

## HIV MONITORING QUARTERLY REPORT

FOR BRITISH COLUMBIA

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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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/ $\mu$ L

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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 British Columbia

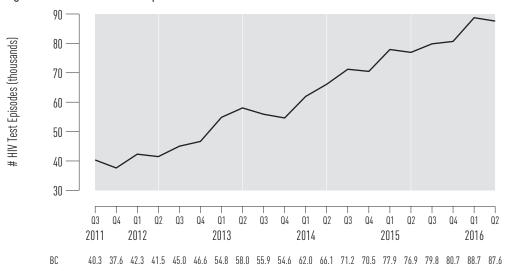


Figure 1.2 HIV Test Episodes by Gender for British Columbia 1,2

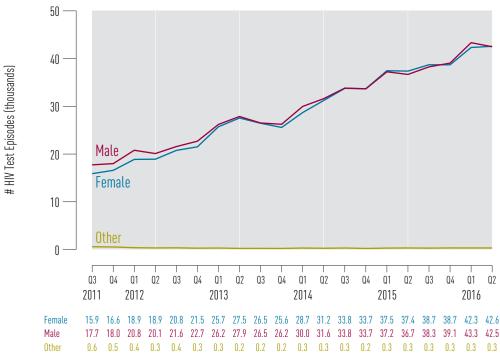
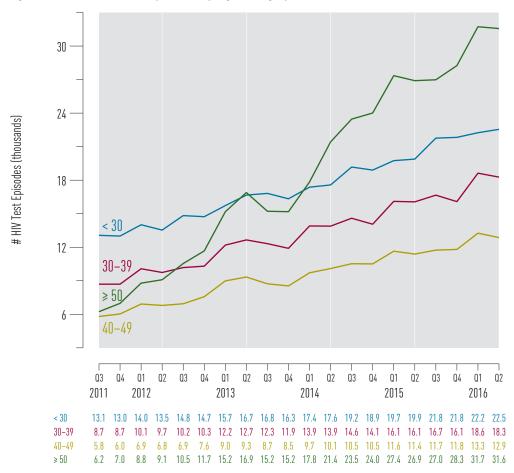
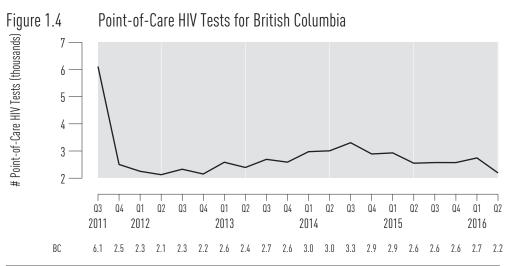


Figure 1.3 HIV Test Episodes by Age Category for British Columbia 1,2



Point-of-Care HIV Tests for British Columbia



Data Source: The BC Public Health Microbiology and Reference Laboratory (BCPHMRL) courtesy of the BC Centre for Disease Control (BCCDC). HIV screening tests conducted by the VIHA Laboratory are not included.

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

Testing does not include point of care tests.

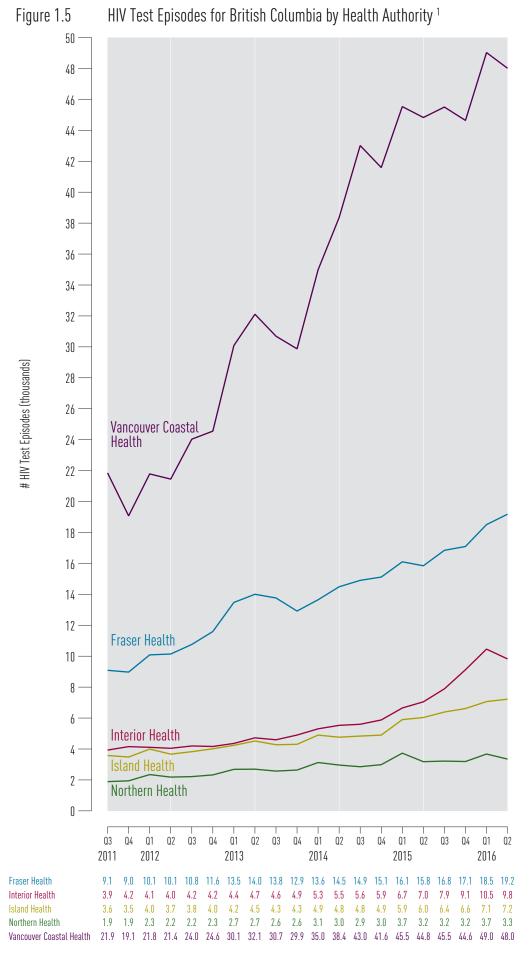


Figure 1.6 HIV Test Episodes for Non-prenatal Females in BC by Health Authority 1.2

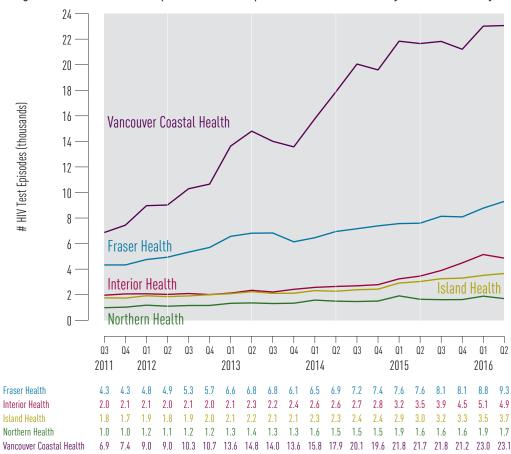
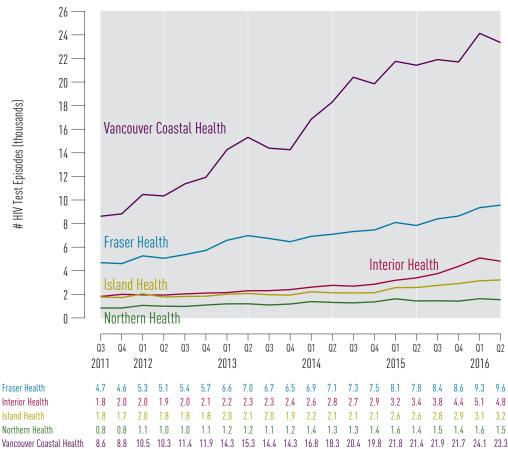


