

# HIV MONITORING QUARTERLY REPORT

FOR INTERIOR HEALTH

THIRD QUARTER 2013

UPDATED VERSION: NOV 28, 2014 \*

\* See foreword

















#### 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 HAS. 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.

<sup>\*</sup> Please note that for Q2 and Q3 2013 reports, a coding revision resulted in data display errors in Indicator 5, the Cascade of Care (in Figures 5.1–5.7 on pp. 20–25 in this report), which has been updated; and, only for Q3 2013 reports, Appendix Table for Indicator 5 (on p. 35 in this report). All other figures and reports remain accurate. Please discard any previous reports and use this updated version. If you have any questions, please contact Irene Day at iday@cfenet.ubc.ca.

# List of Indicators

Indicator 1. 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 Starts

Indicator 8. CD4 Cell Count at ART Initiation

Indicator 9. Active and Inactive Drug Treatment Program Participants

Indicator 10. Antiretroviral Adherence Level

Indicator 11. Resistance Testing Results by Resistance Category

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
Figure 1.1	HIV Test Episodes for Interior Health, 2009 Q1–2013 Q3
Figure 1.2	HIV Test Episodes for Interior Health by Gender and Prenatal Status, 2009 Q1–2013 Q3
Figure 1.3	HIV Test Episodes for Interior Health by Age Category, 2009 Q1–2013 Q3
Figure 1.4	Point-of-Care HIV Tests for Interior Health, 2010 Q4–2013 Q3
Figure 1.5	HIV Test Episodes by HSDA for Interior Health, 2009 Q1–2013 Q3
Indicator 2	HIV Testing Rates
Figure 2.1	Rate of HIV Testing for Interior Health and HSDA's, 2009–2012
Figure 2.2	Rate of HIV Testing for Interior Health by Gender, 2009–2012
Figure 2.3	Rate of HIV Testing for Interior Health by Age Category, 2009–2012
Indicator 3	New HIV Diagnoses
Figure 3.1	New HIV Diagnoses for Interior Health, 2009 Q1–2013 Q3
Figure 3.2	New HIV Diagnoses for Interior Health by Gender, 2009 Q1–2013 Q3
Figure 3.3	New HIV Diagnoses for Interior Health by Age Category, 2009 Q1–2013 Q3
Figure 3.4	New HIV Diagnoses for Interior Health by Exposure Category, 2009 Q1–2012 Q4
Figure 3.5	New HIV Diagnoses for Interior Health by HSDA, 2009 Q1–2012 Q4
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
Figure 4.1	Stage of HIV Infection at Diagnosis for Interior Health, 2010–2012
Figure 4.2	Stage of HIV Infection at Diagnosis for Interior Health by Gender, 2010–2012
Figure 4.3	Stage of HIV Infection at Diagnosis for Interior Health by Age Category, 2010–2012
Figure 4.4	Stage of HIV Infection at Diagnosis for Interior Health by Exposure Category, 2010–2012
Indicator 5	HIV Cascade of Care
Figure 5.1	Estimated Cascade of Care for Interior Health, Year Ending 2013 Q3
Figure 5.2	Estimated Cascade of Care for Interior Health by Gender, Year Ending 2013 Q3
Figure 5.3	Estimated Cascade of Care for Interior Health by Age Category, Year Ending 2013 Q3
Figure 5.4	Estimated Cascade of Care for Interior Health by Msm Status, Year Ending 2013 Q3

Figure 5.5 Estimated Cascade of Care for Interior Health by Age Category and Msm Status, Year Ending 2013 Q3 Figure 5.6 Estimated Cascade of Care for Interior Health by History of IDU, Year Ending 2013 Q3 Figure 5.7 Estimated Cascade of Care for Interior Health by HSDA, Year Ending 2013 Q3 **Indicator 6 Programmatic Compliance Score (PCS)** Probability of Mortality Based on the Programmatic Compliance Score Table 2 Figure 6.1 Pcs Components for Interior Health, 2011 Q4-2013 Q3 First-Year CD4 Measurement First-Year VL measurement Baseline Resistance Testing Recommended Highly Active Antiretroviral Therapy (HAART) Baseline CD<sub>4</sub>  $\geq$  200 cells/ $\mu$ L Suppression at 9 Months Historical Trends for Pcs Score for Interior Health, 2011 Q4-2013 Q3 Figure 6.2 **Indicator 7** New Antiretroviral Therapy Starts in Interior Health Figure 7 BC-CfE Drug Treatment Program Enrollment: New Antiretroviral Participants for Interior Health, 2011 Q4-2013 Q3 **Indicator 8 CD4 Cell Count at ART Initiation** Figure 8 CD4 Cell Count at ART Initiation for Interior Health, 2011 Q4-2013 Q3 Indicator 9 Active and Inactive Drug Treatment Program (DTP) Participants Table 3 Distribution of People on ART for Interior Health, 2013 Q3 Figure 9 Active and Inactive DTP Participants for Interior Health, 2011 Q4-2013 Q3 Indicator 10 **Antiretroviral Adherence** Distribution of Individuals by Adherence Level in 1st Year of Therapy, Figure 10 Based on Pharmacy Refill Compliance for Interior Health, 2011 Q4-2013 Q3 Indicator 11 **Resistance Testing and Results** Figure 11 Cumulative Resistance Testing Results by Resistance Category for Interior Health, 2011 Q4-2013 Q3 Indicator 12 **AIDs-Defining Illness** Figure 12 AIDS Case Rate and Reports for Interior Health, 2005–2012 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. Lillian Lourenco writes and compiles the monitoring report. Guillaume Colley, Dr. Viviane Lima and Nada Gataric perform analysis of Indicators 5–13. James Nakagawa is responsible for publishing and editing. This report was conceived and guided by Dr. Julio Montaner.



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

The expansion to a province-wide programme was announced on November 30th 2012 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 in Interior Health, 2009 Q1–2013 Q3

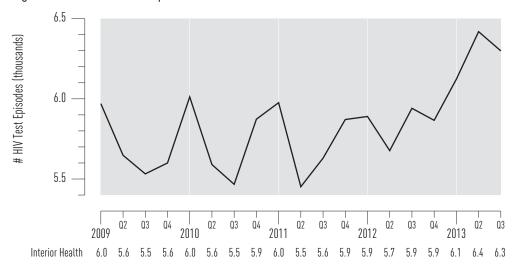
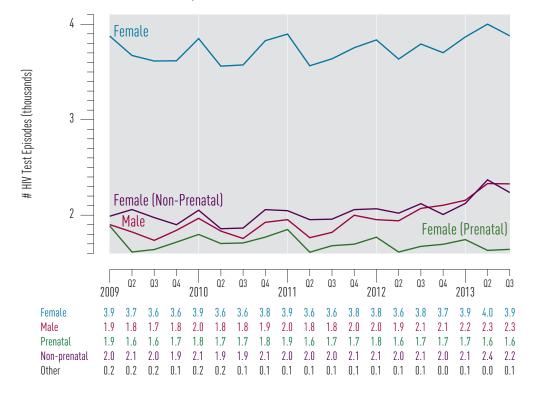


Figure 1.2 HIV Test Episodes by Gender and Prenatal Status in Interior Health, 2009 Q1–2013 Q3 <sup>1</sup>

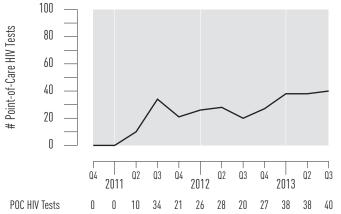


<sup>1</sup> NB: Testing does not include point of care tests.

