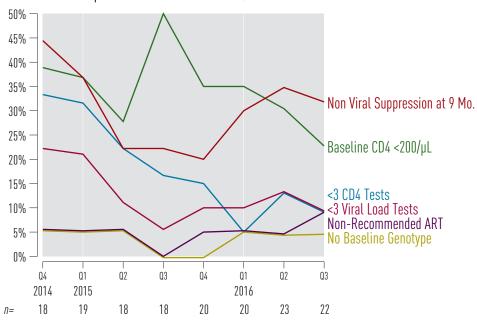
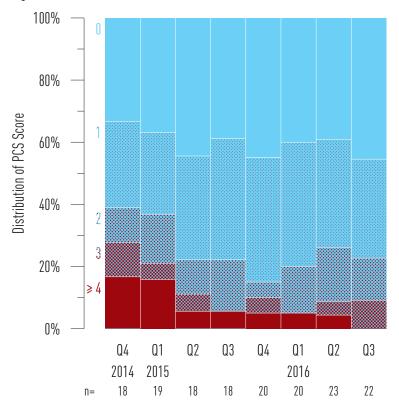


Figure 6.2 Historical Trends for PCS Score for Interior Health, 2014 Q4-2016 Q3 10,11



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

¹¹ 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 o 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 Interior Health

Figure 7 BC-CfE Drug Treatment Program Enrollment: New ART Participants in Interior Health, 2014 Q4-2016 Q3 12



Indicator 8. CD4 Cell Count at ART Initiation

Figure 8 CD4 Cell Count at ART Initiation of ART-Naïve DTP Participants in Interior Health, 2014 Q4-2016 Q3 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.

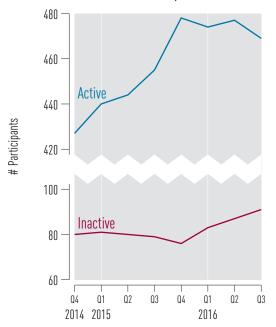
¹³ Data Source: Drug Treatment Program Database Limitations: CD4 cell count data is approximately 80% complete.

Indicator 9. Active and Inactive DTP Participants

Table 3. Distribution of People on ART for Interior Health, 2016 Q3 14

Age	< 30	21
	30-39	46
	40-49	102
	≥ 50	300
Gender	Male	373
	Female	96
Exposure	MSM	153
	PWID	146
Total		469

Figure 9 Active and Inactive DTP Participants for Interior Health, 2014 Q4-2016 Q3 15



Definition:

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

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.

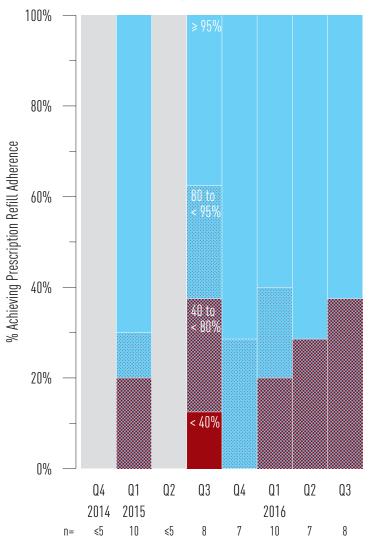
Data Source: Drug Treatment Program Database
Limitation: DTP participants are designated to an HA based on most current residence provided by the participant.

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 Interior Health, 2014 Q4–2016 Q3 ¹⁶



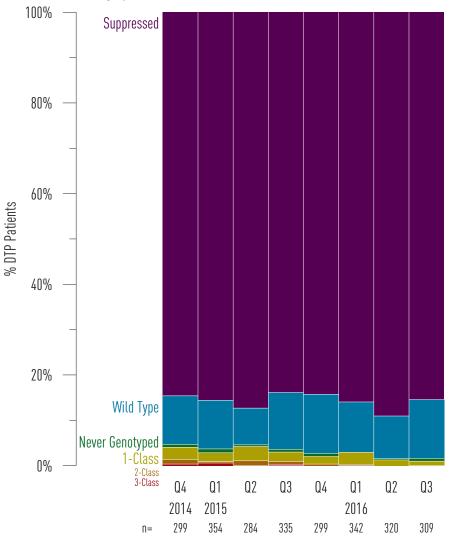
¹⁶ Data Source: Drug Treatment Program Database Limitation: Prescription refill adherence is used as a proxy for patient adherence.

