

HIV MONITORING QUARTERLY REPORT

FOR BRITISH COLUMBIA

FIRST QUARTER 2014

















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. Motoi Matsukura writes and compiles the monitoring report. Guillaume Colley, Dr. Viviane Lima and Nada Gataric perform analysis of Indicators 5–13. James Nakagawa is responsible for publishing and editing. This report was conceived and guided by Dr. Julio Montaner.



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

Other Data Sources:

The above databases were supplemented with:

- (I) The BC Vital Statistics database which was used to calculate Indicator 5. The HIV Cascade of Care and Indicator 13. HIV-Related Mortality.
- (II) Linkage and preparation of the de-identified individual-level database used for calculating Indicator 5. The HIV Cascade of Care was facilitated by the British Columbia Ministry of Health.
- (III) The Statistics Canada database: BC and HIV-positive population counts were acquired through the statistics Canada website to calculate HIV-specific mortality rates for Indicator 13. HIV-Related Mortality.

Membership of the STOP HIV/AIDS Technical Monitoring Committee-BC-CfE

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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.

NB: HIV screening tests conducted by the VIHA Laboratory are not included.

Indicator 1. HIV Testing Episodes

Figure 1.1 HIV Test Episodes for BC, 2009 Q2-2014 Q1

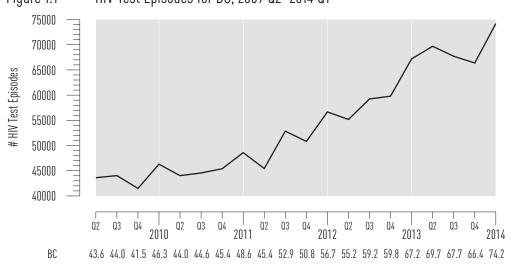


Figure 1.2 HIV Test Episodes by Gender and Prenatal Status for BC, 2009 Q2–2014 Q1 ¹



¹ NB: Testing does not include point of care tests.

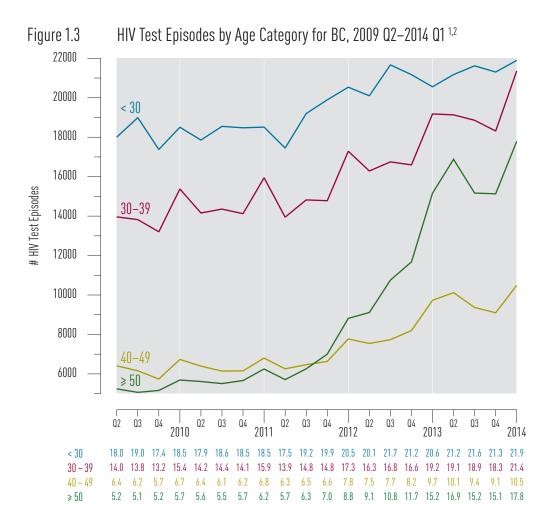


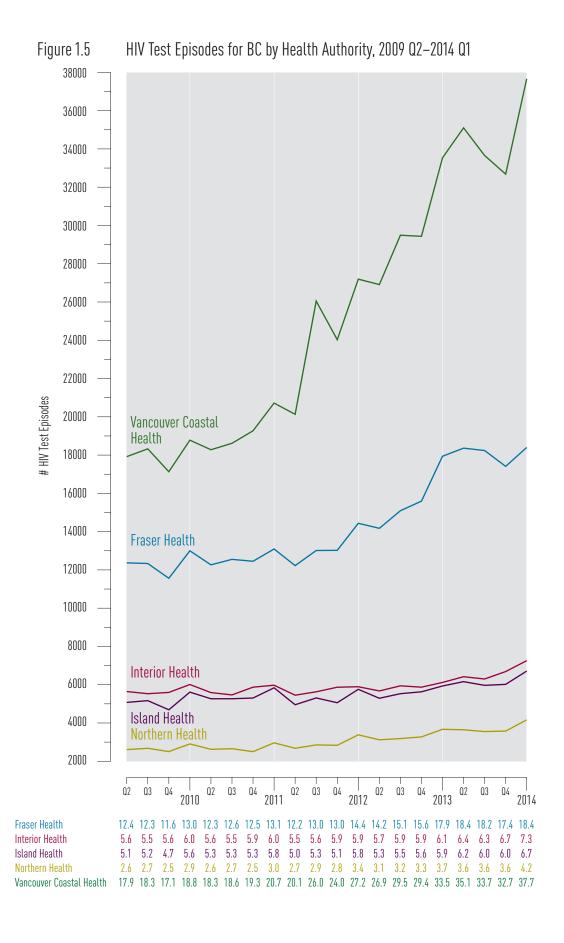
Figure 1.4 Point-of-Care HIV Tests for BC, 2010 Q4-2014 Q1

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2000 & \\
\text{2010} & \text{2000} & \\
\text{2000} & \text{2000} & \\
\text{2011} & \text{20} & \text{2012} & \text{20} & \text{20}

Limitations:

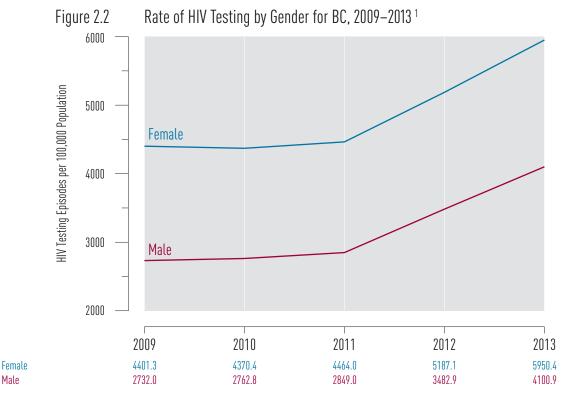
- 1 Repeat tests in individuals who test using various identifiers may not be identified and these individuals may be counted more than once.
- 2 Poc testing data is available from the fourth quarter of 2010 and onwards.

