

## HIV MONITORING QUARTERLY REPORT

FOR INTERIOR HEALTH

SECOND 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 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

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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 Interior Health

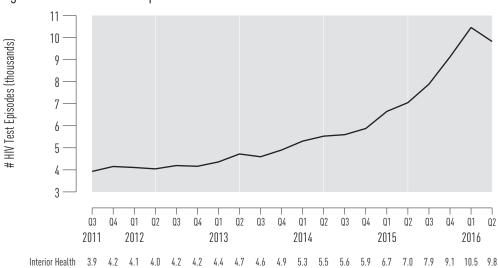


Figure 1.2 HIV Test Episodes by Gender for Interior Health 1.2

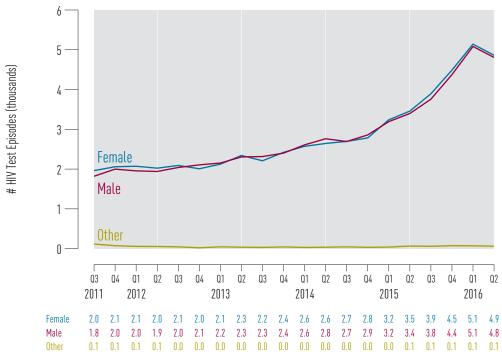


Figure 1.3 HIV Test Episodes by Age Category for Interior Health 1,2

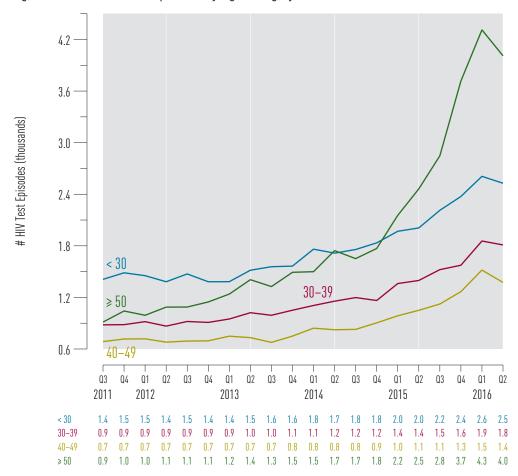
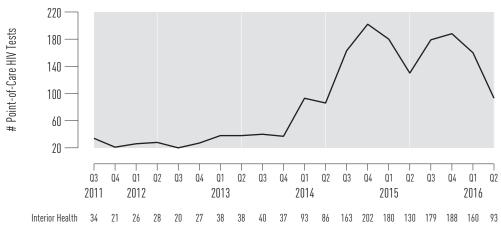


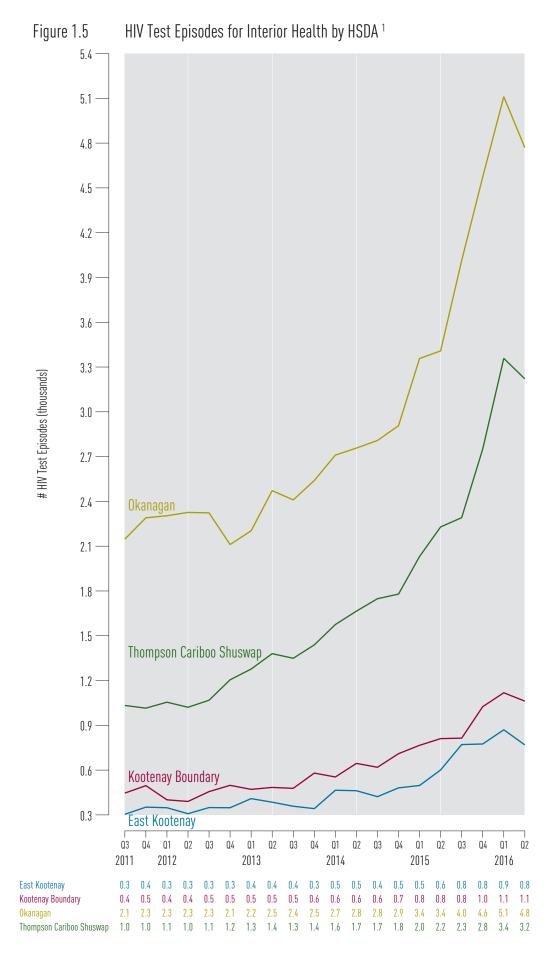
Figure 1.4 Point-of-Care HIV Tests for Interior Health

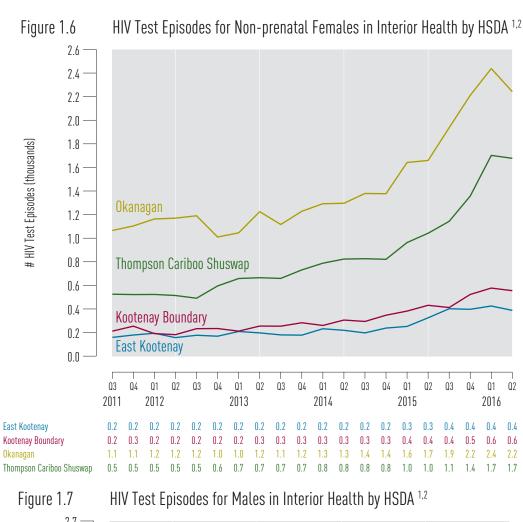


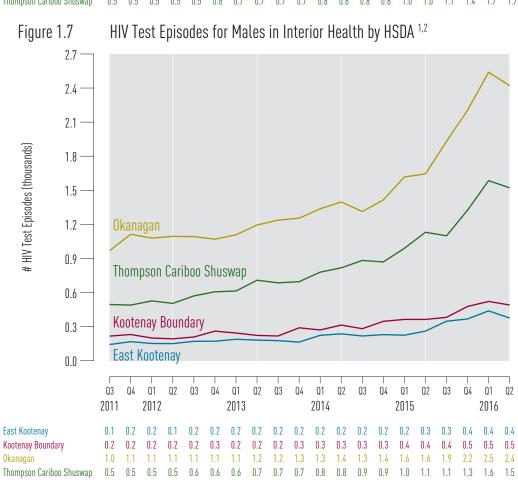
<sup>1</sup> Data Source: The BC Public Health Microbiology and Reference Laboratory (BCPHMRL) courtesy of the BC Centre for Disease Control (BCCDC).