Figure 1.7 HIV Test Episodes for Males in British Columbia by Health Authority 1.2



#### Indicator 2. HIV Testing Rates

Figure 2.1 Rate of HIV Testing for British Columbia and Health Authorities <sup>2</sup>

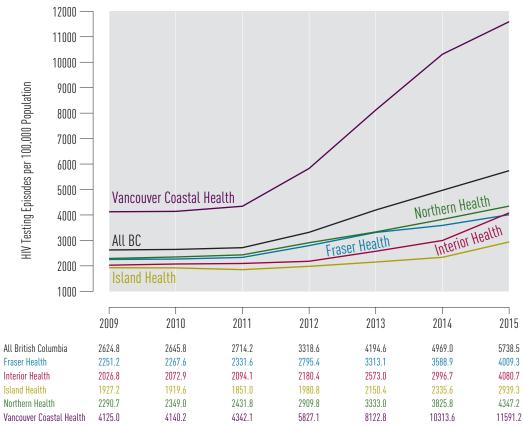
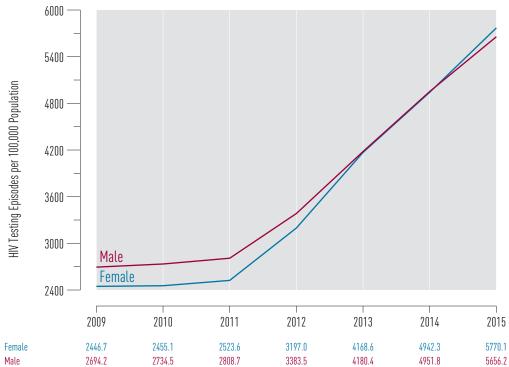


Figure 2.2 Rate of HIV Testing by Gender for British Columbia <sup>2</sup>



9000 8000 7000 HIV Testing Episodes per 100,000 Population 6000 30-39 5000 4000 40-49 3000 < 30 2000 ≥ 50 1000 2010 2012 2013 2009 2011 2014 2015 2795.0 5088.9 2802.4 5225.7 2854.7 5252.5 3231.5 6074.0 < 30 3686.6 4047.2 4542.7 8108.6 5705.4 7206.3 30-39 9020.6 3123.2 40-49 3832.7 4933.5 3027.3 3025.7 6512.6 1395.3 3303.0 4415.1 1240.3 2168.5 5355.5 ≥ 50 1280.6

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

 $Testing\ does\ not\ include\ point\ of\ care\ tests.$ 

## 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 3.1 New HIV Diagnoses for British Columbia <sup>3</sup>

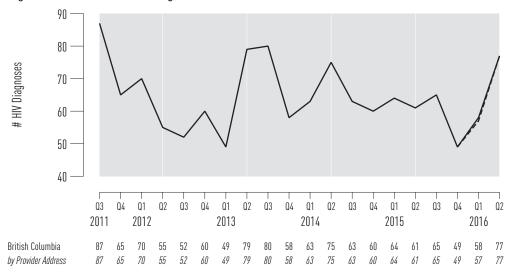


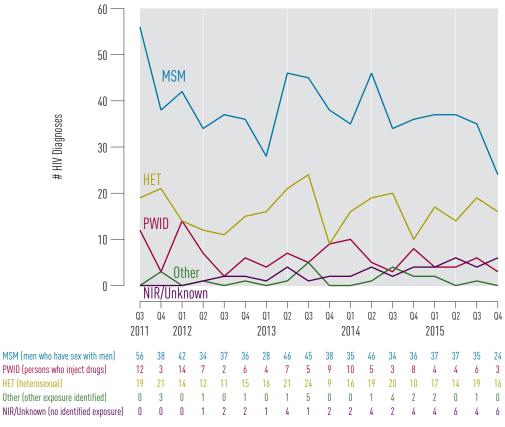
Figure 3.2 New HIV Diagnoses for BC by Gender <sup>3</sup>



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

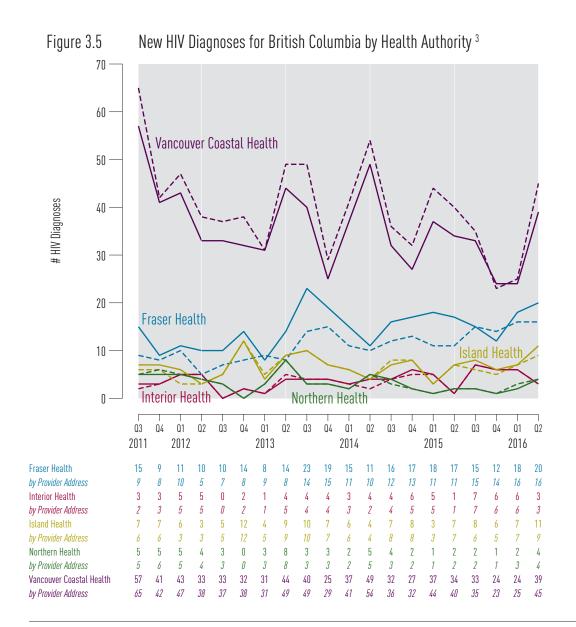
Figure 3.3 New HIV Diagnoses for British Columbia by Age Category <sup>3</sup> ≥ 50 # HIV Diagnoses 10 -< 30 Q4 Q4 Q2 Q3 Q4 Q1 Q1 Q3 Q1 Q2 Q3 Q4 Q2 Q3 Q1 Q2 Q3 Q2 Q1 < 30 30-39 40-49 ≥ 50 

Figure 3.4 New HIV Diagnoses for BC by Exposure Category 3,4



<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.



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

## 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			CD4 350-499										
2b	Stage 0 not met	and	CD4 200-349										
3	HOLHIEL		CD4 <200										
Unknown			No available CD4										

Updated 2016 Q1: AIDS diagnosis date is no longer used in this indicator.

