

HIV MONITORING QUARTERLY REPORT

FOR BRITISH COLUMBIA

THIRD QUARTER 2013

UPDATED VERSION: NOV 28, 2014 *

* See foreword

















Foreword

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

The objectives of these reports are to:

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

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

^{*} Please note that for Q2 and Q3 2013 reports, a coding revision resulted in data display errors in Indicator 5, the Cascade of Care (in Figures 5.1–5.7 on pp. 22–27 in this report), which has been updated; and, only for Q3 2013 reports, Appendix Table for Indicator 5 (on p. 37 in this report). All other figures and reports remain accurate. Please discard any previous reports and use this updated version. If you have any questions, please contact Irene Day at iday@cfenet.ubc.ca.

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Recommended Highly Active Antiretroviral Therapy (HAART)

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Suppression at 9 Months

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



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

Other Data Sources:

The above databases were supplemented with:

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

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

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The Seek and Treat for Optimal Prevention (STOP) HIV/AIDS BC Provincial Program: A Note on Monitoring and Interpreting HIV Indicators

The Seek and Treat for Optimal Prevention (STOP) of HIV/AIDS programme is a provincial initiative to improve HIV diagnosis and care delivery in BC through increased HIV-specific funding to all HSDA's across BC. The STOP provincial programme is an expansion of a four-year STOP pilot project which was implemented in two Health Service Delivery Areas in March 2010; the Vancouver HSDA which bears the largest burden of the HIV epidemic in the province and the Northern Interior HSDA which bears a high burden of HIV-related mortality. The STOP pilot project demonstrated the urgent need for improved efforts in early diagnosis of HIV and timely initiation of highly active antiretroviral therapy (HAART) initiation.

The expansion to a province-wide programme was announced on November 30th 2012 by the BC Ministry of Health with roll out of funding beginning on April 1st, 2013. This funding is intended to be used in the implementation and evaluation of HIV-related diagnosis and care initiatives within individual HA's. Goals of the project include: 1. A reduction in the number of new HIV infections in BC; 2. Improvements in the quality, effectiveness, and reach of HIV prevention services; 3. An increase in early diagnosis of HIV; 4. A reduction in AIDS cases and HIV-related mortality.

The goals of HA-led STOP-funded initiatives are to work toward achieving these goals. To these ends some outcome measures or indicators of progress have been drafted that should be considered in the design and implementation phases of these initiatives.

HIV Testing Episodes and Rates

In this section, the number of HIV test episodes and point of care (POC) HIV tests conducted each quarter in BC is shown. In general terms the goal is to increase the number of tests performed and to maximize testing efficiency. Test episodes are allocated by region according to where the test is performed.

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

Indicator 1. HIV Testing Episodes

Figure 1.1 HIV Test Episodes in BC, 2009 Q1-2013 Q3

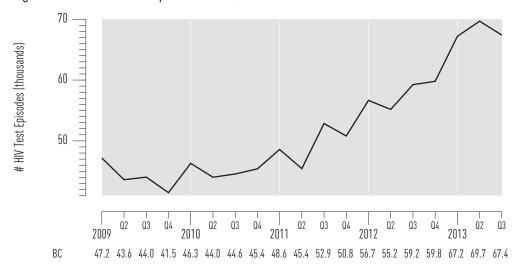
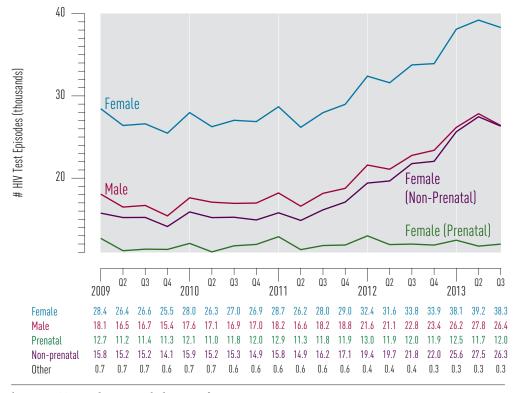
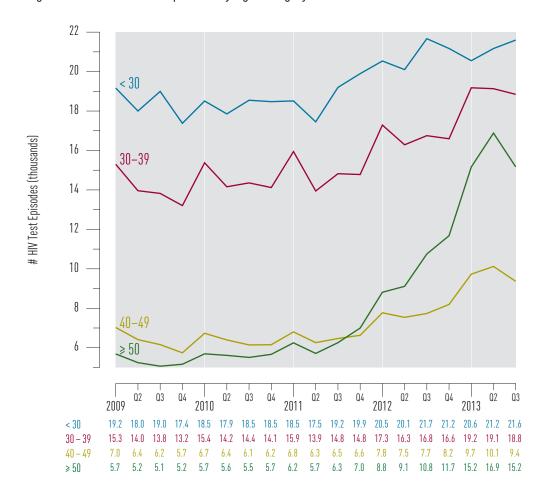


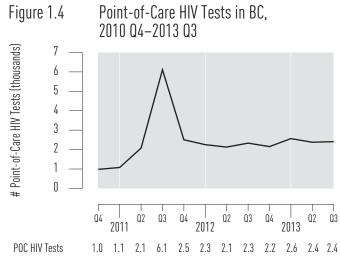
Figure 1.2 HIV Test Episodes by Gender and Prenatal Status in BC, 2009 Q1–2013 Q3 ¹



NB: Testing does not include point of care tests.