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.







## Indicator 2. HIV Testing Rates

Figure 2.1 Rate of HIV Testing for Interior Health and HSDAs <sup>2</sup>

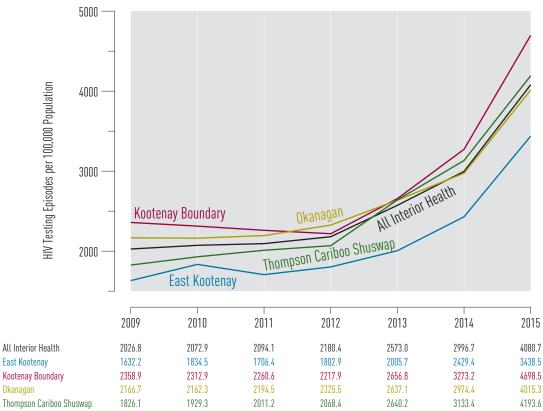
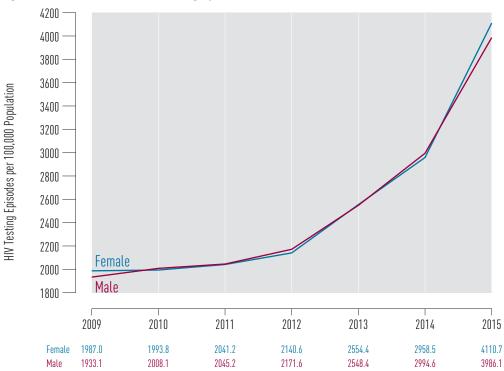


Figure 2.2 Rate of HIV Testing by Gender for Interior Health <sup>2</sup>



7200 -6800 6400 6000 5600 5200 -4800 -HIV Testing Episodes per 100,000 Population 4400 30-39 4000 3600 -3200 -2800 -40-49 2400 < 30 2000 1600 1200 ≥ 50 800 2009 2010 2012 2013 2011 2014 2015 2360.8 4319.5 2774.7 2333.3 2343.6 2389.3 2656.2 3071.6 3687.3 < 30 4131.2 2521.0 4196.1 2592.0 4944.8 3185.0 5595.2 3724.9 6841.9 4904.9 3925.4 30-39 40-49 2479.2 1340.1 ≥ 50 1107.8 1164.0 1175.9 1757.5 2114.1 3444.8

Figure 2.3 Rate of HIV Testing by Age Category for Interior Health  $^{\rm 2}$ 

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

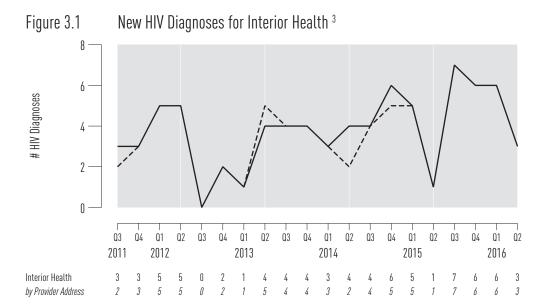


Figure 3.2 New HIV Diagnoses for Interior Health by Gender <sup>3</sup> 8 . # HIV Diagnoses Male Female Q4 Q1 Q2 Q3 Q4 Q1 Q2 Q3 Q4 Q1 Q2 Q3 Q4 Q1 Q2 Q3 Q4 Q1 Q2 2011 2012 2013 2014 2015 2016

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

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

Figure 3.4 New HIV Diagnoses for Interior Health by Exposure Category 3.4

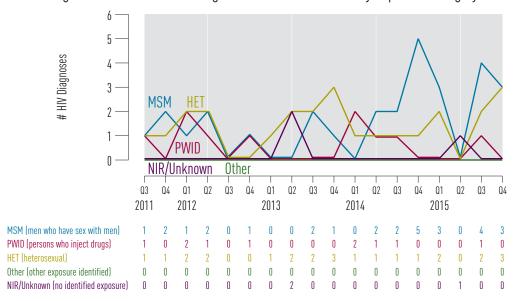
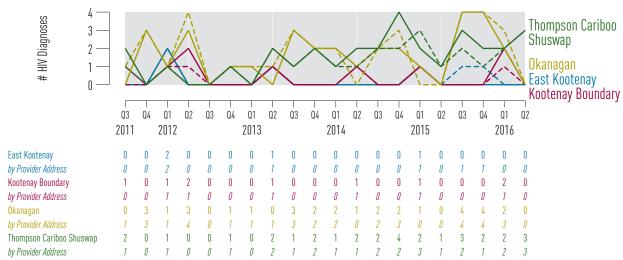


Figure 3.5 New HIV Diagnoses for Interior Health by HSDA <sup>3</sup>



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

<sup>4</sup> 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	Laboratory criteria met for acute HIV infection, or previous negative or indeterminate HIV test within 180 days of first confirmed positive HIV test.											
1			CD4 ≥500										
2a		and	CD4 350-499										
2b	Stage 0 not met		CD4 200-349										
3	HOL HIEL		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 <sup>5</sup>