Figure 4.1 Stage of HIV Infection at Diagnosis for BC, 2011–2015 <sup>5</sup>

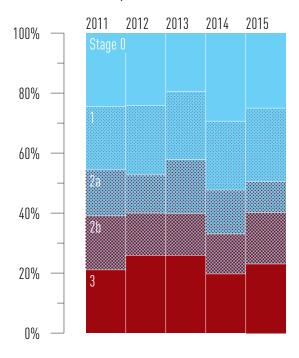
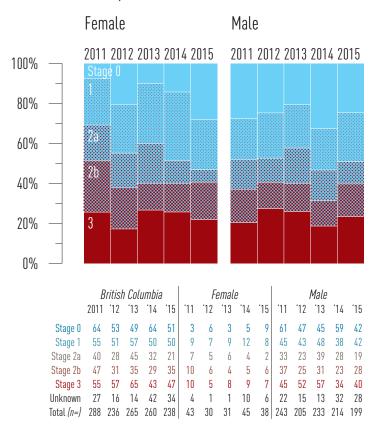


Figure 4.2 Stage of HIV Infection at Diagnosis by Gender for BC, 2011–2015 <sup>5</sup>



Data Source: вссьс

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

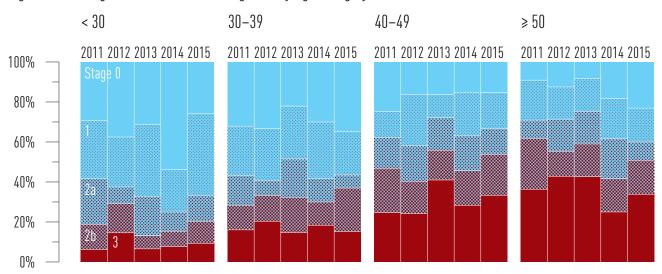
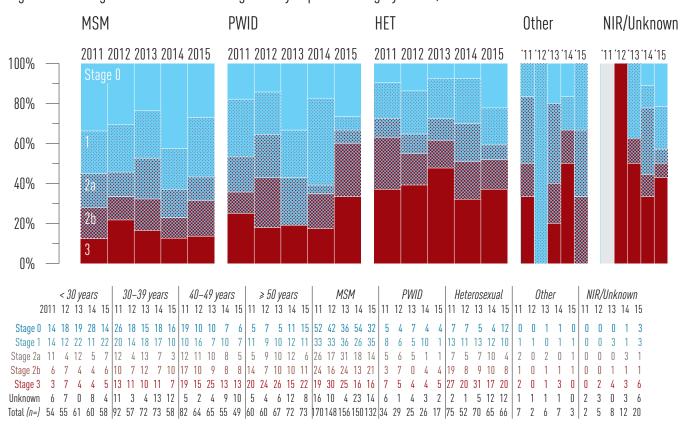


Figure 4.4 Stage of HIV Infection at Diagnosis by Exposure Category for BC, 2011–2015 5.6



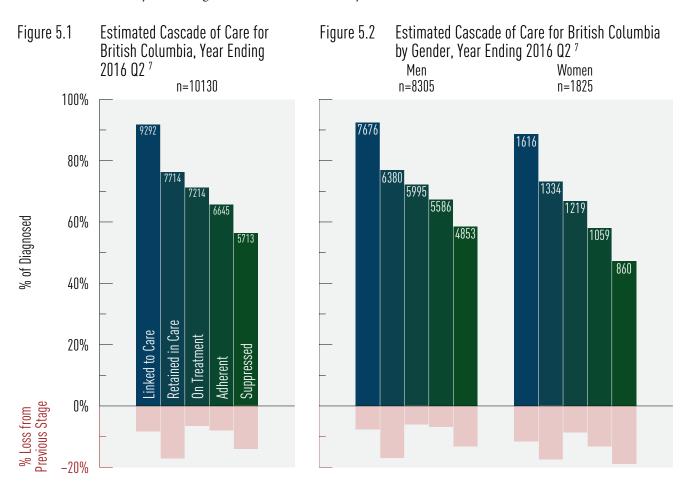
<sup>5</sup> Data Source: BCCDC

<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 BC overall and stratified by sex and age for each Health Authority.



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

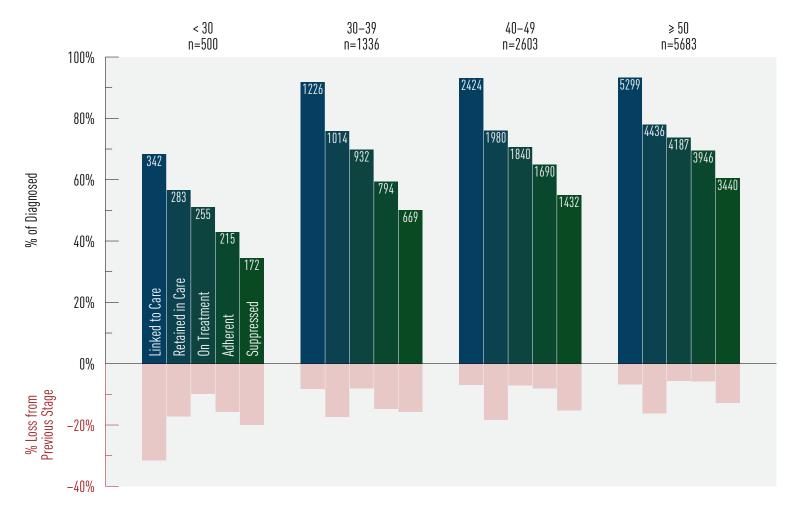
Limitations: HA assignment is based on the most recent HA of residence of the patient, if not available of the HIV-care provider. If the most recent HA of residence is not updated then the designated HA may be incorrect.

NB: Transgender have been assigned to their biological sex.

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)).



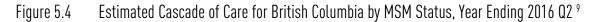


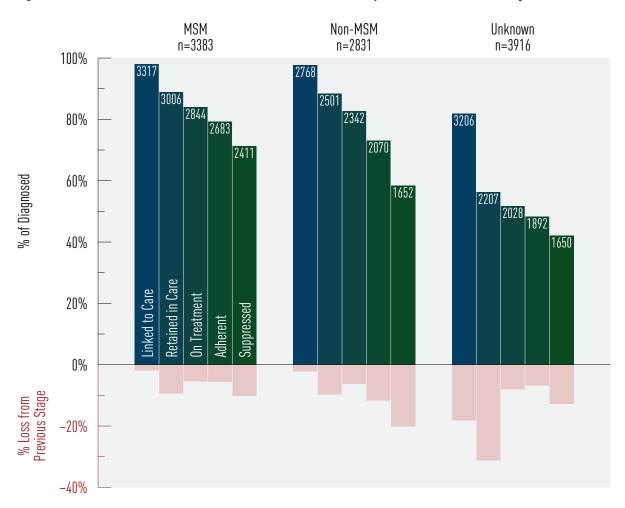
Limitations: HA assignment is based on the most recent HA of residence of the patient, if not available of the HIV-care provider. If the most recent HA of residence is not updated then the designated HA may be incorrect.

