

## HIV MONITORING QUARTERLY REPORT

FOR BRITISH COLUMBIA

FOURTH QUARTER 2013

















#### 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.

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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

Dr. Rolando Barrios, Chair, BC-CFE

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Melanie Rusch, VIHA

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

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

## **HIV Testing Episodes and Rates**

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

#### Indicator 1. HIV Testing Episodes

Figure 1.1 HIV Test Episodes in British Columbia, 2009 Q1–2013 Q4

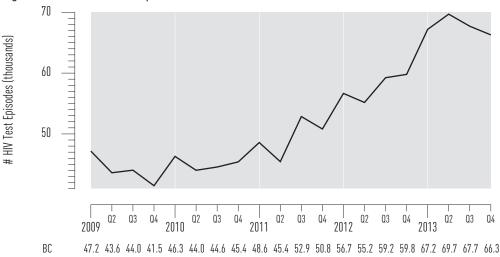
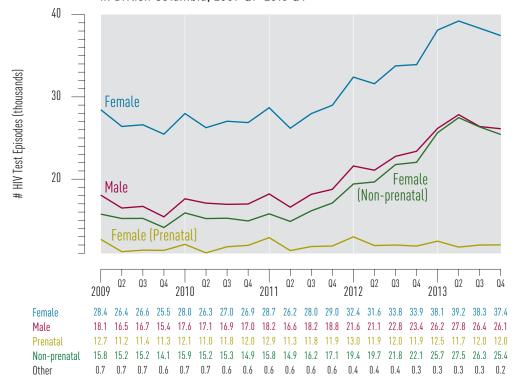
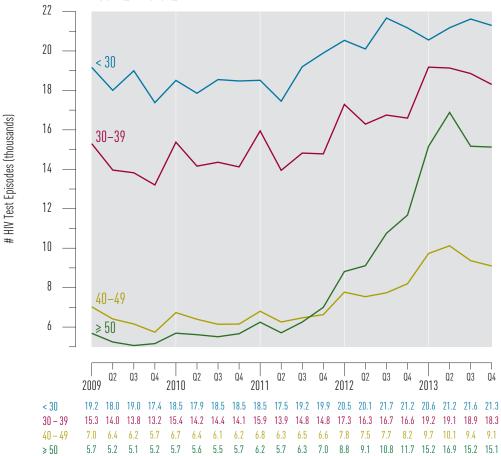


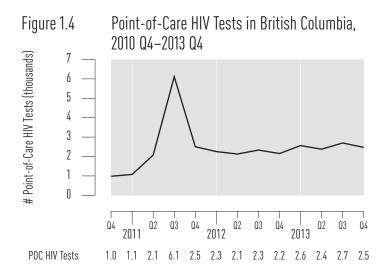
Figure 1.2 HIV Test Episodes by Gender and Prenatal Status in British Columbia, 2009 Q1–2013 Q4 <sup>1</sup>



<sup>1</sup> NB: Testing does not include point of care tests.

Figure 1.3 HIV Test Episodes by Age Category in British Columbia, 2009 Q1–2013 Q4  $^{1.2}$ 





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

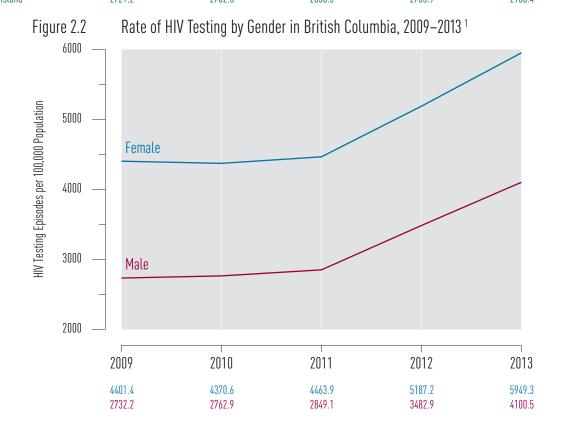
#### Limitations:

- 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 in British Columbia by Health Authority, 2009 Q1–2013 Q4 36 34 32 30 28 26 24 22 # HIV Test Episodes (thousands) 20 Vancouver Coastal 18 16 14 Fraser 12 10 8 Interior Vancouver Island Northern Q4 Q2 Q4 Q2 Q4 2010 2011 2012 2009 13.0 12.4 12.3 11.6 13.0 12.3 12.5 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 Fraser Interior 6.0 5.6 5.5 5.6 6.0 5.6 5.5 5.9 6.0 5.5 5.9 5.9 5.7 5.9 6.1 5.6 5.9 3.1 2.6 2.7 2.5 2.9 2.6 2.7 2.5 3.0 2.7 2.9 2.8 3.1 Northern 3.4 3.2 3.3 3.7 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 33.7 32.7 Vancouver Coastal Vancouver Island  $6.2 \quad 5.1 \quad 5.2 \quad 4.7 \quad 5.6 \quad 5.3 \quad 5.3 \quad 5.3 \quad 5.8 \quad 5.0 \quad 5.3 \quad 5.1 \quad 5.8 \quad 5.3 \quad 5.5 \quad 5.6 \quad 5.9 \quad 6.2 \quad 6.0 \quad 6.0$ 12