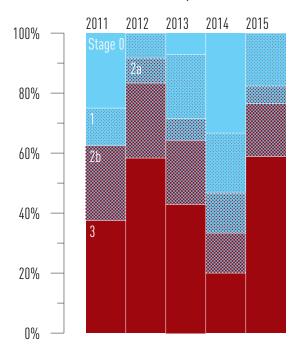
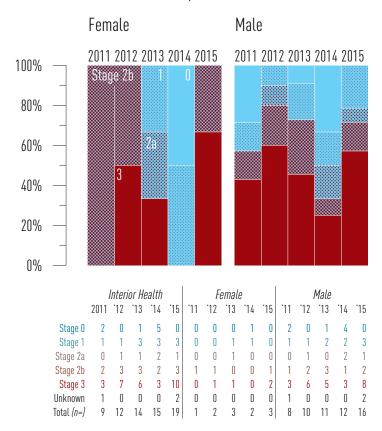


Figure 4.2 Stage of HIV Infection at Diagnosis by Gender for Interior Health, 2011–2015 <sup>5</sup>



Data Source: вссьс

Figure 4.3 Stage of HIV Infection at Diagnosis by Age Category for Interior Health, 2011–2015 <sup>5</sup>

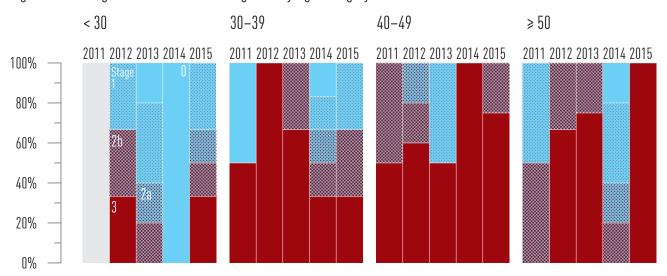
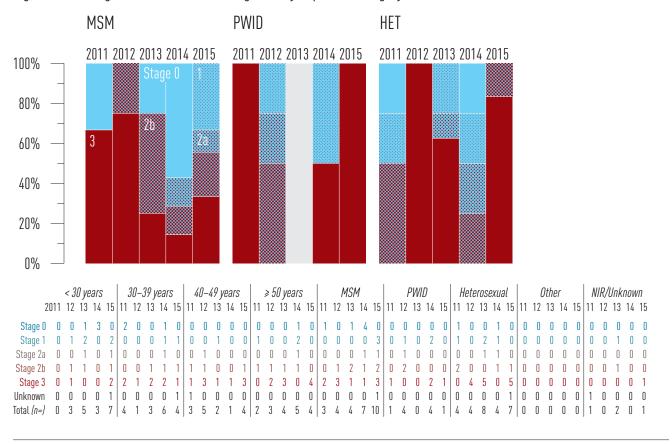


Figure 4.4 Stage of HIV Infection at Diagnosis by Exposure Category for Interior Health, 2011–2015 5,6



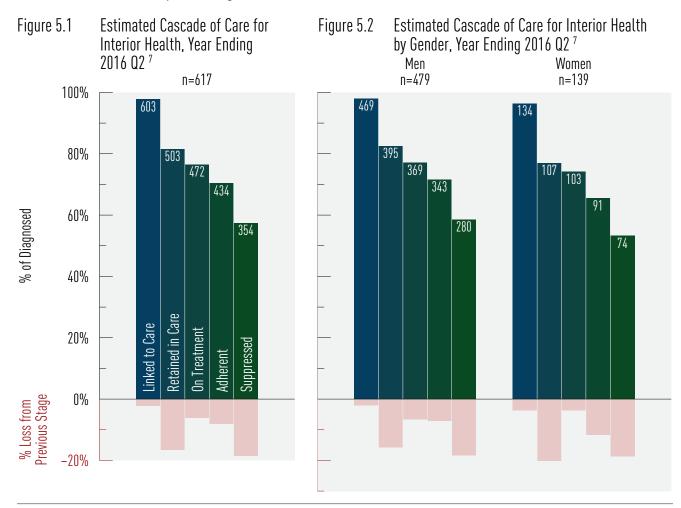
<sup>5</sup> Data Source: BCCDC

<sup>6</sup> 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 Q3–2016 Q2 in Interior Health and stratified by sex and age.



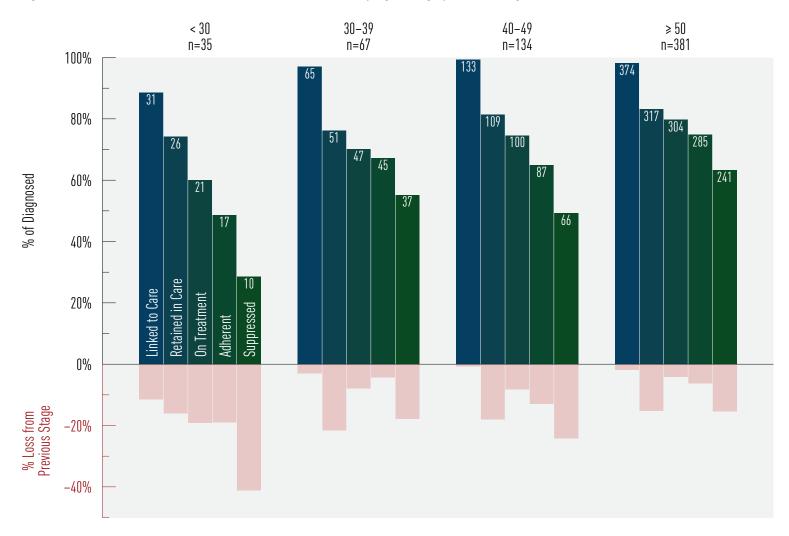
<sup>7</sup> Data is for the period 2015 Q3-2016 Q2. Data Sources:

- 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.