<sup>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)).





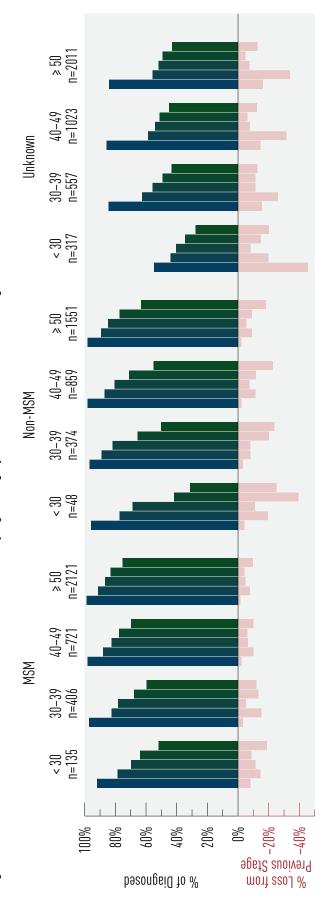
Limitations: HA assignment is based on the most recent HA of residence of the patient, if not available of the HIV-care provider. If the most recent HA of residence is not updated then the designated HA may be incorrect.

<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)).

Estimated Cascade of Care for British Columbia by Age Category and MSM Status, Year Ending 2016  $02^{\,9}$ Figure 5.5



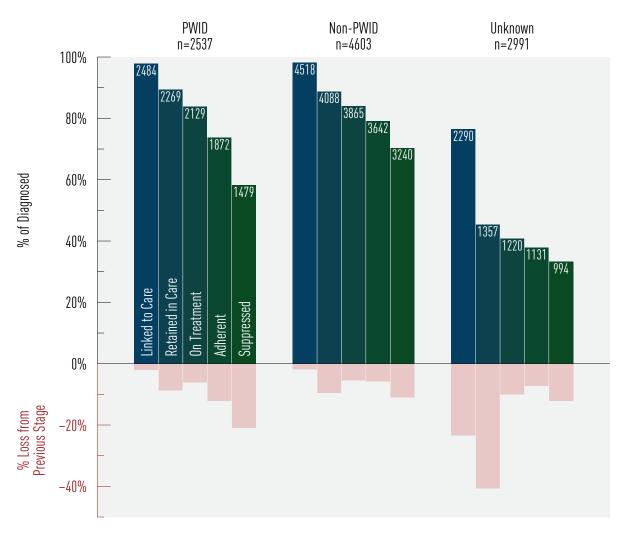
Data is for the period 2015 Q3-2016 Q2. Data Sources:

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).

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





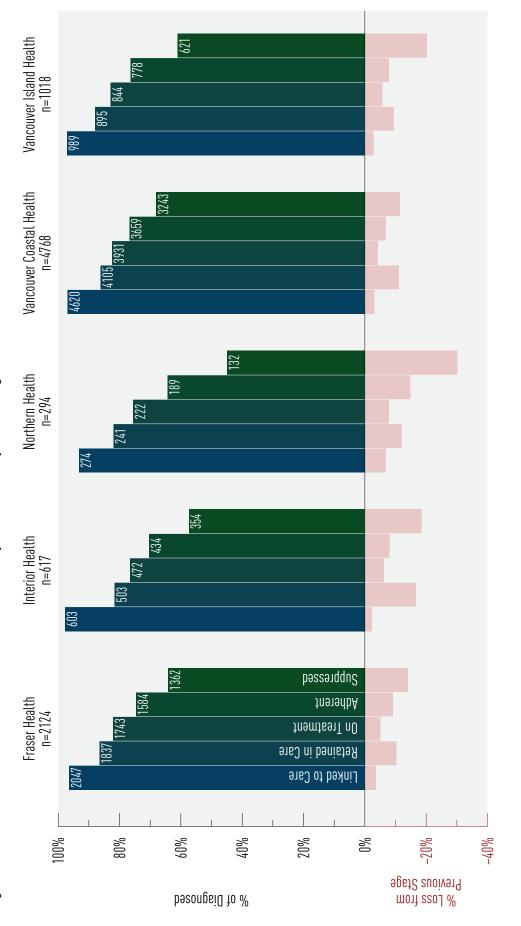
Limitations: HA assignment is based on the most recent HA of residence of the patient, if not available of the HIV-care provider. If the most recent HA of residence is not updated then the designated HA may be incorrect.

<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)).

Estimated Cascade of Care for British Columbia by Health Authority, Year Ending 2016 Q2  $^{\it 9}$ Figure 5.7



9 Data is for the period 2015 Q3-2016 Q2.

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)).

## 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 BC, 2014 Q3-2016 Q2  $^{10}$ 

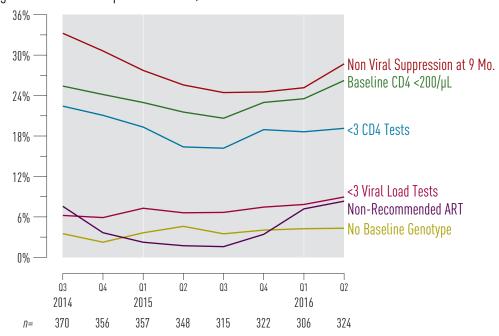
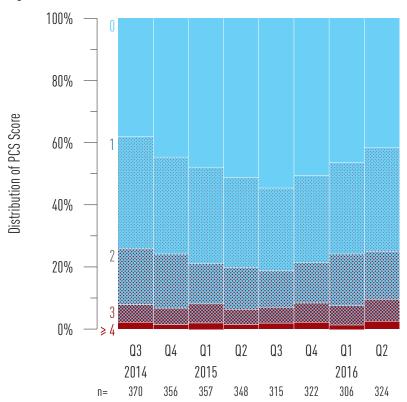


Figure 6.2 Historical Trends for PCS Score for BC, 2014 Q3-2016 Q2 10,11



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

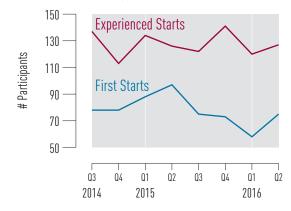
<sup>11</sup> Each quarter's data is calculated as the sum of the 4 quarters leading up to it. e.g. 2013 Q1 is calculated from 2012 Q2 – 2013 Q1. NB: A score of o is the best score and a score of 4 or more is the worst score.

