

HIV MONITORING QUARTERLY REPORT

FOR BRITISH COLUMBIA

SECOND 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.

For Indicator 5 (page 22), recent data have allowed for more comprehensive death information. As a result, 2015 Q2 data for the Diagnosed and Linked to Care steps may be slightly lower than previously reported.

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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

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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 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

Q3

2010

ВС

Q1 Q2 Q3

2011

Q1 Q2 Q3 Q4 Q1 Q2

2013

44.6 45.4 48.6 45.4 52.9 50.8 56.7 55.2 59.3 59.8 67.3 69.8 67.9 66.7 74.9 78.0 83.5 82.9 91.0 88.1

2014

Q1 Q2

2015

Q1 Q2 Q3

2012

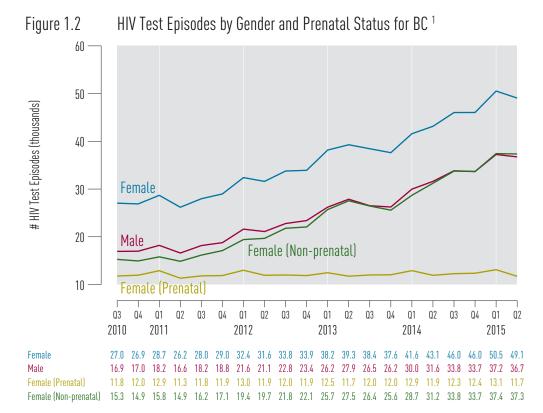
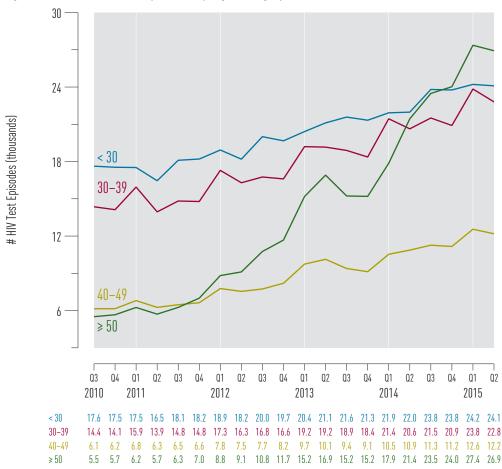
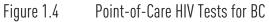
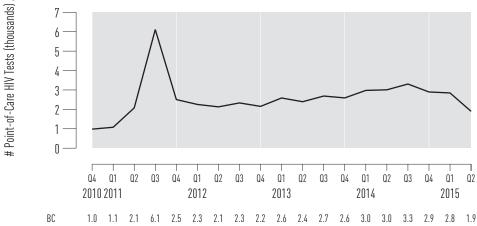


Figure 1.3 HIV Test Episodes by Age Category for BC ^{1,2}







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.

Limitations:

- 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.

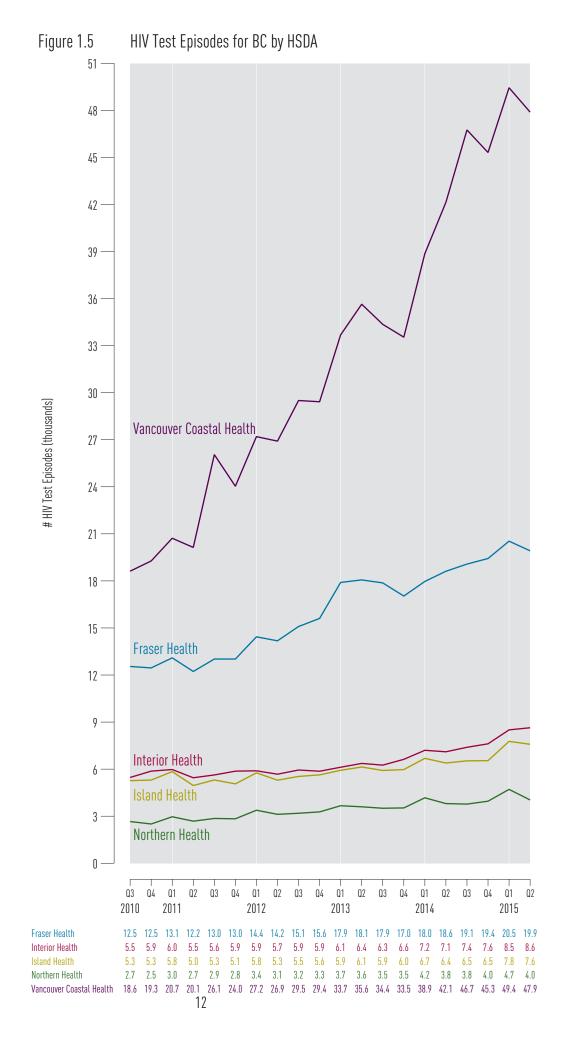


Figure 1.6 HIV Test Episodes for Non-prenatal Females in BC by Health Authority ¹