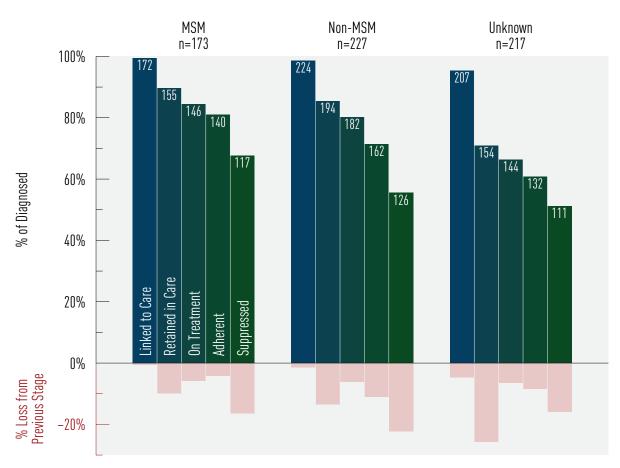
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<sup>8</sup> Data is for the period 2015 Q3-2016 Q2. Data Sources:

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)).





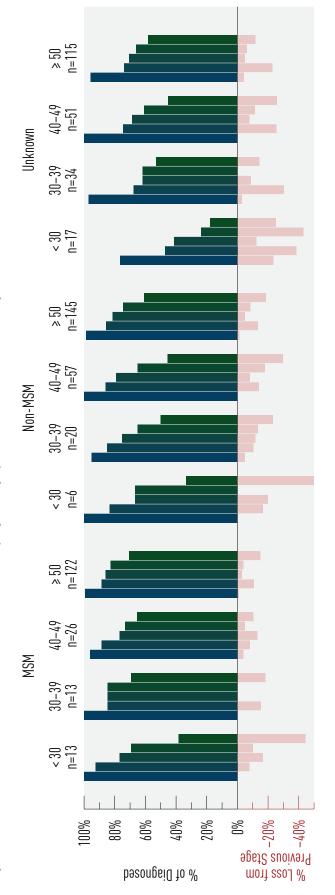
Limitations: HA assignment is based on the most recent HA of residence of the patient, if not available of the HIV-care provider. If the most recent HA of residence is not updated then the designated HA may be incorrect.

<sup>9</sup> Data is for the period 2015 Q3–2016 Q2. 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 Q2  $^{9}$ 

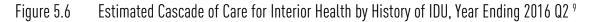


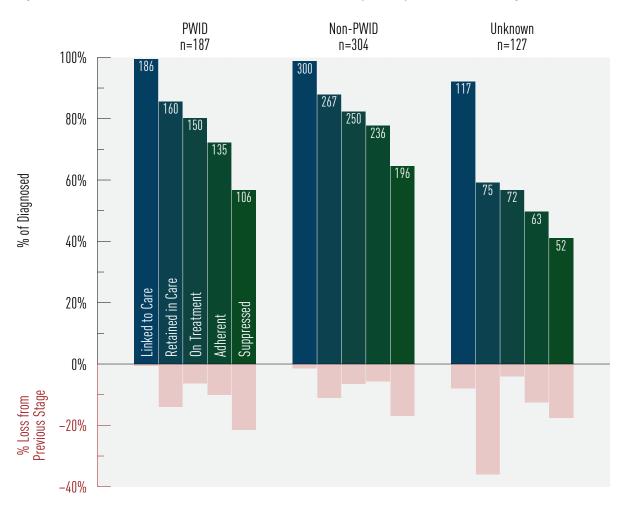
Data is for the period 2015 Q3-2016 Q2. Data Sources:

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)).





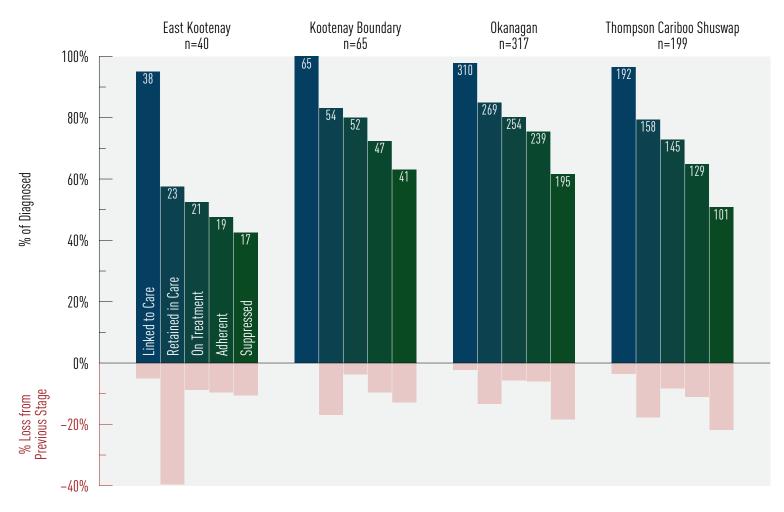
Limitations: HA assignment is based on the most recent HA of residence of the patient, if not available of the HIV-care provider. If the most recent HA of residence is not updated then the designated HA may be incorrect.

<sup>9</sup> Data is for the period 2015 Q3-2016 Q2. 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 5.7 Estimated Cascade of Care for Interior Health by HSDA, Year Ending 2016 Q2  $^{9}$ 



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.

<sup>9</sup> Data is for the period 2015 Q3-2016 Q2.

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 Q3-2016 Q2 10

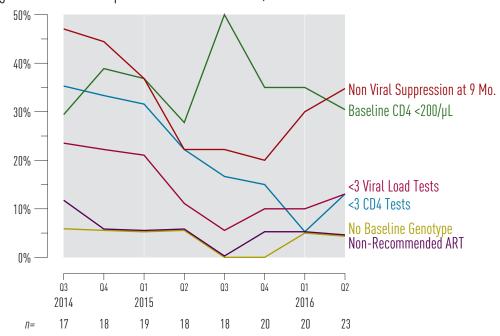
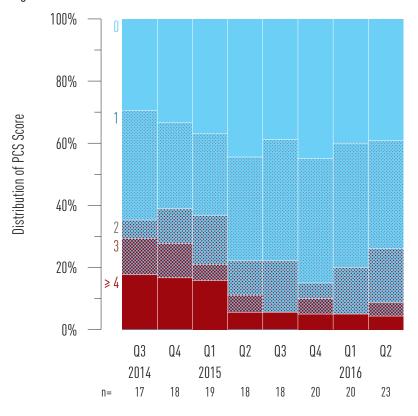


Figure 6.2 Historical Trends for PCS Score for Interior Health, 2014 Q3-2016 Q2 10,11



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 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 Q3-2016 Q2 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 Q3-2016 Q2 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.

