

HIV MONITORING QUARTERLY REPORT

FOR BRITISH COLUMBIA

FOURTH QUARTER 2015

















Foreword

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

The objectives of these reports are to:

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

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

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Acknowledgements and Contributions



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

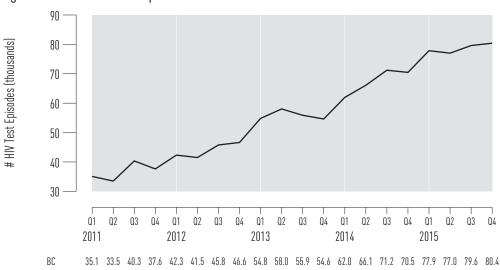


Figure 1.2 HIV Test Episodes by Gender for BC ^{1,2}

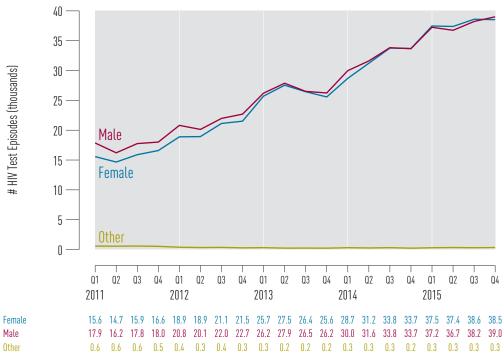
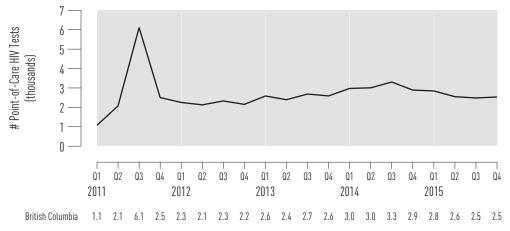


Figure 1.3 HIV Test Episodes by Age Category for BC 1,2 30 28 26 -24 -22 -# HIV Test Episodes (thousands) 20 -18 -16 14 < 30 12 10 30-39 8 -≥ 50 6 Q1 Q2 Q3 Q4 Q1 04 Q1 Q2 Q3 Q4 Q1 02 Q3 04 01 Q2 Q3 Q2 Q3 2012 2013 2014 2015 2011 < 30 12.3 11.8 13.1 13.0 14.0 13.5 15.1 14.7 15.7 16.7 16.8 16.3 17.4 17.6 19.2 18.9 19.7 19.9 21.7 21.8 30-39 10.4 10.3 12.2 12.7 12.3 11.9 13.9 13.9 14.6 14.1 16.1 16.1 16.6 16.0

Figure 1.4 Point-of-Care HIV Tests for BC



7.1

9.0 9.3 8.7 8.5 9.7

7.6

10.1 10.5 10.5 11.6 11.4 11.7 11.8

9.1 10.7 11.7 15.2 16.9 15.2 15.2 17.8 21.4 23.5 24.0 27.4 26.9 26.9 28.2

Limitations:

40-49

≥ 50

- i Repeat tests in individuals who test using various identifiers may not be identified and these individuals may be counted more than once.
- ii In Fraser Health, POC testing data are available from March 2011 forward. In Interior Health, POC testing data are available for May 2011 forward. For all other health authorities, POC testing data are available from the fourth quarter of 2010 forward.
- 2 Testing does not include point of care tests.

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

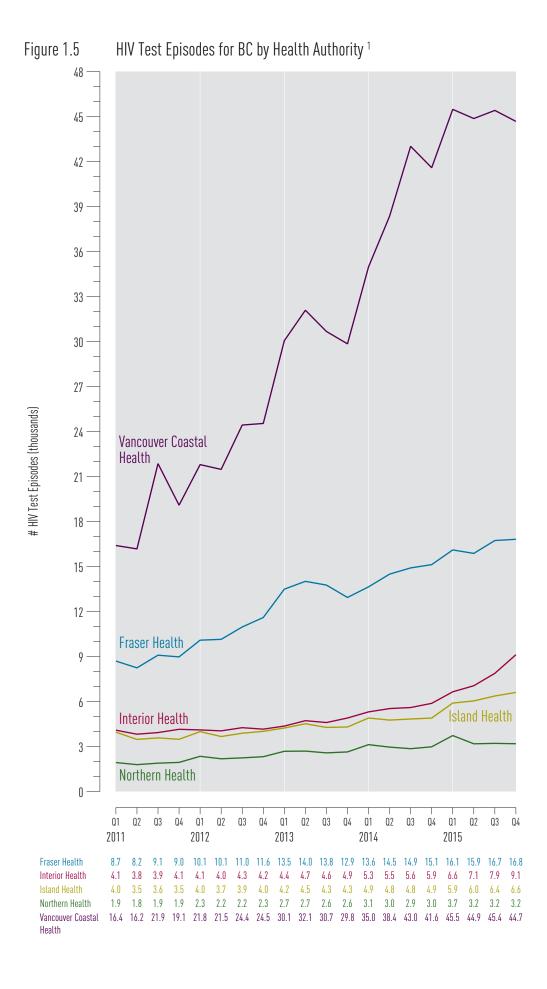


Figure 1.6 HIV Test Episodes for Non-prenatal Females in BC by Health Authority 1.2

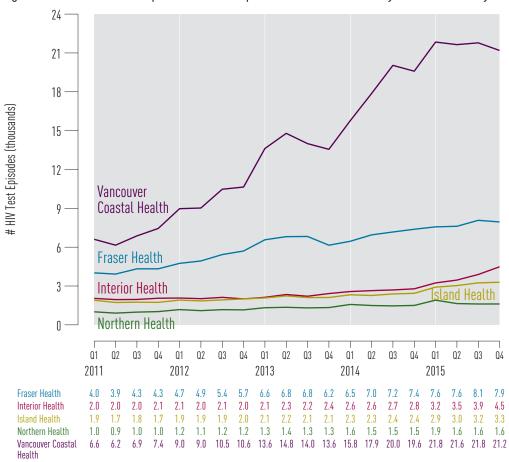
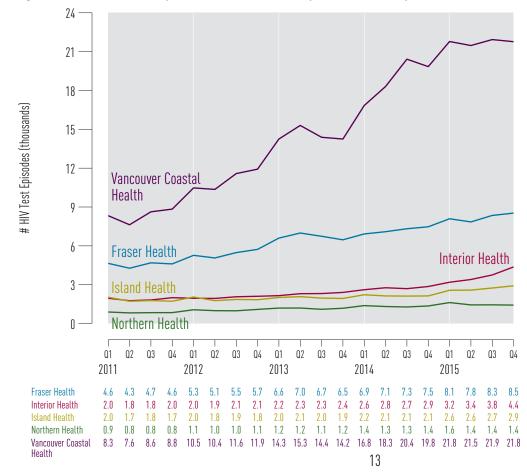