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

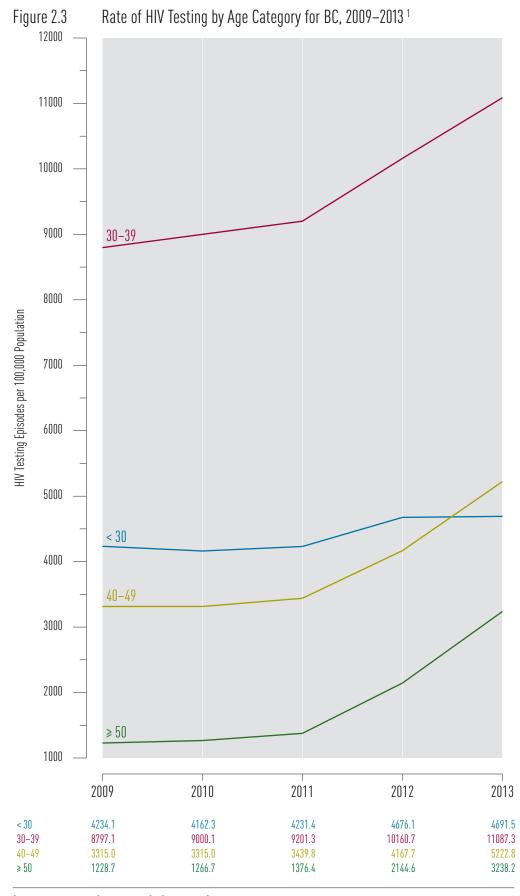


Indicator 2. HIV Testing Rates

Figure 2.1 Rate of HIV Testing for BC and Health Authorities, 2009–2013 ¹ 9000 8000 HIV Testing Episodes per 100,000 Population 7000 6000 Vancouver Coastal Health 5000 Northern Health Fraser Health 4000 All British Columbia Interior Health 3000 Vancouver Island Health 2000 2010 2011 2012 2013 2009 3633.3 3627.9 3714.3 4373.3 5056.0 BC 4262.7 3386.7 3383.1 3438.2 3893.6 Fraser Health Interior Health 2888.9 2925.8 2926.2 2988.8 3265.1 Island Health 2721.3 2703.2 2658.7 2783.9 2902.8 4088.3 4383.5 Northern Health 3515.8 3436.5 3583.1 Vancouver Coastal Health 5115.5 5087.9 5338.2 7023.7 8875.0



Male

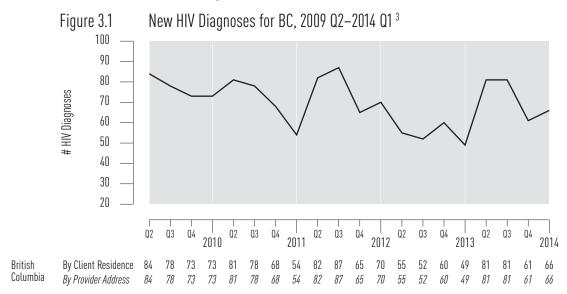


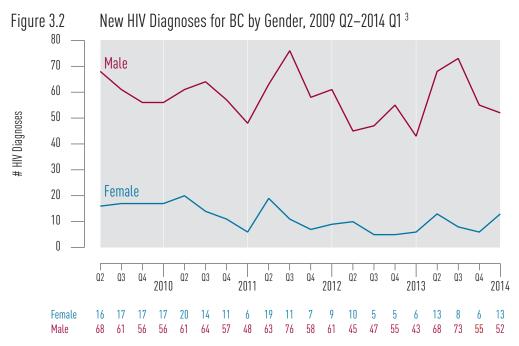
¹ NB: Testing does not include point of care tests.

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

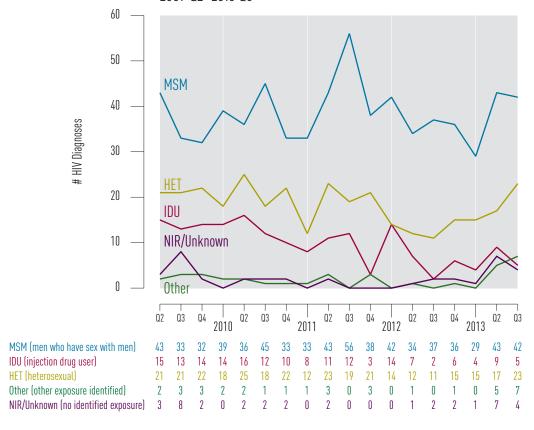




³ Data Source: BCCDC. "By Provider Address" is graphed as dashed line in same colour.

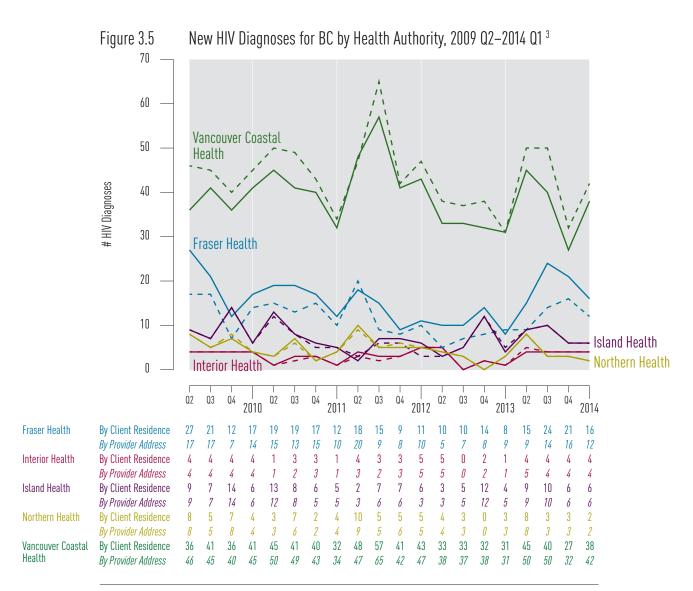
Figure 3.3 New HIV Diagnoses for BC by Age Category, 2009 Q2-2014 Q1 3 # HIV Diagnoses < 30 30–39 ≥ 50 40-49 < 30 years 30-39 years 40-49 years ≥ 50 years

Figure 3.4 New HIV Diagnoses for BC by Exposure Category, 2009 Q2–2013 Q3 3.4



³ 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.



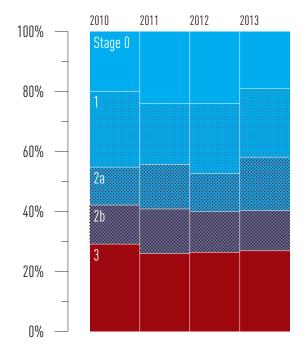
Data Source: BCCDC. "By Provider Address" is graphed as dashed line in same colour.

