

BRITISH COLUMBIA CENTRE for EXCELLENCE in HIV/AIDS

# HIV MONITORING QUARTERLY REPORT FOR VANCOUVER COASTAL HEALTH

FIRST QUARTER 2015

















# Foreword

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

The objectives of these reports are to:

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

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

# List of Indicators

Indicator 1. Testing Episodes Indicator 2. HIV Testing Rate Indicator 3. New HIV Diagnoses Indicator 4. Stage of HIV Infection at Diagnosis Indicator 5. HIV Cascade of Care Indicator 6. Programmatic Compliance Score (PCS) Indicator 7. New Antiretroviral Starts Indicator 8. CD4 Cell Count at ART Initiation Indicator 9. Active and Inactive Drug Treatment Program Participants Indicator 10. Antiretroviral Adherence Level Indicator 11. Resistance Testing Results by Resistance Category Indicator 12. AIDS-Defining Illness Indicator 13. HIV-Related Mortality

# Table of Contents

#### Acknowledgements and Contributions

#### BC Provincial STOP Program:

#### A Note on Monitoring and Interpreting HIV Indicators

Indicator 1	HIV Testing Episodes
Figure 1.1	H1v Test Episodes for Vancouver Coastal Health, 2010 Q2–2015 Q1
Figure 1.2	H1v Test Episodes for Vancouver Coastal Health by Gender and Prenatal Status, 2010 Q2–2015 Q1
Figure 1.3	HIV Test Episodes for Vancouver Coastal Health by Age Category, 2010 Q2–2015 Q1
Figure 1.4	Point-of-Care HIV Tests for Vancouver Coastal Health, 2010 Q4–2015 Q1
Figure 1.5	HIV Test Episodes by HSDA for Vancouver Coastal Health, 2010 Q2–2015 Q1
Figure 1.6	HIV Test Episodes for Non-Prenatal Females in Vancouver Coastal Health by HSDA, 2010 Q2–2015 Q1
Figure 1.7	HIV Test Episodes for Males in Vancouver Coastal Health by HSDA, 2010 Q2–2015 Q1
Indicator 2	HIV Testing Rates
Figure 2.1	Rate of HIV Testing for Vancouver Coastal Health and HSDA's, 2009–2014
Figure 2.2	Rate of HIV Testing for Vancouver Coastal Health by Gender, 2009–2014
Figure 2.3	Rate of HIV Testing for Vancouver Coastal Health by Age Category, 2009–2014
Indicator 3	New HIV Diagnoses
Figure 3.1	New HIV Diagnoses for Vancouver Coastal Health, 2010 Q2–2015 Q1
Figure 3.2	New HIV Diagnoses for Vancouver Coastal Health by Gender, 2010 Q2–2015 Q1
Figure 3.3	New HIV Diagnoses for Vancouver Coastal Health by Age Category, 2010 Q2–2015 Q1
Figure 3.4	New HIV Diagnoses for Vancouver Coastal Health by Exposure Category, 2010 Q1–2014 Q2
Figure 3.5	New HIV Diagnoses for Vancouver Coastal Health by HSDA, 2010 Q2–2015 Q1
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
Figure 4.1	Stage of HIV Infection at Diagnosis for Vancouver Coastal Health, 2010–2014
Figure 4.2	Stage of H1v Infection at Diagnosis for Vancouver Coastal Health by Gender, 2010–2014
Figure 4.3	Stage of HIV Infection at Diagnosis for Vancouver Coastal Health by Age Category, 2010–2014
Figure 4.4	Stage of H1v Infection at Diagnosis for Vancouver Coastal Health by Exposure Category, 2010–2013
Indicator 5	HIV Cascade of Care
Figure 5.1	Estimated Cascade of Care for Vancouver Coastal Health, Year Ending 2015 Q1
Figure 5.2	Estimated Cascade of Care for Vancouver Coastal Health by Gender, Year Ending 2015 Q1
Figure 5.3	Estimated Cascade of Care for Vancouver Coastal Health by Age Category, Year Ending 2015 Q1