Figure 1.7 HIV Test Episodes for Males in BC by Health Authority 1,2



Indicator 2. HIV Testing Rates

Figure 2.1 Rate of HIV Testing for British Columbia and Health Authorities ²

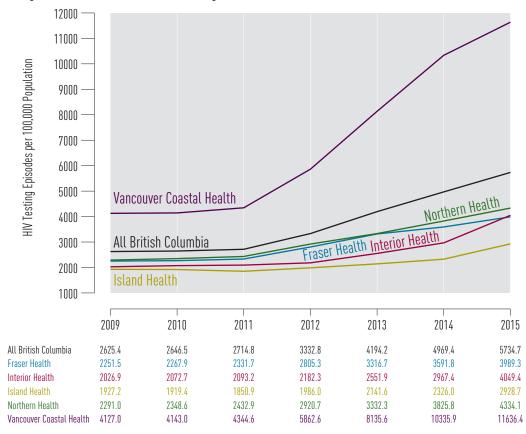
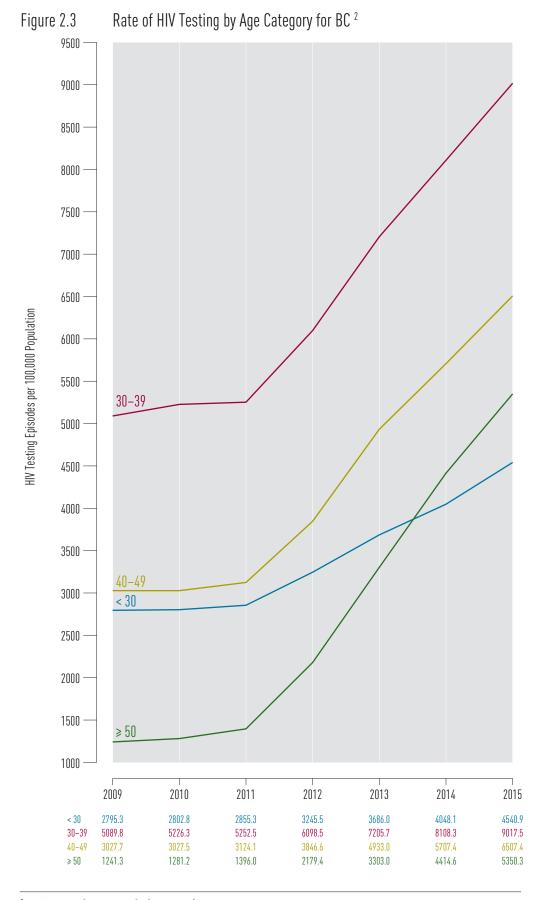


Figure 2.2 Rate of HIV Testing by Gender for BC $^{\rm 2}$





Testing does not include point of care tests.

New HIV Diagnoses

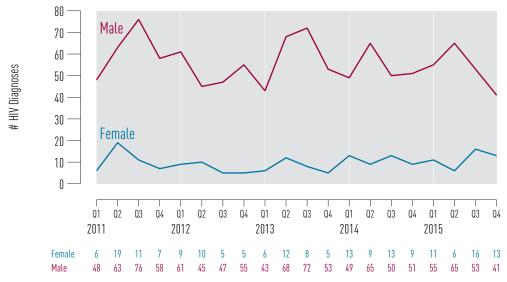
Trends in HIV diagnoses by gender and exposure category are described. Interpreting HIV diagnoses must be done with consideration that trends are influenced by both changes in testing rate as well as changes in transmission rates. It is important to note that new HIV diagnoses cases and rates are not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





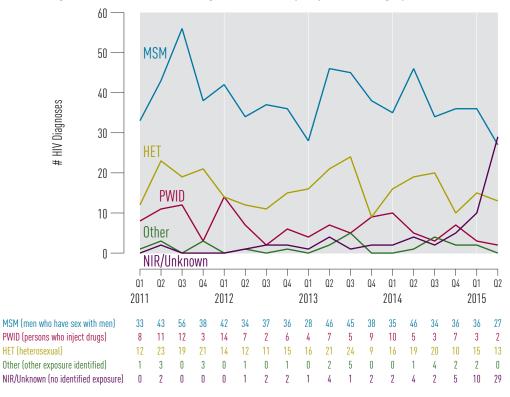




З Data Source: вссьс. When present, "By Provider Address" is graphed as dashed line in same colour.

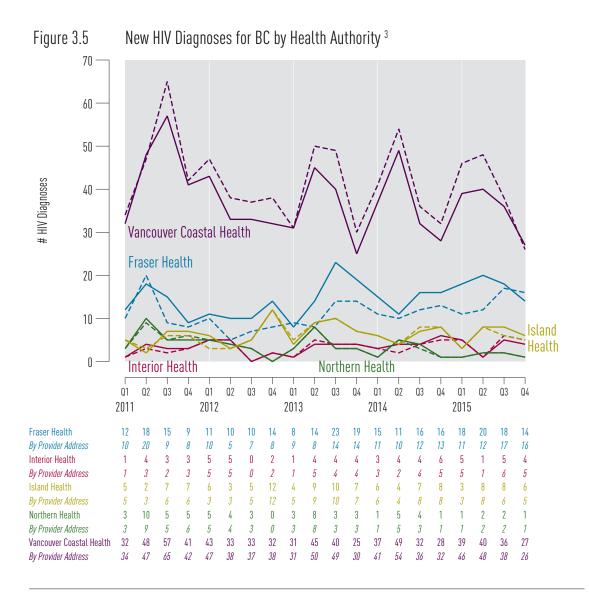
Figure 3.3 New HIV Diagnoses for BC by Age Category ³ 30-39 # HIV Diagnoses 40-49 ≥ 50 < 30 0 -Q2 Q3 Q2 Q4 Q1 Q2 Q1 Q3 Q4 Q2 Q3 Q4 < 30 30-39 40-49 ≥ 50