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.

Figure 4.1 Stage of HIV Infection at Diagnosis for BC, 2010–2013 ⁵

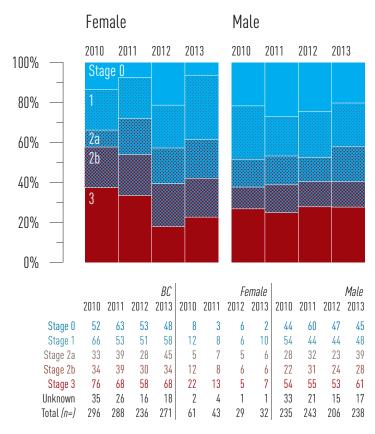


Indicator 4. Stage of HIV Infection at Diagnosis

Table 1 Staging Classifications of Infection at Time of HIV Diagnosis Based on CDC HIV Surveillance Case Definitions

Stage	Criteria																							
0	previous																							
1			CD4 ≥500		N. AIDC																			
2a			CD4 350-499	and	No AIDS case report																			
2b	Stage 0		CD4 200-349		торого																			
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, 2010–2013 ⁵



Data Source: вссьс

Figure 4.3 Stage of HIV Infection at Diagnosis by Age Category for BC, 2010–2013 ⁵

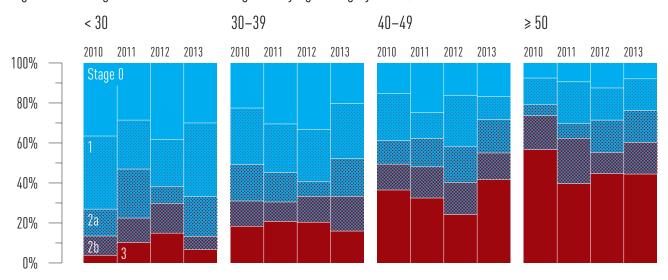
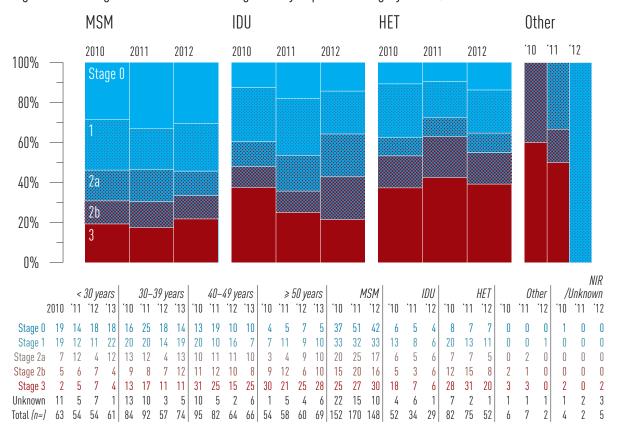


Figure 4.4 Stage of HIV Infection at Diagnosis by Exposure Category for BC, 2010–2012 5.6

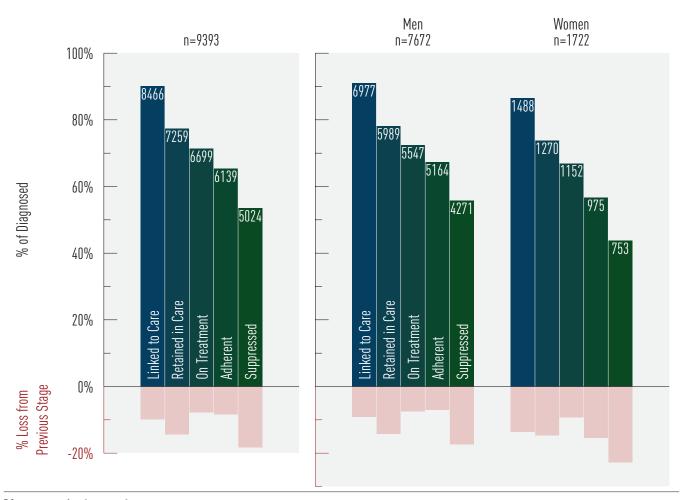


⁶ MSM=men who have sex with men; IDU= injection drug user; HET=heterosexual. NIR=No identified risk/exposure.

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. Linkage to HIV care, 3. Retention in HIV care, 4. On ART and 5. Achieving a suppressed VL; collectively, they are referred to as the cascade of care. Leakage between any of these stages of HIV-care means a reduction in the potential of ART as a benefit to the HIV-positive individual and as an HIV transmission prevention method on a population level. Thus, when interpreting trends in the cascade of care, we strive to see increases along each step of the cascade of care (ie. reduced attrition) with the ultimate goal being 100% within each stage of the cascade. Monitoring the Cascade of Care provides a picture as to where deficiencies lie in the delivery and uptake of HIV-care. In this section we present the cascade of care for the year 2012 in BC overall and stratified by sex and age for each Health Authority.

Figure 5.1 Estimated Cascade of Care for BC, Figure 5.2 Estimated Cascade of Care for BC by Gender, Year Ending 2014 Q1 7 Year Ending 2014 Q1 8



7,8 Data is for the period 2013 Q2-2014 Q1.

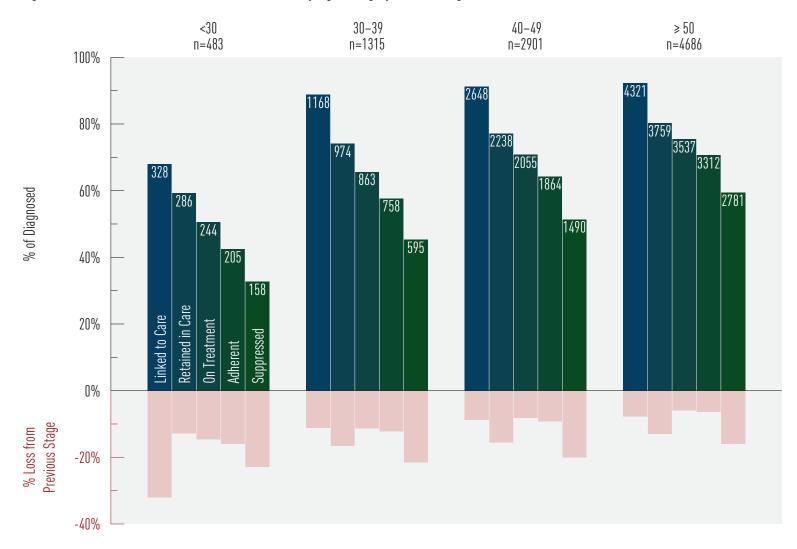
Data Sources:

- 1 British Columbia Centre for Excellence Drug Treatment Program (DTP) Database (ARV use, VL and CD4 count).
- 2 Administrative data (ex. MSP billings; hospitalization data from the Discharge Abstract Database (DAD)).