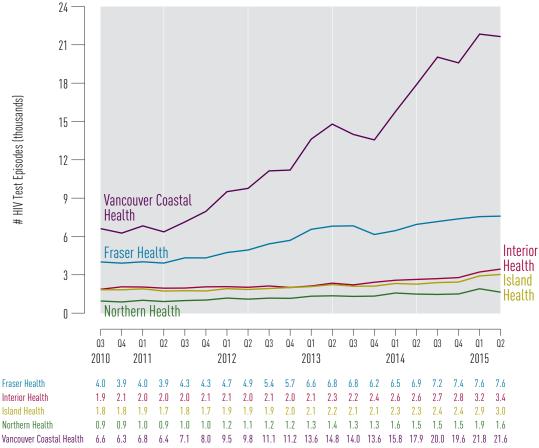
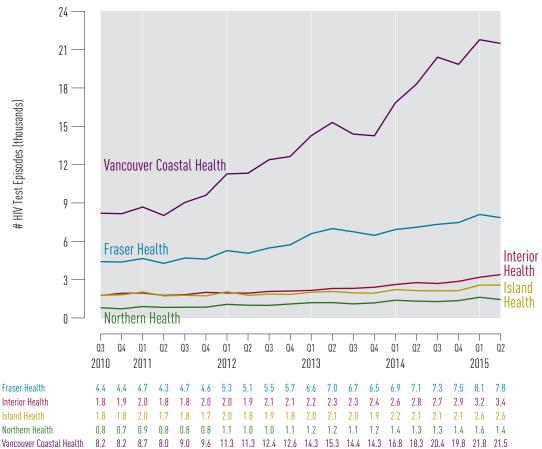
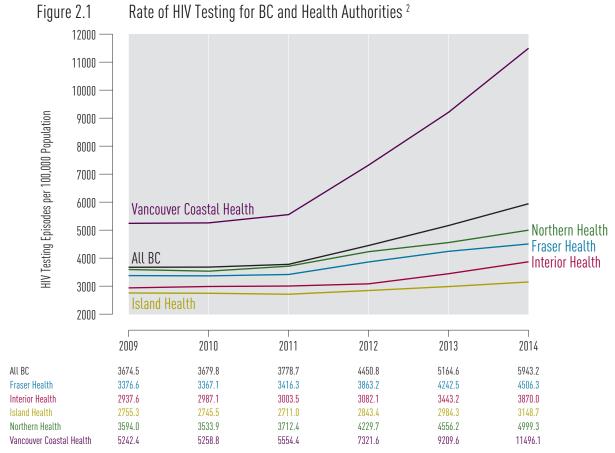


Figure 1.7 HIV Test Episodes for Males in BC by HSDA ¹





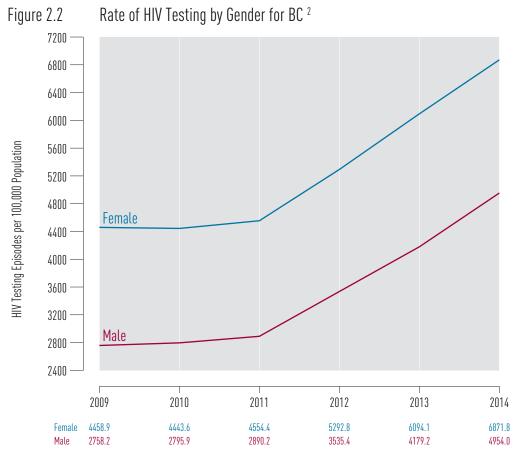


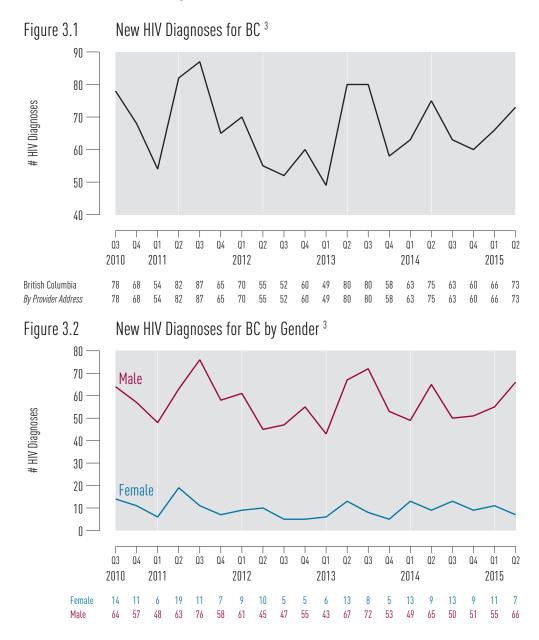
Figure 2.3 Rate of HIV Testing by Age Category for BC $^{\rm 2}$ 13000 12000 11000 10000 30-39 9000 8000 HIV Testing Episodes per 100,000 Population 7000 6000 5000 < 30 4000 40-49 3000 2000 ≥ 50 1000 2009 2010 2012 2011 2013 2014 4006.5 4039.0 < 30 4043.5 4383.1 4784.0 5116.5 30-39 8807.2 9006.0 9211.3 10184.9 11249.1 12228.4 **5321.7** 3306.1 40-49 3356.5 3369.5 3506.4 4241.6 6109.8 1243.0 1284.8 1399.0 2181.7 4418.8 ≥ 50

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.

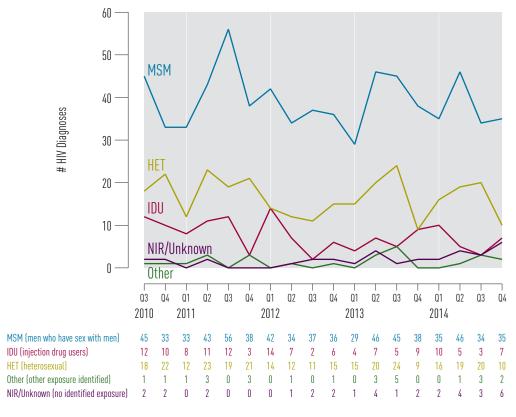
Indicator 3. New HIV Diagnoses



З Data Source: всссьс. "By Provider Address" is graphed as dashed line in same colour.

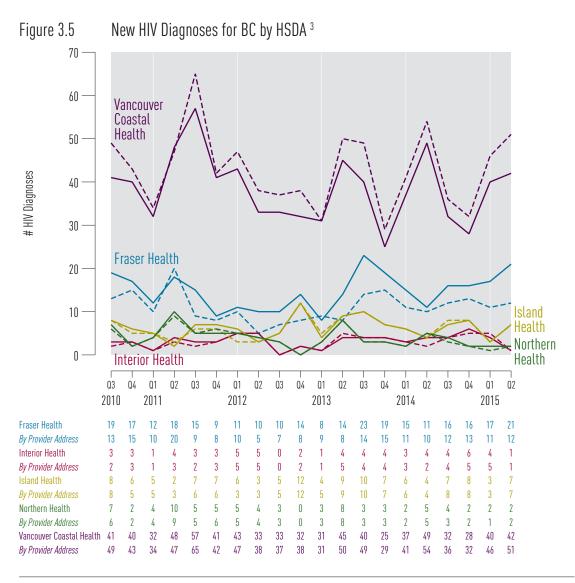
Figure 3.3 New HIV Diagnoses for BC by Age Category 3 30 -30-39 40-49 ≥ 50 # HIV Diagnoses 0 -Q3 Q4 Q1 Q2 Q3 Q4 Q2 Q3 Q3 Q4 Q1 Q2 Q1 Q1 Q1 < 30 30-39 40-49 ≥ 50

Figure 3.4 New HIV Diagnoses for BC by Exposure Category 3.4



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

⁴ MSM=men who have sex with men; IDU= injection drug user; HET=heterosexual. NIR=No identified risk/exposure. "By Provider Address" is graphed as dashed line in same colour.