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.

Figure 11 Cumulative Resistance Testing Results by Resistance Category for Interior Health, 2014 Q4–2016 Q3 ¹⁷



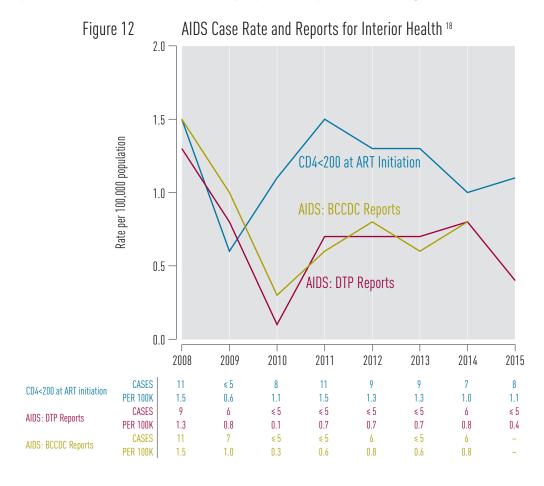
¹⁷ Data Source: Drug Treatment Program Database

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.



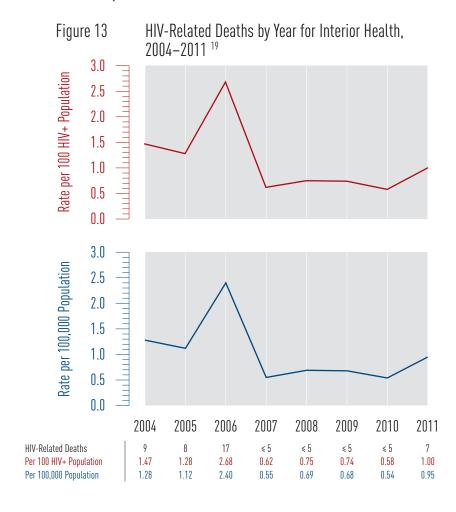
Data Source: DTP AIDS cases are obtained from the Drug Treatment Program Database; BCCDC AIDS cases are obtained from the BC-CDC; CD4<200 at ART initiation data came from the DTP database. Indicator 12 also reflects information from BC Vital Statistics. As this information is made available to BC-CFE, we use it to inform the development and refinement of this indicator.