### 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.

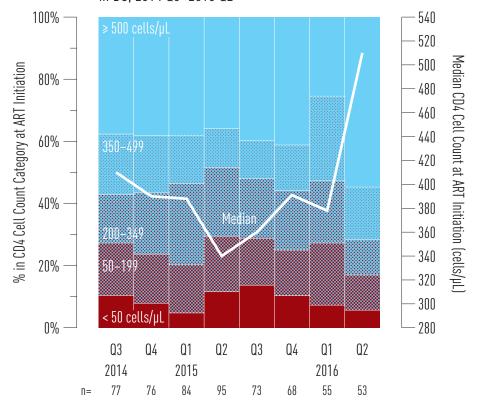
#### Indicator 7. New Antiretroviral Therapy Starts in BC

Figure 7 BC-CfE Drug Treatment Program Enrollment: New ART Participants in BC, 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 BC. 2014 Q3-2016 Q2 13



<sup>12</sup> Data Source: Drug Treatment Program Database
Limitation: DTP participants are designated to an HA based on most current residence provided by the participant.

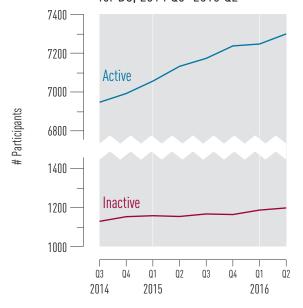
<sup>13</sup> Data Source: Drug Treatment Program Database Limitations: CD4 cell count data is approximately 80% complete.

## Indicator 9. Active and Inactive DTP Participants

Table 3. Distribution of People on ART for BC, 2016 Q2  $^{14}$ 

		Fraser	Interior	Island	Northern	Vancouver Coastal	Total BC
Age	< 30	84	22	30	7	127	270
	30-39	262	53	97	50	541	1005
	40-49	508	108	217	64	1049	1946
	≥ 50	913	294	523	99	2248	4078
Gender	Male	1369	374	709	137	3494	6085
	Female	397	103	158	83	471	1213
Exposure	MSM	573	153	255	30	1898	2910
	PWID	442	149	276	115	1121	2104
Total		1767	477	867	220	3965	7299

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



Definition:

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

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

Active DTP participants: An individual who has had medication prescribed at least once in the preceding quarter.

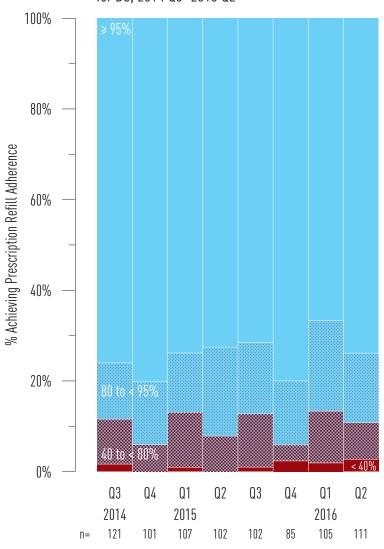
Inactive DTP participants: Persons no longer prescribed drugs through the HIV/AIDS Drug Treatment Program in the last quarter.

#### Antiretroviral Adherence Level

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

#### Indicator 10. Antiretroviral Adherence

Figure 10 Distribution of Individuals by Adherence Level in 1st Year of Therapy, Based on Pharmacy Refill Compliance for BC, 2014 Q3–2016 Q2 16



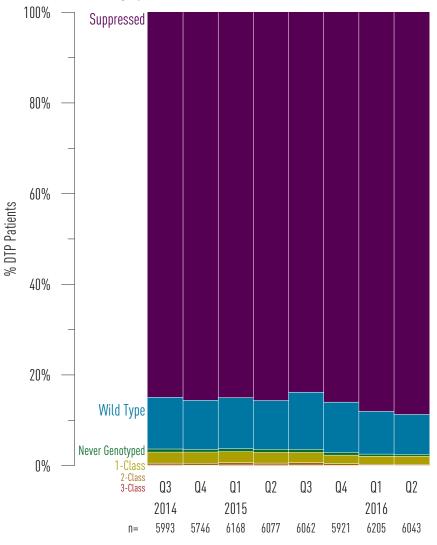
<sup>16</sup> Data Source: Drug Treatment Program Database Limitation: Prescription refill adherence is used as a proxy for patient adherence.

## 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.





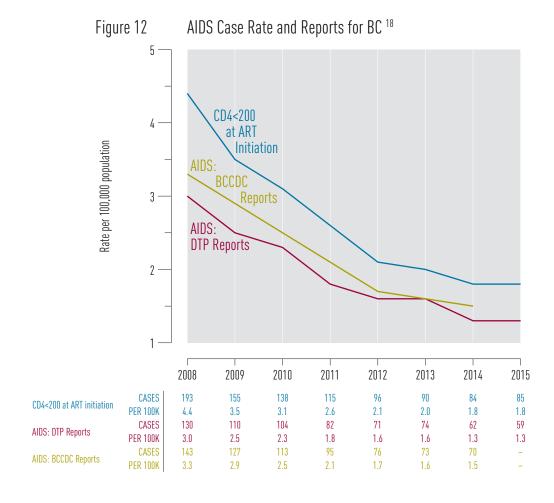
<sup>17</sup> Data Source: Drug Treatment Program Database

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.