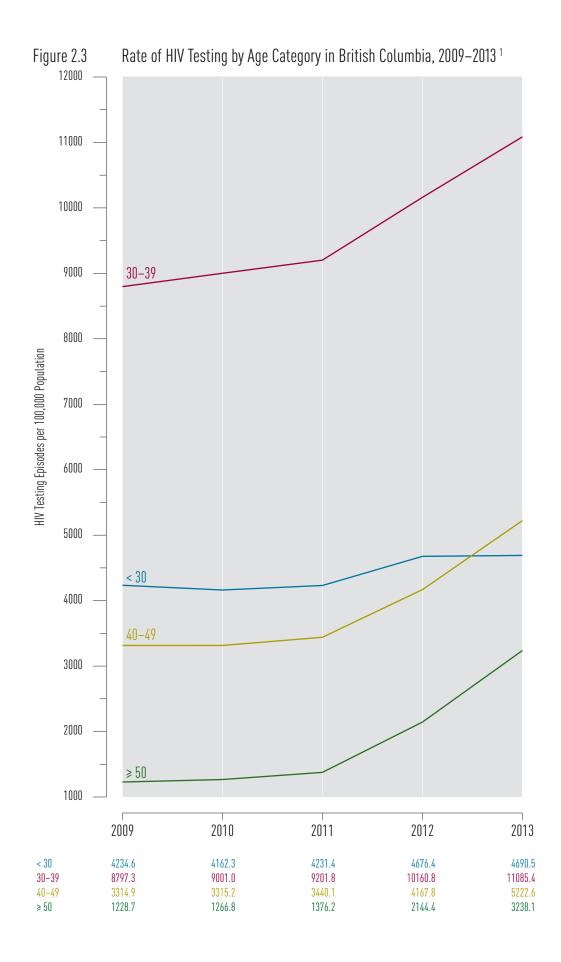
## Indicator 2. HIV Testing Rates

Figure 2.1 Rate of HIV Testing in British Columbia and HAs, 2009–2013 <sup>1</sup> 9000 8000 HIV Testing Episodes per 100,000 Population 7000 6000 Vancouver Coastal 5000 Northern 4000 All BC Fraser Interior 3000 Vancouver Island 2000 2010 2011 2012 2013 2009 3633.5 3628.0 3714.3 4373.4 5055.3 British Columbia 3386.3 3382.4 3437.4 3893.5 4253.2 Fraser Interior 2890.3 2926.6 2927.5 2988.1 3261.1 Northern 3519.6 3437.6 3585.2 4090.0 4381.1 5088.8 Vancouver Coastal 5115.0 5338.4 7024.1 8890.4 Vancouver Island 2721.2 2702.8 2658.3 2783.9 2900.4



Female

Male



## New HIV Diagnoses

Trends in HIV diagnoses by gender and exposure category are described. Interpreting HIV diagnoses must be done with consideration that trends are influenced by both changes in testing rate as well as changes in transmission rates. It is important to note that new HIV diagnoses cases and rates are not synonymous with HIV incidence as a person may have become infected with HIV long before they tested positive for HIV. However, as there is no reliable method for measuring HIV incidence we follow trends in HIV diagnoses.

#### Indicator 3. New HIV Diagnoses

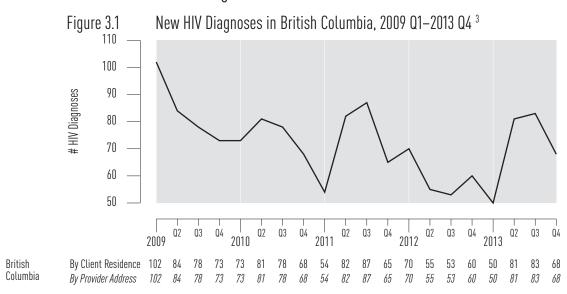


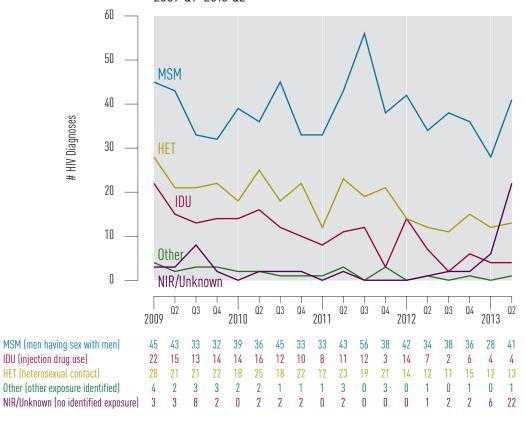
Figure 3.2 New HIV Diagnoses in British Columbia by Gender, 2009 Q1–2013 Q4



<sup>3</sup> Data Source: BCCDC

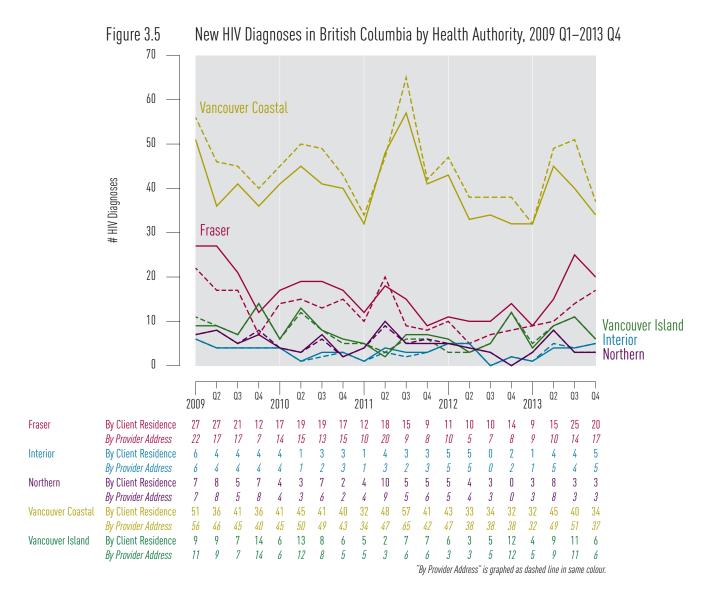
Figure 3.3 New HIV Diagnoses in British Columbia by Age Category, 2009 Q1-2013 Q4 40-49 30 - 39# HIV Diagnoses < 30 ≥ 50 < 30 years 30-39 years 40-49 years ≥ 50 years 

Figure 3.4 New HIV Diagnoses in British Columbia by Exposure Category, 2009 Q1–2013 Q2  $^4$ 



<sup>4</sup> BCCDC: Data lags by 6 months.