Figure 1.3 HIV Test Episodes by Age Category for BC, 2009 Q1–2013 Q3 1.2



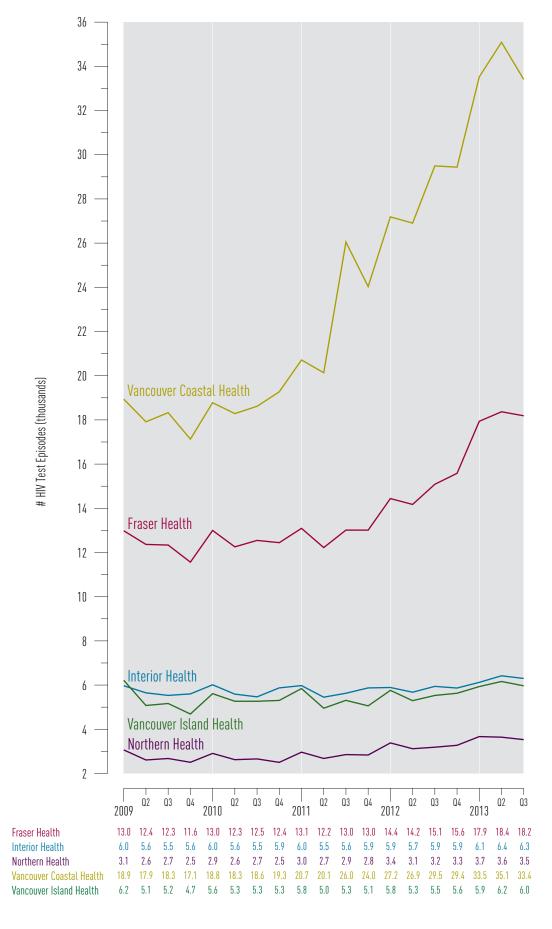


Data Source: The BC Public Health Microbiology and Reference Laboratory (BCPHMRL) courtesy of the BC Centre for Disease Control (BCCDC).

Limitations:

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

Figure 1.5 HIV Test Episodes by Health Authority in BC, 2009 Q1–2013 Q3



Indicator 2. HIV Testing Rates

Figure 2.1 Rate of HIV Testing in BC and Health Authorities, 2009–2012 ¹

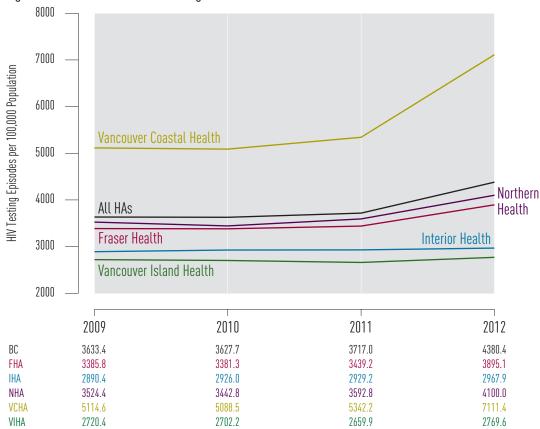


Figure 2.2 Rate of HIV Testing by Gender in BC, 2009–2012 ¹

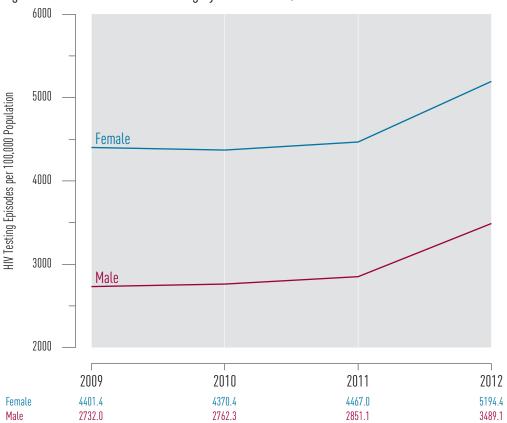
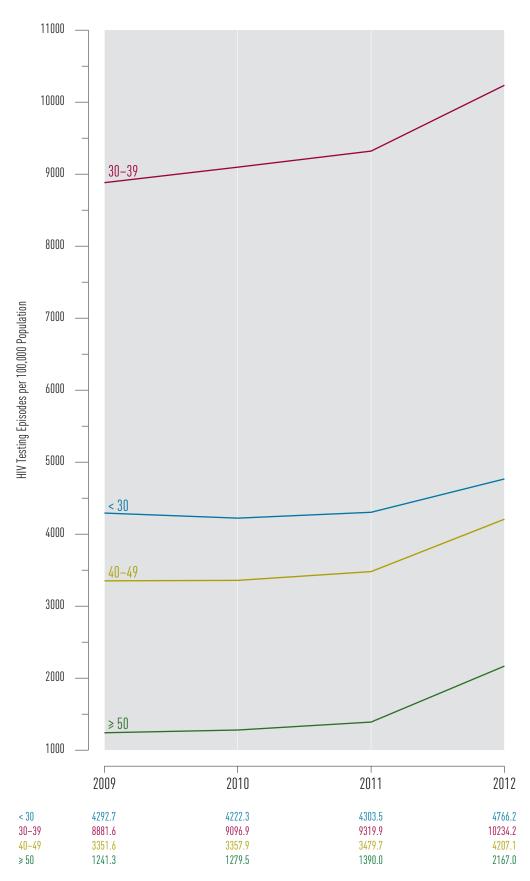


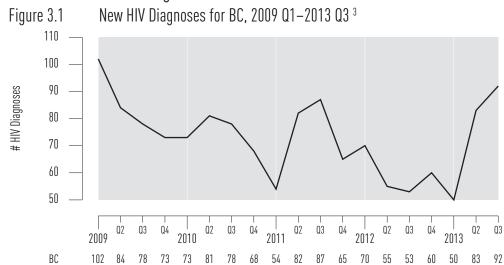
Figure 2.3 Rate of HIV Testing by Age Category in BC, 2009–2012 ¹