Figure 5.4	Estimated Cascade of Care for Vancouver Coastal Health by Msм Status, Year Ending 2015 Q1
Figure 5.5	Estimated Cascade of Care for Vancouver Coastal Health by Age Category and Мѕм Status, Year Ending 2015 Q1
Figure 5.6	Estimated Cascade of Care for Vancouver Coastal Health by History of IDU, Year Ending 2015 Q1
Figure 5.7	Estimated Cascade of Care for Vancouver Coastal Health by HSDA, Year Ending 2015 Q1
Indicator 6	Programmatic Compliance Score (PCS)
Table 2	Probability of Mortality Based on the Programmatic Compliance Score
Figure 6.1	Pcs Components for Vancouver Coastal Health, 2013 Q2–2015 Q1
	First-Year CD4 Measurement
	First-Year VL measurement
	Baseline Resistance Testing
	Recommended Antiretroviral Therapy (ART)
	Baseline CD4 $\ge$ 200 cells/ $\mu$ L
	Suppression at 9 Months
Figure 6.2	Historical Trends for Pcs Score for Vancouver Coastal Health, 2013 Q2–2015 Q1
Indicator 7	New Antiretroviral Therapy Starts in Vancouver Coastal Health
Figure 7	BC-CfE Drug Treatment Program Enrollment: New Antiretroviral Participants for Vancouver Coastal Health, 2013 Q2–2015 Q1
Indicator 8	CD4 Cell Count at ART Initiation
Figure 8	CD4 Cell Count at ART Initiation for Vancouver Coastal Health, 2013 Q2–2015 Q1
Indicator 9	Active and Inactive Drug Treatment Program (DTP) Participants
Table 3	Distribution of People on ART in Vancouver Coastal Health, 2015 Q1
Figure 9	Active and Inactive DTP Participants for Vancouver Coastal Health, 2013 Q2–2015 Q1
Indicator 10	Antiretroviral Adherence
Figure 10	Distribution of Individuals by Adherence Level in 1st Year of Therapy, Based on Pharmacy Refill Compliance for Vancouver Coastal Health, 2013 Q2–2015 Q1
Indicator 11	Resistance Testing and Results
Figure 11	Cumulative Resistance Testing Results by Resistance Category for Vancouver Coastal Health, 2013 Q2–2015 Q1
Indicator 12	AIDS-Defining Illness
Figure 12	AIDs Case Rate and Reports for Vancouver Coastal Health, 2007–2014
Indicator 13	HIV-Related Mortality
Figure 13	H1v-Related Deaths by Year for Vancouver Coastal Health, 2004–2011

# Acknowledgements and Contributions



BRITISH COLUMBIA CENTRE for EXCELLENCE in HIV/AIDS

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



BC Centre for Disease Control An agency of the Provincial Health Services Authority

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

#### **Other Data Sources:**

The above databases were supplemented with:

(I) The BC Vital Statistics database which was used to calculate Indicator 5. The HIV Cascade of Care and Indicator 13. HIV-Related Mortality.

(II) Linkage and preparation of the de-identified individual-level database used for calculating Indicator 5. The HIV Cascade of Care was facilitated by the British Columbia Ministry of Health.

(III) The Statistics Canada database: BC and HIV-positive population counts were acquired through the statistics Canada website to calculate HIV-specific mortality rates for Indicator 13. HIV-Related Mortality.

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

Dr. Rolando Barrios, *Chair*, BC-CFE Kate Heath, BC-CFE Bohdan Nosyk, BC-CFE Viviane Dias Lima, BC-CFE Irene Day, BC-CFE Dr. Jason Wong, BCCDC Dr. Mel Kradjen, BCCDC Salman Klar, FHA Corey Green, FNHA Jennifer May-Hadford, IHA James Haggerstone, NHA Dr. Neora Pick, PHSA Dr. Reka Gustafson, VCHA 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 antiretroviral therapy (ART) initiation.