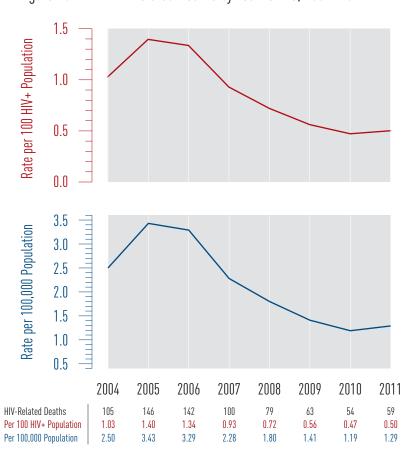


Figure 13 HIV-Related Deaths by Year for BC, 2004–2011 19

#### 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 1 <b>Episodes</b> (	l: Test (thousands)	2011 Q3	l 04	2012 O1	O2	Q3	Q4	2013 Q1		Q3	Q4	2014 Q1	1 Q2	Q3	Q4	2015 O1	Q2	Q3	O4	2016 O1	Q2
British Co	lumbia	_	37.6	42.3	41.5	_	_	_	_	_	_	_	_		_	77.9		_	80.7	88.7	
Gender	Female	15.9	16.6	18.9	18.9	20.8	21.5	25.7	27.5	26.5	25.6	28.7	31.2	33.8	33.7	37.5	37.4	38.7	38.7	42.3	42.6
	Male	17.7	18.0	20.8	20.1	21.6	22.7	26.2	27.9	26.5	26.2	30.0	31.6	33.8	33.7	37.2	36.7	38.3	39.1	43.3	42.5
	Other	0.6	0.5	0.4	0.3	0.4	0.3	0.3	0.2	0.2	0.2	0.3	0.3	0.3	0.2	0.3	0.3	0.3	0.3	0.3	0.3
Age	< 30	13.1	13.0	14.0	13.5	14.8	14.7	15.7	16.7	16.8	16.3	17.4	17.6	19.2	18.9	19.7	19.9	21.8	21.8	22.2	22.5
	30-39	8.7	8.7	10.1	9.7	10.2	10.3	12.2	12.7	12.3	11.9	13.9	13.9	14.6	14.1	16.1	16.1	16.7	16.1	18.6	18.3
	40-49	5.8	6.0	6.9	6.8	6.9	7.6	9.0	9.3	8.7	8.5	9.7	10.1	10.5	10.5	11.6	11.4	11.7	11.8	13.3	12.9
	≥ 50	6.2	7.0	8.8	9.1	10.5	11.7	15.2	16.9	15.2	15.2	17.8	21.4	23.5	24.0	27.4	26.9	27.0	28.3	31.7	31.6
POC HIV	Tests	6.1	2.5	2.3	2.1	2.3	2.2	2.6	2.4	2.7	2.6	3.0	3.0	3.3	2.9	2.9	2.6	2.6	2.6	2.7	2.2
Fraser Hea	alth	9.1	9.0	10.1	10.1	10.8	11.6	13.5	14.0	13.8	12.9	13.6	14.5	14.9	15.1	16.1	15.8	16.8	17.1	18.5	19.2
Female		4.3	4.3	4.8	4.9	5.3	5.7	6.6	6.8	6.8	6.1	6.5	6.9	7.2	7.4	7.6	7.6	8.1	8.1	8.8	9.3
Male		4.7	4.6	5.3	5.1	5.4	5.7	6.6	7.0	6.7	6.5	6.9	7.1	7.3	7.5	8.1	7.8	8.4	8.6	9.3	9.6
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
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
Island Hea	alth	3.6	3.5	4.0	3.7	3.8	4.0	4.2	4.5	4.3	4.3	4.9	4.8	4.8	4.9	5.9	6.0	6.4	6.6	7.1	7.2
Female		1.8	1.7	1.9	1.8	1.9	2.0	2.1	2.2	2.1	2.1	2.3	2.3	2.4	2.4	2.9	3.0	3.2	3.3	3.5	3.7
Male		1.8	1.7	2.0	1.8	1.8	1.8	2.0	2.1	2.0	1.9	2.2	2.1	2.1	2.1	2.6	2.6	2.8	2.9	3.1	3.2
Northern :	Health	1.9	1.9	2.3	2.2	2.2	2.3	2.7	2.7	2.6	2.6	3.1	3.0	2.9	3.0	3.7	3.2	3.2	3.2	3.7	3.3
Female		1.0	1.0	1.2	1.1	1.2	1.2	1.3	1.4	1.3	1.3	1.6	1.5	1.5	1.5	1.9	1.6	1.6	1.6	1.9	1.7
Male		0.8	0.8	1.1	1.0	1.0	1.1	1.2	1.2	1.1	1.2	1.4	1.3	1.3	1.4	1.6	1.4	1.5	1.4	1.6	1.5
Vancouver	Coastal Health	21.9	19.1	21.8	21.4	24.0	24.6	30.1	32.1	30.7	29.9	35.0	38.4	43.0	41.6	45.5	44.8	45.5	44.6	49.0	48.0
Female		6.9	7.4	9.0	9.0	10.3	10.7	13.6	14.8	14.0	13.6	15.8	17.9	20.1	19.6	21.8	21.7	21.8	21.2	23.0	23.1
Male		8.6	8.8	10.5	10.3	11.4	11.9	14.3	15.3	14.4	14.3	16.8	18.3	20.4	19.8	21.8	21.4	21.9	21.7	24.1	23.3

Indicator 2: Rate of HIV Testing per 100,000

		2009	2010	2011	2012	2013	2014	2015
British Co	lumbia	2624.8	2645.8	2714.2	3318.6	4194.6	4969.0	5738.5
Fraser Hea	ılth	2251.2	2267.6	2331.6	2795.4	3313.1	3588.9	4009.3
Interior H	ealth	2026.8	2072.9	2094.1	2180.4	2573.0	2996.7	4080.7
Island Hea	lth	1927.2	1919.6	1851.0	1980.8	2150.4	2335.6	2939.3
Northern	Health	2290.7	2349.0	2431.8	2909.8	3333.0	3825.8	4347.2
Vancouver	Coastal Health	4125.0	4140.2	4342.1	5827.1	8122.8	10313.6	11591.2
Gender	Female	2446.7	2455.1	2523.6	3197.0	4168.6	4942.3	5770.1
	Male	2694.2	2734.5	2808.7	3383.5	4180.4	4951.8	5656.2
Age	< 30	2795.0	2802.4	2854.7	3231.5	3686.6	4047.2	4542.7
	30-39	5088.9	5225.7	5252.5	6074.0	7206.3	8108.6	9020.6
	40-49	3027.3	3025.7	3123.2	3832.7	4933.5	5705.4	6512.6
	≥ 50	1240.3	1280.6	1395.3	2168.5	3303.0	4415.1	5355.5