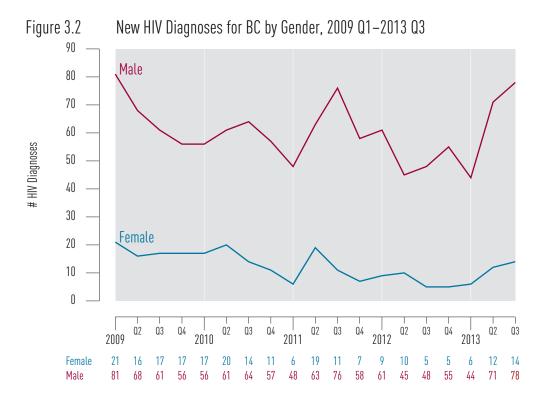


New HIV Diagnoses

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



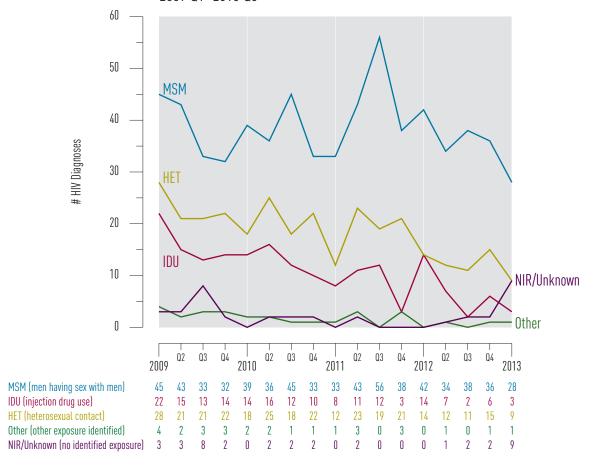


³ Data Source: BCCDC

Figure 3.3 New HIV Diagnoses for BC by Age Category, 2009 Q1–2013 Q3



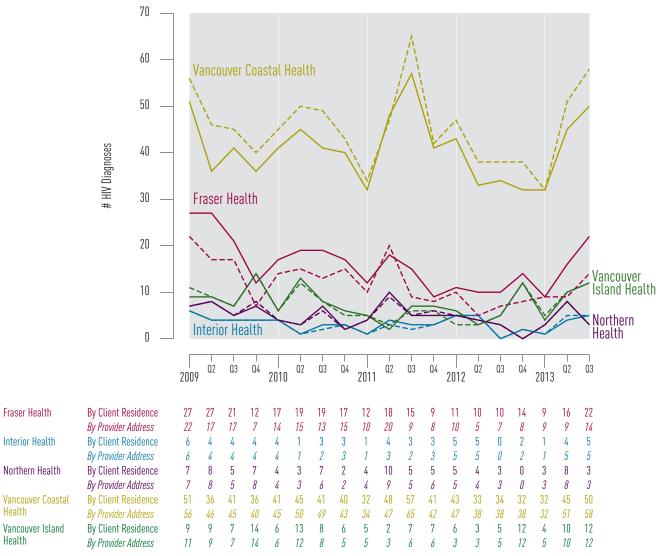
Figure 3.4 New HIV Diagnoses for BC by Exposure Category, 2009 Q1–2013 Q3 4



⁴ BCCDC: Data lags by 6 months.

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

Figure 3.5 New HIV Diagnoses for BC by Health Authority, 2009 Q1-2013 Q3



"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	ńegativ	ria met for acute ve or indetermina Ifirmed positive H	te HIV	test within 180		
1			CD4 ≥500		N AIDO		
2a			CD4 350-499	and	No AIDS case report		
2b	Stage 0		CD4 200-349		τομοιτ		
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, 2010–2012 ⁵

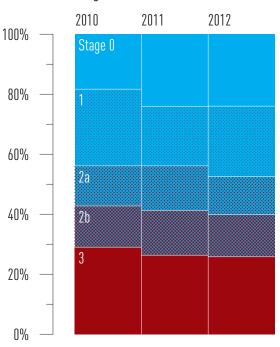
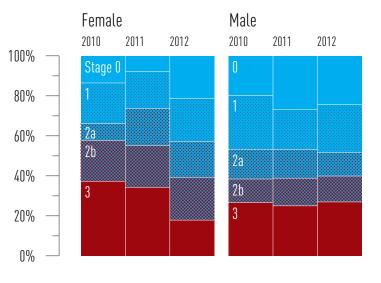


Figure 4.2 Stage of HIV Infection at Diagnosis by Gender for BC, 2010–2012 ⁵



			BL		re	emale			маιе
	2010	2011	2012	2010	2011	2012	2010	2011	2012
Stage 0	48	63	53	8	3	6	40	60	47
Stage 1	67	52	52	12	7	6	55	45	46
Stage 2a	35	39	28	5	7	5	30	32	23
Stage 2b	36	39	31	12	8	6	24	31	25
Stage 3	76	69	57	22	13	5	54	56	52
Unknown	38	26	17	3	5	1	35	21	16
Total (n=)	300	288	238	62	43	29	238	245	209

Famala

Mala

nc

Data Source: вссьс

Figure 4.3 Stage of HIV Infection at Diagnosis by Age Category for BC, 2010–2012 ⁵

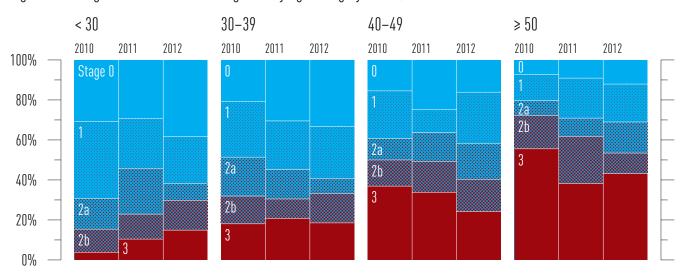
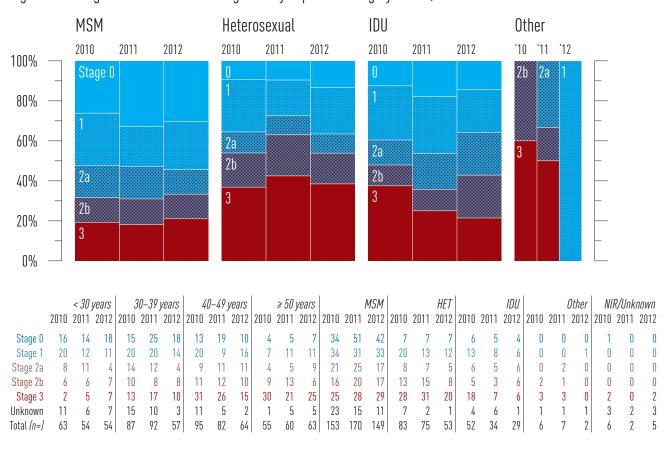