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



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

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



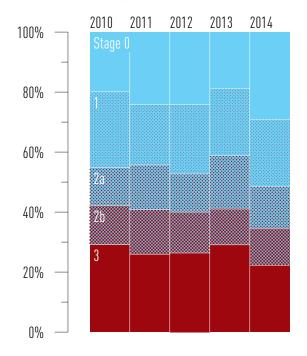
³ Data Source: BCCDC. When present, "By Provider Address" is graphed as dashed line in same colour.

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, 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.

Figure 4.1 Stage of HIV Infection at Diagnosis for BC, 2010–2014 ⁵

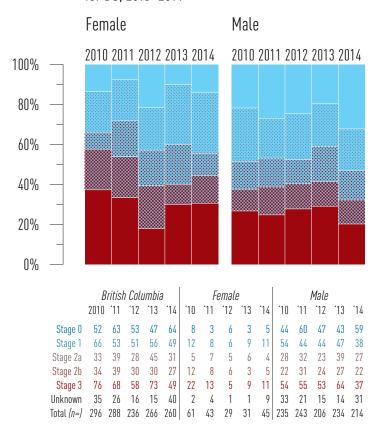


Indicator 4. Stage of HIV Infection at Diagnosis

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		N AIDO																			
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 4.2 Stage of HIV Infection at Diagnosis by Gender for BC, 2010–2014 ⁵



Data Source: BCCDC

Figure 4.3 Stage of HIV Infection at Diagnosis by Age Category for BC, 2010–2014 ⁵

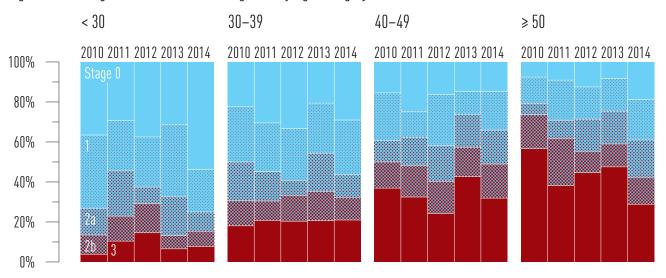
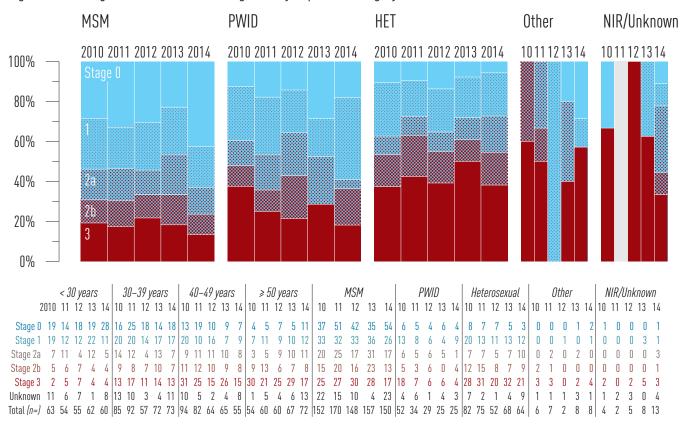


Figure 4.4 Stage of HIV Infection at Diagnosis by Exposure Category for BC, 2010–2014 5.6



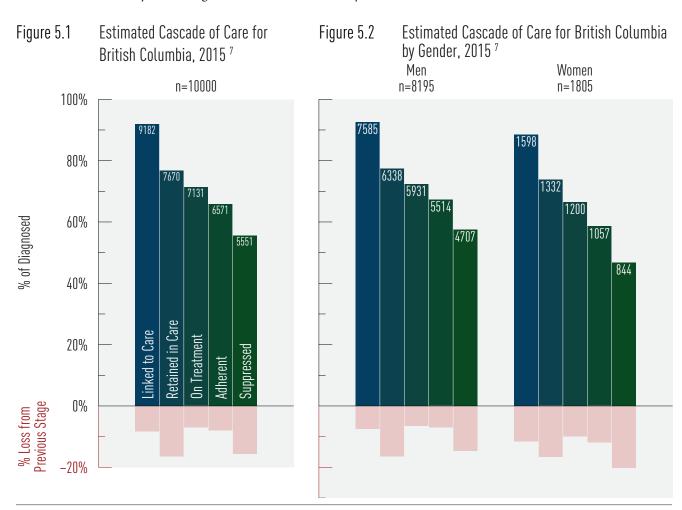
⁵ Data Source: BCCDC

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

HIV Cascade of Care

Indicator 5. HIV Cascade of Care

The success of seek, test, treat and retain (STTR) strategies like STOP is reliant on early diagnosis of HIV, linking newly diagnosed HIV-positive persons with ongoing care, retaining persons in HIV-care; initiating ART based on best evidenced practices and maintaining optimal ART adherence to ensure a suppressed viral load. These stages of HIV-care can be summarized as: 1. HIV diagnosis, 2. Linked to HIV care, 3. Retained in HIV care, 4. On ART, 5. Adherent to ART and 6. Achieving a suppressed VL; collectively, they are referred to as the cascade of care. 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 (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 Q1–2015 Q4 in BC overall and stratified by sex and age for each Health Authority.