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	: Test Episod	les		2012				2013					2014	l.			2015				2.0	016		
(thousand			Q4	Q1	Q2	Q3	Q4	Q1	Q	2 (Q3	Q4	Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q ²		Q1	Q2	Q3
Interior He	*		4.2	4.1	4.0	4.2	4.2	4.4	4.		4.6	4.9	5.3	5.5	5.6	5.9	6.6	7.0	7.9).5	9.8	9.0
Gender	Female		2.1	2.1	2.0	2.1	2.0	2.1	2.		2.2	2.4	2.6	2.6		2.8	3.2	3.4	3.9			5.1	4.9	4.4
	Male		2.0	2.0	1.9	2.0	2.1	2.2	2.		2.3	2.4	2.6	2.8	2.7	2.9	3.2	3.4	3.8			5.1	4.8	4.4
	Other	(0.1	0.1	0.1	0.0	0.0	0.0	0.	.0 (0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.1	0.1	0.	1 ().1	0.1	0.0
Age	< 30		1.5	1.5	1.4	1.5	1.4	1.4	1.	.5	1.6	1.6	1.8	1.7	1.8	1.8	2.0	2.0	2.2	2.4	4 2	2.6	2.5	2.6
0	30-39	(0.9	0.9	0.9	0.9	0.9	1.0	1.	.0	1.0	1.1	1.1	1.2	1.2	1.2	1.3	1.4	1.5	1.6	6 1	1.9	1.8	1.7
	40-49	(0.7	0.7	0.7	0.7	0.7	0.7	0.	.7 (0.7	0.7	0.8	0.8	0.8	0.9	1.0	1.0	1.1	1.3	3 1	1.5	1.4	1.2
	≥ 50		1.0	1.0	1.1	1.1	1.1	1.2	1.	.4	1.3	1.5	1.5	1.7	1.6	1.8	2.2	2.5	2.8	3.7	7 4	1.3	4.0	3.3
POC Tests	(not in thousan	nds)	21	26	28	20	27	38	3	8	40	37	93	86	163	202	180	130	179	188	8 1	60	113	163
East Koote	nay	(0.4	0.3	0.3	0.3	0.3	0.4	0.	.4 (0.4	0.3	0.5	0.5	0.4	0.5	0.5	0.6	0.8	0.8	8 ().9	0.8	0.7
Female		(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	0.4	0.4	4 ().4	0.4	0.3
Male			0.2	0.2	0.1	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	0.4	4 ().4	0.4	0.3
Kootenay I	Boundary		0.5	0.4	0.4	0.5	0.5	0.5	0.	.5 (0.5	0.6	0.6	0.6	0.6	0.7	0.8	0.8	0.8	1.0	0 1	1.1	1.1	1.0
Female		(0.3	0.2	0.2	0.2	0.2	0.2	0.	.3 (0.3	0.3	0.3	0.3	0.3	0.3	0.4	0.4	0.4	0.5	5 ().6	0.6	0.5
Male		(0.2	0.2	0.2	0.2	0.3	0.2	0.	.2 (0.2	0.3	0.3	0.3	0.3	0.3	0.4	0.4	0.4	0.5	5 ().5	0.5	0.5
Okanagan		:	2.3	2.3	2.3	2.3	2.1	2.2	2.	.5 2	2.4	2.5	2.7	2.8	2.8	2.9	3.3	3.4	4.0	4.6	5 5	5.1	4.8	4.5
Female			1.1	1.2	1.2	1.2	1.0	1.0	1.		1.1	1.2	1.3	1.3	1.4	1.4	1.6	1.7	1.9			2.4	2.2	2.2
Male			1.1	1.1	1.1	1.1	1.1	1.1	1.	.2	1.2	1.3	1.3	1.4	1.3	1.4	1.6	1.6	1.9	2.2	2 2	2.5	2.4	2.2
•	ı Cariboo Shu	swap	1.0	1.1	1.0	1.1	1.2	1.3	1.		1.3	1.4	1.6	1.7	1.7	1.8	2.0	2.2	2.3	2.8	8 3	3.4	3.2	2.8
Female			0.5	0.5	0.5	0.5	0.6	0.7	0.		0.7	0.7	0.8	0.8	0.8	0.8	1.0	1.0	1.1	1.4		1.7	1.7	1.4
Male		(0.5	0.5	0.5	0.6	0.6	0.6	0.	.7 (0.7	0.7	0.8	0.8	0.9	0.9	1.0	1.1	1.1	1.3	3 1	1.6	1.5	1.4
T., J:	D-4£1113	7 T4:		. 100 (200		2000			2010		,	0011		2012		201	12		2014		,	0015	