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

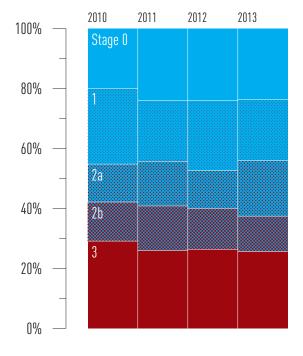


## Stage of HIV infection at diagnosis

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 British Columbia, 2010–2013 <sup>5</sup>

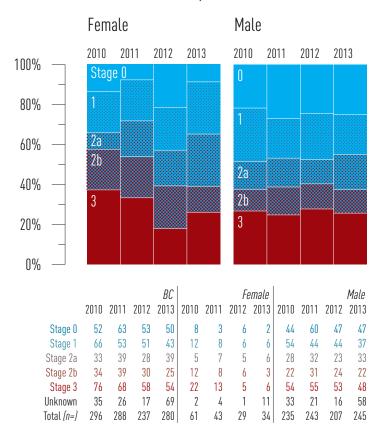


#### Indicator 4. Stage of HIV Infection at Diagnosis

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

Stage	Criteria				
0	previous r	, negativ	ria met for acute ve or indetermina firmed positive H	te HIV	test within 180
1			CD4 ≥500		N. AIDC
2a			CD4 350-499	and	No AIDS case report
2b	Stane N		CD4 200-349		торого
3	Stage 0 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 British Columbia, 2010–2013 <sup>5</sup>



Data Source: BCCDC

Figure 4.3 Stage of HIV Infection at Diagnosis by Age Category for British Columbia, 2010–2013 <sup>5</sup>

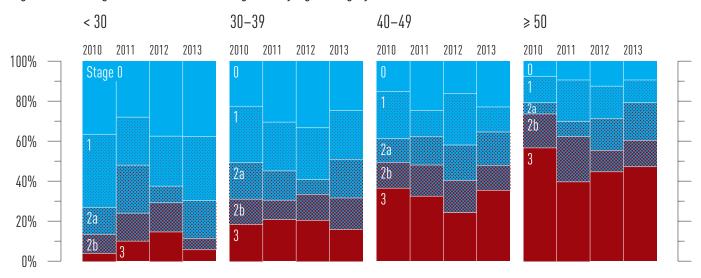
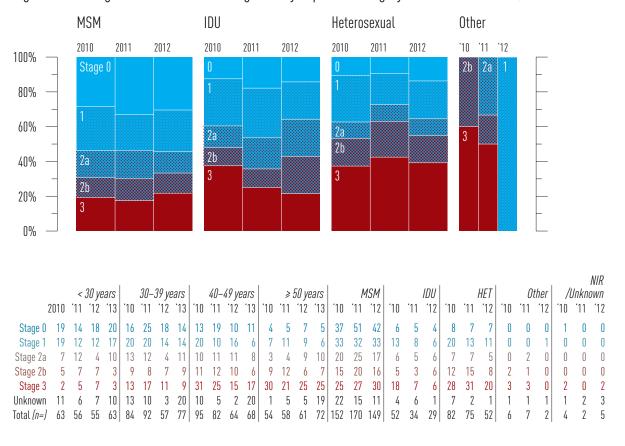


Figure 4.4 Stage of HIV Infection at Diagnosis by Exposure Category for British Columbia, 2010–2012 5.6



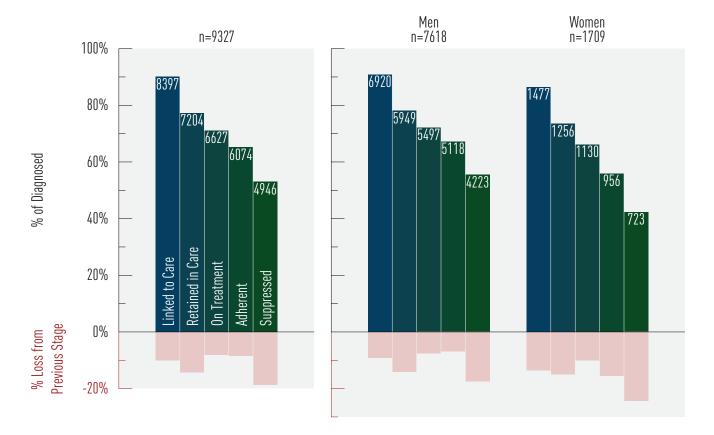
<sup>6</sup> MSM=men who have sex with men; IDU= injection drug user; HET=heterosexual. NIR=No identified risk/exposure.

#### 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 British Columbia, 2013 7

Figure 5.2 Estimated Cascade of Care for British Columbia by Gender, 2013 8



<sup>7,8</sup> Data is for the period 2013 Q1–2013 Q4.

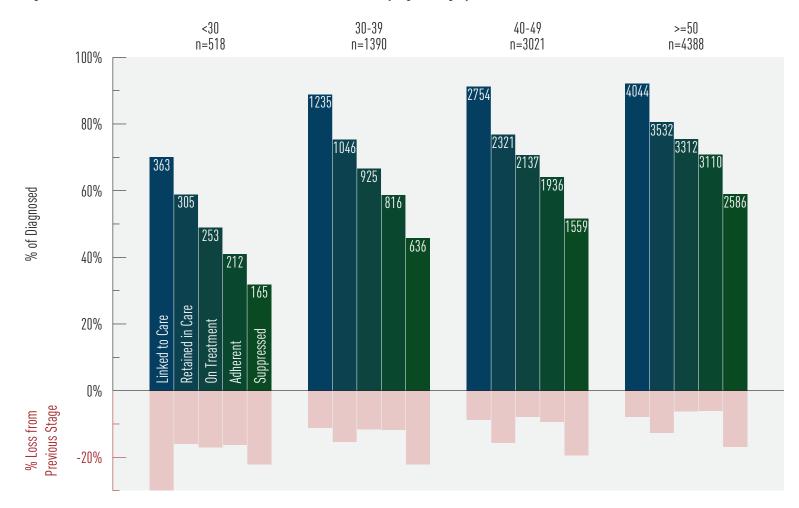
#### Data Sources:

- 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 British Columbia by Age Category, 2013  $^9$ 



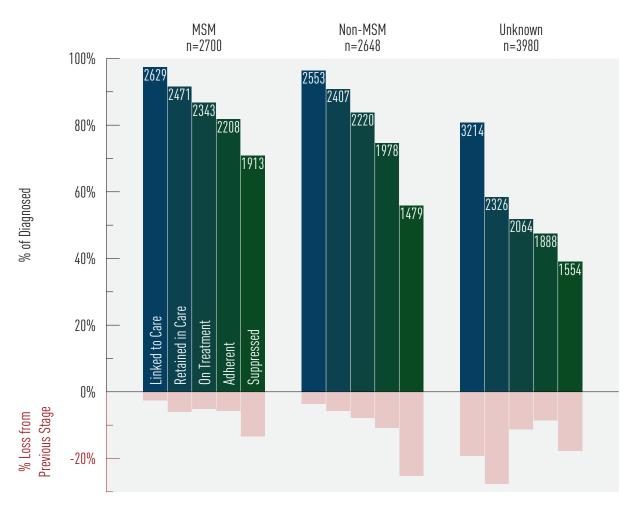
Limitations: на assignment is based on the most recent на of residence of the patient, if not available of the нiv-care provider. If the most recent на of residence is not updated then the designated на may be incorrect.

<sup>9</sup> Data is for the period 2013 Q1–2013 Q4. Data Sources:

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

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





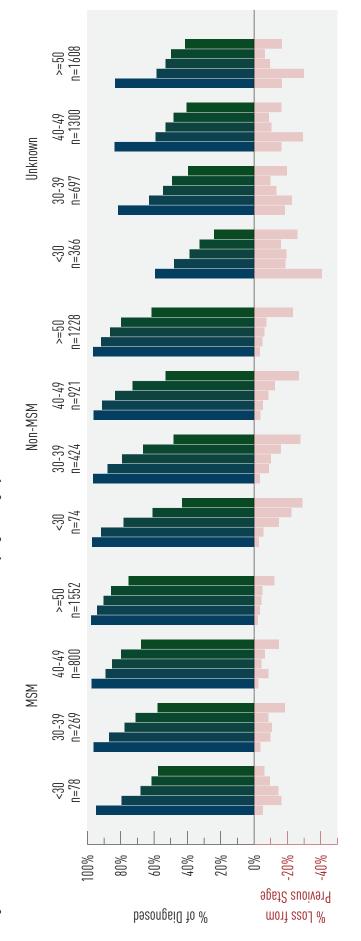
Limitations: HA assignment is based on the most recent HA of residence of the patient, if not available of the HIV-care provider. If the most recent HA of residence is not updated then the designated HA may be incorrect.

<sup>10</sup> Data is for the period 2013 Q1–2013 Q4. Data Sources:

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

<sup>2</sup> 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, 2013 <sup>11</sup> Figure 5.5



Data Sources:

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

Where  $n \le 5$ , data has been withheld for concerns of statistical significance as well as privacy.

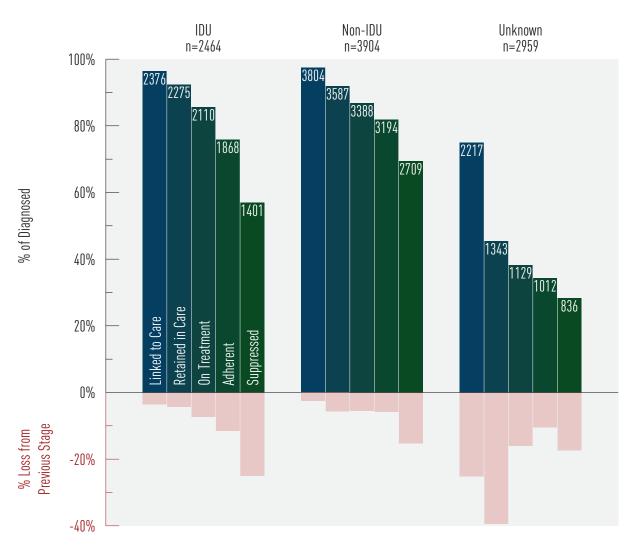
Authorized parties may contact the British Columbia Centre for Excellence in HIV/AIDS to obtain this information.

<sup>11</sup> Data is for the period 2013 Q1-2013 Q4.

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)).





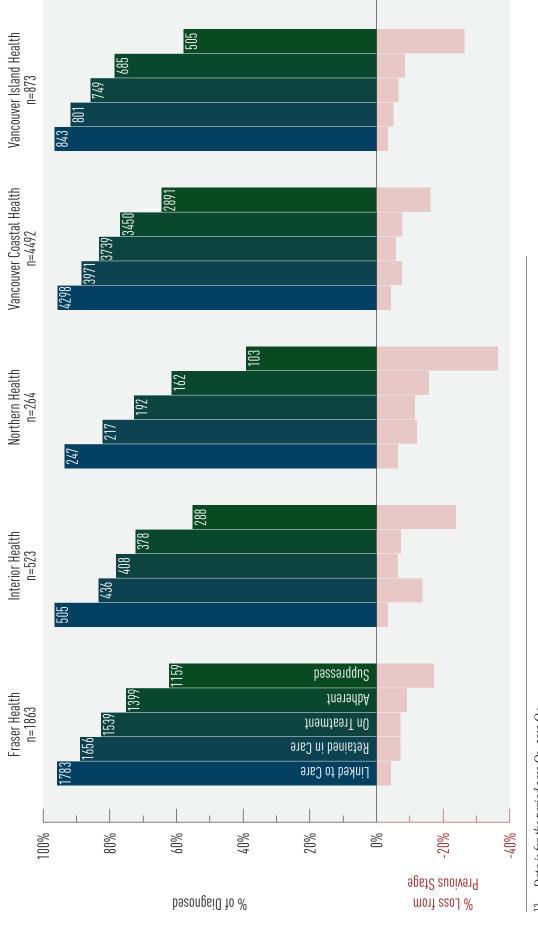
Limitations: на assignment is based on the most recent на of residence of the patient, if not available of the нiv-care provider. If the most recent на of residence is not updated then the designated на may be incorrect.