Figure 1.3 HIV Test Episodes by Age Category for Interior Health, 2009 Q1–2013 Q3  $^{1.2}$ 



Figure 1.4 Point-of-Care HIV Tests in Interior Health, 2010 Q4–2013 Q3



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

#### Limitations:

- 1 Repeat tests in individuals who test using various identifiers may not be identified and these individuals may be counted more than once.
- 2 Poc testing data is available from the fourth quarter of 2010 and onwards.

HIV Test Episodes by Health Service Delivery Area in Interior Health, 2009 Q1–2013 Q3  $\,$ 3.4 -Okanagan 3.2 3.0 -2.8 2.6 2.4 # HIV Test Episodes (thousands) 2.2 2.0 1.8 Thompson Cariboo Shuswap 1.6 1.4 1.2 1.0 **Kootenay Boundary** 0.8 0.6 East Kootenay Q2 Q3 04 Q2 Q3 Q4 2010 2012 2011 2009 **Kootenay Boundary** 0.7 0.6 0.6 0.6 8.0 0.6 0.6 0.8 0.7 0.6 0.6 0.5 East Kootenay 0.6 0.5 0.5 0.6 0.5 0.5 0.6 Okanagan 3.0 2.9 3.1 2.9 2.8 2.9 3.1 2.8 3.0 3.2 3.2 3.1 3.2 3.0 3.1

1.6 1.5 1.6

1.5 1.6

Figure 1.5

Thompson Cariboo Shuswap 1.5 1.5 1.4 1.3 1.7 1.5 1.5 1.6 1.7 1.6

# Indicator 2. HIV Testing Rates

Figure 2.1 Rate of HIV Testing in Interior Health and HSDAs, 2009–2012 <sup>1</sup> 3200 Kootenay Boundary HIV Testing Episodes per 100,000 Population Okanagan 3000 All HSDAs 2800 Thompson Cariboo Shuswap East Kootenay 2600 2011 2009 2010 2012 2890.4 2926.0 2929.2 2967.9 Interior Health 3078.0 **Kootenay Boundary** 3096.6 2984.2 2910.6 East Kootenay 2641.5 2764.6 2621.2 2661.1 3040.6 3128.9 Okanagan 3001.2 3029.9

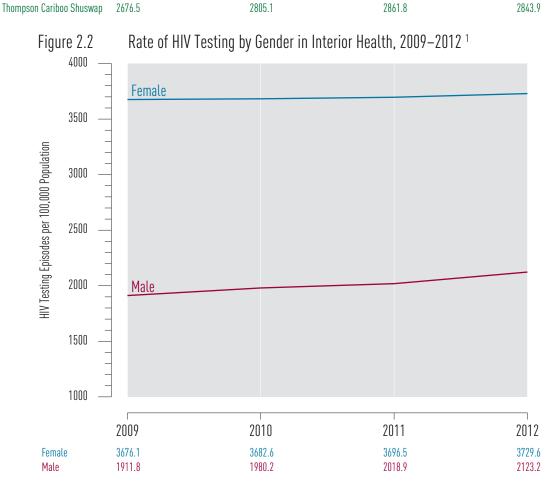
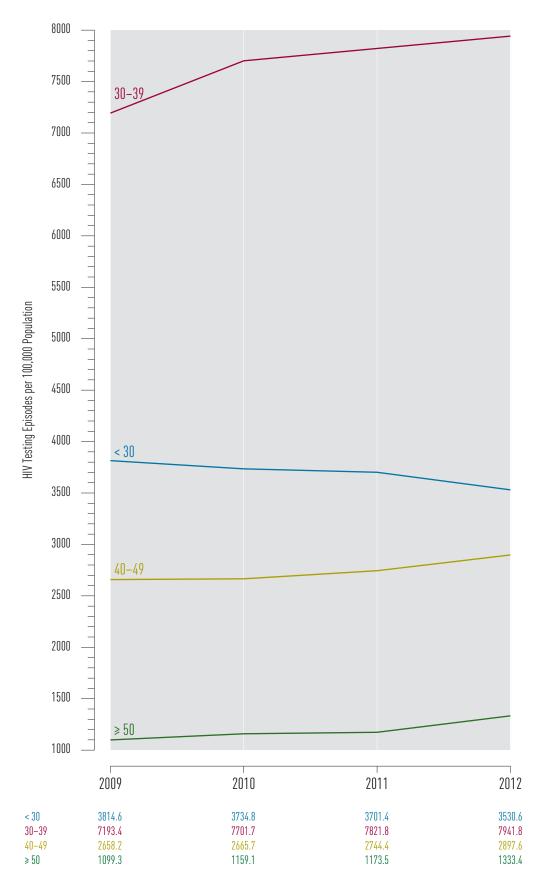


Figure 2.3 Rate of HIV Testing by Age Category in Interior Health, 2009–2012 <sup>1</sup>

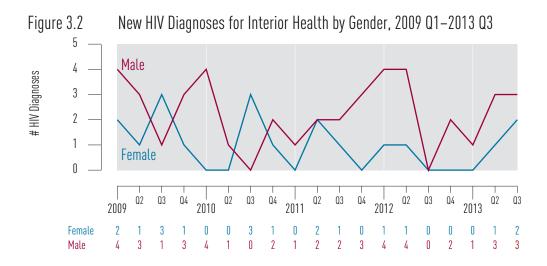


# New HIV Diagnoses

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 3.1 New HIV Diagnoses for Interior Health, 2009 Q1-2013 Q3 3 # HIV Diagnoses Q3 By Client Residence Interior Health By Provider Address