³ Data Source: BCCDC. "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.

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. AIDC									
2a			CD4 350-499	and	No AIDS case report									
2b	Stage 0		CD4 200-349		ισμοιτ									
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 BC, 2010–2014 ⁵

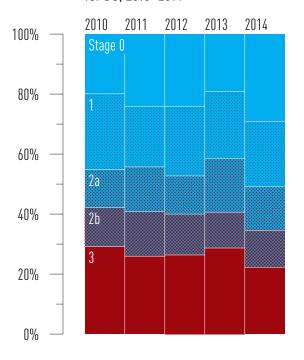


Figure 4.2 Stage of HIV Infection at Diagnosis by Gender for BC, 2010–2014 $^{\rm 5}$

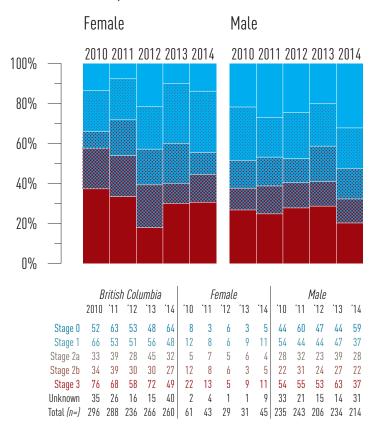


Figure 4.3 Stage of HIV Infection at Diagnosis by Age Category for BC, 2010–2014 ⁵

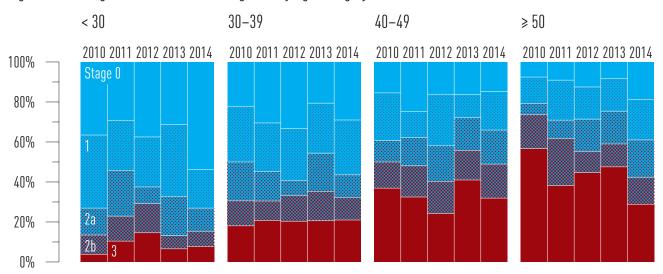
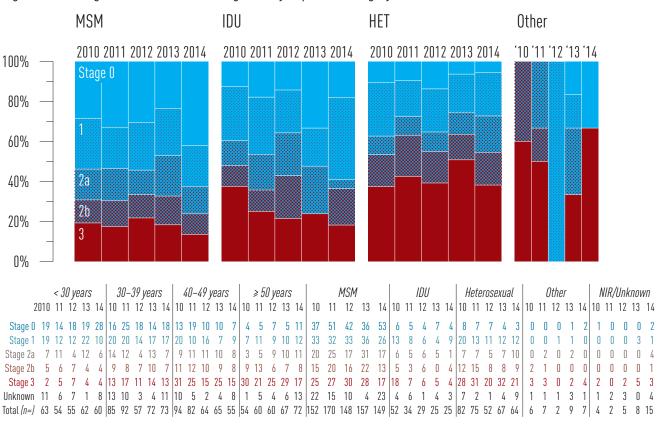


Figure 4.4 Stage of HIV Infection at Diagnosis by Exposure Category for BC, 2010–2014 5.6

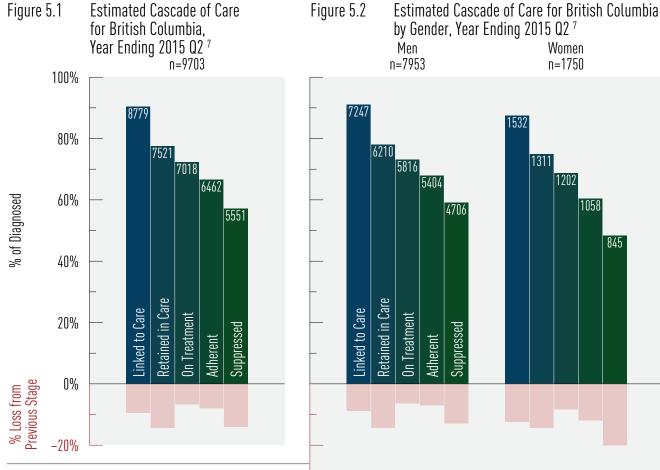


⁶ MSM=men who have sex with men; IDU= injection drug user; HET=heterosexual. NIR=No identified risk/exposure.

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. 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 2014 Q3–2015 Q2 in BC overall and stratified by sex and age for each Health Authority.

Recent data have allowed for more comprehensive death information. As a result, data for the Diagnosed and Linked to Care steps may be slightly lower than previously reported.

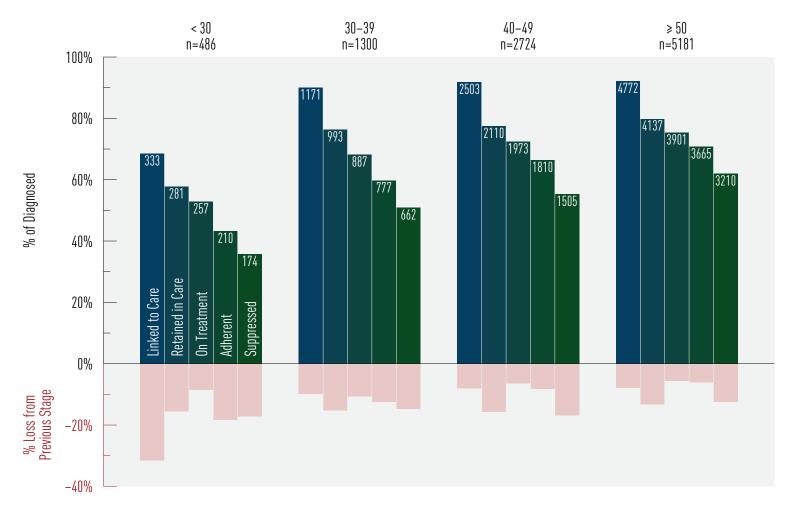


- 7 Data is for the period 2014 Q3-2015 Q2. Data Sources:
 - i British Columbia Centre for Excellence Drug Treatment Program (DTP) Database (ARV use, VL and CD4 count).
 - ii Administrative data (ex. MSP billings; hospitalization data from the Discharge Abstract Database (DAD)).