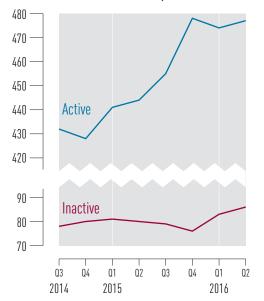
<sup>13</sup> Data Source: Drug Treatment Program Database Limitations: CD4 cell count data is approximately 80% complete.

## Indicator 9. Active and Inactive DTP Participants

Table 3. Distribution of People on ART for Interior Health, 2016 Q2  $^{\rm 14}$ 

Age	< 30	22
	30-39	53
	40-49	108
	≥ 50	294
Gender	Male	374
	Female	103
Exposure	MSM	153
	PWID	149
Total		477

Figure 9 Active and Inactive DTP Participants for Interior Health, 2014 Q3-2016 Q2  $^{15}$ 



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.

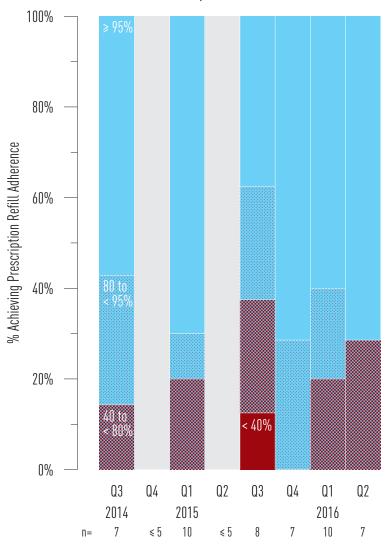
<sup>14</sup> Data Source: Drug Treatment Program Database Limitation: DTP participants are designated to an HA based on most current residence provided by the participant.

#### 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 Q3–2016 Q2 <sup>16</sup>



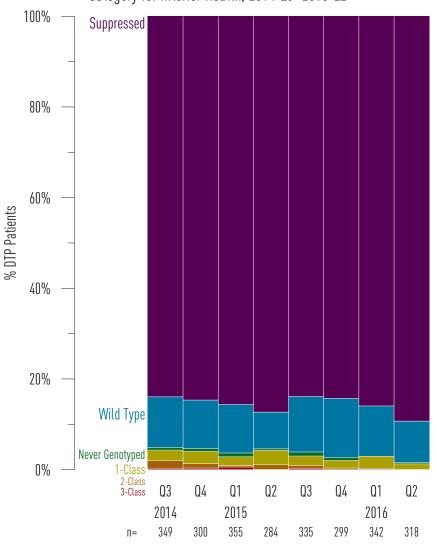
<sup>16</sup> Data Source: Drug Treatment Program Database Limitation: Prescription refill adherence is used as a proxy for patient adherence.

## 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 Q3–2016 Q2 <sup>17</sup>



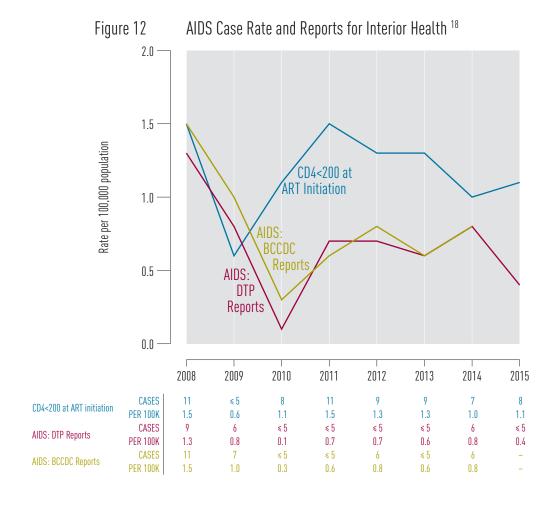
<sup>17</sup> Data Source: Drug Treatment Program Database

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.



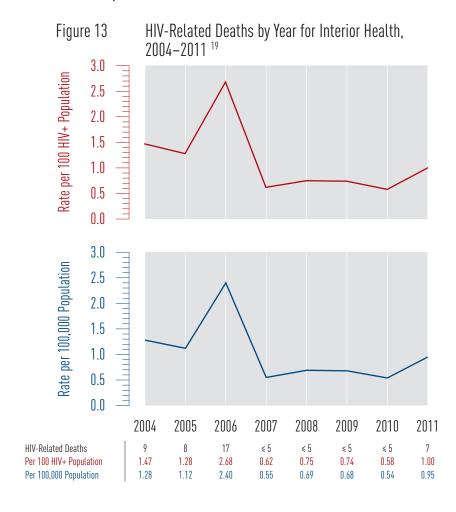
Data Source: DTP AIDS cases are obtained from the Drug Treatment Program Database; BCCDC AIDS cases are obtained from the BC-CDC; CD4<200 at ART initiation data came from the DTP database.

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:

<sup>19</sup> Data Source: BC Vital Statistics

<sup>1.</sup> DTP participants are designated to an HA based on most current residence provided by the participant.

<sup>2.</sup> Mortality data is updated annually.

<sup>3.</sup> The most recent available data was used.