<sup>3</sup> Data Source: BCCDC

Figure 3.3 New HIV Diagnoses for Interior Health by Age Category, 2009 Q1-2013 Q3 HIV Diagnoses 4 3 30-39 2 < 30 ≥ 50 40-49 Q3 Q3 Q3 Q4 2009 2010 2011 2012 2013 < 30 years 0 2 30-39 years 3 0 2 40-49 years 2 0 3 0 3 0 0 ≥ 50 years 0 0 0 0 0 3 0 New HIV Diagnoses for Interior Health by Exposure Category, 2009 Q1-2013 Q3  $^4$ Figure 3.4 HET # HIV Diagnoses NIR/Unknown Other MSM Q2 Q3 Q2 Q3 Q2 Q2 Q4 2009 2010 2011 2012 2013 MSM (men having sex with men) 0 0 IDU (injection drug use) 0 0 0 0 0 0 0 2 0 HET (heterosexual contact) 0 0 0 0 Other (other exposure identified) 0 0 0 0 0 0 0 0 0 0 0 0 0 0 NIR/Unknown (no identified exposure) New HIV Diagnoses for Interior Health by HSDA, 2009 Q1-2013 Q3 Figure 3.5 Thompson Cariboo Shuswap **HIV Diagnoses** 3 0 Kootenay Boundary East Kootenay Q2Q3 Q4 Q2Q3 Q3 Q3 2009 2010 2011 By Client Residence 0

Kootenay Boundary

Okanagan

Shuswap

East Kootenay

Thompson Cariboo

By Provider Address

By Client Residence
By Provider Address

By Client Residence
By Provider Address

By Client Residence

By Provider Address

3 0

0 0

0

2

"By Provider Address" is graphed as dashed line in same colour.

<sup>4</sup> BCCDC: Data lags by 6 months.

MSM=men who have sex with men; IDU= injection drug user; 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, laboratory results suggestive of acute HIV infection, and clinical presentation with an AIDS-defining illness (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 r	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		N. AIDC							
2a			CD4 350-499	and	No AIDS case report							
2b	Stage 0		CD4 200-349		торогс							
3	not met	and	( CD4 <200	or	AIDS case report							
Unknown			No available CD4	and	No AIDS case report							

Figure 4.1 Stage of HIV Infection at Diagnosis for Interior Health, 2010–2012 <sup>5</sup>

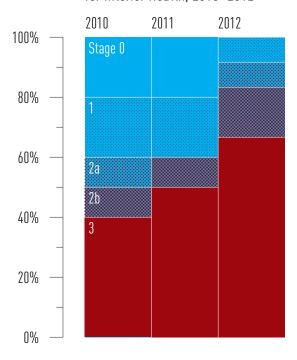
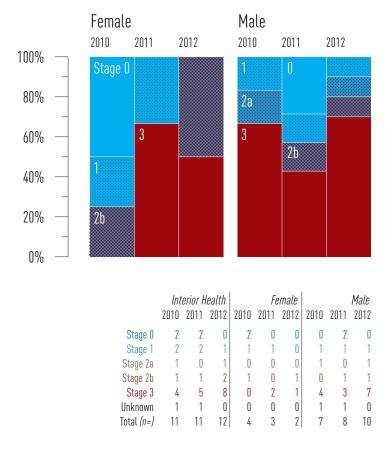


Figure 4.2 Stage of HIV Infection at Diagnosis by Gender for Interior Health, 2010–2012 <sup>5</sup>



Data Source: вссьс

Figure 4.3 Stage of HIV Infection at Diagnosis by Age Category for Interior Health, 2010–2012 <sup>5</sup>

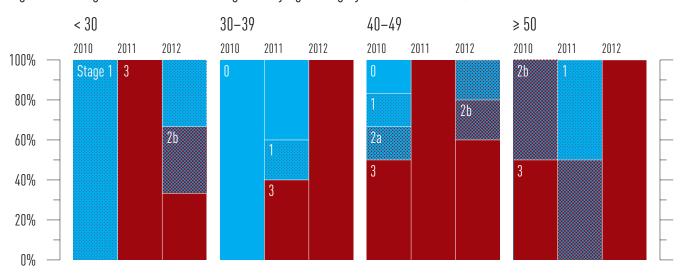
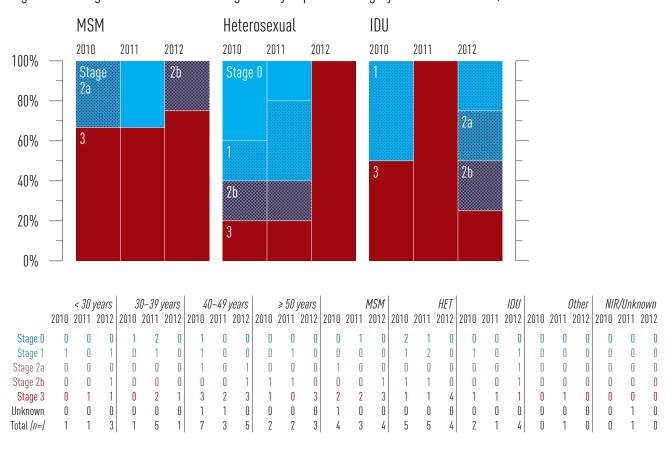


Figure 4.4 Stage of HIV Infection at Diagnosis by Exposure Category for Interior Health, 2010–2012 5,6



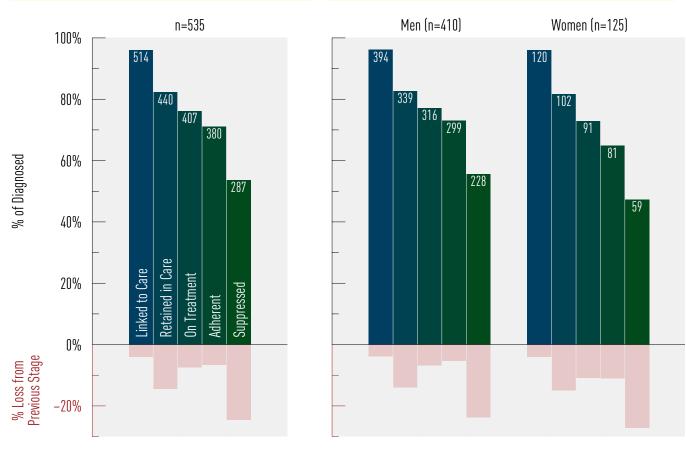
<sup>6</sup> MSM=men who have sex with men; IDU= injection drug user; HET=heterosexual. NIR=No identified risk/exposure.

#### 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. Linkage to HIV care, 3. Retention in HIV care, 4. On ART and 5. Achieving a suppressed VL; collectively, they are referred to as the cascade of care. Leakage 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 (ie. 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 year 2012 in BC overall and stratified by sex and age for each Health Authority.

Figure 5.1 Estimated Cascade of Care for Interior Health, Year Ending 2013 Q3 7

Figure 5.2 Estimated Cascade of Care for Interior Health by Gender, Year Ending 2013 Q3 8



5,6 Data is for the period 2012 Q4-2013 Q3.