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

NB: Transgender has been assigned to their biological sex.





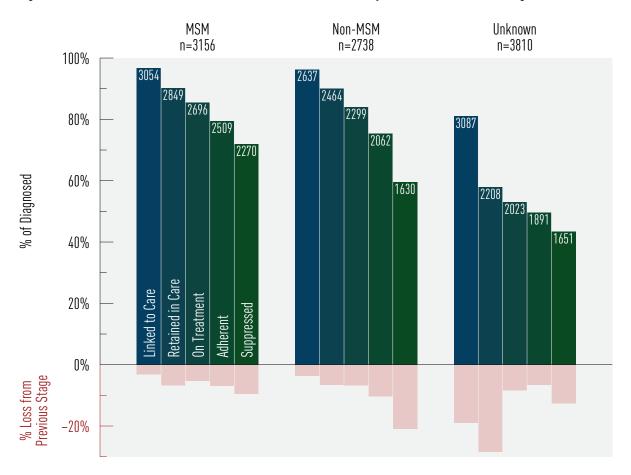
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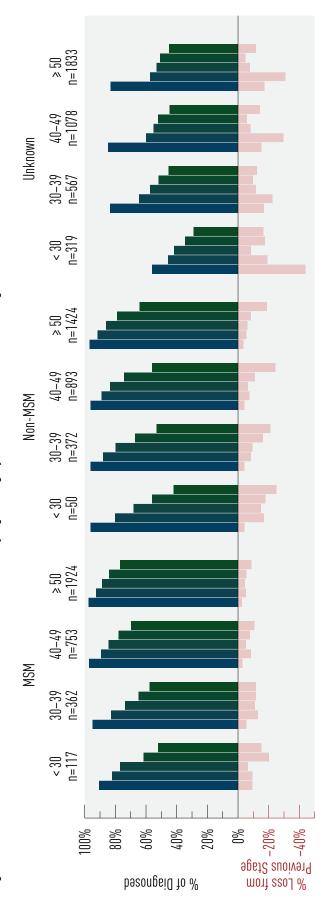
Recent updates to the DTP database have allowed for more comprehensive information on HIV risk group category. As a result, 2014 Q4 data may differ significantly from preceding reports in terms of total numbers ascribed to each risk group.

⁹ Data is for the period 2014 Q3–2015 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 2015 Q2 $^{\it 9}$ Figure 5.5



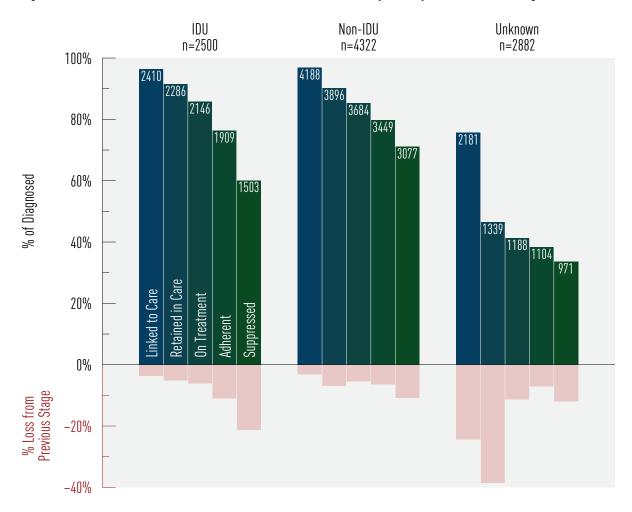
Data is for the period 2014 Q3-2015 Q2. Data Sources:

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. 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.

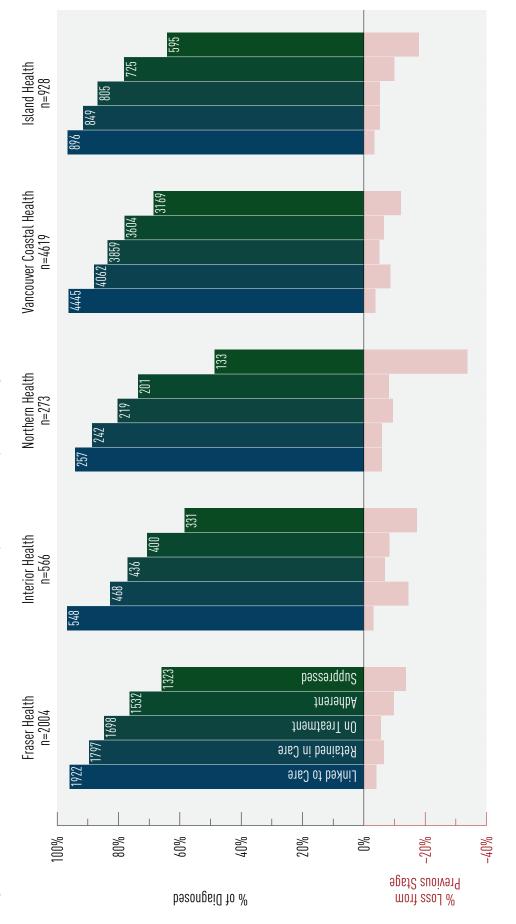
Recent updates to the DTP database have allowed for more comprehensive information on HIV risk group category. As a result, 2014 Q4 data may differ significantly from preceding reports in terms of total numbers ascribed to each risk group.