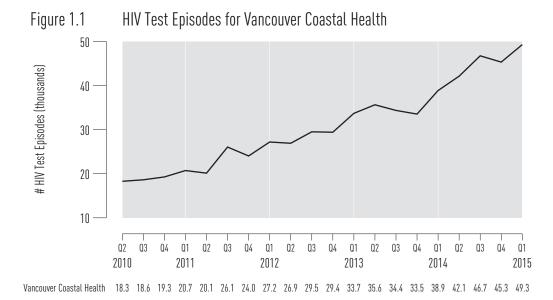
The expansion to a province-wide programme was announced on November 30th 2013 by the BC Ministry of Health with roll out of funding beginning on April 1st, 2013. This funding is intended to be used in the implementation and evaluation of HIV-related diagnosis and care initiatives within individual HA's.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

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

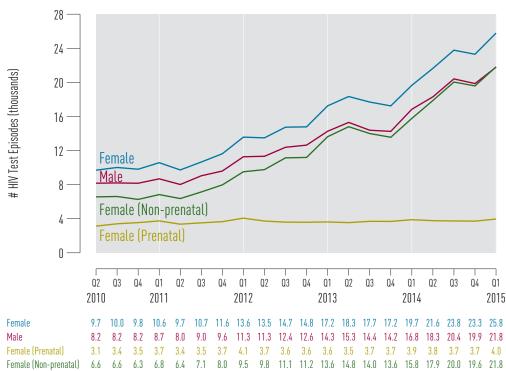
## **HIV Testing Episodes and Rates**

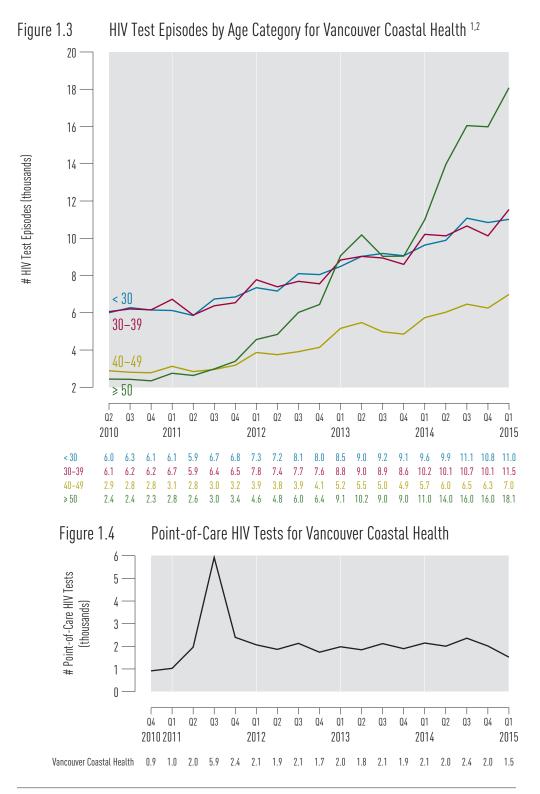
In this section, the number of HIV test episodes and point of care (POC) HIV tests conducted each quarter in BC is shown. In general terms the goal is to increase the number of tests performed and to maximize testing efficiency. Test episodes are allocated by region according to where the test is performed.