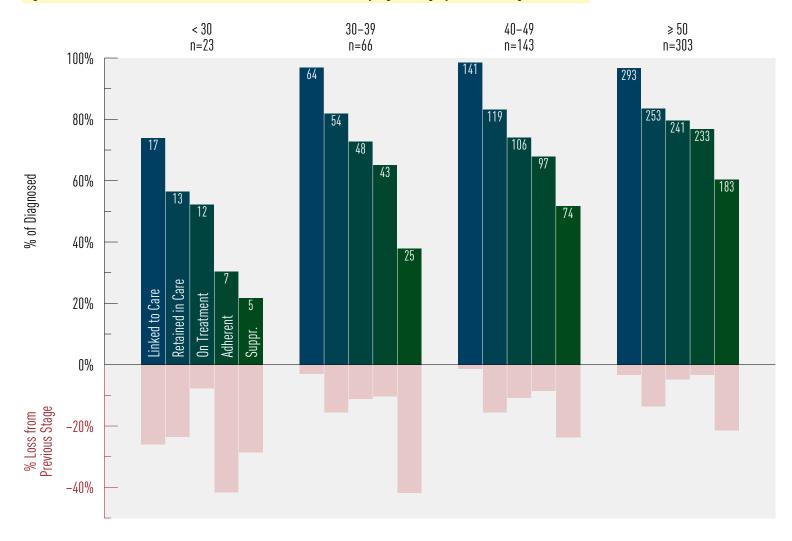
#### Data Sources:

- 1 British Columbia Centre for Excellence Drug Treatment Program (DTP) Database (ARV use, VL and CD4 count).
- 2 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 has been assigned to their biological sex.



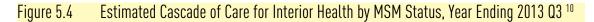


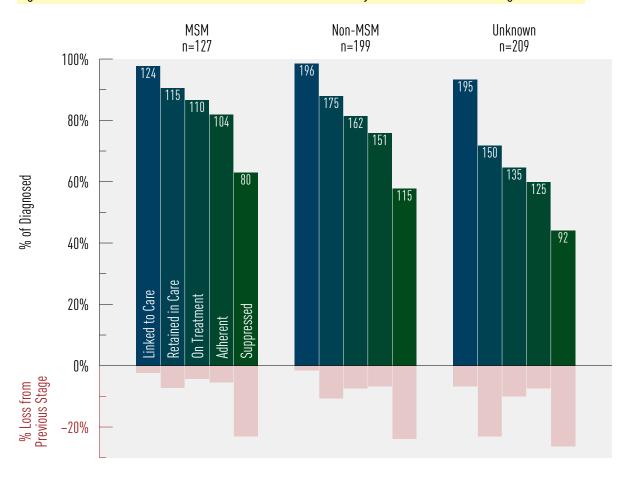
Limitations: HA assignment is based on the most recent HA of residence of the patient, if not available of the HIV-care provider. If the most recent HA of residence is not updated then the designated HA may be incorrect.

<sup>7</sup> Data is for the period 2012 Q4–2013 Q3. Data Sources:

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

<sup>2</sup> Administrative data (ex. MSP billings; hospitalization data from the Discharge Abstract Database (DAD)).





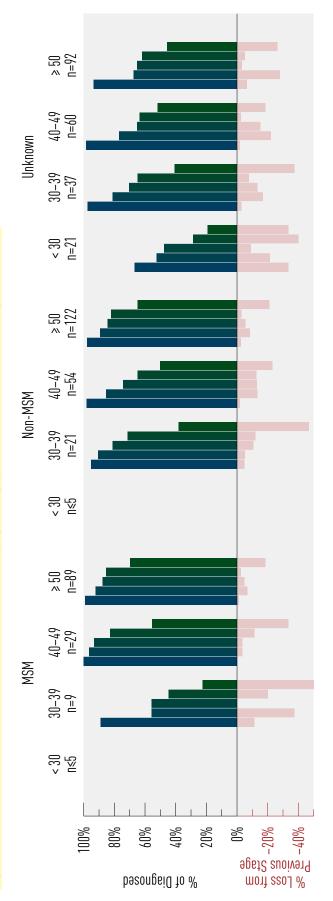
Limitations: HA assignment is based on the most recent HA of residence of the patient, if not available of the HIV-care provider. If the most recent HA of residence is not updated then the designated HA may be incorrect.

<sup>8</sup> Data is for the period 2012 Q4-2013 Q3. Data Sources:

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

<sup>2</sup> 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 2013 Q3 <sup>11</sup>



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.

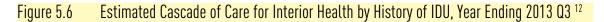
Where  $n \le 5$ , data has been withheld for concerns of statistical significance as well as privacy.

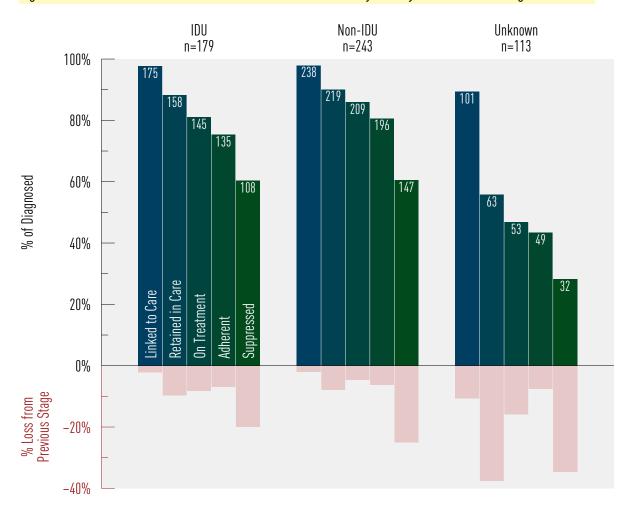
Authorized parties may contact the British Columbia Centre for Excellence in HIV/AIDS to obtain this information.

Data is for the period 2012 Q4-2013 Q3.