⁹ Data is for the period 2014 Q3–2015 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 2015 Q2 $^{\it 9}$ Figure 5.7



9 Data is for the period 2014 Q3-2015 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. 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)).

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 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;
- 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 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, 2013 Q3–2015 Q2 10

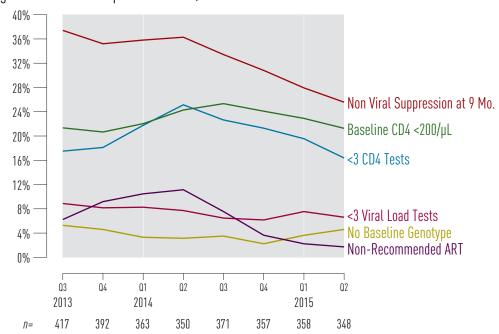
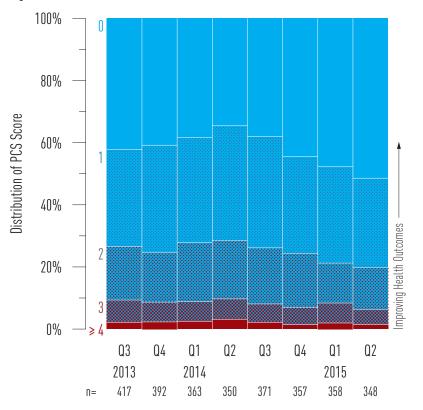


Figure 6.2 Historical Trends for PCS Score for BC, 2013 Q3-2015 Q2 10,11



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

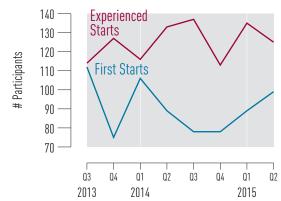
¹¹ Each quarter's data is calculated as the sum of the 4 quarters leading up to it. e.g. 2013 Q1 is calculated from 2012 Q2 – 2013 Q1. NB: A score of o is the best score and a score of 4 or more is the worst score.

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.

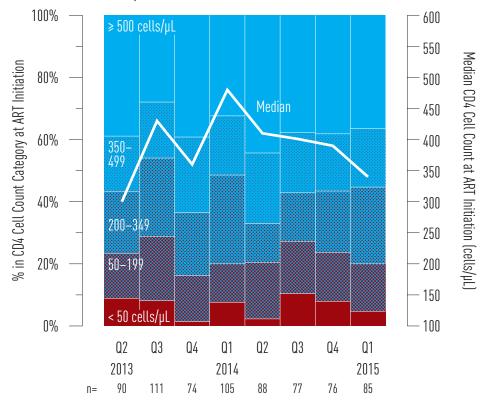
Indicator 7. New Antiretroviral Therapy Starts in BC

Figure 7 BC-CfE Drug Treatment Program Enrollment: New ART Participants in BC, 2013 Q3-2015 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, 2013 Q3-2015 Q2 ¹³



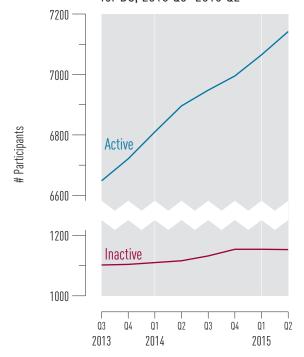
- Data Source: Drug Treatment Program Database Limitation: DTP participants are designated to an HA based on most current residence provided by the participant.
- 3 Data Source: Drug Treatment Program Database Limitations: CD4 cell count data is approximately 80% complete.

Indicator 9. Active and Inactive DTP Participants

Table 3. Distribution of People on ART for BC, 2015 Q2 14

		Fraser	Interior	Island	Northern	Vancouver Coastal	Total BC
Age	< 30	89	15	26	13	143	286
	30-39	257	54	88	53	524	977
	40-49	534	104	229	68	1129	2065
	≥ 50	839	272	479	92	2131	3814
Gender	Male	1322	350	663	143	3459	5940
	Female	397	95	159	83	468	1202
Exposure	MSM	535	128	217	30	1853	2765
	IDU	456	147	276	124	1137	2141
Total		1719	445	822	226	3927	7142

Figure 9 Active and Inactive DTP Participants for BC, 2013 Q3-2015 Q2 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.

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

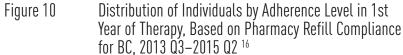
15 Active DTP participants: An individual who has had medication prescribed at least once in the preceding quarter.

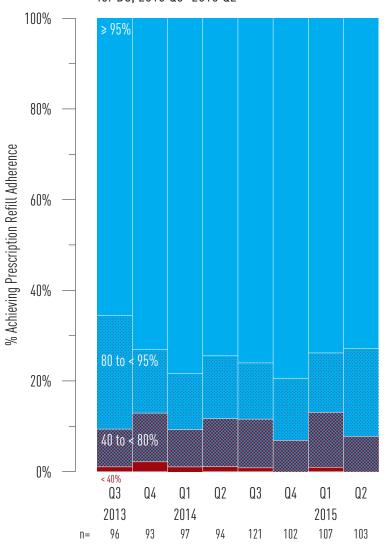
Inactive DTP participants: Persons no longer prescribed drugs through the HIV/AIDS Drug Treatment Program in the last quarter.

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

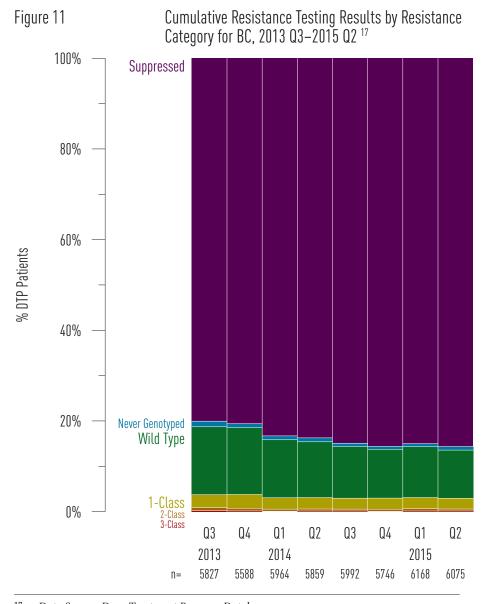




¹⁶ Data Source: Drug Treatment Program Database Limitation: Prescription refill adherence is used as a proxy for patient adherence.