#### Indicator 1. HIV Testing Episodes





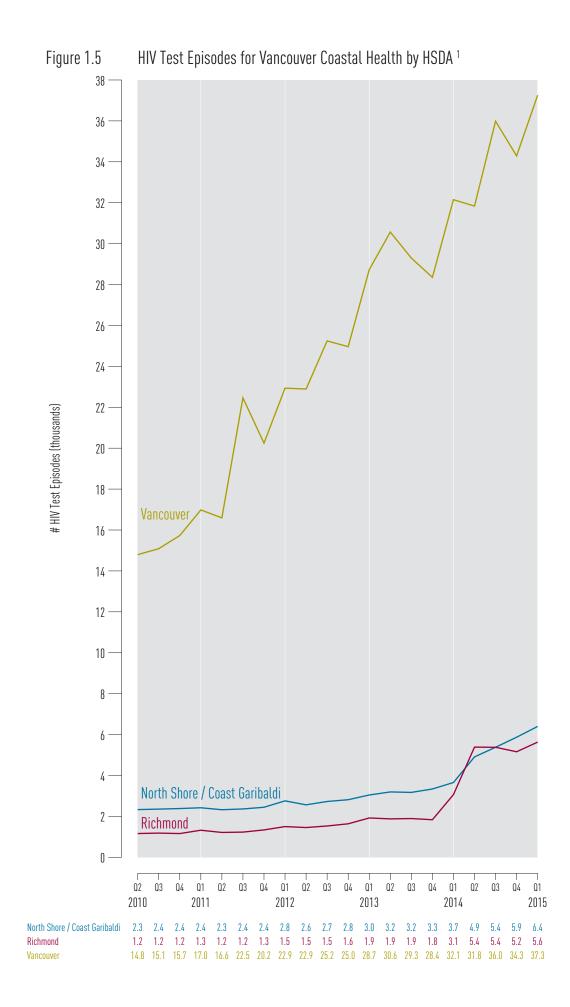


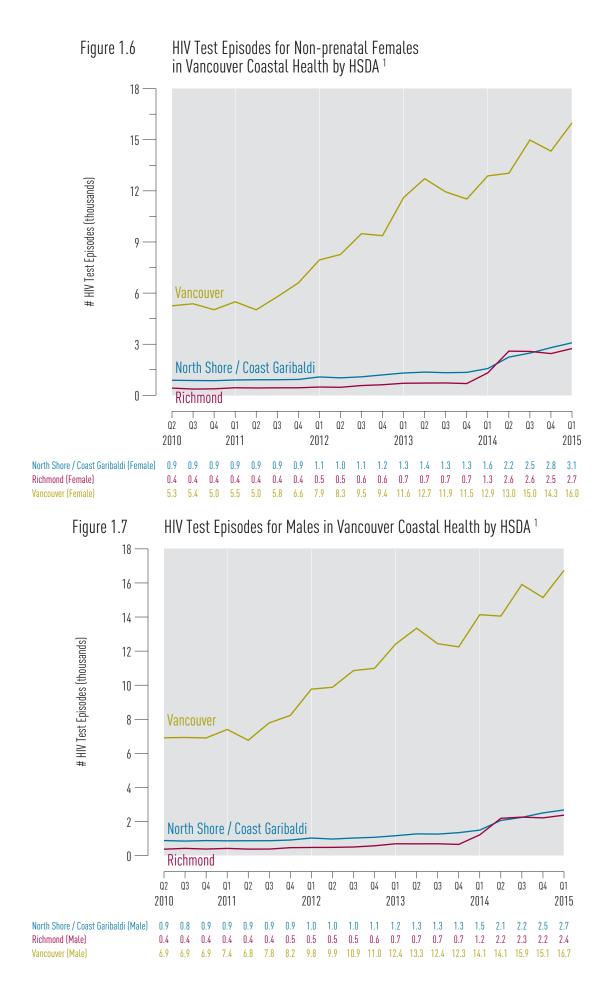


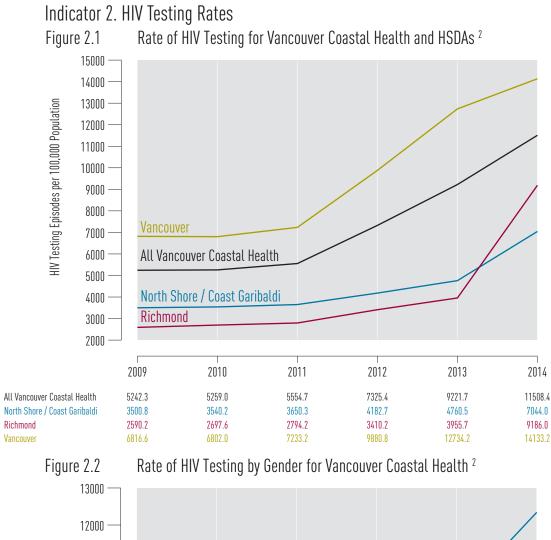
1 Data Source: The BC Public Health Microbiology and Reference Laboratory (BCPHMRL) courtesy of the BC Centre for Disease Control (BCCDC).