Figure 4.4 Stage of HIV Infection at Diagnosis by Exposure Category for BC, 2010–2012 5,6

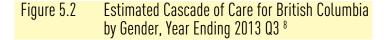


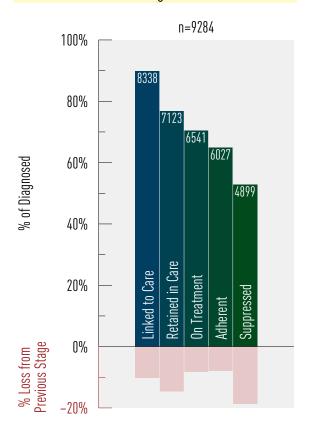
⁶ MSM=men who have sex with men; IDU= injection drug user; HET=heterosexual. NIR=No identified risk/exposure.

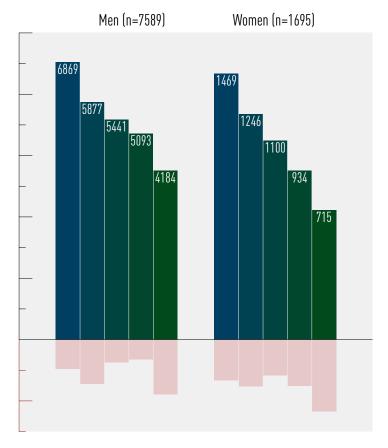
Indicator 5. HIV Cascade of Care

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, Year Ending 2013 Q3 7







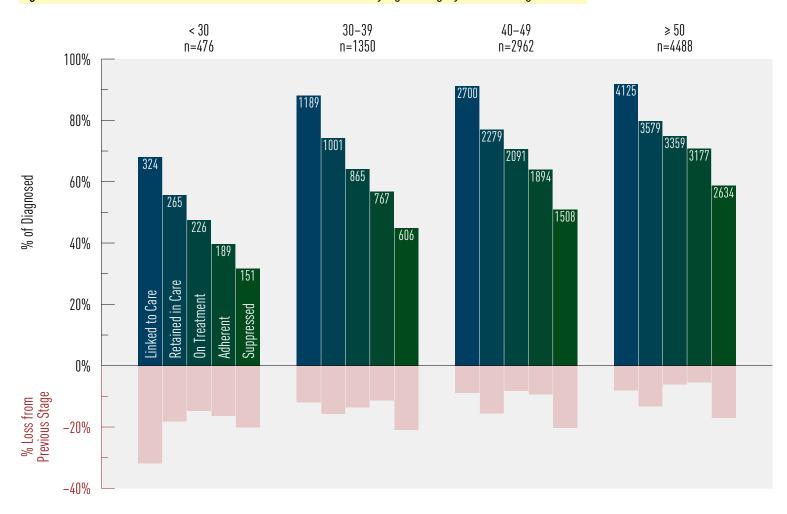
NB: Transgender has been assigned to their biological sex.

^{7,8} 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)).

Figure 5.3 Estimated Cascade of Care for British Columbia by Age Category, Year Ending 2013 Q3 9

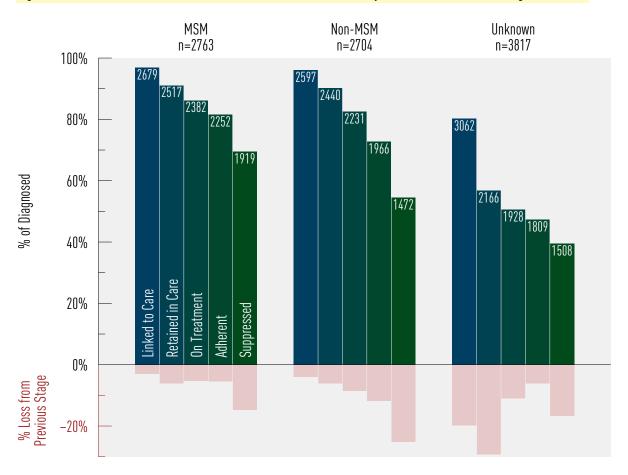


⁹ Data Sources:

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)).



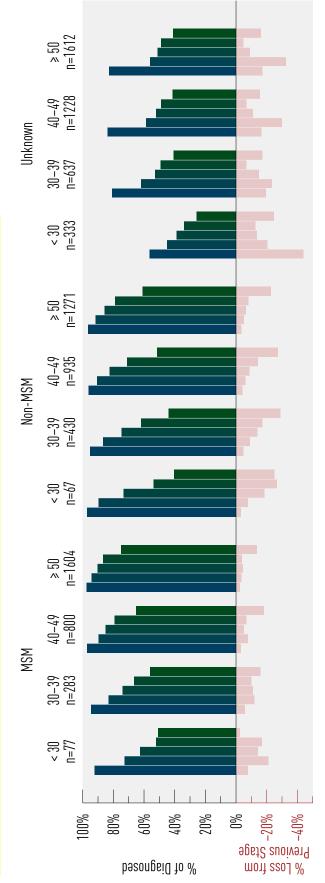


¹⁰ Data Sources:

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 British Columbia by Age Category and MSM Status, Year Ending 2013 Q3 ¹¹ Figure 5.5