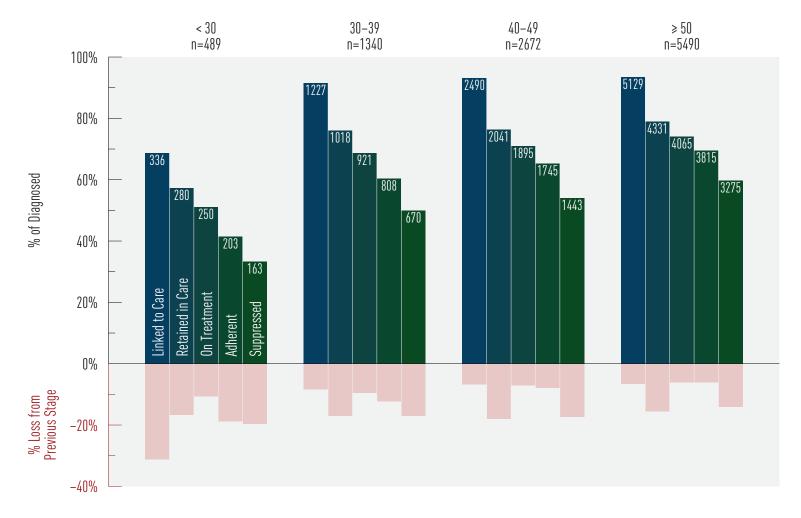


- 7 Data is for the period 2015 Q1–2015 Q4. Data Sources:
 - i British Columbia Centre for Excellence Drug Treatment Program (DTP) Database (ARV use, VL and CD4 count).
 - ii Administrative data (ex. MSP billings; hospitalization data from the Discharge Abstract Database (DAD)).

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

NB: Transgender have been assigned to their biological sex.





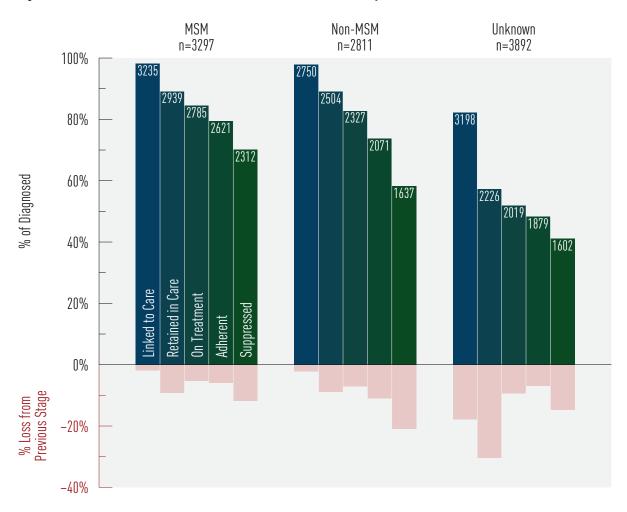
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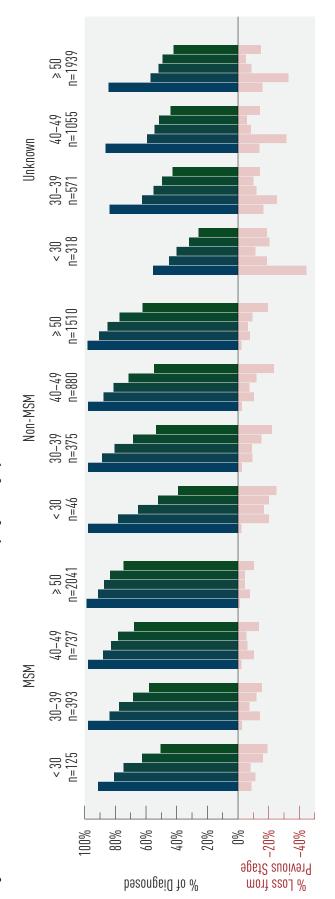
Recent updates to the DTP database have allowed for more comprehensive information on HIV risk group category. As a result, 2014 Q4 data may differ significantly from preceding reports in terms of total numbers ascribed to each risk group.

⁹ Data is for the period 2015 Q1-2015 Q4. 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, 2015 $^{\it 9}$ Figure 5.5

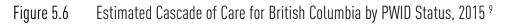


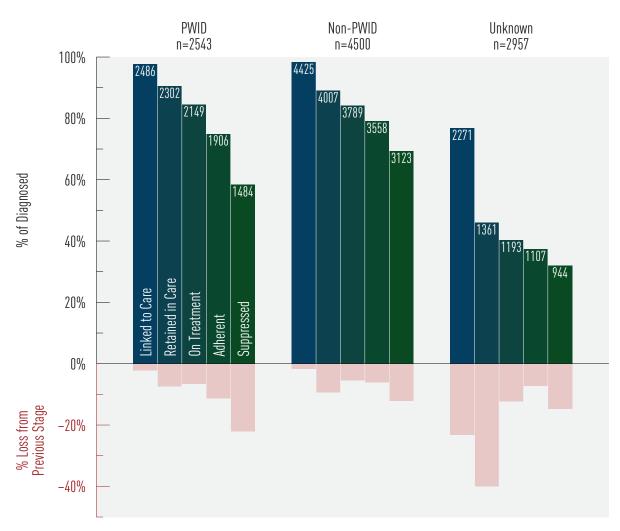
Data Sources:

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. Recent updates to the DTP database have allowed for more comprehensive information on HIV risk group category. As a result, 2014 Q4 data may differ significantly from preceding reports in terms of total numbers ascribed to each risk group.

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.

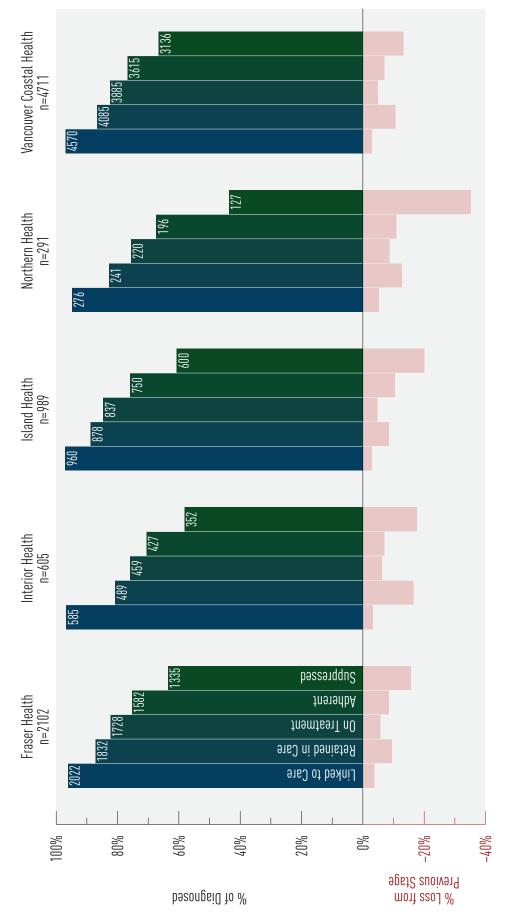
Recent updates to the DTP database have allowed for more comprehensive information on HIV risk group category. As a result, 2014 Q4 data may differ significantly from preceding reports in terms of total numbers ascribed to each risk group.