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

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



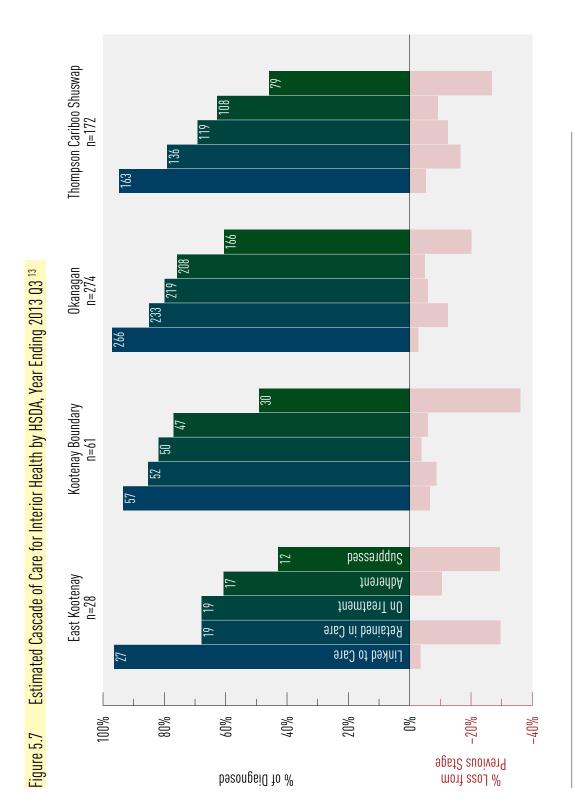


Limitations: HA assignment is based on the most recent HA of residence of the patient, if not available of the HIV-care provider. If the most recent HA of residence is not updated then the designated HA may be incorrect.

<sup>10</sup> Data is for the period 2012 Q4-2013 Q3. Data Sources:

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

<sup>2</sup> Administrative data (ex. MSP billings; hospitalization data from the Discharge Abstract Database (DAD)).



11 Data is for the period 2012 Q4–2013 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).

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

# Indicator 6. The Programmatic Compliance Score (PCS)

The Programmatic Compliance Score (PCS) is a summary measure of risk of future death, immunologic failure and virologic failure from all causes for people who are starting ART for the first time. It is composed of patient-and physician-driven effects. PCS scores range from o−6 with higher scores indicative of poorer health outcomes and greater risk of death. Table 1 provides mortality, immunologic failure and virologic failure probabilities for given PCS scores. We interpret an individual with a PCS≥4 as being 22 times more likely to die, almost 10 times more likely to have immunologic failure and nearly 4 times as likely to demonstrate virologic failure compared to those individuals with a PCS score of o. A detailed description of how the PCS score is calculated and its valida¬tion 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:

- 1. 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. The Probability of Mortality, Immunologic Failure and Virologic Failure based on the Programmatic Compliance Score

Programmatic	Mortality Risk Ratio	Immunologic Failure Risk	Virologic Failure Risk Ratio
Compliance Score	(95% Confidence Interval)	Ratio (95% CI)	(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

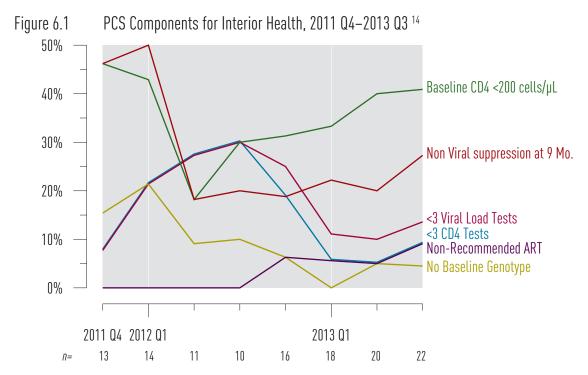


Figure 6.2 Historical Trends for PCS Score for Interior Health, 2011 Q4-2013 Q3  $^{15}$ 100% 80% Distribution of PCS Score 60% 40% 2 Improving Health Outcomes 20% 3 ≥ 4 0% Q4 Q1 Q2 Q3 Q4 Q1 Q2Q3 2012 2011 2013 16 18 20 22 13 14 11 10 NB: A score of o is the best score and a score of 4 or more is the worst score.

<sup>12</sup> Data Source: British Columbia Centre for Excellence Drug Treatment Program (DTP) Database. Limitations: CD4 cell count capture is approximately 80%.

Data Source: British Columbia Centre for Excellence Drug Treatment Program (DTP) Database. Each quarter's data is calculated as the sum of the 4 quarters leading up to it. e.g. 2012 Q1 is calculated from 2011 Q2 – 2012 Q1.

# 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.

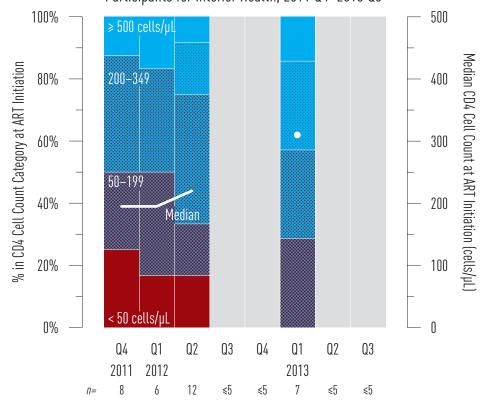
# Indicator 7. New Antiretroviral Therapy Starts in Interior Health

Figure 7 BC-CfE Drug Treatment Program Enrollment: New Antiretroviral Participants for Interior Health, 2011 Q4-2013 Q3 16



Indicator 8. CD4 Cell Count at ART Initiation

Figure 8 CD4 Cell Count at ART Initiation of ART-Naïve DTP Participants for Interior Health, 2011 Q4–2013 Q3 <sup>17</sup>



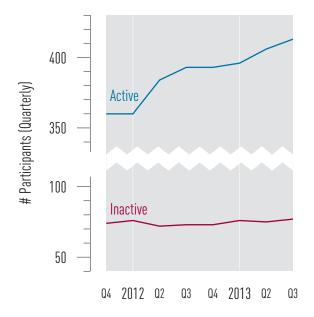
- 14 Data Source: Drug Treatment Program Database Limitation: DTP participants are designated to an HA based on most current residence provided by the participant.
- Data Source: Drug Treatment
  Program Database
  Limitations: CD4 cell count data
  is approximately 80% complete.
  Where n ≤ 5, data has been
  withheld for concerns of statistical significance as well as
  privacy. Authorized parties may
  contact the British Columbia
  Centre for Excellence in HIV/
  AIDS to obtain this information.

# Indicator 9. Active and Inactive DTP Participants

Table 3. Distribution of People on ART for Interior Health, 2013 Q3  $^{16}$ 

Age	< 30	9
	30-39	52
	40-49	113
	≥ 50	239
Gender	Male	321
	Female	92
Exposure	MSM	112
	IDU	144
Total		413

Figure 9 Active and Inactive DTP Participants for Interior Health, 2011 Q4–2013 Q3 <sup>19</sup>



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

#### Definitions:

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

'Unknown/not stated' defined as being on treatment in the current quarter, and city of residence unknown

Active DTP participants: are those who are prescribed one or more drugs in the last six months.

Inactive DTP Participants: Persons no longer prescribed drugs through the HIV/AIDS Drug Treatment Program in the last quarter.