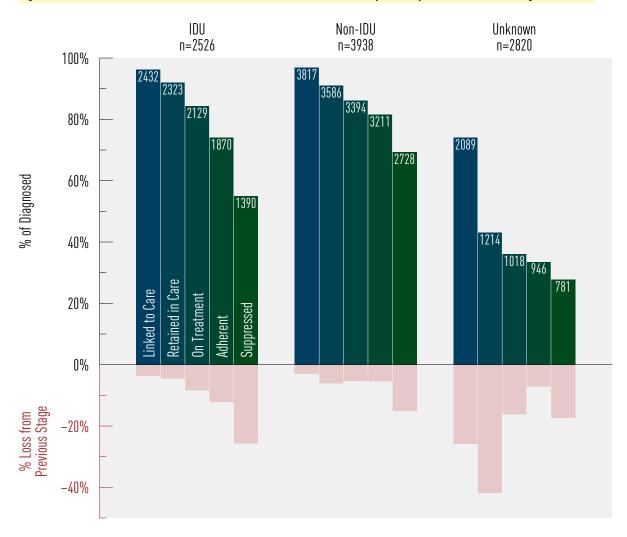


11 Data Sources:

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)).



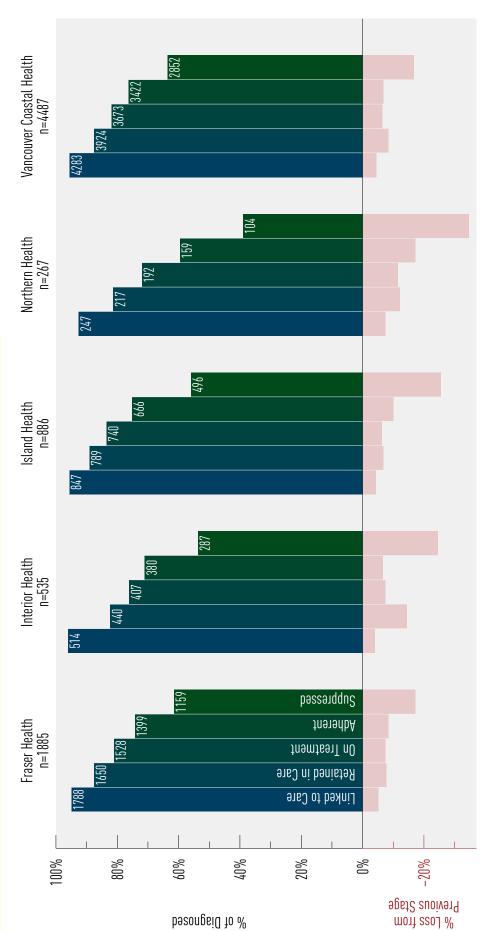


¹² Data Sources:

¹ 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)).

Estimated Cascade of Care for British Columbia by Health Authority, Year Ending 2013 Q3 13 Figure 5.7



13 Data Sources:

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

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

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

Indicator 6. The Programmatic Compliance Score (PCS)

The Programmatic Compliance Score (PCS) is a summary measure of risk of future death, immunologic failure and virologic failure from all causes for people who are starting ART for the first time. It is composed of patient- and physician-driven effects. PCs scores range from o−6 with higher scores indicative of poorer health outcomes and greater risk of death. Table 2 provides mortality, immunologic failure and virologic failure probabilities for given PCs scores. We interpret an individual with a PCs≥4 as being 22 times more likely to die, almost 10 times more likely to have immunologic failure and nearly 4 times as likely to demonstrate virologic failure compared to those individuals with a PCs score of o. A detailed description of how the PCs score is calculated and its valida¬tion can be found in the technical report. In short, PCs scores are calculated by summing the results (yes=1, no=0) of six un-weighted non-performance indicators based on IAS−USA treatment guidelines:

- 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

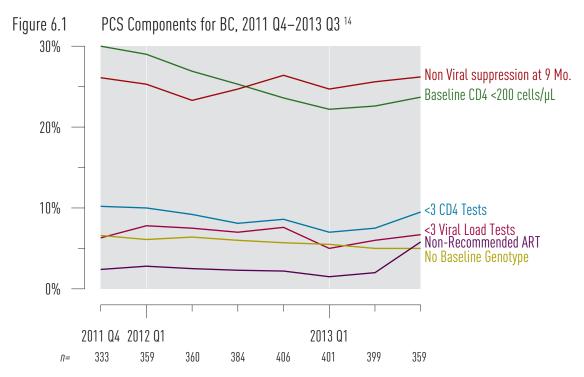


Figure 6.2 Historical Trends for PCS Score for BC, 2011 Q4-2013 Q3 15 100% 80% Distribution of PCS Score 60% 40% Improving Health Outcomes 20% 2 0% Q1 Q4 Q2Q3 Q4 Q1 Q2Q3 2012 2011 2013 333 359 406 401 399 360 384 359 NB: A score of o is the best score and a score of 4 or more is the worst score.

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

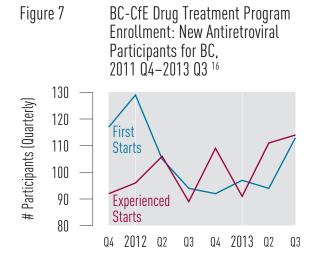
Data Source: British Columbia Centre for Excellence Drug Treatment Program (DTP) Database.

Each quarter's data is calculated as the sum of the 4 quarters leading up to it. e.g. 2012 Q1 is calculated from 2011 Q2 – 2012 Q1.