## **Appendices**

Indicator <b>Episodes</b>	1: Test (thousands)	2011 Q3	Q4	2012 Q1	Q2	Q3	Q4	2013 Q1	Q2	Q3	Q4	2014 Q1	Q2	Q3	Q4	2015 Q1	Q2	Q3	Q4	2016 Q1	Q2
Interior H	ealth	3.9	4.2	4.1	4.0	4.2	4.2	4.4	4.7	4.6	4.9	5.3	5.5	5.6	5.9	6.7	7.0	7.9	9.1	10.5	9.8
Gender	Female	2.0	2.1	2.1	2.0	2.1	2.0	2.1	2.3	2.2	2.4	2.6	2.6	2.7	2.8	3.2	3.5	3.9	4.5	5.1	4.9
	Male	1.8	2.0	2.0	1.9	2.0	2.1	2.2	2.3	2.3	2.4	2.6	2.8	2.7	2.9	3.2	3.4	3.8	4.4	5.1	4.8
	Other	0.1	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	0.1	0.1
Age	< 30	1.4	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	2.6	2.5
	30-39	0.9	0.9	0.9	0.9	0.9	0.9	0.9	1.0	1.0	1.1	1.1	1.2	1.2	1.2	1.4	1.4	1.5	1.6	1.9	1.8
	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	0.8	0.9	1.0	1.1	1.1	1.3	1.5	1.4
	≥ 50	0.9	1.0	1.0	1.1	1.1	1.1	1.2	1.4	1.3	1.5	1.5	1.7	1.7	1.8	2.2	2.5	2.8	3.7	4.3	4.0
POC HIV (not in the		34	21	26	28	20	27	38	38	40	37	93	86	163	202	180	130	179	188	160	93
East Koote	enay	0.3	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	0.9	0.8
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	0.2	0.3	0.3	0.4	0.4	0.4	0.4
Male		0.1	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	0.4	0.4
Kootenay	Boundary	0.4	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	1.1	1.1
Female		0.2	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	0.6	0.6
Male		0.2	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	0.5	0.5
Okanagan	1	2.1	2.3	2.3	2.3	2.3	2.1	2.2	2.5	2.4	2.5	2.7	2.8	2.8	2.9	3.4	3.4	4.0	4.6	5.1	4.8
Female		1.1	1.1	1.2	1.2	1.2	1.0	1.0	1.2	1.1	1.2	1.3	1.3	1.4	1.4	1.6	1.7	1.9	2.2	2.4	2.2
Male		1.0	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	2.5	2.4
Thompson Shuswap	n Cariboo	1.0	1.0	1.1	1.0	1.1	1.2	1.3	1.4	1.3	1.4	1.6	1.7	1.7	1.8	2.0	2.2	2.3	2.8	3.4	3.2
Female		0.5	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	1.7
Male		0.5	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	1.6	1.5

Indicator 2: Rate of HIV Testing per 100,000

		2009	2010	2011	2012	2013	2014	2015
All Interior	Health	2026.8	2072.9	2094.1	2180.4	2573.0	2996.7	4080.7
East Kooten	East Kootenay		1834.5	1706.4	1802.9	2005.7	2429.4	3438.5
Kootenay Bo	oundary	2358.9	2312.9	2260.6	2217.9	2656.8	3273.2	4698.5
Okanagan		2166.7	2162.3	2194.5	2325.5	2637.1	2974.4	4015.3
Thompson (	Cariboo Shuswap	1826.1	1929.3	2011.2	2068.4	2640.2	3133.4	4193.6
Gender	Female	1987.0	1993.8	2041.2	2140.6	2554.4	2958.5	4110.7
	Male	1933.1	2008.1	2045.2	2171.6	2548.4	2994.6	3986.1
Age	< 30	2333.3	2343.6	2389.3	2360.8	2656.2	3071.6	3687.3
	30-39	3925.4	4131.2	4196.1	4319.5	4944.8	5595.2	6841.9
	40-49	2479.2	2521.0	2592.0	2774.7	3185.0	3724.9	4904.9
	≥ 50	1107.8	1164.0	1175.9	1340.1	1757.5	2114.1	3444.8