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

NB: Transgender has been assigned to their biological sex.

Figure 5.3 Estimated Cascade of Care for BC by Age Category, Year Ending 2014 Q1 9

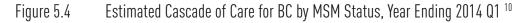


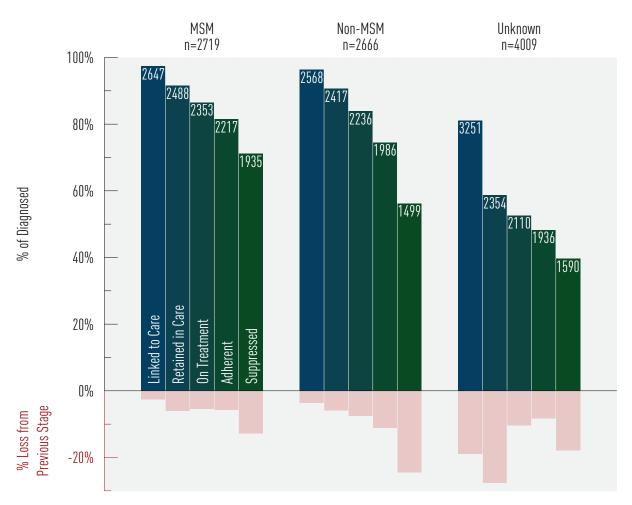
Limitations: на assignment is based on the most recent на of residence of the patient, if not available of the нiv-care provider. If the most recent на of residence is not updated then the designated на may be incorrect.

⁹ Data is for the period 2013 Q2-2014 Q1. Data Sources:

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

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





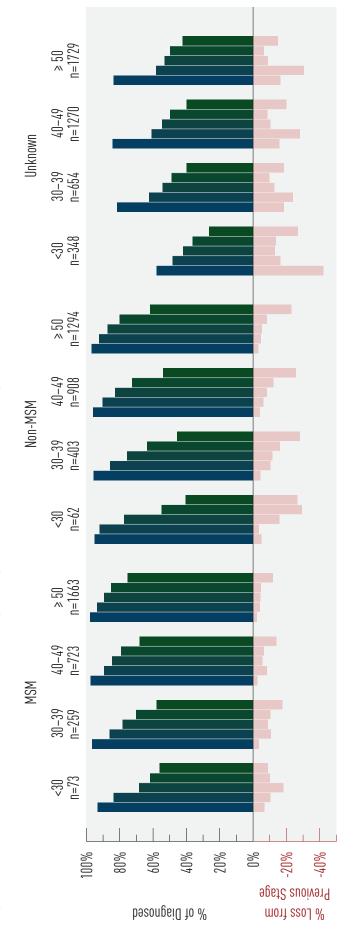
Limitations: HA assignment is based on the most recent HA of residence of the patient, if not available of the HIV-care provider. If the most recent HA of residence is not updated then the designated HA may be incorrect.

¹⁰ Data is for the period 2013 Q2-2014 Q1. Data Sources:

¹ British Columbia Centre for Excellence Drug Treatment Program (DTP) Database (ARV use, VL and CD4 count).

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

Estimated Cascade of Care for BC by Age Category and MSM Status, Year Ending 2014 Q1 ¹¹ Figure 5.5



11 Data is for the period 2013 Q2-2014 Q1.

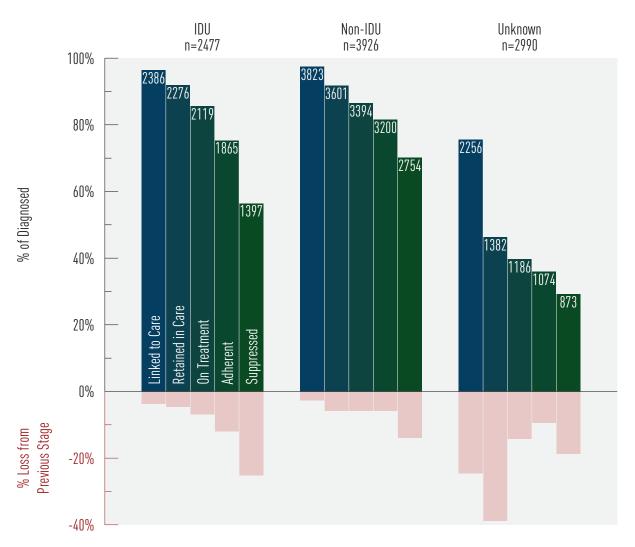
Data Sources:

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

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

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





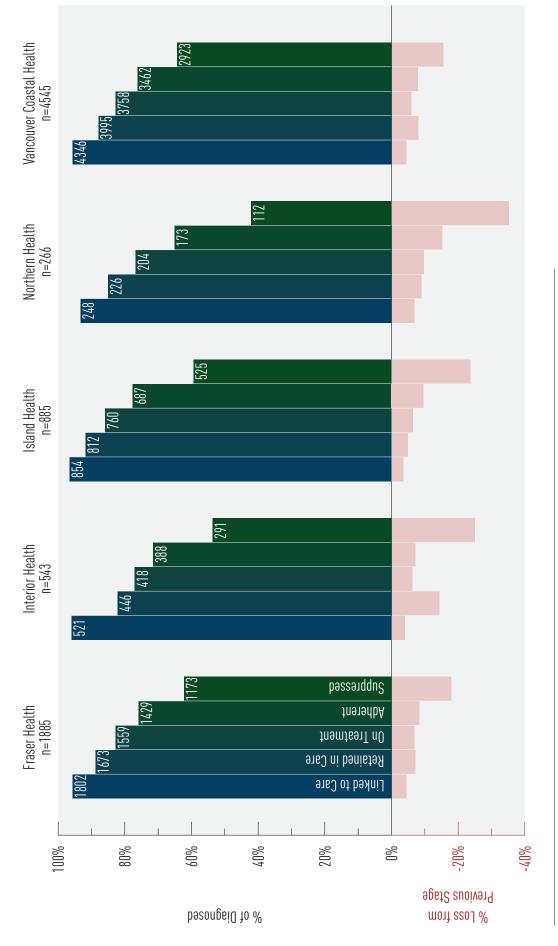
Limitations: на assignment is based on the most recent на of residence of the patient, if not available of the нiv-care provider. If the most recent на of residence is not updated then the designated на may be incorrect.