<sup>12</sup> Data is for the period 2013 Q1–2013 Q4. Data Sources:

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

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





Data is for the period 2013 Q1-2013 Q4. 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 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 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 6.1 PCS Components for British Columbia, 2011–2013 <sup>14</sup>

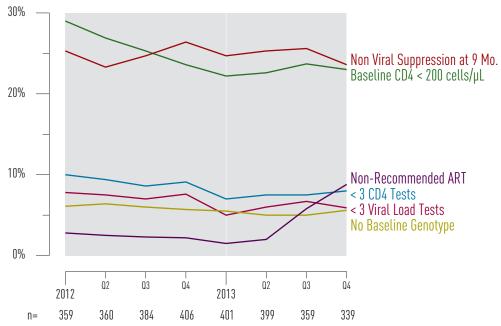
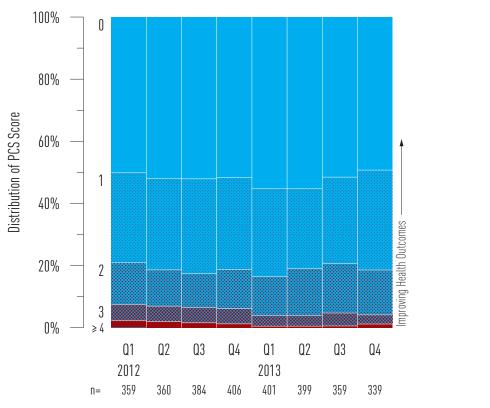


Figure 6.2 Historical Trends for PCS Score for British Columbia, 2011 Q1-2013 Q4 <sup>15</sup>



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

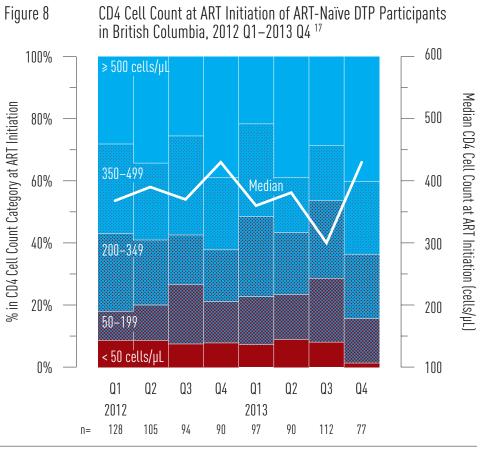
In this section we present trends in ART uptake, the number and proportion of new HIV treatment initiations and the number of active and inactive DTP participants. Trends in ART uptake should be interpreted under the consideration of changing BC HIV treatment guidelines. BC HIV treatment guidelines are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

Figure 7 BC-CfE Drug Treatment Program Enrollment: New ART Participants in BC, 2012 Q1-2013 Q4 16



Indicator 8. CD4 Cell Count at ART Initiation



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

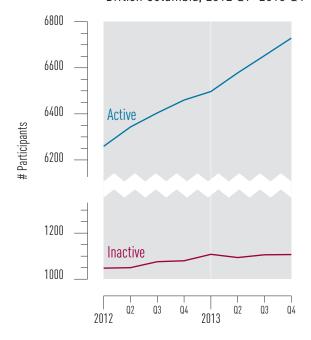
<sup>17</sup> Data Source: Drug Treatment Program Database Limitations: CD4 cell count data is approximately 80% complete.

## Indicator 9. Active and Inactive DTP Participants

Table 3. Distribution of People on ART in British Columbia, 2013 Q4  $^{18}$ 

		Fraser	Interior	Northern	Vancouver Coastal	Vancouver Island	Total BC
Age	< 30	68	10	16	132	28	254
	30-39	261	51	42	523	91	968
	40-49	530	111	63	1242	224	2170
	≥ 50	710	236	86	1884	421	3337
Gender	Male	1212	315	123	3321	617	5588
	Female	357	93	84	460	147	1141
Exposure	MSM	457	109	22	1589	176	2353
	IDU	444	143	124	1116	277	2104
Total		1569	408	207	3781	764	6729

Figure 9 Active and Inactive DTP Participants in British Columbia, 2012 Q1-2013 Q4 19



<sup>18</sup> 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.

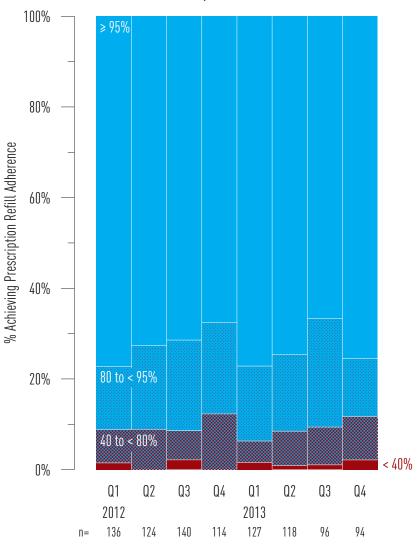
Inactive DTP Participants: Persons no longer prescribed drugs through the HIV/AIDS Drug Treatment Program in the last quarter.