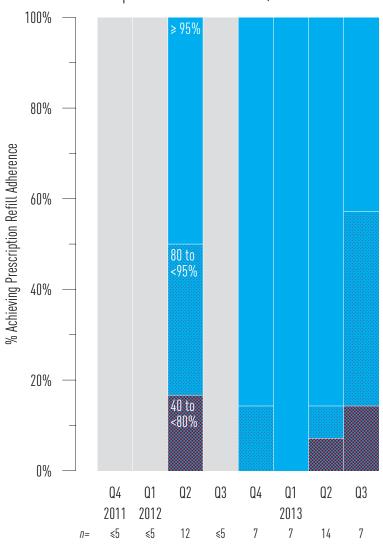
<sup>16</sup> Data Source: Drug Treatment Program Database

## 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

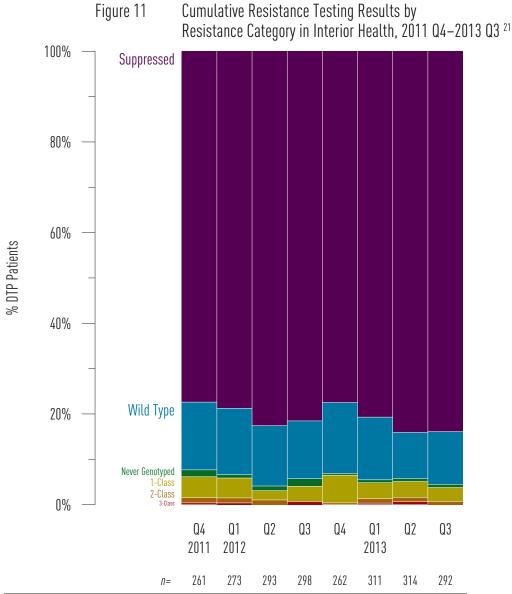
Figure 10 Distribution of Individuals by Adherence Level in 1st Year of Therapy, Based on Pharmacy Refill Compliance for Interior Health, 2011 Q4–2013 Q3 <sup>20</sup>



Data Source: Drug Treatment Program Database Limitation: Prescription refill adherence is used as a proxy for patient adherence. Where n ≤ 5, data has been withheld for concerns of statistical significance as well as privacy. Authorized parties may contact the British Columbia Centre for Excellence in HIV/AIDS to obtain this information.

# 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 or three 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.



19 Data Source: Drug Treatment Program Database

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

# 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; as such, we have plotted DTP reported AIDS cases as well as the proportion of persons initiating ART with a CD4<200 cells/µL.

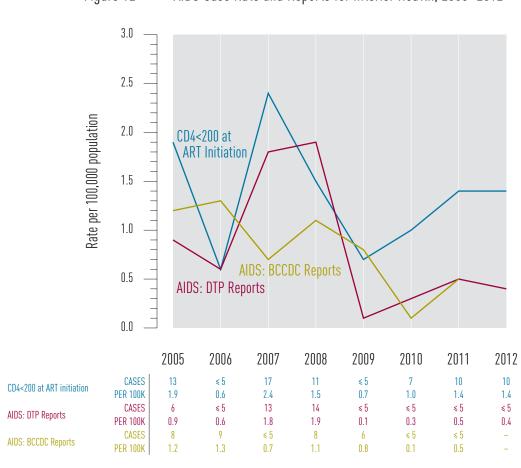


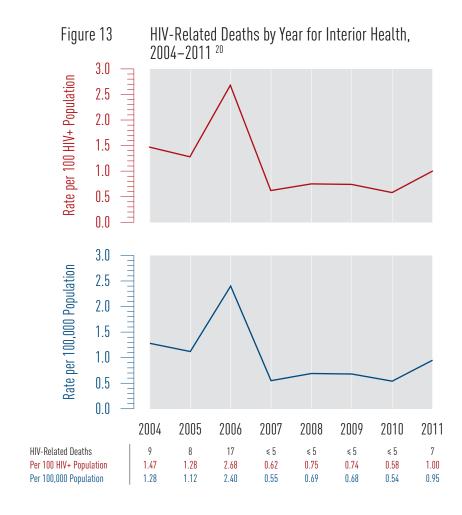
Figure 12 AIDS Case Rate and Reports for Interior Health, 2005–2012 <sup>22</sup>

Limitation: AIDS case reporting was investigated using 2 definitions: First, using AIDS cases reported in AIDS case report forms from the DTP, and second, using a CD4 cell count of <200 cells/ $\mu$ L 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. 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.

<sup>20</sup> Data Source: Drug Treatment Program Database

# 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>21</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		2009				2010				2011				2012				2013		
Episodes	(thousands)	Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4	Q1	Q2	Q3
Interior H	lealth	6.0	5.6	5.5	5.6	6.0	5.6	5.5	5.9	6.0	5.5	5.6	5.9	5.9	5.7	5.9	5.9	6.1	6.4	6.3
Gender	Female	3.9	3.7	3.6	3.6	3.9	3.6	3.6	3.8	3.9	3.6	3.6	3.8	3.8	3.6	3.8	3.7	3.9	4.0	3.9
	Male	1.9	1.8	1.7	1.8	2.0	1.8	1.8	1.9	2.0	1.8	1.8	2.0	2.0	1.9	2.1	2.1	2.2	2.3	2.3
	Other	0.2	0.2	0.2	0.1	0.2	0.2	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.0	0.1	0.0	0.1
Female (P	renatal)	1.9	1.6	1.6	1.7	1.8	1.7	1.7	1.8	1.9	1.6	1.7	1.7	1.8	1.6	1.7	1.7	1.7	1.6	1.6
Female (N	Ion-prenatal)	2.0	2.1	2.0	1.9	2.1	1.9	1.9	2.1	2.0	2.0	2.0	2.1	2.1	2.0	2.1	2.0	2.1	2.4	2.2
Age	< 30	2.7	2.4	2.5	2.3	2.5	2.4	2.4	2.4	2.5	2.2	2.3	2.5	2.3	2.2	2.4	2.3	2.2	2.3	2.4
	30-39	1.7	1.5	1.5	1.5	1.8	1.6	1.6	1.6	1.8	1.6	1.6	1.6	1.8	1.6	1.7	1.6	1.8	1.8	1.8
	40-49	0.8	0.8	0.7	0.7	0.8	0.8	0.7	0.7	0.7	0.7	0.8	0.8	0.8	0.8	0.8	0.7	0.8	0.8	0.7
	≥ 50	0.8	0.9	0.8	1.1	0.9	0.9	0.9	1.2	1.0	0.9	0.9	1.0	1.0	1.1	1.1	1.1	1.3	1.5	1.4
POC HIV	Tests (number	r not in	thous	sands)					0	0	10	34	21	26	28	20	27	38	38	40
Kootenay	Boundary	0.6	0.6	0.6	0.8	0.7	0.6	0.6	0.8	0.7	0.6	0.6	0.6	0.6	0.5	0.6	0.7	0.6	0.6	0.6
East Koote	enay	0.5	0.5	0.5	0.6	0.6	0.5	0.5	0.6	0.5	0.5	0.5	0.6	0.6	0.5	0.6	0.6	0.6	0.6	0.5
Okanagan	l	3.3	3.0	3.0	2.9	3.1	2.9	2.8	2.9	3.1	2.8	3.0	3.2	3.2	3.1	3.2	3.0	3.1	3.3	3.3
Thompson Shuswap	n Cariboo	1.5	1.5	1.4	1.3	1.7	1.5	1.5	1.6	1.7	1.6	1.6	1.5	1.6	1.5	1.6	1.7	1.8	1.9	1.8