¹² Data is for the period 2013 Q2-2014 Q1. Data Sources:

¹ British Columbia Centre for Excellence Drug Treatment Program (DTP) Database (ARV use, VL and CD4 count).

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





Data is for the period 2013 Q2–2014 Q1.

Data Sources:

13

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

British Columbia Centre for Excellence Drug Treatment Program (DTP) Database (ARV use, VL and CD4 count). Administrative data (ex. MSP billings; hospitalization data from the Discharge Abstract Database (DAD)).

Indicator 6. The Programmatic Compliance Score (PCS)

The Programmatic Compliance Score (PCS) is a summary measure of risk of future death, immunologic failure and virologic failure from all causes for people who are starting ART for the first time. It is composed of patient- and physician-driven effects. PCS scores range from o−6 with higher scores indicative of poorer health outcomes and greater risk of death. Table 1 provides mortality, immunologic failure and virologic failure probabilities for given PCS scores. We interpret an individual with a PCS≥4 as being 22 times more likely to die, almost 10 times more likely to have immunologic failure and nearly 4 times as likely to demonstrate virologic failure compared to those individuals with a PCS score of o. A detailed description of how the PCS score is calculated and its 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. 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, 2012 Q2-2013 Q4 14 40% Non Viral Suppression at 9 Mo. 30% <3 CD4 Tests Baseline CD4 <200 cells/μL 20% Non-Recommended ART 10% <3 Viral Load Tests No Baseline Genotype 0% Q3 Q4 Q2 Q2 Q3 Q4 2013 2014

461

419

396

366

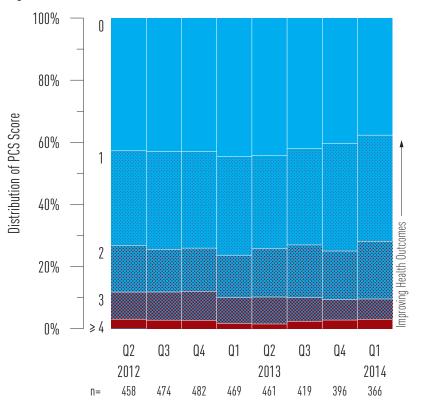
Figure 6.2 Historical Trends for PCS Score for BC, 2012 Q2-2014 Q1 14,15

469

458

474

482



Data Source: British Columbia Centre for Excellence Drug Treatment Program (DTP) Database. Limitations: CD4 cell count capture is approximately 80%. Due to improvements in the automated system, some changes in data representation are expected compared to previous reports.

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 0 is the best score and a score of 4 or more is the worst score.

Antiretroviral Uptake

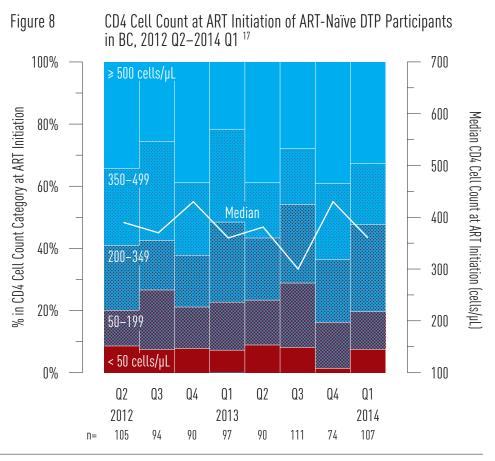
In this section we present trends in ART uptake, the number and proportion of new HIV treatment initiations and the number of active and inactive DTP participants. Trends in ART uptake should be interpreted under the consideration of changing BC HIV treatment guidelines. BC HIV treatment guidelines are 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, 2012 Q2-2014 Q1 16



Indicator 8. CD4 Cell Count at ART Initiation



¹⁶ Data Source: Drug Treatment Program Database
Limitation: DTP participants are designated to an HA based on most current residence provided by the participant.

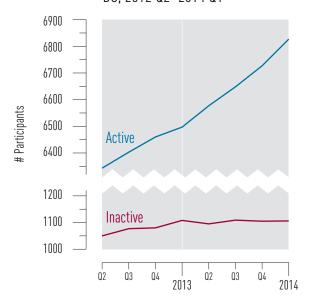
¹⁷ Data Source: Drug Treatment Program Database Limitations: CD4 cell count data is approximately 80% complete.

Indicator 9. Active and Inactive DTP Participants

Table 3. Distribution of People on ART for BC, 2014 Q1 18

		Fraser	Interior	Island	Northern	Vancouver Coastal	Total BC
Age	< 30	64	14	26	15	138	257
	30-39	264	57	92	41	516	970
	40-49	534	115	226	67	1258	2200
	≥ 50	725	243	428	89	1915	3400
Gender	Male	1230	327	624	124	3363	5668
	Female	357	102	148	88	464	1159
Exposure	MSM	457	109	181	23	1601	2371
	IDU	443	145	274	126	1144	2132
Total		1587	429	772	212	3827	6827

Figure 9 Active and Inactive DTP Participants in BC, 2012 Q2-2014 Q1 19



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

Definitions:

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

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

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

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

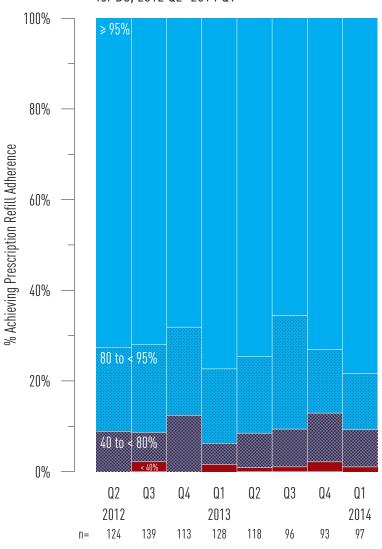
¹⁸ Data Source: Drug Treatment Program Database