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 θ Data is for the period 2015 Q1-2015 Q4.

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. Recent updates to the DTP database have allowed for more comprehensive information on HIV risk group category. As a result, 2014 Q4 data may differ significantly from preceding reports in terms of total numbers ascribed to each risk group.

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)).

Programmatic Compliance Score

Indicator 6. Programmatic Compliance Score (PCS)

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

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

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

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

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

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

Figure 6.1 PCS Components for BC, 2014 Q1-2015 Q4 10

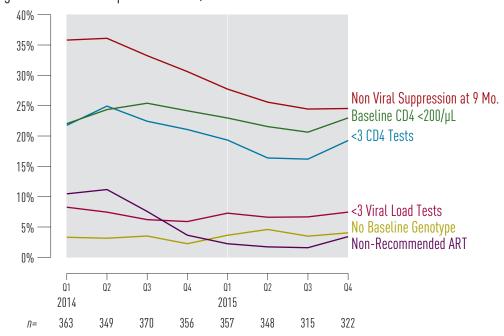
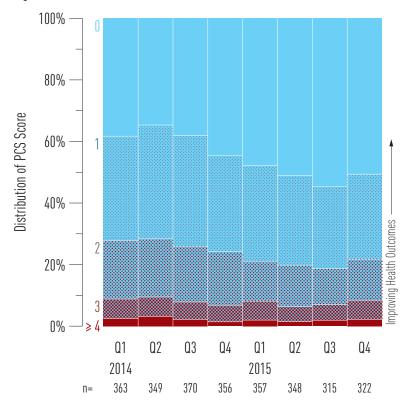


Figure 6.2 Historical Trends for PCS Score for BC, 2014 Q1-2015 Q4 10,11



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

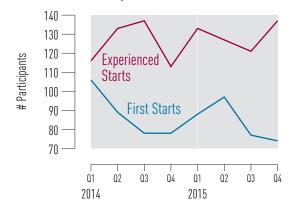
¹¹ Each quarter's data is calculated as the sum of the 4 quarters leading up to it. e.g. 2013 Q1 is calculated from 2012 Q2 – 2013 Q1. NB: A score of o is the best score and a score of 4 or more is the worst score.

Antiretroviral Uptake

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

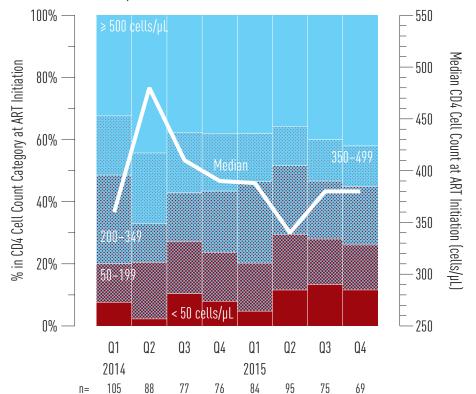
Indicator 7. New Antiretroviral Therapy Starts in BC

Figure 7 BC-CfE Drug Treatment Program Enrollment: New ART Participants in BC, 2014 Q1–2015 Q4 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 Q1–2015 Q4 ¹³



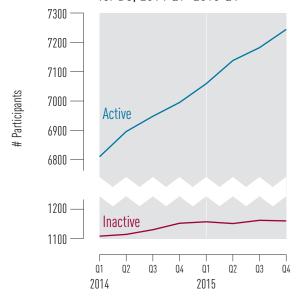
- Data Source: Drug Treatment Program Database Limitation: DTP participants are designated to an HA based on most current residence provided by the participant.
- 3 Data Source: Drug Treatment Program Database Limitations: CD4 cell count data is approximately 80% complete.

Indicator 9. Active and Inactive DTP Participants

Table 3. Distribution of People on ART for BC, 2015 Q4 14

		Fraser	Interior	Island	Northern	Vancouver Coastal	Total BC
Age	< 30	88	18	26	10	143	285
	30-39	257	59	94	46	519	976
	40-49	538	111	223	64	1085	2022
	≥ 50	874	290	496	98	2203	3961
Gender	Male	1362	380	680	138	3481	6043
	Female	395	98	159	80	469	1201
Exposure	MSM	560	143	228	30	1883	2846
	PWID	455	153	274	117	1134	2133
Total		1757	478	839	218	3950	7244

Figure 9 Active and Inactive DTP Participants for BC, 2014 Q1–2015 Q4 ¹⁵



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

Recent updates to the DTP database provides for improved classification allowing some individuals previously classified as 'unknown' to be reclassified into specific risk groups. This update is in effect from 2014Q4 and may result in noticeable changes of numbers in each risk group category compared to previous reports.

Definition:

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

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

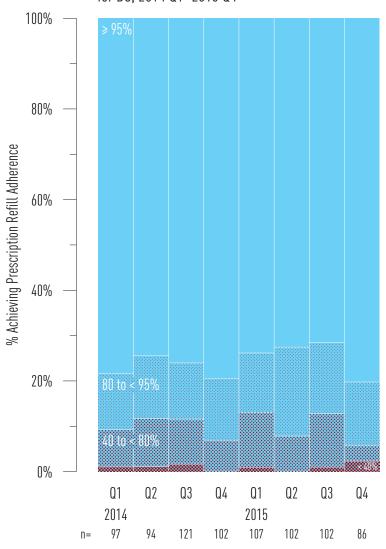
Inactive DTP participants: Persons no longer prescribed drugs through the HIV/AIDS Drug Treatment Program in the last quarter.