All Interior	2: Rate of HIV	resting	g per	100,0	JUU		2009 2027.			2010 072.6			2011 93.6		$\frac{2012}{2181.3}$		201			$\frac{2014}{000.2}$			$\frac{2015}{77.4}$	
East Koote							2027 1634.8			372.0 334.5			07.6		1804.2		2013			124.2			46.2	
Kootenay I							2360.2			312.9			61.9		2212.8		2661			277.0			03.6	
Okanagan	Doundary						2366.2 2166.2			162.0			92.7		2325.5		2644			978.7			05.6	
_	Cariboo Shu	ewan					1827.9			928.8			11.7		2072.9		2642			38.8			93.6	
Gender	Female	Swap					1987.0			992.4			40.3		2141.1		2559			962.5			99.1	
Gerraer	Male						1934.			008.9			45.2		2173.0		2553			97.9			91.3	
Age	< 30						2332.9			341.9			88.4		2363.0		2662			77.9			83.2	
8	30-39						3927.9			132.4			96.1		4322.0		4961			595.2			11.9	
	40-49						2481.	1	25	521.0)	25	91.0		2775.7		3190	.4	37	734.8		49	07.2	
	≥ 50						1108.	4	11	164.3	3	11	75.9		1339.8		1759	0.0	21	15.0		34	46.9	
Indicator 3	3: New HIV D	iagnoses	S				2012				2013				2014			2015				201		
Total and The	1(1.	D. Cli	. (D .	1						Q4		Q2			Q1 Q2			Q1			Q4	Q1		Q3
Interior He	ealth	By Clien				3	5	5	0	2	1	4	4	4		4 4		5	1	7	6	6	2	4
C 1		By Provi	iaer	Aaare	255	3	5	5	0	2	1	5	4	4		2 4		5	1	7	6	6	2	3
Gender		Female Male				0	1 4	1 4	0	0 2	0	1	1	1 3		0 1 4 3		0 5	0 1	0 7	2 4	1 5	1 1	0 4
Age		< 30				0	1	1	0	1	0	0	2	2		1 1	2	2	0	5	2	2	0	1
rige		30-39				1	1	1	0	0	0	0	2	1		1 3		0	0	0	1	1	0	1
		40-49				1	2	1	0	1	0	1	0	1		1 0		1	1	2	1	3	0	1
		≥ 50				1	1	2	0	0	1	3	0	0		1 0		2	0	0	2	0	2	1
Exposure		MSM				2	1	2	0	1	0	0	2	1		2 2		3	0	4	3	3		
1		PWID				0	2	1	0	1	0	0	0	0		1 1		0	0	1	0	0		
		HET				1	2	2	0	0	1	2	2	3	1	1 1	1	2	0	2	3	1		
		Other				0	0	0	0	0	0	0	0	0	0 (0 0	0	0	0	0	0	0		
		NIR/Un	kno	wn		0	0	0	0	0	0	2	0	0	0 (0 0	0	0	1	0	0	2		
East Koote	nay	By Clien	ıt Re	esiden	ce	0	2	0	0	0	0	1	0	0	0 (0 0	0	1	0	0	0	0	0	0
		By Provi	ider	Addre	ess	0	2	0	0	0	0	1	0	0	0 (0 0	0	1	0	1	1	0	0	0
Kootenay I	Boundary	By Clien				0	1	2	0	0	0	1	0	0	0	1 0	0	1	0	0	0	2	0	0
		By Provi				0	1	1	0	0	0	1	0	0		1 0		1	0	0	0	1	0	0
Okanagan		By Clien				3	1	3	0	1	1	0	3	2		1 2		1	0	4	4	2	0	2
		By Provi				3	1	4	0	1	1	1	3	2		0 2		0	0	4	4	3	0	2
Thompson	Cariboo	By Clien				0	1	0	0	1	0	2	1	2		2 2		2	1	3	2	2	2	2
Shuswap		By Provi	ıder	Addre	ess	0	1	0	0	1	0	2	1	2	1 .	1 2	2	3	1	2	1	2	2	1