Antiretroviral Adherence Level

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

Indicator 10. Antiretroviral Adherence

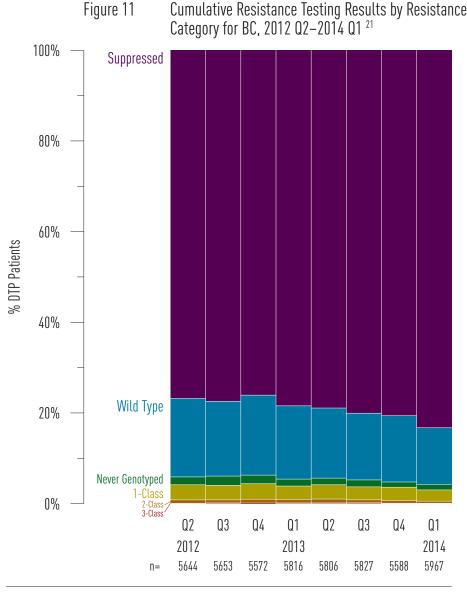
Figure 10 Distribution of Individuals by Adherence Level in 1st Year of Therapy, Based on Pharmacy Refill Compliance for BC, 2012 Q2–2014 Q1 ²⁰



²⁰ 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.

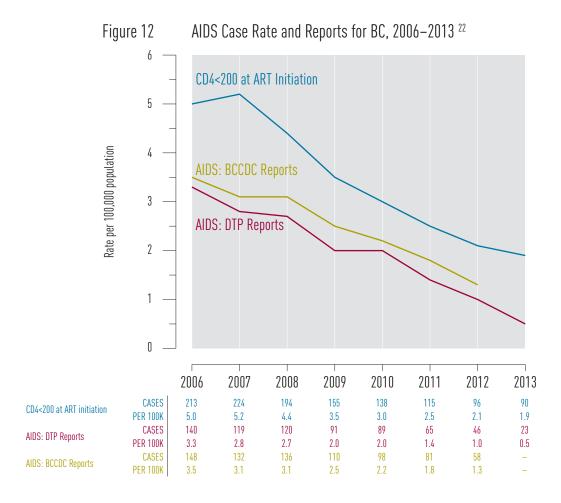


²¹ Data Source: Drug Treatment Program Database

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.

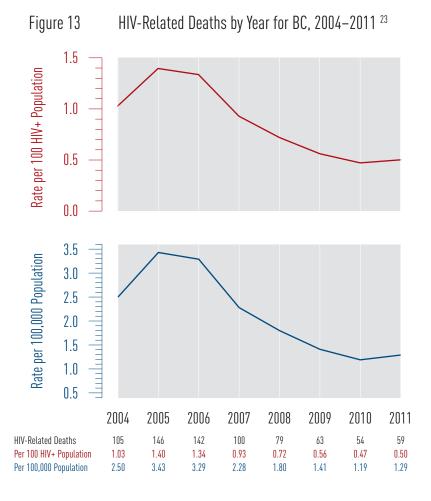


Data Source: DTP AIDS cases are obtained from the Drug Treatment Program Database; BCCDC AIDS cases are obtained from the BCCDC; CD4<200 at ART initiation data came from the DTP database.

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:

²³ 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		2009	9		2010)			201	l			2012	2			2013	3			2014
Episodes ((thousands)	Q2	Q3	Q4	Q1																
British Co	lumbia	43.6	44.0	41.5	46.3	44.0	44.6	45.4	48.6	45.4	52.9	50.8	56.7	55.2	59.2	59.8	67.2	69.7	67.7	66.4	74.2
Gender	Female	26.4	26.6	25.5	28.0	26.3	27.0	26.9	28.7	26.2	28.0	29.0	32.4	31.6	33.8	33.9	38.1	39.2	38.3	37.5	41.4
	Male	16.5	16.7	15.4	17.6	17.1	16.9	17.0	18.2	16.6	18.2	18.8	21.6	21.1	22.8	23.4	26.2	27.8	26.4	26.1	29.8
	Other	0.7	0.7	0.6	0.7	0.7	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
Female (Pr	renatal)	11.2	11.4	11.3	12.1	11.0	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
Female (N	on-prenatal)	15.2	15.2	14.1	15.9	15.2	15.3	14.9	15.8	14.9	16.2	17.1	19.4	19.7	21.8	22.1	25.7	27.5	26.3	25.4	28.5
Age	< 30	18.0	19.0	17.4	18.5	17.9	18.6	18.5	18.5	17.5	19.2	19.9	20.5	20.1	21.7	21.2	20.6	21.2	21.6	21.3	21.9
	30-39	14.0	13.8	13.2	15.4	14.2	14.4	14.1	15.9	13.9	14.8	14.8	17.3	16.3	16.8	16.6	19.2	19.1	18.9	18.3	21.4
	40-49	6.4	6.2	5.7	6.7	6.4	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
	≥ 50	5.2	5.1	5.2	5.7	5.6	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	15.1	17.8
POC HIV	Tests							1.0	1.1	2.1	6.1	2.5	2.3	2.1	2.3	2.2	2.6	2.4	2.7	2.5	2.7
Fraser Hea	alth	12.4	12.3	11.6	13.0	12.3	12.6	12.5	13.1	12.2	13.0	13.0	14.4	14.2	15.1	15.6	17.9	18.4	18.2	17.4	18.4
Interior H	ealth	5.6	5.5	5.6	6.0	5.6	5.5	5.9	6.0	5.5	5.6	5.9	5.9	5.7	5.9	5.9	6.1	6.4	6.3	6.7	7.3
Island Hea	ılth	5.1	5.2	4.7	5.6	5.3	5.3	5.3	5.8	5.0	5.3	5.1	5.8	5.3	5.5	5.6	5.9	6.2	6.0	6.0	6.7
Northern 1	Health	2.6	2.7	2.5	2.9	2.6	2.7	2.5	3.0	2.7	2.9	2.8	3.4	3.1	3.2	3.3	3.7	3.6	3.6	3.6	4.2
Vancouver	Coastal Health	17.9	18.3	17.1	18.8	18.3	18.6	19.3	20.7	20.1	26.0	24.0	27.2	26.9	29.5	29.4	33.5	35.1	33.7	32.7	37.7