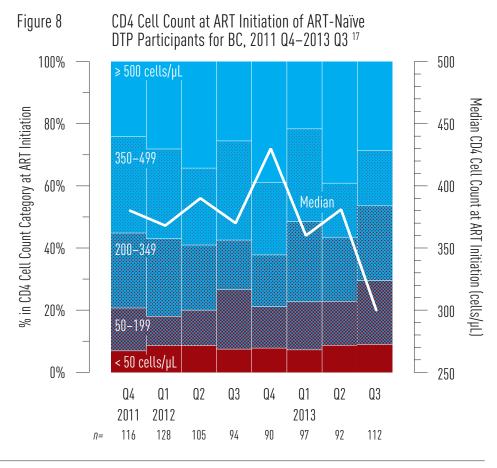
Antiretroviral Uptake

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

Indicator 7. New Antiretroviral Therapy Starts in BC



Indicator 8. CD4 Cell Count at ART Initiation



¹⁶ Data Source: Drug Treatment Program Database

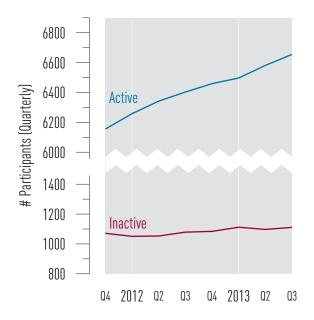
¹⁷ Data Source: Drug Treatment Program Database Limitations: CD4 cell count data is approximately 80% complete.

Indicator 9. Active and Inactive DTP Participants

Table 3. Distribution of People on ART in BC by Health Authority, 2013 Q3 18

		Interior	Fraser	Vancouver Coastal	Vancouver Island	Northern	Total BC
Age	< 30	9	64	141	29	15	258
	30-39	52	259	508	96	43	958
	40-49	113	516	1256	222	61	2168
	≥ 50	239	702	1838	410	81	3271
Gender	Male	321	1191	3293	615	117	5538
	Female	92	350	450	142	83	1117
Exposure	MSM	112	451	1587	180	20	2350
	IDU	144	456	1094	272	122	2088
Total		413	1541	3743	757	200	6655

Figure 9 Active and Inactive DTP Participants for BC, 2011 Q4–2013 Q3 19



¹⁸ Data Source: Drug Treatment Program Database

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.

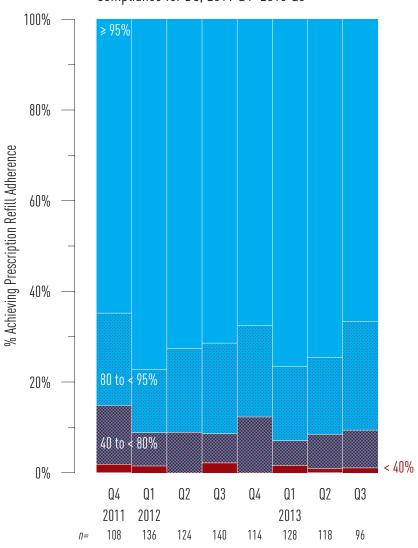
Inactive DTP Participants: Persons no longer prescribed drugs through the HIV/AIDS Drug Treatment Program in the last quarter.

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

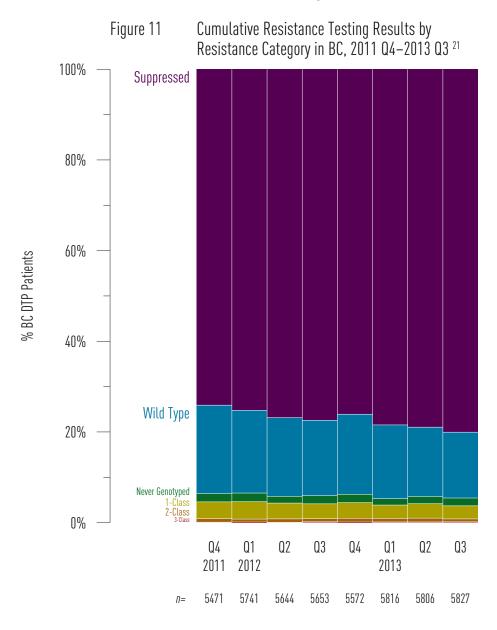
Figure 10 Distribution of Individuals by Adherence Level in 1st Year of Therapy, Based on Pharmacy Refill Compliance for BC, 2011 Q4–2013 Q3 ²⁰



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

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.

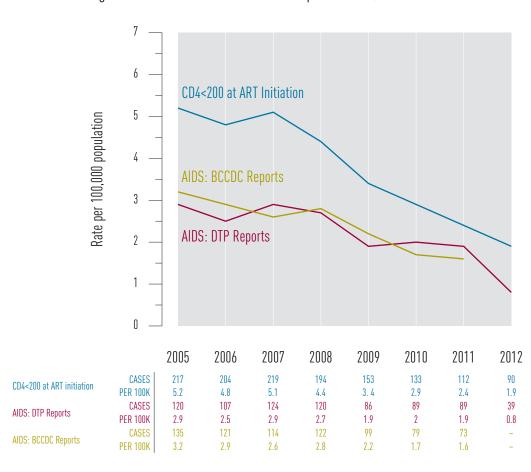


Figure 12 AIDS Case Rate and Reports for BC, 2005–2012 ²²

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.

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.



²³ Data Source: BC Vital Statistics
Limitation: Mortality data is updated annually. The most recent available data was used.