Indicator 3	: Nev	w HI	VΓ	Diagi	nose	es			2011 Q3		201 Q1		Q	3 (		013 Q1	Q2	Q3		2014 Q1		Q3		2015 Q1	Q2	Q3	Q4	2016 Q1	Q2
Interior He	alth		]	By C	llien	t Re	side	nce	3	3	5	5 5		0	2	1	4	4	4	3	4	4	6	5	1	7	6	6	3
			1	Ву Р	rovi	der 1	Addr	ess	2	3	5	5 5		0	2	1	5	4	4	3	2	4	5	5	1	7	6	6	3
Gender			]	Fem	ale				1	0	1	. 1		0	0	0	1	1	1	1	0	1	0	0	0	0	2	1	1
			]	Male	9				2	3	4	4	:	0	2	1	3	3	3	2	4	3	6	5	1	7	4	5	2
Age				< 30					0	0	1	. 1		0	1	0	0	2	2	0	1	1	2	2	0	5	2	2	0
			;	30-3	39				3	1	1	. 1		0	0	0	0	2	1	0	1	3	2	0	0	0	1	1	0
			4	40-4	19				0	1	2	2 1		0	1	0	1	0	1	1	1	0	1	1	1	2	1	3	1
				≥ 50					0	1	1	. 2	,	0	0	1	3	0	0	2	1	0	1	2	0	0	2	0	2
Exposure			1	MSN	Л				1	2	. 1	. 2		0	1	0	0	2	1	0	2	2	5	3	0	4	3	-	_
				PWI	D				1	0	2	1		0	1	0	0	0	0	2	1	1	0	0	0	1	0	-	_
				HET					1	1	2	2 2	2	0	0	1	2	2	3	1	1	1	1	2	0	2	3	-	_
			(	Othe	er				0	0	(	0	)	0	0	0	0	0	0	0	0	0	0	0	0	0	0	-	_
				NIR.	/Un	knov	wn		0	0	(	0	)	0	0	0	2	0	0	0	0	0	0	0	1	0	0	-	_
East Kooter	nay		1	By C	Clien	ıt Re	side	nce	0	0	2	2 0	)	0	0	0	1	0	0	0	0	0	0	1	0	0	0	0	0
				Ву Р	rovi	der 1	Addı	ress	0	0	) 2	2 0	)	0	0	0	1	0	0	0	0	0	0	1	0	1	1	0	0
Kootenay B	Sound	dary		By C	Clien	ıt Re	side	nce	1	0	]	. 2	2	0	0	0	1	0	0	0	1	0	0	1	0	0	0	2	0
				Ву Р	rovi	ider 1	Addı	ress	0	0	j	! 1		0	0	0	1	0	0	0	1	0	0	1	0	0	0	1	0
Okanagan				By C	Clien	ıt Re	side	nce	0	3	1	. 3	;	0	1	1	0	3	2	2	1	2	2	1	0	4	4	2	0
				Ву Р	rovi	der 1	Addr	ress	1	3	1	4	!	0	1	1	1	3	2	2	0	2	3	0	0	4	4	3	0
Thompson	Caril	000	]	By C	llien	ıt Re	side	nce	2	0	1	. 0	)	0	1	0	2	1	2	1	2	2	4	2	1	3	2	2	3
Shuswap				Ву Р	rovi	der 1	Addr	ress	1	0	1	0	)	0	1	0	2	1	2	1	1	2	2	3	1	2	1	2	3
	_						_																						
Indicator 4:						ion a										ı		• •		ı							4.0		
		iterio '12				'11		emal '13		<b>'</b> 15	<b>'</b> 11	'12	Iale '13		<b>'</b> 15	<b>'11</b>		30 ye '13		<b>'</b> 15			9 yea: 13 '1	rs .4 '1!	5 '1		-49 y		<b>'</b> 15
Stage 0	2	0	1	5	0	0	0	0	1	0	2	0	1	4	0	0			3	0	2	0	0		_	0 0		0	0
Stage 1	1	1	3	3	3	0	0	1	1	0	1	1	2	2		0			0	2	0	0	0			0 0		0	0
Stage 2a	0	1	1	2	1	0	0	1	0	0	0	1	0	2		0			0	1	0	0	0			0 1		0	0
Stage 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
Stage 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
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	5 2	1	4
	1					1																							
	<b>'11</b>	≥ 50 '12			<b>'</b> 15	'11		MSM '13		<b>'</b> 15	'11		NII '13		'15			erose					Expos	ure 4 '1!			Unki		
Stage 0	0	0	0	1	0	1	0	1	4	0	0	0	0	0					1	0	0	0	0			0 0		0	0
Stage 1	1	0	0	2	0	0	0	0	0	3	0	1	0	2					1	0	0	0	0			0 0		0	0
Stage 2a	0	0	0	1	0	0	0	0	1	1	0	1	0	0					1	0	0	0	0			0 (			0
Stage 2b	1	1	1	1	0	0	1	2	1	2	0	2	0	0					1	1	0	0	0			0 (		0	0
Stage 20 Stage 3	0	2	3	0	2	2	3	1	1	3	1	0	0	2						5	0	0	0			0 (		0	1
Unknown	0	0	0	0	1		0		0	1	0	0	0	0						1	0	0	0			1 (			0
Total	2	3	4	5	3		4		7			4	0	4						7	0	0	0			1 (			1
10(a)	4	3	7	3	5	3	4	4	,	10	1	7	U	4	1	4	. 4	. 0	4	/	U	J	v	U	۱	. (	, 4	U	1