#### 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

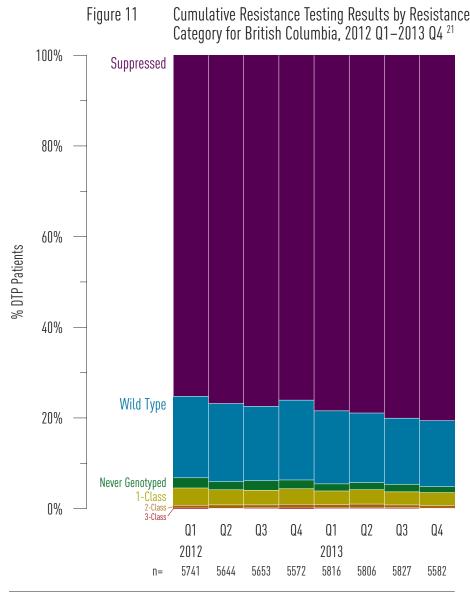




<sup>20</sup> Data Source: Drug Treatment Program Database Limitation: Prescription refill adherence is used as a proxy for patient adherence.

#### 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.



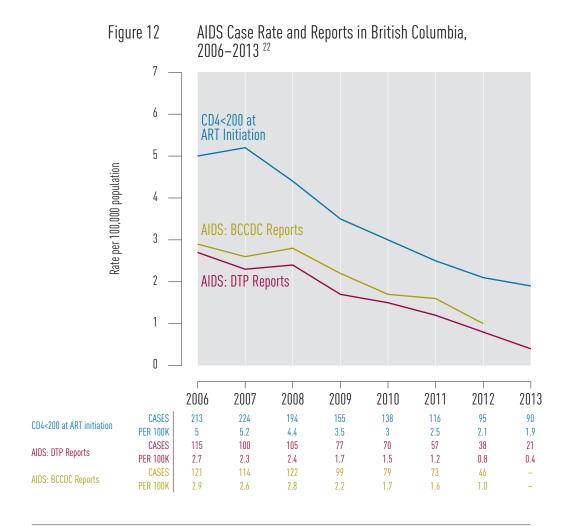
<sup>21</sup> Data Source: Drug Treatment Program Database

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; as such, we have plotted DTP reported AIDS cases as well as the proportion of persons initiating ART with a CD4<200 cells/µL.

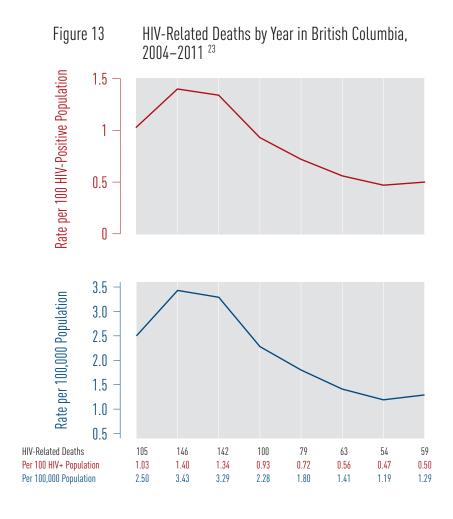


#### 22 Data Source: Drug Treatment Program Database

Limitation: AIDs case reporting was investigated using 2 definitions: First, using AIDs cases reported in AIDs case report forms from the DTP, and second, 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. 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:

<sup>23</sup> Data Source: BC Vital Statistics

<sup>1.</sup> DTP participants are designated to an HA based on most current residence provided by the participant.

<sup>2.</sup> Mortality data is updated annually.

<sup>3.</sup> The most recent available data was used.

## **Appendices**

Indicator		200	9			2010	0			201	1			2012	2			2013	3		
Episodes	(thousands)	Q1	Q2	Q3	Q4																
British Co	lumbia	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	67.7	66.3
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	37.4
	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	26.1
	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	0.2
Female (Pr	renatal)	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	12.0
Female (N	Ion-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	22.1	25.7	27.5	26.3	25.4
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	21.3
	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	16.7	16.6	19.2	19.1	18.9	18.3
	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	9.1
	≥ 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	15.1
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
Fraser Hea	alth	13.0	12.4	12.3	11.6	13.0	12.3	12.5	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
Interior H	ealth	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	6.7
Northern 1	Health	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	3.6	3.6
Vancouver	r Coastal 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	33.7	32.7
Vancouver	r 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	6.0

Indicator 2: Rate of HIV Testing per 100,000

		2009	2010	2011	2012	2013
British Col	umbia	3633.5	3628.0	3714.3	4373.4	5055.3
Fraser Heal	lth	3386.3	3382.4	3437.4	3893.5	4253.2
Interior He	alth	2890.3	2926.6	2927.5	2988.1	3261.1
Northern F	Iealth	3519.6	3437.6	3585.2	4090.0	4381.1
Vancouver	Coastal Health	5115.0	5088.8	5338.4	7024.1	8890.4
Vancouver	Island Health	2721.2	2702.8	2658.3	2783.9	2900.4
Gender	Female	4401.4	4370.6	4463.9	5187.2	5949.3
	Male	2732.2	2762.9	2849.1	3482.9	4100.5
Age	< 30	4234.6	4162.3	4231.4	4676.4	4690.5
	30-39	8797.3	9001.0	9201.8	10160.8	11085.4
	40-49	3314.9	3315.2	3440.1	4167.8	5222.6
	≥ 50	1228.7	1266.8	1376.2	2144.4	3238.1