Indicator 3: 1	Nev	v HI	V D	Diag	nose	es		2	2011 Q3		2012 Q1		Q3	Q4	2013 Q1		Q3		2014 Q1		Q3		2015 Q1		Q3		2016 Q1	Q2
British Colum						t Res	sider	nce	87	65	70	55	52	60		79	80	58	63	75	63	60	64	61	65	49	58	77
			i	By P	rovi	der A	Addr	ess	87	65	70	55	52	60	49	79	80	58	63	75	63	60	64	61	65	49	57	77
Gender			]	Fem	ale				11	7	9	10	5	5	6	12	8	5	13	9	13	9	10	4	12	11	10	11
			]	Male	e				76	58	61	45	47	55	43	67	72	53	49	65	50	51	54	57	53	38	47	66
Age				< 30					17	18	18	14	9	18	9	17	23	15	17	15	13	18	14	14	21	13	19	15
			3	30-3	39				30	13	16	17	11	10	16	25	18	11	17	21	25	15	15	17	17	8	19	17
			4	40-4	19				22	19	20	11	19	19	12	14	21	20	14	14	7	13	11	19	10	11	11	15
				≥ 50					18	15	16	13	13	13	12	23	18	12	15	25	18	14	24	11	17	17	9	30
Exposure			]	MSN	A				56	38	42	34	37	36	28	46	45	38	35	46	34	36	37	37	35	24	-	_
			]	PWI	ID				12	3	14	7	2	6	4	7	5	9	10	5	3	8	4	4	6	3	-	_
			]	НЕТ					19	21	14	12	11	15	16	21	24	9	16	19	20	10	17	14	19	16	-	_
			(	Oth	er				0	3	0	1	0	1	. 0	1	5	0	0	1	4	2	2	0	1	0	-	_
			]	NIR	/Un	knov	vn		0	0	0	1	2	2	1	4	1	2	2	4	2	4	4	6	4	6	-	_
Fraser Health	h		]	Ву С	Clien	t Re	sider	nce	15	9	11	10	10	14	8	14	23	19	15	11	16	17	18	17	15	12	18	20
			1	Ву Р	rovi	der A	Addr	ess	9	8	10	5	7	8	9	8	14	15	11	10	12	13	11	11	15	14	16	16
Interior Heal	lth		]	ВуС	Clien	t Re	sider	nce	3	3	5	5	0	2	1	4	4	4	3	4	4	6	5	1	7	6	6	3
				Ву Р	rovi	der 1	Addr	ess	2	3	5	5	0	2	2 1	5	4	4	3	2	4	5	5	1	7	6	6	3
Island Health	h		]	ВуС	Clien	t Re	sidei	nce	7	7	6	3	5	12	4	9	10	7	6	4	7	8	3	7	8	6	7	11
			i	Ву Р	rovi	der A	Addr	ess	6	6	3	3	5	12	5	9	10	7	6	4	8	8	3	7	6	5	7	9
Northern He	ealth	ı	]	ВуС	Clien	t Res	sider	nce	5	5	5	4	3	0	3	8	3	3	2	5	4	2	1	2	2	1	2	4
			i	Ву Р	rovi	der A	Addr	ess	5	6	5	4	3	0	3	8	3	3	2	5	3	2	1	2	2	1	3	4
Vancouver C	Coas	tal	]	ВуС	Clien	t Res	sider	nce	57	41	43	33	33	32	31	44	40	25	37	49	32	27	37	34	33	24	24	39
Health			i	Ву Р	rovi	der A	Addr	ess	65	42	47	38	37	38	31	49	49	29	41	54	36	32	44	40	35	23	25	45
<b>7</b> 10	0.				٠.		. 5																					
Indicator 4: S	_					ion a		_		ı			1				20		ı						40	40		
		tish '12				<b>'</b> 11		emal '13		<b>'</b> 15	'11 '		ale 13'1	4 '	15 1		30 ye : '13		<b>'</b> 15		30-39 12. 1			5 1		-49 y 2. '13		<b>'</b> 15
		53	49	64	51	3	6	3	5	9	61					1 18			14					6 1				6
	55	51	57	50	50	9	7	9	12	8	45						2 22		22					0 1				7
	40	28	45	32	21	7	5	6	4	2	33				19 1			5	7	12		13	7		2 11			5
	47	31	35	29	35		6	4	5	6	37					5 7			6	10	7			0 1				8
-	55	57	65	43	47	10	5	8	9	7	45	52	57 3	4	40	3 7	7 4	4	5	13	11	10	11		9 15	5 25	13	13
		16					1						13 3				7 0			11	3		13 1			2 4		10
Total 2	2882	236	265	260	238	43	30								99 5						57				2 64	4 65	55	49
c	<b>'</b> 11		0 ye: '13		<b>'</b> 15	<b>'</b> 11		MSM '13		<b>'</b> 15	<b>'</b> 11		VID 13'1	4 '	15 1		erose 2 '13				her E '12'				NIR/ 1 '12			
Stage 0	5	7				52					5	4		4		7 7			12	0	0	1			0 0			3
	11	9				33					8	6	5 1	0	1 13	3 11	13	12	10	1	1	0	1	1	0 0	) 3	1	3
Stage 2a	5	9	10	12	6	26	17	31	18	14	5	6	5	1	1	7 5	5 7	10	4	2	0	2	0	1	0 (	0 (	3	1
Stage 2b	14	7	10	10	11	24	16	24	13	21	3	7	0	4	4 19	9 8	9	10	8	1	0	1	1	1	0 0	) 1	1	1
-				15		19	30	25	16	16	7	5	4	4	5 2	7 20	31	17	20	2	0	1	3	0	0 2	2 4	3	6
-			26	15 12	22	19 16					7 6	5 1	4	4		7 20 2 1				2	0	1 1				<ul><li>2 4</li><li>3 0</li></ul>		6 6