Indicator 4: Stage of HIV Infe	ction at Baseline
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Indicator 4	l: Sta	ge of	HI	V In	fecti	ion a	t Ba	selin	e																					
	In	terio	or H	ealth			Fe	male				N	1ale				< 30) yea	rs			30-3	9 yea	ars			40-4	9 yea	ars	
	'11	'12	'13	'14	'15	'11	'12	'13	'14	'15	'11	'12	'13	'14	'15	'11	'12	'13	'14	'15	'11	'12	'13	'14	'15	'11	'12	'13	'14	'15
Stage 0	2	0	1	5	0	0	0	0	1	0	2	0	1	4	0	0	0	1	3	0	2	0	0	1	0	0	0	0	0	0
1	1	1	3	3	3	0	0	1	1	0	1	1	2	2	3	0	1	2	0	2	0	0	0	1	1	0	0	1	0	0
2a	0	1	1	2	1	0	0	1	0	0	0	1	0	2	1	0	0	1	0	1	0	0	0	1	0	0	1	0	0	0
2b	2	3	3	2	3	1	1	0	0	1	1	2	3	1	2	0	1	1	0	1	0	0	1	1	1	1	1	0	0	1
3	3	7	6	3	10	0	1	1	0	2	3	6	5	3	8	0	1	0	0	2	2	1	2	2	1	1	3	1	1	3
Unknown	1	0	0	0	2	0	0	0	0	0	1	0	0	0	2	0	0	0	0	1	0	0	0	0	1	1	0	0	0	0
Total	9	12	14	15	19	1	2	3	2	3	8	10	11	12	16	0	3	5	3	7	4	1	3	6	4	3	5	2	1	4
					•										•															
		≥ 50) yea	rs			N	ISM			ŀ	leter	osex	kual			PΛ	WID)		O	ther I	Expo	sure	2	N	IR/U	Jnkn	own	
	'11	'12	'13	'14	'15	'11	'12	'13	'14	'15	'11	'12	'13	'14	'15	'11	'12	'13	'14	'15	'11	'12	'13	'14	'15	'11	'12	'13	'14	'15
Stage 0	0	0	0	1	0	1	0	1	4	0	0	0	0	0	0	1	0	0	1	0	0	0	0	0	0	0	0	0	0	0
1	1	0	0	2	0	0	0	0	0	3	0	1	0	2	0	1	0	2	1	0	0	0	0	0	0	0	0	1	0	0
2a	0	0	0	1	0	0	0	0	1	1	0	1	0	0	0	0	0	1	1	0	0	0	0	0	0	0	0	0	0	0
2b	1	1	1	1	0	0	1	2	1	2	0	2	0	0	0	2	0	0	1	1	0	0	0	0	0	0	0	1	0	0
3	0	2	3	0	4	2	3	1	1	3	1	0	0	2	1	0	4	5	0	5	0	0	0	0	0	0	0	0	0	1
Unknown	0	0	0	0	0	0	0	0	0	1	0	0	0	0	0	0	0	0	0	1	0	0	0	0	0	1	0	0	0	0
Total	2	3	4	5	4	3	4	4	7	10	1	4	0	4	1	4	4	8	4	7	0	0	0	0	0	1	0	2	0	1