Indicator 2: Rate of HIV Testing per 100,000

		2009	2010	2011	2012
Interior He	alth	2 890.4	2 926.0	2 929.2	2 967.9
Kootenay I	Boundary	3 078.0	3 096.6	2 984.2	2 910.6
East Koote	nay	2 641.5	2 764.6	2 621.2	2 661.1
Okanagan		3 040.6	3 001.2	3 029.9	3 128.9
Thompson	Cariboo Shuswap	2 676.5	2 805.1	2 861.8	2 843.9
Gender	Female	3 676.1	3 682.6	3 696.5	3 729.6
	Male	1 911.8	1 980.2	2 018.9	2 123.2
Age	< 30	3 814.6	3 734.8	3 701.4	3 530.6
	30-39	7 193.4	7 701.7	7 821.8	7 941.8
	40-49	2 658.2	2 665.7	2 744.4	2 897.6
	≥ 50	1 099.3	1 159.1	1 173.5	1 333.4

		2009				2010				2011				2012				2013	,	
Indicator 3: New HI	V Diagnoses	Q1	Q2	Q3	Q4		Q2	Q3												
Interior Health	By Client Residence	6	4	4	4	4	1	3	3	1	4	3	3	5	5	0	2	1	4	5
	By Provider Address	6	4	4	4	4	1	2	3	1	3	2	3	5	5	0	2	1	5	5
Gender	Female	2	1	3	1	0	0	3	1	0	2	1	0	1	1	0	0	0	1	2
	Male	4	3	1	3	4	1	0	2	1	2	2	3	4	4	0	2	1	3	3
Age	< 30	0	1	2	1	0	0	0	1	0	1	0	0	1	1	0	1	0	0	2
	30-39	3	0	1	2	0	0	1	0	1	0	3	1	1	1	0	0	0	0	2
	40-49	1	1	1	1	3	1	1	2	0	3	0	1	2	1	0	1	0	1	0
	≥ 50	2	2	0	0	1	0	1	0	0	0	0	1	1	2	0	0	1	3	1
Exposure	MSM	1	2	0	1	2	1	0	1	0	0	1	2	1	2	0	1	0	_	_
•	IDU	1	0	0	0	1	0	0	1	0	0	1	0	2	1	0	1	0	_	_
	HET	4	1	4	2	1	0	3	1	1	2	1	1	2	2	0	0	0	_	_
	Other	0	1	0	1	0	0	0	0	0	1	0	0	0	0	0	0	0	_	_
	NIR	0	0	0	0	0	0	0	0	0	1	0	0	0	0	0	0	1	_	_
Kootenay Boundary	By Client Residence	0	1	0	0	1	0	0	0	0	0	1	0	1	2	0	0	0	1	0
	By Provider Address	0	1	0	0	1	0	0	0	0	0	0	0	1	1	0	0	0	1	0
East Kootenay	By Client Residence	0	1	0	0	3	0	0	0	0	0	0	0	2	0	0	0	0	1	0
•	By Provider Address	0	1	0	0	3	0	0	0	0	0	0	0	2	0	0	0	0	1	0
Okanagan	By Client Residence	2	2	2	2	0	1	1	0	1	3	0	3	1	3	0	1	1	0	3
-	By Provider Address	2	2	2	2	0	1	1	0	1	2	1	3	1	4	0	1	1	1	3
Thompson Cariboo	By Client Residence	4	0	2	2	0	0	2	3	0	1	2	0	1	0	0	1	0	2	2
Shuswap	By Provider Address	4	0	2	2	0	0	1	3	0	1	1	0	1	0	0	1	0	2	2

Indicator 4: Stag	e of HI	V Infe	ction :	at Base	line													
	Inter 2010	rior He 2011	ealth 2012	2010	Female 2011	2012	2010	Male 2011	2012	< 2010	30 yea 2011	rs 2012	30- 2010	-39 yea 2011	rs 2012	40- 2010	-49 yea	ars 2012
Stage 0	2	2	0	2	0	0	0	2	0	0	0	0	1	2	0	1	0	0
Stage 1	2	2	1	1	1	0	1	1	1	1	0	1	0	1	0	1	0	0
Stage 2a	1	0	1	0	0	0	1	0	1	0	0	0	0	0	0	1	0	1
Stage 2b	1	1	2	1	0	1	0	1	1	0	0	1	0	0	0	0	0	1
Stage 3	4	5	8	0	2	1	4	3	7	0	1	1	0	2	1	3	2	3
Unknown	1	1	0	0	0	0	1	1	0	0	0	0	0	0	0	1	1	0
Total	11	11	12	4	3	2	7	8	10	1	1	3	1	5	1	7	3	5
	≥ 2010	50 yea 2011	rs 2012	2010	MSM 2011	2012	Het 2010	erosex	ual 2012	2010	IDU 2011	2012	Other	r Expo	sure 2012	NIR 2010	/Unkn	own 2012
Stage 0	0	0	0	0	1	0	2	1	0	0	0	0	0	0	0	0	0	0
Stage 1	0	1	0	0	0	0	1	2	0	1	0	1	0	0	0	0	0	0
Stage 2a	0	0	0	1	0	0	0	0	0	0	0	1	0	0	0	0	0	0
Stage 2b	1	1	0	0	0	1	1	1	0	0	0	1	0	0	0	0	0	0
Stage 3	1	0	3	2	2	3	1	1	4	1	1	1	0	1	0	0	0	0
Unknown	0	0	0	1	0	0	0	0	0	0	0	0	0	0	0	0	1	0
Total	2	2	3	4	3	4	5	5	4	2	1	4	0	1	0	0	1	0
Indicator 5: HIV	/ Casca	ide of	Care		DIAGN	OSED		LINKE	D	RETAI	NED	(	ON ART	A	DHERE	NT	SUPPR	ESSED
Interior Health						535		51			440		407			380		287
Age Category	< 30					23		1	7		13		12			7		5