Age Category Mand MSM Status	< 30 30–39 40–49 ≥ 50 MSM Non-MSM	$< 30$ $30-39$ $40-49$ $\geq 50$ $< 30$ $30-39$ $40-49$ $\geq 50$ $< 30$	617 35 67 134 381 13 13 26 122 6 20 57 145	603 31 65 133 374 13 13 25 121 6 19 57	503 26 51 109 317 12 11 23 108 5		472 21 47 100 304 10 11 20 105 4	434 17 45 87 285 9 11 19	354 10 37 66 241 5 9 17
3 4 ≥ Age Category Mand MSM Status	60–39 40–49 ≥ 50 MSM Non-MSM	$30-39$ $40-49$ $\geq 50$ $< 30$ $30-39$ $40-49$ $\geq 50$	67 134 381 13 13 26 122 6 20 57	65 133 374 13 13 25 121 6	51 109 317 12 11 23 108 5		47 100 304 10 11 20 105	45 87 285 9 11 19	37 66 241 5 9 17
4 ≥ Age Category M and MSM Status	10–49 ≥ 50 MSM Non-MSM	$30-39$ $40-49$ $\geq 50$ $< 30$ $30-39$ $40-49$ $\geq 50$	134 381 13 13 26 122 6 20 57	133 374 13 13 25 121 6 19	109 317 12 11 23 108		100 304 10 11 20 105	87 285 9 11 19	666 241 5 9
≥ Age Category Mand MSM Status	≥ 50 MSM Non-MSM	$30-39$ $40-49$ $\geq 50$ $< 30$ $30-39$ $40-49$ $\geq 50$	381 13 13 26 122 6 20 57	374 13 13 25 121 6 19	317 12 11 23 108 5		304 10 11 20 105	285 9 11 19 101	241 5 9 17
Age Category Mand MSM Status	MSM Non-MSM	$30-39$ $40-49$ $\geq 50$ $< 30$ $30-39$ $40-49$ $\geq 50$	13 13 26 122 6 20 57	13 13 25 121 6 19	12 11 23 108 5		10 11 20 105	9 11 19 101	5 9 17
and MSM Status	Non-MSM	$30-39$ $40-49$ $\geq 50$ $< 30$ $30-39$ $40-49$ $\geq 50$	13 26 122 6 20 57	13 25 121 6 19	11 23 108 5		11 20 105	11 19 101	9 17
Status		$40-49$ $\geq 50$ < 30 $30-39$ $40-49$ $\geq 50$	26 122 6 20 57	25 121 6 19	23 108 5		20 105	19 101	17
Ν		≥ 50 < 30 30-39 40-49 ≥ 50	122 6 20 57	121 6 19	108 5		105	101	
		< 30 30-39 40-49 ≥ 50	6 20 57	6 19	5				86
		30-39 40-49 ≥ 50	20 57	19			4		
U	Jnknown	40–49 ≥ 50	57		17			4	2
U	Jnknown	≥ 50		57			15	13	10
U	Jnknown		145		49		45	37	26
τ	Jnknown	< 30		143	124		118	108	88
			17	13	8		7	4	3
		30-39	34	33	23		21	21	18
		40-49	51	51	38		35	31	23
		≥ 50	115	110	85		81	76	67
Gender M	Male		479	469	395		369	343	280
F	Female		139	134	107		103	91	74
Injection P	PWID		187	186	160		150	135	106
Drug Use	Non-PWID		304	300	267		250	236	196
U	Jnknown		127	117	75		72	63	52
MSM Status M	MSM		173	172	155		146	140	117
N	Non-MSM		227	224	194		182	162	126
U	Jnknown		217	207	154		144	132	111
Health E	East Kootenay	7	40	38	23		21	19	17
A (1 *)	Kootenay Bou		65	65	54		52	47	41
	) Okanagan	,	317	310	269		254	239	195
	Thompson Ca	riboo	199	192	158		145	129	101
	Shuswap								
Indicator 6: <b>Prog</b>	rammatic C	ompliance Sco	re (PCS)						
	,	2014		2015				2016	
		Q3	Q4	Q1	Q2	Q3	Q4		
< 3 CD4 Tests		35.3%	33.3%	31.6%	22.2%	16.7%	15.0%		
< 3 Viral Load Te		23.5%	22.2%	21.1%	11.1%	5.6%	10.0%	10.0%	
No Baseline Gen		5.9%	5.6%	5.3%	5.6%	0.0%	0.0%	5.0%	
Baseline CD4 < 2	•	29.4%	38.9%	36.8%	27.8%	50.0%	35.0%	35.0%	
Non-Recommend		11.8%	5.6%	5.3%	5.6%	0.0%	5.0%	5.0%	
Non Viral suppre	ession at 9 Mo		44.4%	36.8%	22.2%	22.2%	20.0%	30.0%	
PCS Score: 0		5	6	7	8	7	9	8	
PCS Score: 1		6	5	5	6	7	8	8	
PCS Score: 2		1	2	3	2	3	1	3	
PCS Score: 3		2	2	1	1	0	1	0	
PCS Score: 4 or n Total (n=)	nore	3 17	3 18	3 19	1 18	1 18	1 20	1 20	

Indicator 7: New DT	P ARV Participants									
	2014 Q3	Q4		2015 Q1	Q2	Q3	Q	4	2016 Q1	Q2
First Starts	5	5		8	5	5		7	4	6
Experienced Starts	5	6		13	8	11	1	5	5	11
Indicator 8: CD4 Ce	ll Count at ART Initiati	on for ARV	-Naïve	DTP Partie	rinants					
CD4 ≥ 500		-	THUITE	1		_		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 Median (cells/µ)	L) –	_		310	_	_	19		_	545
Total (n=)	≤ 5	≤ 5		7	≤ 5	≤ 5		7	≤ 5	6
Indicator 9: Active a	nd Inactive DTP Partici	inants								
Active DTP Participa		428		441	444	455	47	8	474	477
Inactive DTP Partici		80		81	80	79	7		83	86
Indicator 10: Antiret	troviral Adherence									
≥ 95%	4	_		7	_	3		5	6	5
80% to < 95%	2	_		1	_	2		2	2	0
40% to < 80%	1	_		2	_	2		0	2	2
< 40%	0	_		0	_	1		0	0	0
Total (n=)	7	≤ 5		10	≤ 5	8		7	10	7
T 1: 4 11 D 14	Tract In to									
	nce Testing and Results 293	254		304	248	281	25	2	294	284
Suppressed				38	23				38	29
Wild Type	39	32				41	3			
Never Genotyped 1-Class	2	2		3 7	1	3 7		2	0	1
2-Class	8	8		1	9	2		5 1	9	4
3-Class	6 1	1		2	3	1		0	1 0	0
Total (n=)	349	300		355	284	335	29		342	318
	D C 1 711	•••	•		2010		2012	2012		2015
Indicator 12: AIDS-I	Cases	2007 18	2008	2009 ≤ 5	2010	2011	2012	2013	2014 7	2015 8
ART initiation	Rate per 100,000	2.6	1.5	≥ 5 0.6	8 1.1	1.5	1.3	1.3	1.0	1.1
AIDS Cases	Cases	9	9	6	≤ 5	≤ 5	≤ 5	≤ 5	6	≤ 5
(DTP Reports)	Rate per 100,000	1.3	1.3	0.8	0.1	0.7	0.7	0.6	0.8	0.4
AIDS Cases	Cases	6	11	7	≤ 5	≤ 5	6	≤ 5	6	_
(BCCDC Reports)	Rate per 100,000	0.9	1.5	1.0	0.3	0.6	0.8	0.6	0.8	-
Indicator 13: HIV-R	elated Mortality	2004	2005	2006	2007	2008	2009	2010	2011	
Interior Health	ciated Mioriality	9	8	17	≤ 5	<u>2008</u> ≤ 5	≤ 5	≤ 5	7	
Per 100 HIV+ Popul	ation	1.47	1.28	2.68	0.62	0.75	0.74	0.58	1.00	
Per 100,000 Populati		1.28	1.12	2.40	0.55	0.69	0.68	0.54	0.95	
				2.10	2.00	2.02			3.70	