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.

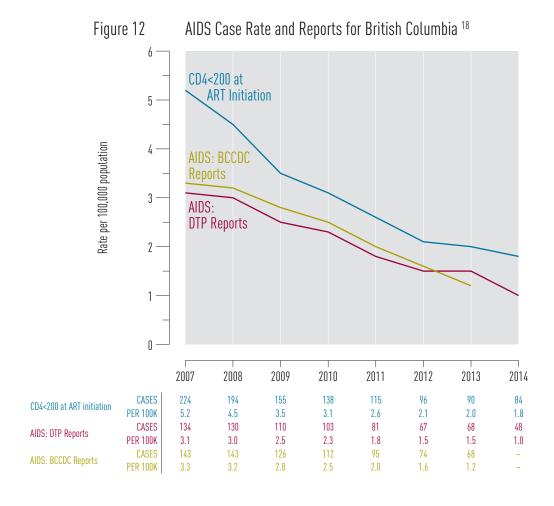


¹⁷ Data Source: Drug Treatment Program Database

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 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.

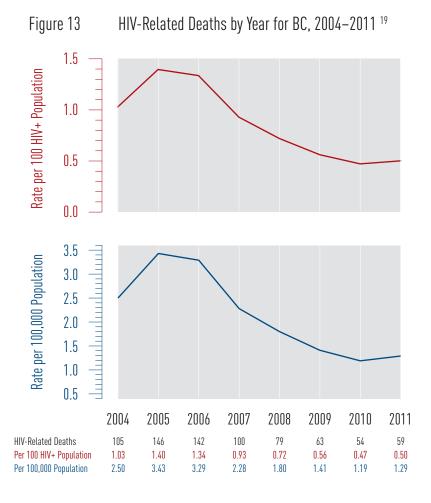


Data Source: DTP AIDS cases are obtained from the Drug Treatment Program Database; BCCDC AIDS cases are obtained from the BCCDC; CD4<200 at ART initiation data came from the DTP database.

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.

Indicator 13. HIV-Related Mortality

Evidence indicates that individuals who initiate treatment with recommended ART in a timely fashion may live near normal lifespans. Excess mortality among HIV positive persons is, therefore, an important measure of HIV care with a goal of minimizing HIV-related mortality in British Columbia.



Limitation:

¹⁹ Data Source: BC Vital Statistics

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

^{2.} Mortality data is updated annually.

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

Appendices

Indicator 1: Test Episodes (thousands)		2010 Q3	Q4	2011 Q1	Q2	Q3	Q4	2012 Q1	Q2	Q3	Q4	2013 Q1	Q2	Q3	Q4	2014 Q1	Q2	Q3	Q4	2015 Q1	Q2
British Col	44.6	45.4	48.6	45.4	52.9	50.8	56.7	55.2	59.3	59.8	67.3	69.8	67.9	66.7	74.9	78.0	83.5	82.9	91.0	88.1	
Gender	Female	27.0	26.9	28.7	26.2	28.0	29.0	32.4	31.6	33.8	33.9	38.2	39.3	38.4	37.6	41.6	43.1	46.0	46.0	50.5	49.1
	Male	16.9	17.0	18.2	16.6	18.2	18.8	21.6	21.1	22.8	23.4	26.2	27.9	26.5	26.2	30.0	31.6	33.8	33.7	37.2	36.7
	Other	0.6	0.6	0.6	0.6	0.6	0.6	0.4	0.4	0.4	0.3	0.3	0.3	0.3	0.2	0.3	0.3	0.3	0.2	0.3	0.4
Female (Pr	renatal)	11.8	12.0	12.9	11.3	11.8	11.9	13.0	11.9	12.0	11.9	12.5	11.7	12.0	12.0	12.9	11.9	12.3	12.4	13.1	11.7
Female (Non-prenatal)		15.3	14.9	15.8	14.9	16.2	17.1	19.4	19.7	21.8	22.1	25.7	27.5	26.4	25.6	28.7	31.2	33.8	33.7	37.4	37.3
Age	< 30	17.6	17.5	17.5	16.5	18.1	18.2	18.9	18.2	20.0	19.7	20.4	21.1	21.6	21.3	21.9	22.0	23.8	23.8	24.2	24.1
	30-39	14.4	14.1	15.9	13.9	14.8	14.8	17.3	16.3	16.8	16.6	19.2	19.2	18.9	18.4	21.4	20.6	21.5	20.9	23.8	22.8
	40-49	6.1	6.2	6.8	6.3	6.5	6.6	7.8	7.5	7.7	8.2	9.7	10.1	9.4	9.1	10.5	10.9	11.3	11.2	12.6	12.2
	≥ 50	5.5	5.7	6.2	5.7	6.3	7.0	8.8	9.1	10.8	11.7	15.2	16.9	15.2	15.2	17.9	21.4	23.5	24.0	27.4	26.9
POC HIV	Tests	0.0	1.0	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	1.9
Fraser Hea	alth	12.5	12.5	13.1	12.2	13.0	13.0	14.4	14.2	15.1	15.6	17.9	18.1	17.9	17.0	18.0	18.6	19.1	19.4	20.5	19.9
Female (Non-prenatal)	4.0	3.9	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	6.9	7.2	7.4	7.6	7.6
Male		4.4	4.4	4.7	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
Interior H	ealth	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	6.6	7.2	7.1	7.4	7.6	8.5	8.6
Female (Non-prenatal)	1.9	2.1	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	3.4
Male		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	2.4	2.6	2.8	2.7	2.9	3.2	3.4
Island Hea	alth	5.3	5.3	5.8	5.0	5.3	5.1	5.8	5.3	5.5	5.6	5.9	6.1	5.9	6.0	6.7	6.4	6.5	6.5	7.8	7.6
Female (.	Non-prenatal)	1.8	1.8	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
Male		1.8	1.8	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
Northern l	Health	2.7	2.5	3.0	2.7	2.9	2.8	3.4	3.1	3.2	3.3	3.7	3.6	3.5	3.5	4.2	3.8	3.8	4.0	4.7	4.0
Female (Non-prenatal)	0.9	0.9	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
Male		0.8	0.7	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
Vancouver	Coastal Health	18.6	19.3	20.7	20.1	26.1	24.0	27.2	26.9	29.5	29.4	33.7	35.6	34.4	33.5	38.9	42.1	46.7	45.3	49.4	47.9
Female (Non-prenatal)	6.6	6.3	6.8	6.4	7.1	8.0	9.5	9.8	11.1	11.2	13.6	14.8	14.0	13.6	15.8	17.9	20.0	19.6	21.8	21.6
Male		8.2	8.2	8.7	8.0	9.0	9.6	11.3	11.3	12.4	12.6	14.3	15.3	14.4	14.3	16.8	18.3	20.4	19.8	21.8	21.5