Antiretroviral Adherence Level

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

Indicator 10. Antiretroviral Adherence

Figure 10 Distribution of Individuals by Adherence Level in 1st Year of Therapy, Based on Pharmacy Refill Compliance for BC, 2014 Q1–2015 Q4 16

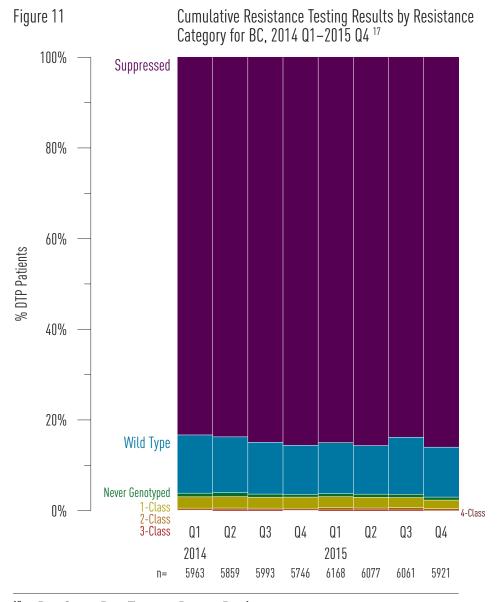


¹⁶ Data Source: Drug Treatment Program Database Limitation: Prescription refill adherence is used as a proxy for patient adherence.

Resistance Testing and Results

Indicator 11. Resistance Testing and Results

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



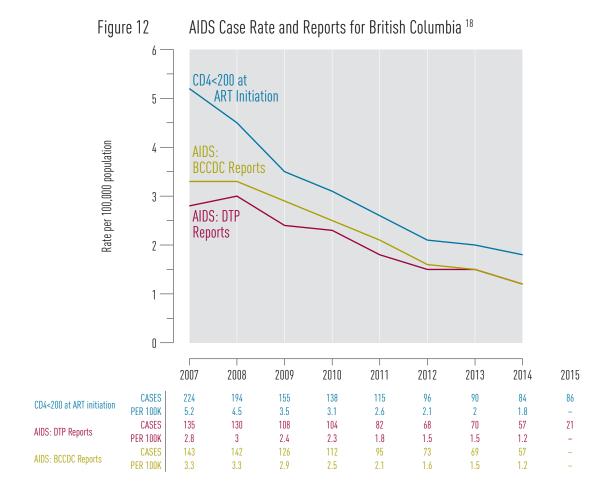
¹⁷ Data Source: Drug Treatment Program Database

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

AIDS-Defining Illness

Indicator 12. AIDS-Defining Illness

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



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

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.

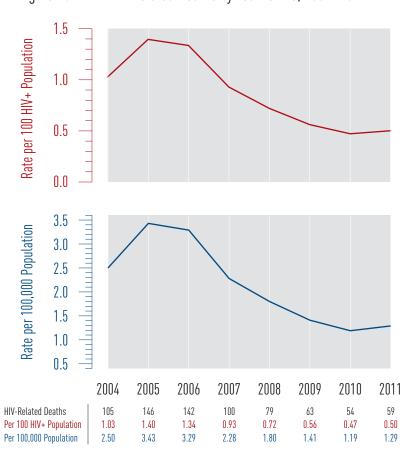


Figure 13 HIV-Related Deaths by Year for BC, 2004–2011 19

Limitation:

¹⁹ Data Source: BC Vital Statistics

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

^{2.} Mortality data is updated annually.

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

Appendices

Indicator : Episodes	l: Test (thousands)	2011 Q1	l Q2	Q3	Q4	2012 Q1	_	Q3	Q4	2013 Q1	3 Q2	Q3	Q4	2014 Q1	Q2	Q3	Q4	2015 Q1	Q2	Q3	O4
British Co	lumbia	35.1	33.5	40.3	37.6	42.3	41.5	45.8	46.6	54.8	58.0	55.9	54.6	62.0	66.1	71.2	70.5	77.9	77.0	79.6	80.4
Gender	Female	15.6	14.7	15.9	16.6	18.9	18.9	21.1	21.5	25.7	27.5	26.4	25.6	28.7	31.2	33.8	33.7	37.5	37.4	38.6	38.5
	Male	17.9	16.2	17.8	18.0	20.8	20.1	22.0	22.7	26.2	27.9	26.5	26.2	30.0	31.6	33.8	33.7	37.2	36.7	38.2	39.0
	Other	0.6	0.6	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
Age	< 30	12.3	11.8	13.1	13.0	14.0	13.5	15.1	14.7	15.7	16.7	16.8	16.3	17.4	17.6	19.2	18.9	19.7	19.9	21.7	21.8
	30-39	9.0	8.0	8.7	8.7	10.1	9.7	10.4	10.3	12.2	12.7	12.3	11.9	13.9	13.9	14.6	14.1	16.1	16.1	16.6	16.0
	40-49	6.0	5.5	5.8	6.0	6.9	6.8	7.1	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
	≥ 50	6.2	5.7	6.2	7.0	8.8	9.1	10.7	11.7	15.2	16.9	15.2	15.2	17.8	21.4	23.5	24.0	27.4	26.9	26.9	28.2
POC HIV	Tests	1.1	2.1	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	2.8	2.6	2.5	2.5
Fraser Hea	alth	8.7	8.2	9.1	9.0	10.1	10.1	11.0	11.6	13.5	14.0	13.8	12.9	13.6	14.5	14.9	15.1	16.1	15.9	16.7	16.8
Female		4.0	3.9	4.3	4.3	4.7	4.9	5.4	5.7	6.6	6.8	6.8	6.2	6.5	7.0	7.2	7.4	7.6	7.6	8.1	7.9
Male		4.6	4.3	4.7	4.6	5.3	5.1	5.5	5.7	6.6	7.0	6.7	6.5	6.9	7.1	7.3	7.5	8.1	7.8	8.3	8.5
Interior H	ealth	4.1	3.8	3.9	4.1	4.1	4.0	4.3	4.2	4.4	4.7	4.6	4.9	5.3	5.5	5.6	5.9	6.6	7.1	7.9	9.1
Female		2.0	2.0	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
Male		2.0	1.8	1.8	2.0	2.0	1.9	2.1	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
Island Hea	alth	4.0	3.5	3.6	3.5	4.0	3.7	3.9	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
Female		1.9	1.7	1.8	1.7	1.9	1.9	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
Male		2.0	1.7	1.8	1.7	2.0	1.8	1.9	1.8	2.0	2.1	2.0	1.9	2.2	2.1	2.1	2.1	2.6	2.6	2.7	2.9
Northern	Health	1.9	1.8	1.9	1.9	2.3	2.2	2.2	2.3	2.7	2.7	2.6	2.6	3.1	3.0	2.9	3.0	3.7	3.2	3.2	3.2
Female		1.0	0.9	1.0	1.0	1.2	1.1	1.2	1.2	1.3	1.4	1.3	1.3	1.6	1.5	1.5	1.5	1.9	1.6	1.6	1.6
Male		0.9	0.8	0.8	0.8	1.1	1.0	1.0	1.1	1.2	1.2	1.1	1.2	1.4	1.3	1.3	1.4	1.6	1.4	1.4	1.4
Vancouver	r Coastal Health	16.4	16.2	21.9	19.1	21.8	21.5	24.4	24.5	30.1	32.1	30.7	29.8	35.0	38.4	43.0	41.6	45.5	44.9	45.4	44.7
Female		6.6	6.2	6.9	7.4	9.0	9.0	10.5	10.6	13.6	14.8	14.0	13.6	15.8	17.9	20.0	19.6	21.8	21.6	21.8	21.2
Male		8.3	7.6	8.6	8.8	10.5	10.4	11.6	11.9	14.3	15.3	14.4	14.2	16.8	18.3	20.4	19.8	21.8	21.5	21.9	21.8