Indicator 2: Rate of HIV Testing per 100,000

		2009	2010	2011	2012	2013
British Colu	ımbia	3633.3	3627.9	3714.3	4373.3	5056.0
Fraser Heal	th	3386.7	3383.1	3438.2	3893.6	4262.7
Interior Hea	alth	2888.9	2925.8	2926.2	2988.8	3265.1
Island Heal	th	2721.3	2703.2	2658.7	2783.9	2902.8
Northern H	Iealth	3515.8	3436.5	3583.1	4088.3	4383.5
Vancouver	Coastal Health	5115.5	5087.9	5338.2	7023.7	8875.0
Gender	Female	4401.3	4370.4	4464.0	5187.1	5950.4
	Male	2732.0	2762.8	2849.0	3482.9	4100.9
Age	< 30	4234.1	4162.3	4231.4	4676.1	4691.5
	30-39	8797.1	9000.1	9201.3	10160.7	11087.3
	40-49	3315.0	3315.0	3439.8	4167.7	5222.8
	≥ 50	1228.7	1266.7	1376.4	2144.6	3238.2

		2009			2010	1			2011				2012				2013			2	2014
Indicator 3: New HIV	Diagnoses	Q2	Q3	Q4	Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4	Q1
British Columbia	By Client Residence	84	78	73	73	81	78	68	54	82	87	65	70	55	52	60	49	81	81	61	66
	By Provider Address	84	78	73	73	81	78	68	54	82	87	65	70	55	52	60	49	81	81	61	66
Gender	Female	16	17	17	17	20	14	11	6	19	11	7	9	10	5	5	6	13	8	6	13
	Male	68	61	56	56	61	64	57	48	63	76	58	61	45	47	55	43	68	73	55	52
Age	< 30	16	16	24	17	20	22	9	5	18	17	18	18	14	8	18	9	18	23	15	18
	30-39	27	17	20	25	18	23	20	18	30	30	13	16	17	11	10	16	25	18	13	17
	40-49	25	30	21	23	29	19	24	18	22	22	19	20	11	19	19	12	14	21	21	15
	≥ 50	16	15	8	8	14	14	15	13	12	18	15	16	13	14	13	12	24	19	12	16
Exposure	MSM	43	33	32	39	36	45	33	33	43	56	38	42	34	37	36	29	43	42	-	_
	IDU	15	13	14	14	16	12	10	8	11	12	3	14	7	2	6	4	9	5	-	_
	HET	21	21	22	18	25	18	22	12	23	19	21	14	12	11	15	15	17	23	-	_
	Other	2	3	3	2	2	1	1	1	3	0	3	0	1	0	1	0	5	7	-	_
	NIR/Unknown	3	8	2	0	2	2	2	0	2	0	0	0	1	2	2	1	7	4	-	_
Fraser Health	By Client Residence	27	21	12	17	19	19	17	12	18	15	9	11	10	10	14	8	15	24	21	16
	By Provider Address	17	17	7	14	15	13	15	10	20	9	8	10	5	7	8	9	9	14	16	12
Interior Health	By Client Residence	4	4	4	4	1	3	3	1	4	3	3	5	5	0	2	1	4	4	4	4
	By Provider Address	4	4	4	4	1	2	3	1	3	2	3	5	5	0	2	1	5	4	4	4
Island Health	By Client Residence	9	7	14	6	13	8	6	5	2	7	7	6	3	5	12	4	9	10	6	6
	By Provider Address	9	7	14	6	12	8	5	5	3	6	6	3	3	5	12	5	9	10	6	6
Northern Health	By Client Residence	8	5	7	4	3	7	2	4	10	5	5	5	4	3	0	3	8	3	3	2
	By Provider Address	8	5	8	4	3	6	2	4	9	5	6	5	4	3	0	3	8	3	3	2
Vancouver Coastal	By Client Residence	36	41	36	41	45	41	40	32	48	57	41	43	33	33	32	31	45	40	27	38
Health	By Provider Address	46	45	40	45	50	49	43	34	47	65	42	47	38	37	38	31	50	50	32	42