Indicator 2: Rate of HIV Testing per 100,000

		2009	2010	2011	2012	2013	2014
British Colu	ımbia	3674.5	3679.8	3778.7	4450.8	5164.6	5943.2
Fraser Heal	th	3376.6	3367.1	3416.3	3863.2	4242.5	4506.3
Interior He	alth	2937.6	2987.1	3003.5	3082.1	3443.2	3870.0
Island Heal	th	2755.3	2745.5	2711.0	2843.4	2984.3	3148.7
Northern F	Iealth	3594.0	3533.9	3712.4	4229.7	4556.2	4999.3
Vancouver	Coastal Health	5242.4	5258.8	5554.4	7321.6	9209.6	11496.1
Gender	Female	4458.9	4443.6	4554.4	5292.8	6094.1	6871.8
	Male	2758.2	2795.9	2890.2	3535.4	4179.2	4954.0
Age	< 30	4043.5	4006.5	4039.0	4383.1	4784.0	5116.5
	30-39	8807.2	9006.0	9211.3	10184.9	11249.1	12228.4
	40-49	3356.5	3369.5	3506.4	4241.6	5321.7	6109.8
	≥ 50	1243.0	1284.8	1399.0	2181.7	3306.1	4418.8

									2010		201				201				201				201					2015	
Indicator 3				_								Q2							Q1										Q2
British Colu	British Columbia By Client Residenc							78	68	54	82	87								80	58	63				60	66	73	
		By Provider Address					ess	78	68	54	82	87								80	58	63				60	66	73	
Gender								14	11	6	19	11	7							8	5	13			13	9	11	7	
	Male						64	57	48	63	76								72	53	49	65	; 5	50	51	55	66		
Age	Age < 30						22	9	5	18	17								23	15	17	15		13	18	15	16		
	30–39						23	20	18	30	30	13	16	17	7 11				18	11	17	21	ι 2	25	15	16	17		
			4	40-4	19				19	24	18	22	22	19	20						21	20	14	14	ł	7	13	11	24
				≥ 50					14	15	13	12	18		16	13	3 13			23	18	12	15	25	5]	18	14	24	16
Exposure				MSN	M				45	33	33	43	56	38	3 42	34	1 37	36	29	46	45	38	35	46	5 3	34	35	-	-
				IDU	ſ				12	10	8	11	12	. 3	14	: 7	7 2	2 6	4	7	5	9	10	5	5	3	7	-	_
				HET	Γ				18	22	12	23	19	21	14	: 12	2 11	. 15	15	20	24	9	16	19) 2	20	10	-	-
			(Oth	er				1	1	1	3	0	3	()]	l ()]	0	3	5	0	0	1	1	3	2	-	-
				NIR	/Un	knov	vn		2	2	0	2	C	C) ()]	l 2	2 2	1	4	1	2	2	4	1	3	6	-	_
Fraser Heal	lth			By C	Clien	t Res	sider	ıce	19	17	12	18	15	9	11	10) 1(14	. 8	14	23	19	15	11	1 1	16	16	17	21
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Interior He	alth			By C	Clien	t Res	sider	ıce	3	3	1	4	3	3	5		5 () 2	1	4	4	4	3	4	4	4	6	4	1
				Ву Р	rovi	der A	Addr	ess	2	3	1	3	2	3	3 5	5 5	5 () 2	2 1	5	4	4	3	2	2	4	5	5	1
Island Heal	lth			By C	Clien	t Res	sider	nce	8	6	5	2	7	7	' (3	3 5	12	4	9	10	7	6	4	4	7	8	3	7
			i	Ву Р	rovi	der A	Addro	ess	8	5	5	3	6	6	5 3	3	3 5	12	5	9	10	7	6	4	4	8	8	3	7
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Stage 0	52	63		48	64	8	3	6	3	5					_		4 1			16						19	10	10	7
Stage 1	66	53	51		48	12	8	6	9	11								2 22								10	16	7	9
Stage 2a	33	39	28	45	32	5	7	5	6	4			23		28	7 1		4 12				4		7	9	11	11	10	8
Stage 2b		39		30		12	8	6	3	5			24		22	5		7 4		9	8	7		7		12		9	8
Stage 3		68				22						55				2		7 4			17			.				-	
Unknown		26							1			21									10							4	8
Total																			2 60										
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Stage 0	4	5 11	7 0			37					6	5	4	7	4			7 4 1 11		0	0	0	1	2	1	0	0	0	2
Stage 1		11				33						8	6	4					12		0	1	1	0	0	0	0	3	1
Stage 2a	3					20					6	5	6	5	1	7		5 7			2	0	2	0	0	0	0	0	4
Stage 2b		13	6	7		15					5	3	6	0		.2 1				2	1	0	0	0	0	0	0	0	1
Stage 3		21				25						7	6	5	.				2 21	3	3	0	2	4	2	0		5	3
Unknown	1		4			22					4	6	1	4	3	7		1 4		1	1	1	3	1	1	2		0	4
Total	54	60	60	6/	/2	152	1/0	148	157	149	52	54	29	25	25 8	52 7	5 5	2 67	64	6	7	2	9	7	4	2	5	8	15