		2009	,			2010	)			2011				2012				2013			
Indicator 3: New HIV	Diagnoses	Q1	Q2	Q3	Q4																
British Columbia	By Client Residence	102	84	78	73	73	81	78	68	54	82	87	65	70	55	53	60	50	81	83	68
	By Provider Address	102	84	78	73	73	81	78	68	54	82	87	65	70	55	53	60	50	81	83	68
Gender	Female	21	16	17	17	17	20	14	11	6	19	11	7	9	10	5	5	6	12	9	7
	Male	81	68	61	56	56	61	64	57	48	63	76	58	61	45	48	55	44	69	74	61
Age	< 30	18	16	16	24	17	20	22	9	5	18	17	18	18	14	8	18	9	18	23	16
	30-39	26	27	17	20	25	18	23	20	18	30	30	13	16	17	11	10	17	25	19	15
	40-49	35	25	30	21	23	29	19	24	18	22	22	19	20	11	19	19	12	14	21	23
	≥ 50	23	16	15	8	8	14	14	15	13	12	18	15	16	13	15	13	12	24	20	14
Exposure	MSM	45	43	33	32	39	36	45	33	33	43	56	38	42	34	38	36	28	41	-	_
	IDU	22	15	13	14	14	16	12	10	8	11	12	3	14	7	2	6	4	4	_	_
	HET	28	21	21	22	18	25	18	22	12	23	19	21	14	12	11	15	12	13	_	_
	Other	4	2	3	3	2	2	1	1	1	3	0	3	0	1	0	1	0	1	_	_
	NIR/Unknown	3	3	8	2	0	2	2	2	0	2	0	0	0	1	2	2	6	22	_	_
Fraser Health	By Client Residence	27	27	21	12	17	19	19	17	12	18	15	9	11	10	10	14	9	15	25	20
	By Provider Address	22	17	17	7	14	15	13	15	10	20	9	8	10	5	7	8	9	10	14	17
Interior Health	By Client Residence	6	4	4	4	4	1	3	3	1	4	3	3	5	5	0	2	1	4	4	5
	By Provider Address	6	4	4	4	4	1	2	3	1	3	2	3	5	5	0	2	1	5	4	5

Northern He	alth		Ву С	lient	Reside	nce	7	8	5	7	4	3	7	2	4	10	5	5	5	4	3	0	3	8	3	3
			By P	rovide	er Add	ress	7	8	5	8	4	3	6	2	4	9	5	6	5	4	3	0	3	8	3	3
Vancouver Co Health	oastal		,		Reside er Add				41 45	36 40				40 43	32 34	48 47	57 65	41 42	43 47	33 38	34 38	32 38	32 32	45 49	40 51	34 37
Vancouver Isl Health	land		,		Reside er Add		9 11	9	7 <i>7</i>	14 14	6 6	13 12	8	6 5	5 5	2	7 6	7 6	6	3	5 5	12 12	4 5	9 9	11 11	6 6
Indicator 4: S	taga		,						,		Ü						Ü									
mulcator 4: 3	lage (	л гп В		lectio	ni at d	Fema				M	ale				20 **			2	0.20			1	40	40.		
	'10	'11	°12	<b>'</b> 13	'10			3	'10	'11	°12	'13	'10		30 yo 11	'12	<b>'</b> 13	'10	ود–0 11'	year 12		3 '		-49 '11	'12	<b>'13</b>
Stage 0	52	63	53	50	8	3	6	2	44	60	47	47	19	9	14	18	20	16	25	18	1	4	13	19	10	11
Stage 1	66	53	51	43	12	8	6	6	54	44	44	37	19	9	12	12	17	20	20	14	1	4	20	10	16	6
Stage 2a	33	39	28	39	5	7	5	6	28	32	23	33	7	7	12	4	10	13	12	4	1	1	10	11	11	8
Stage 2b	34	39	30	25	12	8	6	3	22	31	24	22		5	7	7	3	9	8	7		9	11	12	10	6
Stage 3	76	68	58	54	22	13	5	6	54	55	53	48	2	2	5	7	3	13	17	11			31	25	15	17
Unknown	35	26	17	69	2	4	1	1	33	21	16				6	7	10	13	10	3			10	5	2	20
Total	296	288	237	280	61	43	29	34	235	243	207	245	63	3	56	55	63	84	92	57	7	7	95	82	64	68
	'10	≥ 50 '11	years '12	<b>'13</b>	2010	MSN 201		2	2010	IE ) 20	OU 011	2012		Het 10	eros 201	exua	12012	Oth 2010		xpos	ure 201:	2 2	NIF 2010		know	n 012
Stage 0	4	5	7	5	37	5		2			5	4	-	8		7	7		)	0		0	1		0	0
Stage 1	7	11	9	6	33	32	2 3	3	13	3	8	6		20	1	13	11	(	)	0		1	0		0	0
Stage 2a	3	4	9	10	20	25	5	7	6	5	5	6		7		7	5	(	)	2	(	0	0		0	0
Stage 2b	9	12	6	7	15	20	)	6	5	5	3	6		12	1	15	8	2	2	1	(	0	0		0	0
Stage 3	30	21	25	25	25	27	7 (	0	18	8	7	6		28	3	31	20	3	3	3	(	0	2		0	2
Unknown	1	5	5	19	22	15	5	1	4	4	6	1		7		2	1		1	1		1	1		2	3
Total	54	58	61	72	152	170	) 14	9	52	2	34	29		82	7	75	52	(	5	7	:	2	4		2	5
Indicator 5: 1		Casca	de of	f Care	e	DI	AGNO			]	LINKI		R	ETA	INEI			ON A		A	DHE			SUP	PRES	
British Colu		• •						327			83				720			66				607				946
Age Category	•							518				63			30.				53			21				165
		-39 40						390 321			12. 27.				104			21	25			81				636
	40 ≥ !	-49						388			40				<ul><li>232</li><li>353</li></ul>				12			193 311				559 586
Age Categor		SM		< :	30		4.	78				74			6.				53				8		۷.	45
and MSM	y 1V1	31 <b>V</b> 1			-39			269				59			23				09			19				156
Status					-49			300				80			71:				81			63				543
				≥.				552			15				146				00			133				169
	No	on-M	SM	< .				74				72			6	8			58			4	15			32
				30	-39			124			4	09			37	3		3	36			28	32			204
				40	-49		9	921			8	86			839	9		7	67			67	1			490
				$\geq$	50		1:	228			11	87			112	8		10	59			98	80			753
	Ur	ıkno	wn	< .	30			366				17			17	6		1	42			11	9			88
					-39			597				68			439				80			34				276
					-49			300			10				769				89			62				526
				≥ .	50			808			13				94				53			79				664
Gender		ale .						518			69				5949				97			511				223
T		male						709			14				125				30			95				723
Injection	ID		NT T					164			23				227				10			186				401
Drug Use		n-II						904			38				358				88			319				709
MSM Status		nkno SM	WII					959 700			22 26				134: 247				29 43			101 220				836 913
Wisivi Status		on-M	SM					548			25				240				20			197				479
		nkno						980			32				232				64			188				554
Health			wn Iealth	1				363			17				165				39			139				159
Authority			Heal					523				05			430				08			37				288
,			n He					264				47			21				92			16				103
					l Healt	h		192			42				397				39			345				891
					Health			373				43			80				49			68				505