Indicator 4: Stage of HIV Infection at Baseline

indicator 4: St	tage (iectio	n at B	aseme		ı											1				
	'10	В '11	C '12	'13		Female '11 '12	'13	'10	Ma '11	ile '12	'13		30 y '11	ears '12	' 13	30 10)–39 [·]	years '12	' 13	40 10)–49 '11	years '12	' 13
Stage 0	52	63	53	48	8	3 6	2	44	60	47	45	19	14	18	18	16	25	18	14	13	19	10	10
Stage 1	66	53	51	58	12	8 6	10	54	44	44	48	19	12	11	22	20	20	14	19	20	10	16	7
Stage 2a	33	39	28	45	5	7 5	6	28	32	23	39	7	12	4	12	13	12	4	13	10	11	11	10
Stage 2b	34	39	30	34	12	8 6	6	22	31	24	28	5	6	7	4	9	8	7	12	11	12	10	8
Stage 3	76	68	58	68	22	13 5	7	54	55	53	61	2	5	7	4	13	17	11	11	31	25	15	25
Unknown	35	26	16	18	2	4 1	1	33	21	15	17	11	5	7	1	13	10	3	5	10	5	2	6
Total	296	288	236	271	61	43 29	32	235	243	206	238	63	54	54	61	84	92	57	74	95	82	64	66
	'10	≥ 50 '11	years '12	'13	2010	MSM 2011	2012	2010	ID 0 20		2012	He 2010	teros 201	exua 11 2	012	Oth 2010		posui 11 2	re 012	NII 2010	R/Unl 201		n 012
Stage 0	4	5	7	5	37	51	42	(6	5	4	8		7	7	C)	0	0	1		0	0
Stage 1	7	11	9	10	33	32	33	13	3	8	6	20	1	13	11	C)	0	1	0		0	0
Stage 2a	3	4	9	10	20	25	17		6	5	6	7		7	5	C)	2	0	0		0	0
Stage 2b	9	12	6	10	15	20	16		5	3	6	12	1	15	8	2	2	1	0	0		0	0
Stage 3	30	21	25	28	25	27	30	18	8	7	6	28	3	31	20	3	3	3	0	2		0	2
Unknown	1	5	4	6	22	15	10	4	4	6	1	7		2	1	1		1	1	1		2	3
Total	54	58	60	69	152	170	148	52	2	34	29	82	7	75	52	6	Ď	7	2	4		2	5
Indicator 5: H	HIV (Casca	ide of	f Car	e	DIAG	NOSEI)	L	INKE	D	RETA	AINE	D		ON A	RT	AD	HERE	ENT	SUP	PRES	SED
British Colun	nbia						939	3		846	6		725	9		669	99		6	139		5	024
Age Category	< .	30					48.	3		32	8		28	6		2	44		2	205			158
	30	-39					131	5		116	8		97	4		80	63		7	758			595
	40	-49					290			264	8		223	8		20.			18	864		1	490
	≥.						468			432			375			35.			33	312		2	781
Age Category	M	SM			30		7.				8		6	1			50			45			41
and MSM Status					-39		259			25			22				03			182			150
otatus					-49		72.			70			64				12			572			493
	N.T.		.03.4	≥ .			1663			162			155			148			14	418		1:	251
	No	on-M	SM		30		62				9		5				48			34			25
					1–39		403			38			34				05			256			184
					1–49 50		908			87			81				52			560			491
	T T.	alem o			50		129			125			119			11:				036			799
	UI	nkno	wn		30		348			20			16				46 			126			92
)–39)–49		654 1270			53 107			40 77				55 91			320 532			261506
					50		1729			144			100				18			858			731
Gender	М	ale		_	30		767			697			598			554				164			271
Gender		male					172			148			127			11.				975			753
Injection	ID						247			238			227			21				865			397
Drug Use		on-II	Ν				3920			382			360			339				200			754
		nkno					2990			225			138			118				074			873
MSM Status		SM					2719			264			248			23.				217			935
		on-M	SM				266			256			241			22				986			499
		nkno					4009			325			235			21				936			590
Health			Health	1			188			180			167			15				429			173
Authority			Heal				54.			52			44				18			388			291
			Health				88			85			81				60			587			525
			n He				260			24			22				04			173			112
					l Healt	h	454			434	:6		399			37.			34	462		2	923

Indicator 6: Programmatic	-	(PCS)			2012					201
	2012 Q2	Q3	Q4		2013 Q1	Q2	Q3		Q4	201 ⁴ Q1
< 3 CD4 Tests	17.9%	18.1%	19.3%		17.5%	18.4%	17.9%		18.4%	22.1%
< 3 Viral Load Tests	9.2%	9.1%	10.0%		7.7%	8.7%	9.3%		8.6%	8.7%
No Baseline Genotype	6.3%	5.7%	6.0%		6.4%	5.6%	5.5%		5.6%	4.4%
Baseline CD4 < 200 cells/µL		24.3%	22.2%		21.3%	20.6%	21.2%		20.5%	21.9%
•								•		
Non-Recommended ART	3.5%	3.4%	2.9%		2.3%	2.8%	6.2%		9.3%	10.7%
Non Viral suppression at 9 l		36.7%	37.8%		36.0%	37.5%	37.7%		35.1%	35.8%
PCS Score: 0	195	204	207		209	204	176		160	138
PCS Score: 1	140	149	150		149	138	130		137	125
PCS Score: 2	69	65	67		64	72	71		62	68
PCS Score: 3	40	43	45		39	40	32		26	24
PCS Score: 4 or more	14	13	13		8	7	10		11	11
Total (n=)	458	474	482		469	461	419		396	366
Indicator 7: New DTP ARV	Participants									
First Starts	105	94	92		97	92	112		75	108
Experienced Starts	106	89	110		91	111	114		130	117
Indicator 8: CD4 Cell Coun	nt at ART Initiation	for ARV-1	Naïve DTD	Partic	inants					
CD4 ≥ 500	36	24	35	_ u1 t1C	21	35	31		29	35
CD4 2 300 CD4 350-499	26	30	21		29	16	20		18	21
CD4 200-349	22	15	15		25	18	28		15	30
CD4 50–199	12	18	13		15	13	23		11	13
			7							
CD4 < 50	9	7			7	8	9		1	8
CD4 Median (cells/μL)	390	370	430		360	381	300		430	360
Total (n=)	105	94	90		97	90	111		74	107
Indicator 9: Active and Inac	ctive DTP Participa	nts								
Active DTP Participants	6343	6403	6460		6497	6577	6648		6727	6827
Inactive DTP Participants	1050	1077	1080		1108	1095	1109		1105	1106
Indicator 10: Antiretrovira l	l Adherence									
≥ 95%	90	100	77		99	88	63		68	76
80% to < 95%	23	27	22		21	20	24		13	12
40% to < 80%	11	9	14		6	9	8		10	8
< 40%	0	3	0		2	1	1		2	1
Total (n=)	124	139	113		128	118	96		93	97
Indicator 11: Resistance Tes	sting and Results									
Suppressed	4335	4377	4239		4562	4582	4665		4505	4969
Wild Type	977	933	983		940	898	858		817	745
Never Genotyped	98	115	105		89	85	89		64	73
1-Class	188	182	193		173	183	167		164	148
2-Class	43	37	45		43	49	38		33	28
3-Class	3	9	7		9	9	10		5	4
Total (n=)	5644	5653	5572		5816	5806	5827		5588	5967
Indicator 12: AIDS-Definin			2006	2007	2008	2009	2010	2011	2012	2013
	Cases		213	224	194	155	138	115	96	90
	Rate per 100,000		5.0	5.2	4.4	3.5	3.0	2.5	2.1	1.9
	Cases		140	119	120	91	89	65	46	23
	Rate per 100,000		3.3	2.8	2.7	2.0	2.0	1.4	1.0	0.5
	Cases		148	132	136	110	98	81	58	-
(BCCDC Reports)	Rate per 100,000		3.5	3.1	3.1	2.5	2.2	1.8	1.3	-
Indicator 13: HIV-Related 1	Mortality 2004	2005	2006	2007	2008	2009	2010	2011		
British Columbia	105	146	142	100	79	63	54	59		
Per 100 HIV+ Population	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		
1 cr 100,000 ropulation	2.30	5.15	3.27	2.20	1.00	1,71	1.17	1.27		