Appendices

Indicator 1		2009)			2010)			2011				2012				2013	,	
Episodes (thousands)	Q1	Q2	Q3	Q4	Q1	Q2	Q3												
B.C. Overa	11	47.2	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	67.4
Gender	Female	28.4	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
	Male	18.1	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
	Other	0.7	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
Female (Pr	enatal)	12.7	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
Female (No	on-prenatal)	15.8	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	22.0	25.6	27.5	26.3
Age	< 30	19.2	18.0	19.0	17.4	18.5	17.9	18.5	18.5	18.5	17.5	19.2	19.9	20.5	20.1	21.7	21.2	20.6	21.2	21.6
	30-39	15.3	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	18.8
	40-49	7.0	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
	≥ 50	5.7	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
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	2.4
Fraser Hea	lth	13.0	12.4	12.3	11.6	13.0	12.3	12.5	12.4	13.1	12.2	13.0	13.0	14.4	14.2	15.1	15.6	17.9	18.4	18.2
Interior He	alth	6.0	5.6	5.5	5.6	6.0	5.6	5.5	5.9	6.0	5.5	5.6	5.9	5.9	5.7	5.9	5.9	6.1	6.4	6.3
Northern F	Iealth	3.1	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	3.5
Van. Coast	al Health	18.9	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	33.4
Van. Island	Health	6.2	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

Indicator 2	2: Rate of HIV Te	sting per 10	0,000		
		2009	2010	2011	2012
British Co	lumbia Overall	3 633.4	3 627.7	3 717.0	4 380.4
Fraser Hea	ılth	3 385.8	3 381.3	3 439.2	3 895.1
Interior H	ealth	2 890.4	2 926.0	2 929.2	2 967.9
Northern 1	Health	3 524.4	3 442.8	3 592.8	4 100.0
Vancouver	Coastal Health	5 114.6	5 088.5	5 342.2	7 111.4
Vancouver	Island Health	2 720.4	2 702.2	2 659.9	2 769.6
Gender	Female	4401.4	4 370.4	4467.0	5 194.4
	Male	2 732.0	2 762.3	2 851.1	3 489.1
Age	< 30	4 292.7	4 222.3	4 303.5	4 766.2
	30-39	8 881.6	9 096.9	9 319.9	10 234.2
	40-49	3 351.6	3 357.9	3 479.7	4 207.1
	≥ 50	1 241.3	1 279.5	1 390.0	2 167.0

						2010				2011				2012				2013		
Indicator 3: Nev	v HIV Diagnoses	Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4	Q1	Q2	Q3
B.C. Overall		102	84	78	73	73	81	78	68	54	82	87	65	70	55	53	60	50	83	92
Gender	Female	21	16	17	17	17	20	14	11	6	19	11	7	9	10	5	5	6	12	14
	Male	81	68	61	56	56	61	64	57	48	63	76	58	61	45	48	55	44	71	78
Age	< 30	18	16	16	24	17	20	22	9	5	18	17	18	18	14	8	18	9	18	24
	30-39	26	27	17	20	25	18	23	20	18	30	30	13	16	17	11	10	17	25	20
	40-49	35	25	30	21	23	29	19	24	18	22	22	19	20	11	19	19	12	14	23
	≥ 50	23	16	15	8	8	14	14	15	13	12	18	15	16	13	15	13	12	26	25
Exposure	MSM	45	43	33	32	39	36	45	33	33	43	56	38	42	34	38	36	28	-	_
	IDU	22	15	13	14	14	16	12	10	8	11	12	3	14	7	2	6	3	_	_
	HET	28	21	21	22	18	25	18	22	12	23	19	21	14	12	11	15	9	_	_
	Other	4	2	3	3	2	2	1	1	1	3	0	3	0	1	0	1	1	_	_
	NIR	3	3	8	2	0	2	2	2	0	2	0	0	0	1	2	2	9	_	_
Fraser Health	By Client Residence	27	27	21	12	17	19	19	17	12	18	15	9	11	10	10	14	9	16	22
	By Provider Address	22	17	17	7	14	15	13	15	10	20	9	8	10	5	7	8	9	9	14
Interior Health	By Client Residence	6	4	4	4	4	1	3	3	1	4	3	3	5	5	0	2	1	4	5
	By Provider Address	6	4	4	4	4	1	2	3	1	3	2	3	5	5	0	2	1	5	5
Northern	By Client Residence	7	8	5	7	4	3	7	2	4	10	5	5	5	4	3	0	3	8	3
Health	By Provider Address	7	8	5	8	4	3	6	2	4	9	5	6	5	4	3	0	3	8	3
Vancouver	By Client Residence	51	36	41	36	41	45	41	40	32	48	57	41	43	33	34	32	32	45	50
Coastal Health	By Provider Address	56	46	45	40	45	50	49	43	34	47	65	42	47	38	38	38	32	51	58
Vancouver	By Client Residence	9	9	7	14	6	13	8	6	5	2	7	7	6	3	5	12	4	10	12
Island Health	By Provider Address	11	9	7	14	6	12	8	5	5	3	6	6	3	3	5	12	5	10	12

		BC]	Female			Male			30 yea			-39 yea	rs	40	-49 yea	ars
	2010	2011	2012	2010	2011	2012	2010	2011	2012	2010	2011	2012	2010	2011	2012	2010	2011	201
Stage 0	48	63	53	8	3	6	40	60	47	16	14	18	15	25	18	13	19	1
Stage 1	67	52	52	12	7	6	55	45	46	20	12	11	20	20	14	20	9	1
Stage 2a	35	39	28	5	7	5	30	32	23	8	11	4	14	12	4	9	11	1
Stage 2b	36	39	31	12	8	6	24	31	25	6	6	7	10	8	8	11	12	1
Stage 3	76	69	57	22	13	5	54	56	52	2	5	7	13	17	10	31	26	
Unknown	38	26	17	3	5	1	35	21	16	11	6	7	15	10	3	11	5	
Гotal	300	288	238	62	43	29	238	245	209	63	54	54	87	92	57	95	82	(
	≥ 2010	50 year 2011		2010	MSM 2011	2012	Het 2010	erosexi 2011	ual 2012	2010	IDU 2011	2012	Othe 2010	r Expo		NIR 2010	/Unkne 2011	
Stage 0	4	5	7	34	51	42	7	7	7	6	5	4	0	0	0	1	0	
Stage 1	7	11	11	34	31	33	20	13	12	13	8	6	0	0	1	0	0	
Stage 2a	4	5	9	21	25	17	8	7	5	6	5	6	0	2	0	0	0	
Stage 2b	9	13	6	16	20	17	13	15	8	5	3	6	2	1	0	0	0	
Stage 3	30	21	25	25	28	29	28	31	20	18	7	6	3	3	0	2	0	
Unknown	1	5	5	23	15	11	7	2	1	4	6	1	1	1	1	3	2	
Total	55	60	63	153	170	149	83	75	53	52	34	29	6	7	2	6	2	
Indicator 5: HI	V Casca	ade of (Care		DIAGN	OSED		LINKE	D	RETAI	NED	(ON ART	A	DHERE	NT	SUPPR	ESSI
3.C. Overall						9284		833	8	7	123		6541		60)27		48
Age Category	< 30					476		32	4		265		226		1	189		1
	30-39					1350		118	9	1	001		865		7	767		6
	40-49					2962		270	0	2	279		2091		18	394		15
	≥ 50					4488		412	5	3	579		3359		31	177		26
Age Category	MSM		< 30			77		7	1		56		48			40		
and MSM			30-3	9		283		26	7		235		209		1	188		1.
Status			40-4	9		800		77	6		716		680		6	534		5
			≥ 50			1604		156	4	1	510		1445		13	390		12
	Non-M	ISM	< 30			67		6	5		60		49			36		
			30-3	9		430		40	9		372		321		2	266		1
			40-4	9		935		89	8		845		771		6	562		4
			≥ 50			1271		122	6	1	164		1090		10	002		7
Į	Unkno	wn	< 30			333		18	7		149		129		1	113		