< 3 CD4 Tests < 3 Viral Load Tests	Q1					2013				
< 3 Viral Load Tests	Q1	Q2	Q3		Q4	Q1	Q2		Q3	Q <sup>2</sup>
	10.0%	9.4%	8.6%		9.1%	7.0%	7.5%		7.5%	8.09
	7.8%	7.5%	7.0%		7.6%	5.0%	6.0%		6.7%	5.99
No Baseline Genotype	6.1%	6.4%	6.0%		5.7%	5.5%	5.0%		5.0%	5.69
Baseline CD4 < 200 cells/μΙ	L 29.0%	26.9%	25.3%	2	23.6%	22.2%	22.6%	2	3.7%	23.09
Non-Recommended ART	2.8%	2.5%	2.3%		2.2%	1.5%	2.0%		5.8%	8.89
Non Viral suppression at 9	Mo. 25.3%	23.3%	24.7%	2	26.4%	24.7%	25.3%	2	5.6%	23.69
PCS Score: 0	180	187	200		210	221	220		185	16
PCS Score: 1	104	106	117		120	114	103		100	10
PCS Score: 2	48	42	42		51	50	60		57	4
PCS Score: 3	19	18	19		20	14	14		15	1
PCS Score: 4 or more	8	7	6		5	2	2		2	-
Total (n=)	359	360	384		406	401	399		359	33
Indicator 7: New DTP ARV	/ Particinants									
First Starts	129	105	94		92	97	92		113	7
Experienced Starts	96	106	89		109	91	111		113	128
Indicator 8: CD4 Cell Cour	nt at ADT Initiation	for ADV	Naïva DTD	Doutici	nanta					
$\frac{\text{Indicator 8: CD4 Cell Coul}}{\text{CD4} \ge 500}$	at AKT Initiation	36	Naive DTP	r ar ticl	35	21	35		32	3
CD4 ≥ 500 CD4 350–499	36 37	26	30		35 21	21 29	35 16		32 20	1
CD4 200-349	32	22	15		15	25	18		28	1
CD4 50-199	12	12	18		12	15	13		23	1
CD4 < 50	11	9	7		7	7	8		9	
CD4 Median (cells/µL)	368	390	370		430	360	381		300	43
Total (n=)	128	105	94		90	97	90		112	77
Indicator 9: Active and Ina	ctive DTP Participa	ants								
Active DTP Participants	6259	6343	6404		6460	6497	6578		6653	6729
Inactive DTP Participants	1048	1050	1076		1080	1108	1094		1106	1107
Indicator 10: Antiretrovira	d Adherence									
≥ 95%	105	90	100		77	98	88		64	7:
80% to < 95%	19	23	28		23	21	20		23	13
40% to < 80%	10	11	9		14	6	9		8	
< 40%	2	0	3		0	2	1		1	
Total (n=)	136	124	140		114	127	118		96	9,
I. J 11 D										
Indicator 11: <b>Resistance Te</b> Suppressed	4319	4335	4377		4239	4562	4582		4665	449
Wild Type	1032	975	929		981	937	892		851	81
Never Genotyped	129	100	118		108	91	90		96	7.
1-Class	221	188	183		192	174	184		168	15
2-Class	33	43	37		45	43	49		37	3:
	33 7				43 7					
3-Class Total (n=)	5741	3 5644	9 5653		5572	9 5816	9 5806		10 5827	558
Indicator 12: AIDS-Definit			2006	2007	2008	2009	2010	2011	2012	201
	Cases		213	224	194	155	138	116	95	9
ART initiation	Rate per 100,000		5	5.2	4.4	3.5	3	2.5	2.1	1.
	Cases		115	100	105	77	70	57	38	2
AIDS Cases	Rate per 100,000		2.7	2.3	2.4	1.7	1.5	1.2	0.8	0.
AIDS Cases (DTP Reports)			121	114	122	99	79	73	46	
AIDS Cases (DTP Reports) AIDS Cases	Cases					2.2				
AIDS Cases (DTP Reports) AIDS Cases	Cases Rate per 100,000		2.9	2.6	2.8	2.2	1.7	1.6	1.0	
AIDS Cases (DTP Reports) AIDS Cases (BCCDC Reports)	Rate per 100,000	2005	2.9	2.6	2.8	2.2	2010	1.6 2011	1.0	
AIDS Cases (DTP Reports) AIDS Cases (BCCDC Reports)  Indicator 13: HIV-Related	Rate per 100,000         Mortality       2004	2005 146	2006		2008	2009		2011	1.0	
AIDS Cases (DTP Reports) AIDS Cases	Rate per 100,000			2007			2010		1.0	