Indicator 2: Rate of HIV Testing per 100,000

		2009	2010	2011	2012	2013	2014	2015
British Col	umbia	2625.4	2646.5	2714.8	3332.8	4194.2	4969.4	5734.7
Fraser Hea	lth	2251.5	2267.9	2331.7	2805.3	3316.7	3591.8	3989.3
Interior He	ealth	2026.9	2072.7	2093.2	2182.3	2551.9	2967.4	4049.4
Island Hea	lth	1927.2	1919.4	1850.9	1986.0	2141.6	2326.0	2928.7
Northern I	Health	2291.0	2348.6	2432.9	2920.7	3332.3	3825.8	4334.1
Vancouver	Coastal Health	4127.0	4143.0	4344.6	5862.6	8135.6	10335.9	11636.4
Gender	Female	2447.1	2455.6	2524.0	3210.9	4168.0	4942.3	5760.8
	Male	2695.1	2735.4	2809.5	3398.2	4180.1	4952.6	5657.3
Age	< 30	2795.3	2802.8	2855.3	3245.5	3686.0	4048.1	4540.9
	30-39	5089.8	5226.3	5252.5	6098.5	7205.7	8108.3	9017.5
	40-49	3027.7	3027.5	3124.1	3846.6	4933.0	5707.4	6507.4
	≥ 50	1241.3	1281.2	1396.0	2179.4	3303.0	4414.6	5350.3

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Gender				rem Mal					6									6									6	16	13
Λ				viai < 30					48	63	76					47	55	43	68	72	53	49	65	50	51	55		53	41
Age									5	18	17	18				9	18	9	18	23	15	17	15	13	18	15		23	13
				30–3					18	30	30		16			11	10	16	25	18	11	17	21	25	15	16		19	12
				40-4					18	22	22		20			19	19	12	14	21	20	14	14	10	13	11	22	11	13
Г				≥ 50					13	12	18					13	13	12	23	18	12	15	25	18	14	24		18	16
Exposure				MSN					33	43	56					37	36	28	46	45	38	35	46	34	36	36		_	_
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Interior He	alth			•		t Re			1	4	3				5	0	2	1	4	4	4	3	4	4	6	5	1	5	4
				,		der 1			1	3	2				5	0	2	1	5	4	4	3	2	4	5	5		6	5
Island Heal	th					t Re			5	2	7	7	6	5	3	5	12	4	9	10	7	6	4	7	8	3	8	8	6
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Stage 1	66	53		56		12	8	6	9	11	54				38	19			22	11							0 16		9
Stage 2a	33	39		45	31	5	7	5	6	4	28				27	7			12	5		12	4		7	9 1			
Stage 2b	34			30		12	8	6	3	5	22		24		22	5			4	4	9	8	7				2 10		
Stage 3	76					22		5		.		55							4	4		17				31 2			15
Unknown			16				4	1	1									7			13		3		11 1		5 2		
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Stage 2a	3	5				20					6	5	6	5	1	7		5			0	2	0	2	0		0 0		
Stage 2b	9	13		7		15					5	3	6	0			15		7	9	2	1	0	0	0	0	0 0		
Stage 3	30	21	25			25			28	17	18	7	6	6	4	28	31	20	32	21	3	3	0	2	4	2	0 2	5	3
Unknown	1	5				22				23	4	6	1	4	3	7			4	9	1	1	1	3	1	1	2 3	0	4
Total	54	60	60	67	72	152	170	148	157	150	52	34	29	25	25	82	75	52	68	64	6	7	2	8	8	4	2 5	8	13