Indicator 5: H	IV Cascade of	Care	DIAGNOSED	LINKED	RETAINED	Oi	N ART A	DHERENT	SUPPRESSED
British Columb	bia		10130	9292	7714		7214	6645	5713
Age Category	< 30		500	342	283		255	215	172
	30-39		1336	1226	1014		932	794	669
	40-49		2603	2424	1980		1840	1690	1432
	≥ 50		5683	5299	4436		4187	3946	3440
Age Category	MSM	< 30	135	124	106		94	86	70
and MSM		30-39	406	394	334		317	275	242
Status		40-49	721	706	635		595	559	503
		≥ 50	2121	2093	1931		1838	1763	1596
	Non-MSM	< 30	48	46	37		33	20	15
		30-39	374	362	333		306	245	187
		40-49	859	841	746		691	610	471
		≥ 50	1551	1519	1385		1312	1195	979
	Unknown	< 30	317	173	1303		128	109	87
	CIIKIIOWII	30-39	557	470	348		309	274	240
		40-49	1023	876	599		554	521	458
Gender	Male	≥ 50	2011	1687	1120		1037	988	865
Gender			8305	7676	6380		5995	5586	4853
<b>.</b>	Female		1825	1616	1334		1219	1059	860
Injection Drug Use	PWID		2537	2484	2269		2129	1872	1479
Drug Osc	Non-PWID		4603	4518	4088		3865	3642	3240
	Unknown		2991	2290	1357		1220	1131	994
MSM Status	MSM		3383	3317	3006		2844	2683	2411
	Non-MSM		2831	2768	2501		2342	2070	1652
	Unknown		3916	3206	2207		2028	1892	1650
Health	Fraser Health		2124	2047	1837		1743	1584	1362
Authority	Interior Healt		617	603	503		472	434	354
	Island Health		1018	989	895		844	778	621
	Northern Hea	alth	294	274	241		222	189	132
	Vancouver Co	oastal Health	4768	4620	4105		3931	3659	3243
Indicator 6: Pr	ogrammatic C	Compliance S	core (PCS)						
	8	2014	:	2015				2016	, )
		Q3	Q4	Q1	Q2	Q3	Q4	Q1	
< 3 CD4 Tests		22.4%	21.1%	19.3%	16.4%	16.2%	18.9%	18.6%	
< 3 Viral Load	Tests	6.2%	5.9%	7.3%	6.6%	6.7%	7.5%	7.8%	9.0%
No Baseline G	enotype	3.5%	2.2%	3.6%	4.6%	3.5%	4.0%	4.2%	4.3%
Baseline CD4 ·	< 200 cells/μL	25.4%	24.2%	23.0%	21.6%	20.6%	23.0%	23.5%	26.2%
Non-Recomme	ended ART	7.6%	3.7%	2.2%	1.7%	1.6%	3.4%	7.2%	8.3%
Non Viral supp	pression at 9 M	o. 33.2%	30.6%	27.7%	25.6%	24.4%	24.5%	25.2%	28.7%
PCS Score: 0		141	159	171	178	172	163	142	135
PCS Score: 1		133	111	111	101 84 90		90	90	108
PCS Score: 2			62	46	47	37	42	51	50
		67	02						
PCS Score: 3		21	19	22	17	16	20	19	
PCS Score: 3 PCS Score: 4 o	r more					16 6			23

Indicator 7: New DTP A	RV Participants									
	2014 Q3	Q4	2	015 Q1	Q2	Q3	Q	4	2016 Q1	Q2
First Starts	78	78		88	97	75	7:		58	75
Experienced Starts	137	113		134	126	122	14	1	120	127
Indicator 8: CD4 Cell Co	ount at APT Initiati	on for ARV	-Νοΐνο Γ	TD Darti	cinante					
CD4 ≥ 500	29	29	-Naive D	32	34	29	2	R	14	29
CD4 350-499	15	14		13	12	9	10		15	9
CD4 200-349	12	15		22	21	14	1:		11	6
CD4 50–199	13	12		13	17	11	10		11	6
CD4 < 50	8	6		4	11	10		7	4	3
	410	390		388	340	360	39		378	510
CD4 Median (cells/μL) Total (n=)	77	76		84	95	73	6		55	53
				01	,,,	, 0	0.	3	33	
Indicator 9: Active and I				057	7122	7172	F22	7	72.45	7200
Active DTP Participants	6947	6992		056	7132	7173	723		7247	7299
Inactive DTP Participant	rs 1129	1153	1	158	1154	1167	116	4	1187	1198
Indicator 10: Antiretrov	iral Adherence									
≥ 95%	92	81		79	74	73	6	8	70	82
80% to < 95%	15	14		14	20	16	13	2	21	17
40% to < 80%	12	6		13	8	12	:	3	12	9
< 40%	2	0		1	0	1	:	2	2	3
Total (n=)	121	101		107	102	102	8.	5	105	111
Indicator 11: Resistance	Testing and Results	<b>.</b>								
Suppressed	5090	4919	5.	244	5205	5081	5093	3	5463	5361
Wild Type	684	623		694	654	764	65	7	585	532
Never Genotyped	41	33		40	42	42	3.	5	29	25
1-Class	146	141		151	143	137	112	2	110	109
2-Class	25	27		31	25	31	2:	2	15	14
3-Class	7	3		8	8	7		2	3	2
Total (n=)	5993	5746	6	168	6077	6062	5922	2	6205	6043
Indicator 12: AIDS-Defi	ning Illness	2007	2008	2009	2010	2011	2012	2013	2014	2015
CD4 < 200 at	Cases	224	193	155	138	115	96	90	84	85
ART initiation	Rate per 100,000	5.2	4.4	3.5	3.1	2.6	2.1	2.0	1.8	1.8
AIDS Cases	Cases	135	130	110	104	82	71	74	62	59
(DTP Reports)	Rate per 100,000	2.8	3.0	2.5	2.3	1.8	1.6	1.6	1.3	1.3
AIDS Cases	Cases	143	143	127	113	95	76	73	70	_
(BCCDC Reports)	Rate per 100,000	3.3	3.3	2.9	2.5	2.1	1.7	1.6	1.5	_
Indicator 12: IIIV D-1-4-	nd Montality	2004	2005	2006	2007	2009	2000	2010	2011	
Indicator 13: <b>HIV-Relate</b> British Columbia	tu iviortality	2004 105	2005 146	2006 142	2007 100	2008 79	2009 63	2010 54	2011 59	
Per 100 HIV+ Population	2	1.03			0.93					
Per 100 HIV+ Population	1	2.50	1.40	1.34 3.29		0.72 1.80	0.56	0.47	0.50 1.29	
rei 100,000 ropulation		2.30	3.43	3.29	2.28	1.80	1.41	1.19	1.29	