Indicator 5: HI	V Cascade of C	Care	Diagnosed	Linked	Retained	On ARVs	Adherent	Suppressed
Interior Health	1		617	603	498	467	430	345
Gender	Men		481	471	391	367	342	275
	Women		137	132	106	100	88	70
Age Category	< 30		35	30	25	23	18	13
	30-39		62	60	46	42	40	34
	40-49		137	135	113	102	85	65
	≥ 50		383	378	314	300	287	233
MSM Status	MSM		177	175	160	153	139	118
	Non-MSM		225	222	188	174	158	123
	Unknown		215	206	150	140	133	104
Age Category	MSM	< 30	13	12	12	12	9	7
and MSM Stati	18	30-39	14	14	11	10	10	8
		40-49	29	28	28	26	20	17
		≥ 50	122	121	108	105	100	86
	Non-MSM	< 30	≤ 5	≤ 5	≤ 5	≤ 5	≤ 5	≤ 5
		30-39	17	16	14	12	11	9
		40-49	56	55	45	41	34	24
		≥ 50	147	147	124	117	110	88
	Unknown	< 30	18	14	8	7	6	4
		30-39	31	30	21	20	19	17
		40-49	52	52	39	35	31	24
		≥ 50	114	110	82	78	77	59
PWID Status	PWID		184	183	154	142	133	103
	Non-PWID		309	303	271	257	233	190
	Unknown		125	116	73	68	64	52
HSDA	East Kootena	y	40	38	25	22	19	16
	Kootenay Bo	undary	65	65	55	53	47	39
	Okanagan		321	314	267	254	241	192
	Thompson C Shuswap	ariboo	194	187	151	138	123	98