British Columbia 10000 Age Category < 30 489 30-39 1340 40-49 2672 ≥ 50 5490 Age Category and MSM < 30 125 Status 30-39 393 Status 40-49 737 ≥ 50 2041 Non-MSM < 30 46 30-30 275	336 1227 2490 5129 114 384	7670 280 1018 2041 4331 101 329		50 203 21 808 95 1745	5551 163 670 1443 3275
$30-39 & 1340 \\ 40-49 & 2672 \\ \geq 50 & 5490 \\ Age Category & MSM & < 30 & 125 \\ and & MSM & 30-39 & 393 \\ Status & 40-49 & 737 \\ & \geq 50 & 2041 \\ Non-MSM & < 30 & 46 \\ \hline$	1227 2490 5129 114 384	1018 2041 4331 101	92 189 400	21 808 95 1745 65 3815	670 1443
	2490 5129 5 114 5 384	2041 4331 101	189 406	95 1745 65 3815	1443
≥ 50	5129 5 114 5 384	4331 101	406	55 3815	
Age Category and MSM < 30 125 and MSM $30-39$ 393 Status $40-49$ 737 ≥ 50 2041 Non-MSM < 30 46	114 384	101	Ç		3275
and MSM $30-39$ 393 Status $40-49$ 737 ≥ 50 2041 Non-MSM < 30 46	384			93 78	32/3
Status $40-49 737$ $\geq 50 2041$ Non-MSM < 30 46		329	24		63
40-49 737 ≥ 50 2041 Non-MSM < 30 46	721		30)5 269	227
Non-MSM < 30 46		647	60	9 576	498
	2017	1863	177	78 1698	1524
	45	36	3	30 24	18
30–39 375		332	30	256	200
40–49 880		770	71		481
≥ 50 1510		1367	128		938
Unknown < 30 318		143	12		82
30–39 571		357	31		243
40–49 1055		625	57		464
≥ 50 1939		1101	100		813
Gender Male 8195		6338	593		4707
Female 1805		1332	120		844
Injection PWID 2543 Drug Use Non-PWID 4500		2302	214		1484
- 11011 1 11115 1300		4007	378		3123
Unknown 2957		1361	119		944
MSM Status MSM 3297		2939	278		2312
Non-MSM 2811		2504	232		1637
Unknown 3892		2226	201		1602
Health Fraser Health 2102		1832	172		1335
Authority Interior Health 605		489	45		352
Island Health 989		878	83		600
Northern Health 291	276	241	22		127
Vancouver Coastal Health 4711	4570	4085	388	3615	3136
Indicator 6: Programmatic Compliance Score (PCS)					
2014			2015		
Q1 Q		Q4	Q1		Q3 Q4
< 3 CD4 Tests 21.8% 24.99		21.1%	19.3%	16.4% 16.2	
< 3 Viral Load Tests 8.3% 7.49		5.9%	7.3%	6.6% 6.7	
No Baseline Genotype 3.3% 3.29		2.2%	3.6%	4.6% 3.5	
Baseline CD4 < 200 cells/ μ L 22.0% 24.49		24.2%	23.0%	21.6% 20.6	
Non-Recommended ART 10.5% 11.29		3.7%	2.2%	1.7% 1.6	
Non Viral suppression at 9 Mo. 35.8% 36.19	% 33.2%	30.6%	27.7%	25.6% 24.4	24.5%
PCS Score: 0 139 12	1 141	159	171	178 1	72 163
PCS Score: 1 123 12	9 133	111	111	101	84 89
PCS Score: 2 69 6	6 67	62	46	47	37 43
PCS Score: 3 23 2.	2 21	19	22	17	16 20
PCS Score: 4 or more 9 1	1 8	5	7	5	6 7
Total (n=) 363 34	9 370	356	357	348 3	15 322

Indicator 7: New DTP A	RV Participants									
	2014 Q1	Q2		Q3	Q4	2015 Q1	Q2)	Q3	Q4
First Starts	106	89		78	78	88	97		77	74
Experienced Starts	116	133		137	113	133	127		121	137
Indicator 8: CD4 Cell Co	ount at ADT Initiati	on for ADV	Noïvo F	TD Dauti	cinanto					
CD4 ≥ 500	34	39	-Naive L	29	29	32	34	1	30	29
CD4 350-499	20	20		15	14	13	12		10	9
CD4 200-349	30	20 11		12	15	22	21		14	13
CD4 50–199				13						
CD4 50-199	13	16		8	12 6	13	17		11 10	10
	8	2				4	11			300
CD4 Median (cells/μL)	360	480		410	390	388	340		380	380
Total (n=)	105	88		77	76	84	95	•	75	69
Indicator 9: Active and I	nactive DTP Partici	ipants								
Active DTP Participants	6810	6896	6	948	6995	7059	7138	3	7182	7244
Inactive DTP Participant	s 1108	1114	1	130	1152	1157	1151		1162	1160
Indicator 10: Antiretrov	iral Adherence									
≥ 95%	76	70		92	81	79	74	ŀ	73	69
80% to < 95%	12	13		15	14	14	20)	16	12
40% to < 80%	8	10		12	7	13	8	3	12	3
< 40%	1	1		2	0	1	C)	1	2
Total (n=)	97	94		121	102	107	102	2	102	86
In dianton 11. Desistance	Tooting and Deculto									
Indicator 11: Resistance	4968	4906	-	090	4919	5244	5205		5081	5093
Suppressed										
Wild Type	766	722		685	622	693	652		764	652
Never Genotyped	48	50		41	34	41	42		41	39
1-Class	149	148		145	141	151	143		137	112
2-Class	27	26		25	27	31	27		31	23
3-Class Total (n=)	5 5963	7 5859	_	7 993	3 5746	6168	6077		7 6061	2 5922
Total (n=)	3903	3639	3	993	3/40	0108	6077		0001	3922
Indicator 12: AIDS-Defi		2007	2008	2009	2010	2011	2012	2013	2014	2015
CD4 < 200 at	Cases	224	194	155	138	115	96	90	84	86
ART initiation	Rate per 100,000	5.2	4.5	3.5	3.1	2.6	2.1	2.0	1.8	_
AIDS Cases	Cases	135	130	108	104	82	68	70	57	21
(DTP Reports)	Rate per 100,000	2.8	3.0	2.4	2.3	1.8	1.5	1.5	1.2	-
AIDS Cases	Cases	143	142	126	112	95	73	69	57	_
(BCCDC Reports)	Rate per 100,000	3.3	3.3	2.9	2.5	2.1	1.6	1.5	1.2	-
Indicator 13: HIV-Relate	ed Mortality	2004	2005	2006	2007	2008	2009	2010	2011	
British Columbia		105	146	142	100	79	63	54	59	
Per 100 HIV+ Population	1	1.03	1.40	1.34	0.93	0.72	0.56	0.47	0.50	
Per 100,000 Population		2.50	3.43	3.29	2.28	1.80	1.41	1.19	1.29	
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