	V Cascade of C	Care 1	DIAGNOSED	LINKED	RETAINED	Ol		DHERENT	SUPPRESSED
British Columb			9703	8779	7521		7018	6462	5551
Age Category	< 30		486	333	281		257	210	174
	30-39		1300	1171	993		887	777	662
	40-49		2724	2503	2110		1973	1810	1505
	≥ 50		5181	4772	4137			3665	3210
Age Category	MSM	< 30	117	106	96		90	72	61
and MSM Status		30-39	362	343	299		266	235	208
Status		40-49	753	731	671		636	587	524
		≥ 50	1924	1874	1782		1704	1615	1477
	Non-MSM	< 30	50	48	40		34	28	21
		30-39	372	357	327		297	249	197
		40-49	893	858	794		744	663	501
		≥ 50	1424	1376	1303		1224	1122	911
	Unknown	< 30	319	179	145		133	110	92
		30-39	567	472	366		324	293	257
		40-49	1078	914	645		593	560	480
		≥ 50	1833	1522	1052		973	928	822
Gender	Male		7953	7247	6210		5816	5404	4706
	Female		1750	1532	1311		1202	1058	845
Injection	IDU		2500	2410	2286		2146	1909	1503
Drug Use	Non-IDU		4322	4188	3896		3684	3449	3077
	Unknown		2882	2181	1339		1188	1104	971
MSM Status	MSM		3156	3054	2849		2696	2509	2270
Wiowi Status	Non-MSM		2738	2637	2464		2299	2062	1630
	Unknown		3810	3087	2208		2023	1891	1651
Health	Fraser Health		2004	1922	1797		1698	1532	1323
Authority	Interior Health	1	566	548	468		436	400	331
,	Island Health	1	928	896	849		805	725	595
	Northern Heal	lth	273	257	242		219	201	133
	Vancouver Co		4619	4445	4062		3859	3604	3169
				4443	4002		3639	3004	3109
Indicator 6: Pr	ogrammatic Co	•	ore (PCS)	2011					
		2013 Q3	Q4	2014 Q1	Q2	Q3	Q4	2015 Q1	Q2
< 3 CD4 Tests		17.5%	18.1%	21.8%	25.1%	22.6%	21.3%	19.6%	
< 3 Viral Load	Tests	8.9%	8.2%	8.3%	7.7%	6.5%	6.2%	7.5%	
No Baseline Ge		5.3%	4.6%	3.3%	3.1%	3.5%	2.2%	3.6%	
Baseline CD4 <	, ,	21.3%	20.7%	22.0%	24.3%	25.3%	24.1%	22.9%	
Non-Recomme	· ·	6.2%	9.2%	10.5%	11.1%	7.5%	3.6%	2.2%	
	oression at 9 Mc		35.2%	35.8%	36.3%	33.4%	30.8%	27.9%	
PCS Score: 0	ression at 9 Mic	176	160	139	121	141	159	171	179
PCS Score: 0		130	135	123	121	133	111	111	100
PCS Score: 1 PCS Score: 2		72	63	69	66	67	62	46	
PCS Score: 3		30	25	23	23	22	20	23	
PCS Score: 4 or	r more	9	9	9	11	8	5 357	7	
Total (n=)		417	392	363	350	371	357	358	348
	w DTP ARV P								
First Starts		112	75	106	89	78	78	89	
Experienced St	arts	114	127	116	133	137	113	135	125

Indicator 8: CD4 Cell	Count at ART Initiation	on for ARV-	Naïve DTP I	Participants									
	2013 Q3	Q4	2014 Q1	Q2		Q3	Q4	2015 Q1	Q2				
CD4 ≥ 500	31	29	34	39		29	29	33	34				
CD4 350-499	20	18	20	20		15	14	13	13				
CD4 200-349	28	15	30	11		12	15	22	21				
CD4 50-199	23	11	13	16		13	12	13	16				
CD4 < 50	9	1	8	2		8	6	4	11				
CD4 Median (cells/μL)	300	430	360	480	4	410	401	390	340				
Total (n=)	111	74	105	88		77	76	85	95				
Indicator 9: Active and Inactive DTP Participants													
Active DTP Participan	ts 6648	6721	6810	6896	69	948	6995	7065	7142				
Inactive DTP Participa	ints 1102	1104	1110	1116	11	132	1154	1154	1153				
Indicator 10: Antiretro	oviral Adherence												
≥ 95%	63	68	76	70		92	81	79	75				
80% to < 95%	24	13	12	13		15	14	14	20				
40% to < 80%	8	10	8	10		13	7	13	8				
< 40%	1	2	1	1		1	0	1	0				
Total (n=)	96	93	97	94	1	21	102	107	103				
Indicator 11: Resistano	ce Testing and Results												
Suppressed	4665	4505	4968	4906	50)89	4919	5244	5203				
Wild Type	873	824	763	721	(583	619	691	646				
Never Genotyped	72	48	51	51		43	37	42	48				
1-Class	166	170	150	148	1	.45	141	151	144				
2-Class	38	33	27	26		25	27	32	26				
3-Class	13	8	5	7		7	3	8	8				
Total (n=)	5827	5588	5964	5859	59	992	5746	6168	6075				
Indicator 12: AIDS-De	efining Illness	2007	2008	2009	2010	2011	2012	2013	2014				
CD4 < 200 at	Cases	224	194	155	138	115	96	90	84				
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	134	130	110	103	81	67	68	48				
(DTP Reports)	Rate per 100,000	3.1	3.0	2.5	2.3	1.8	1.5	1.5	1.0				
AIDS Cases	Cases	143	143	126	112	95	74	68	_				
(BCCDC Reports)	Rate per 100,000	3.3	3.2	2.8	2.5	2.0	1.6	1.2	_				
Indicator 13: HIV-Rela	ated Mortality	2004	2005	2006	2007	2008	2009	2010	2011				
British Columbia		105	146	142	100	79	63	54	59				
Per 100 HIV+ Populat		1.03	1.40	1.34	0.93	0.72	0.56	0.47	0.50				
Per 100,000 Population	1	2.50	3.43	3.29	2.28	1.80	1.41	1.19	1.29				