30-39

40 - 49

≥ 50

Gender

Injection

Drug Use

MSM Status

Health

Authority

Male

IDU

MSM

Female

Non-IDU

Unknown

Non-MSM

Unknown

Fraser Health

Island Health

Interior Health

Northern Health

Vancouver Coastal Health

Indicator 6: Programmatic	2011	2012					2013		
	Q4	Q1	Q2	Q3	Q4		Q1	Q2	Q3
< 3 CD4 Tests	10.2%	10.0%	9.2%	8.1%	8.6%		7.0%	7.5%	9.5%
< 3 Viral Load Tests	6.3%	7.8%	7.5%	7.0%	7.6%		5.0%	6.0%	6.7%
No Baseline Genotype	6.6%	6.1%	6.4%	6.0%	5.7%		5.5%	5.0%	5.0%
Baseline CD4 < 200 cells/μL		29.0%	26.9%	25.3%	23.6%		22.2%	22.6%	23.7%
Non-Recommended ART	2.4%	2.8%	2.5%	2.3%	2.2%		1.5%	2.0%	5.8%
Non Viral suppression at 9 l		25.3%	23.3%	24.7%	26.4%		24.7%	25.6%	26.2%
PCS Score: 0	165	180	188	202	20.4%		222	220	184
PCS Score: 1	94	104	105	115	118		113	103	98
PCS Score: 2	52	48	42	42	51		49	59	57
PCS Score: 3	15	19	18	19	20		15	15	16
PCS Score: 4 or more	7	8	7	6	5		2	2	4
Total (n=)	333	359	360	384	406		401	399	359
Indicator 7: New DTP	2011	2012					2013		
ARV Participants	Q4	Q1	Q2	Q3	Q4		Q1	Q2	Q3
First Starts	117	129	105	94	92		97	94	113
Experienced Starts	92	96	106	89	109		91	111	113
Experienced starts	,2	70	100	0)	107		71	111	113
Indicator 8: CD4 Cell Coun	nt at ART Initiation	for ARV-N	aïve DTP Pa	articipants					
CD4 ≥ 500	28	36	36	24	35		21	36	32
CD4 350-499	36	37	26	30	21		29	16	20
CD4 200-349	28	32	22	15	15		25	19	26
CD4 50-199	16	12	12	18	12		15	13	23
CD4 < 50	8	11	9	7	7		7	8	10
CD4 Median (cells/µL)	380	368	390	370	430		360	381	300
Total (n=)	116	128	105	94	90		97	92	111
,									
Indicator 9: Active and Inac	ctive DTP Participa	nts							
Active DTP Participants	6155	6258	6342	6403	6459		6497	6581	6653
Inactive DTP Participants	1217	1200	1198	1219	1223		1251	1235	1249
_									
Indicator 10: Antiretroviral	l Adherence								
≥ 95%	70	105	90	100	77		98	88	64
80% to < 95%	22	19	23	28	23		21	20	23
40% to < 80%	14	10	11	9	14		7	9	8
< 40%	2	2	0	3	0		2	1	1
Total (n=)	108	136	124	140	114		128	118	96
Indicator 11: Resistance Tes									
Suppressed	4054	4319	4337	4378	4240		4563	4584	4665
Wild Type	1067	1048	979	939	984		947	888	846
Never Genotyped	100	109	85	102	103		81	92	101
1-Class	203	221	197	186	193		173	185	168
2-Class	42	37	43	38	45		43	48	37
3-Class	5	7	3	10	7		9	9	10
Total (n=)	5471	5741	5644	5653	5572		5816	5806	5827
Indicator 12: AIDS-Definin	_	2005	2006	2007	2008	2009	2010	2011	2012
	Cases	217	204	219	194	153	133	112	90
	Rate per 100,000	5.2	4.8	5.1	4.4	3.4	2.9	2.4	1.9
	Cases	120	107	124	120	86	89	89	39
	Rate per 100,000	2.9	2.5	2.9	2.7	1.9	2	1.9	0.8
	Cases	135	121	114	122	99	79	73	_
(BCCDC Reports)	Rate per 100,000	3.2	2.9	2.6	2.8	2.2	1.7	1.6	_
Indicator 13: HIV-Related 1		2005	2006	2007	2008	2009	2010	2011	
British Columbia Overall	105	146	142	100	79	63	54	59	
Per 100 HIV+ Population	1.03	1.4	1.34	0.93	0.72	0.56	0.47	0.5	
Per 100,000 Population	2.5	3.43	3.29	2.28	1.8	1.41	1.19	1.29	