Indicator 6: Programmati	c		2015					2016		
Compliance Score (PCS)	•	Q4	Q1		Q2	Q3	Q4	Q1	Q2	Q3
< 3 CD4 Tests		3.3%	31.6%	2	22.2%	16.7%	15.0%	5.0%	13.0%	9.1%
< 3 CD4 Tests		2.2%	21.1%		1.1%	5.6%	10.0%	10.0%	13.0%	9.1%
	2			1						
No Baseline Genotype	т -	5.6%	5.3%	2	5.6%	0.0%	0.0%	5.0%	4.3%	4.5%
Baseline CD4 < 200 cells/µ	IL 3	88.9%	36.8%	2	27.8%	50.0%	35.0%	35.0%	30.4%	22.7%
Non-Recommended ART		5.6%	5.3%	_	5.6%	0.0%	5.0%	5.0%	4.3%	9.1%
Non Viral Suppression at	9 Mo. 4	4.4%	36.8%	2	22.2%	22.2%	20.0%	30.0%	34.8%	31.8%
PCS Score: 0		6	7		8	7	9	8	9	10
PCS Score: 1		5	5		6	7	8	8	8	7
PCS Score: 2		2	3		2	3	1	3	4	3
PCS Score: 3		2	1		1	0	1	0	1	2
PCS Score: 4 or more		3	3		1	1	1	1	1	0
Total (n=)		18	19		18	18	20	20	23	22
Indicator 7: New DTP AR	V Participants									
First Starts	_	5	8		5	5	7	4	6	4
Experienced Starts		6	13		8	11	15	5	11	9
Indicator 8: CD4 Cell Cou	ınt Initiation for A	ARV-N	aïve DTP P	articipa	nts					
CD4 ≥ 500		-	1	F	-	-	1	-	4	
CD4 350-499		_	2		-	_	0	_	1	_
CD4 200-349		_	2		_	_	2	_	1	_
CD4 50-199		_	0		_	_	4	_	0	_
CD4 < 50		_	2		_	_	0	_	0	_
CD4 MED		_	310		_	_	190	_	545	_
Total (n=)		- ≤ 5	7		<u>-</u> ≤ 5	≤ 5	7	≤ 5	6	≤ 5
Total (II–)		33	,		33	33	,	23	Ū	33
Indicator 9: Active and In	active DTP Partic	_	440		444	455	470	47.4	477	460
Active DTP Participants		427	440 81		444 80	455 79	478	474	477 87	469 91
Inactive DTP Participants		80	01		80	79	76	83	67	91
Indicator 10: Antiretrovir	al Adherence									
≥ 95%		-	7		-	3	5	6	5	5
80% to < 95%		-	1		-	2	2	2	0	0
40% to < 80%		-	2		-	2	0	2	2	3
< 40%		-	0		-	1	0	0	0	0
Total (n=)		≤ 5	10		≤ 5	8	7	10	7	8
Indicator 11: Resistance T	esting and Result									
Suppressed		253	303		248	281	252	294	285	264
Wild Type		32	38		23	42	39	38	30	40
Never Genotyped		2	3		1	2	2	0	1	2
1-Class		8	7		9	7	5	9	4	3
2-Class		3	1		3	2	1	1	0	0
3-Class		1	2		0	1	0	0	0	0
4-Class		0	0		0	0	0	0	0	0
		200	354		284	335	299	342	320	309
Total (n=)		299	334			333	2//	312	020	
	ing Illness	299								2015
Indicator 12: AIDS-Defin		299	2008	2009	2010	2011	2012	2013	2014	2015
Indicator 12: AIDS-Defin CD4 < 200 at	Cases	299	2008	2009 ≤ 5	2010	2011	2012	2013	2014	8
Indicator 12: AIDS-Defin CD4 < 200 at ART initiation	Cases Rate per 100,000	299	2008 11 1.5	2009 ≤ 5 0.6	2010 8 1.1	2011 3 11 1.5	2012 9 1.3	2013 9 1.3	2014 7 1.0	8 1.1
Indicator 12: AIDS-Defin CD4 < 200 at ART initiation AIDS Cases	Cases Rate per 100,000 Cases	299	2008 11 1.5 9	2009 ≤ 5 0.6 6	2010 8 1.1 ≤ 5	2011 3 11 1.5 ≤ 5	2012 9 1.3 ≤ 5	2013 9 1.3 ≤ 5	2014 7 1.0 6	8 1.1 ≤ 5
Indicator 12: AIDS-Defin CD4 < 200 at ART initiation AIDS Cases (DTP Reports)	Cases Rate per 100,000 Cases Rate per 100,000	299	2008 11 1.5 9 1.3	2009 ≤ 5 0.6 6 0.8	2010 8 1.1 ≤ 5 0.1	2011 3 11 1.5 4 ≤ 5 0.7	2012 9 1.3 ≤ 5 0.7	2013 9 1.3 ≤ 5 0.7	2014 7 1.0 6 0.8	8 1.1
Indicator 12: AIDS-Defin CD4 < 200 at ART initiation AIDS Cases (DTP Reports) AIDS Cases	Cases Rate per 100,000 Cases Rate per 100,000 Cases	299	2008 11 1.5 9 1.3 11	2009 ≤ 5 0.6 6 0.8 7	2010 8 1.1 ≤ 5 0.1 ≤ 5	2011 11 1.5 ≤ 5 0.7 ≤ 5	2012 9 1.3 ≤ 5 0.7 6	2013 9 1.3 ≤ 5 0.7 ≤ 5	2014 7 1.0 6 0.8 6	8 1.1 ≤ 5 0.4
Indicator 12: AIDS-Defin CD4 < 200 at ART initiation AIDS Cases (DTP Reports) AIDS Cases	Cases Rate per 100,000 Cases Rate per 100,000	299	2008 11 1.5 9 1.3	2009 ≤ 5 0.6 6 0.8	2010 8 1.1 ≤ 5 0.1	2011 11 1.5 ≤ 5 0.7 ≤ 5	2012 9 1.3 ≤ 5 0.7	2013 9 1.3 ≤ 5 0.7	2014 7 1.0 6 0.8	8 1.1 ≤ 5
Indicator 12: AIDS-Defin CD4 < 200 at ART initiation AIDS Cases (DTP Reports) AIDS Cases (BCCDC Reports) Indicator 13: HIV-Related	Cases Rate per 100,000 Cases Rate per 100,000 Cases Rate per 100,000	299	2008 11 1.5 9 1.3 11 1.5	2009 ≤ 5 0.6 6 0.8 7 1.0 2005	2010 8 1.1 ≤ 5 0.1 ≤ 5 0.3	$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	2012 9 1.3 ≤ 5 0.7 6 0.8 2008	2013 9 1.3 ≤ 5 0.7 ≤ 5 0.6	2014 7 1.0 6 0.8 6 0.8 2010	8 1.1 ≤ 5 0.4 - - 2011
Indicator 12: AIDS-Defin CD4 < 200 at ART initiation AIDS Cases (DTP Reports) AIDS Cases (BCCDC Reports) Indicator 13: HIV-Related British Columbia	Cases Rate per 100,000 Cases Rate per 100,000 Cases Rate per 100,000	299	2008 11 1.5 9 1.3 11 1.5 2004	2009 ≤ 5 0.6 6 0.8 7 1.0 2005	2010 8 1.1 ≤ 5 0.1 ≤ 5 0.3 2006	$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	2012 9 1.3 ≤ 5 0.7 6 0.8 2008	2013 9 1.3 ≤ 5 0.7 ≤ 5 0.6 2009	2014 7 1.0 6 0.8 6 0.8 2010	8 1.1 ≤ 5 0.4 - - 2011 59
Indicator 12: AIDS-Defin CD4 < 200 at	Cases Rate per 100,000 Cases Rate per 100,000 Cases Rate per 100,000	299	2008 11 1.5 9 1.3 11 1.5	2009 ≤ 5 0.6 6 0.8 7 1.0 2005	2010 8 1.1 ≤ 5 0.1 ≤ 5 0.3	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	2012 9 1.3 ≤ 5 0.7 6 0.8 2008	2013 9 1.3 ≤ 5 0.7 ≤ 5 0.6	2014 7 1.0 6 0.8 6 0.8 2010	8 1.1 ≤ 5 0.4 - - 2011