40- ≥ 5	0–39 0–49 50 SM < 30	535 23 66 143 303	514 17 64 141	440 13 54 119	407 12 48	380 7 43	287 5 25
30- 40- ≥ 5	0–39 0–49 50 SM < 30	66 143 303	64 141	54	48		
40- ≥ 5	1–49 50 SM < 30	143 303	141			43	25
≥ 5	50 SM < 30	303		119			23
	SM < 30		202		106	97	74
			293	253	241	233	183
0 0 1		≤ 5	≤ 5	≤ 5	≤ 5	≤ 5	≤ 5
and MSM	30-	39 9	8	5	5	4	2
Status	40-	19 29	29	28	27	24	16
	≥ 50	89	88	82	78	76	62
No	on-MSM < 30	≤ 5	≤ 5	≤ 5	≤ 5	≤ 5	≤ 5
	30-	39 21	20	19	17	15	8
	40-	19 54	53	46	40	35	27
	≥ 50	122	119	109	103	100	79
Un	nknown < 30	21	14	11	10	6	4
	30-	39 37	36	30	26	24	15
	40-	49 60	59	46	39	38	31
	≥ 50	92	86	62	60	57	42
Gender Ma	ale	410	394	339	316	299	228
Fer	emale	125	120	102	91	81	59
Injection ID	U	179	175	158	145	135	108
Drug Use No	on-IDU	243	238	219	209	196	147
Un	nknown	113	101	63	53	49	32
MSM Status MS	SM	127	124	115	110	104	80
No	on-MSM	199	196	175	162	151	115
Un	nknown	209	195	150	135	125	92
	ist Kootenay	28	27	19	19	17	12
Authority Ko	ootenay Boundary	61	57	52	50	47	30
Ok	kanagan	274	266	233	219	208	166
	nompson Cariboo nuswap	172	163	136	119	108	79

Indicator 6: <b>Programmatic</b>	•					2012		
	2011 Q4	2012 Q1	Q2	Q3	Q4	2013 Q1	Q2	Q3
< 3 CD4 Tests	7.7%	21.4%	27.3%	30.0%	18.8%	5.6%	5.0%	9.1%
< 3 Viral Load Tests	7.7%	21.4%	27.3%	30.0%	25.0%	11.1%	10.0%	13.6%
No Baseline Genotype	15.4%	21.4%	9.1%	10.0%	6.3%	0.0%	5.0%	4.5%
Baseline CD4 < 200 cells/μL		42.9%	18.2%	30.0%	31.3%	33.3%	40.0%	40.9%
Non-Recommended ART	0.0%	0.0%	0.0%	0.0%	6.3%	5.6%	5.0%	9.1%
Non Viral suppression at 9 l		50.0%	18.2%	20.0%	18.8%	22.2%	20.0%	27.3%
PCS Score: 0	40.270	30.0%	7	20.0%	7	8	20.070	7.5%
PCS Score: 1	4	3	1	2	5	7	9	
PCS Score: 1 PCS Score: 2	3		1		2	2		9
		4		1			4	5
PCS Score: 3	2	2	1	1	1	1	0	0
PCS Score: 4 or more	0	1	1	1	1	0	0	1
Total (n=)	13	14	11	10	16	18	20	22
Indicator 7: New DTP	2011	2012				2013		
ARV Participants	Q4	Q1	Q2	Q3	Q4	Q1	Q2	Q3
First Starts	8	6	12	4	3	7	4	5
Experienced Starts	7	3	13	8	7	7	8	7
Indicator 0 CD4 C 11 C	4 o4 A D'T I '4' 4'	for ADVIN	own Dan p	<b>t.i</b>				
Indicator 8: <b>CD4 Cell Coun</b> CD4 ≥ 500	t at ART Initiation		aive DTP Pa	articipants		1		
CD4 ≥ 500 CD4 350–499		1		_	_		-	_
	0	0	2	_	_	2	_	_
CD4 200-349	3	2	5	_	_	2	_	_
CD4 50–199	2	2	2	_	_	2	_	_
CD4 < 50	2	1	2	_	_	0	-	-
CD4 Median (cells/µL)	195	195	220		_	310	_	_
Total (n=)	8	6	12	≤ 5	≤ 5	7	≤ 5	≤ 5
Indicator 9: Active and Inac	tive DTP Particina	ante						
Active DTP Participants	360	360	385	396	396	398	408	416
Inactive DTP Participants	83	85	81	82	82	85	84	86
inactive D 11 Turticipants	00	00	01	02	02	03	01	00
Indicator 10: Antiretroviral	Adherence							
≥ 95%	_	_	6	_	6	7	12	3
80% to < 95%	_	_	4	_	1	0	1	3
40% to < 80%	_	_	1	_	0	0	1	1
< 40%	_	_	0	_	0	0	0	0
Total (n=)	≤ 5	≤ 5	11	≤ 5	7	7	14	7
Indicator 11: Resistance Tes		21.5	2.12	2.12	200	271	251	2.15
Suppressed	202	215	242	243	203	251	264	245
Wild Type	39	40	39	38	41	43	32	34
Never Genotyped	4	2	3	5	1	2	2	2
1-Class	12	12	6	10	16	11	11	9
2-Class	3	3	3	0	1	3	3	2
3-Class	1	1	0	2	0	1	2	0
Total (n=)	261	273	293	298	262	311	314	292
Indicator 12: AIDS-Definin	a Illness	2005	2006	2007	2008	2009 20	10 2011	2012
	Cases	13	<u>2000</u> ≤ 5	17	11	≤ 5	7 10	10
	Rate per 100,000	1.9	0.6	2.4	1.5		1.0 1.4	1.4
	Cases	6	<ul><li>5</li></ul>	13	1.3		≤5 ≤5	1.4 ≤ 5
	Rate per 100,000	0.9	0.6	1.8	1.9		0.5	0.4
_	Cases	8	9	1.0 ≤ 5	8		$\leq 5 \qquad \leq 5$	-
	Cases Rate per 100,000	1.2	1.3	≤ 3 0.7	8 1.1		$\begin{array}{ccc} 2.5 & \leq 5 \\ 0.1 & 0.5 \end{array}$	
(DCCDC Reports)	Kuie pei 100,000	1.2	1.3	0.7	1.1	0.0	0.5	-
Indicator 13: HIV-Related I	Mortality 2004	2005	2006	2007	2008	2009 20	10 2011	
Interior Health	9	8	17	≤ 5	≤ 5		5 7	
Per 100 HIV+ Population	1.47	1.28	2.68	0.62	0.75		58 1.00	
Per 100,000 Population	1.28	1.12	2.40	0.55	0.69		54 0.95	
1								