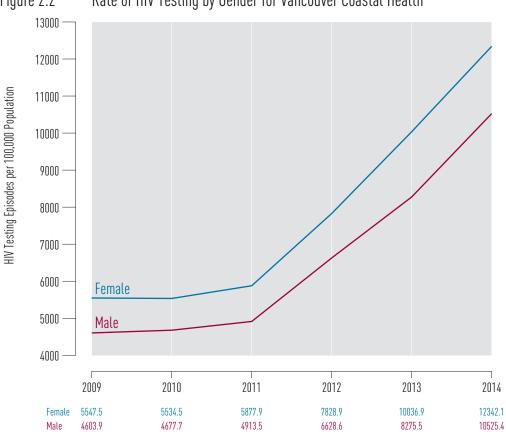
Limitations:

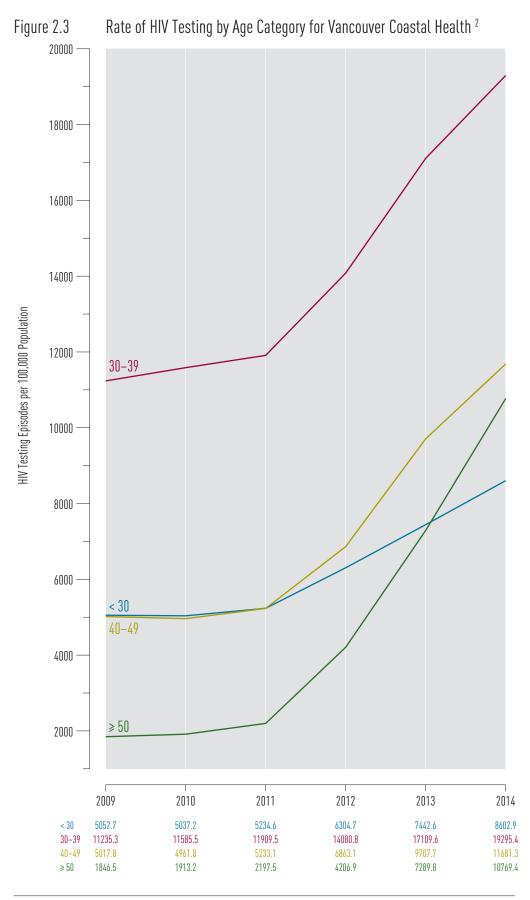
- *i* Repeat tests in individuals who test using various identifiers may not be identified and these individuals may be counted more than once.
- *ii* Poc testing data are available from the fourth quarter of 2010 forward.
- 2 *Testing does not include point of care tests.*









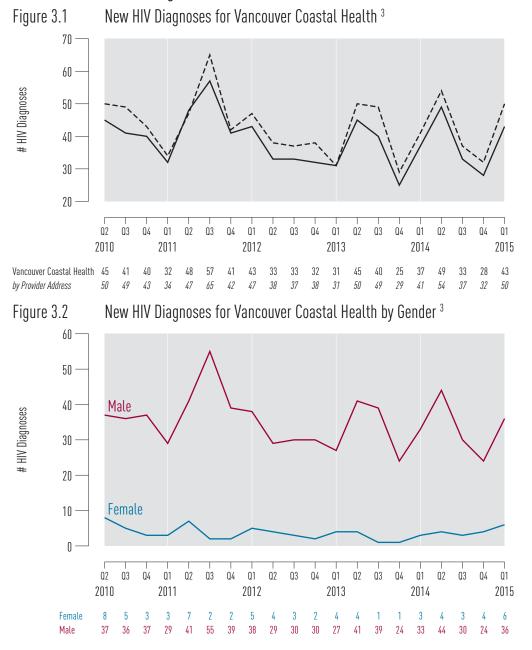


*Testing does not include point of care tests.* 

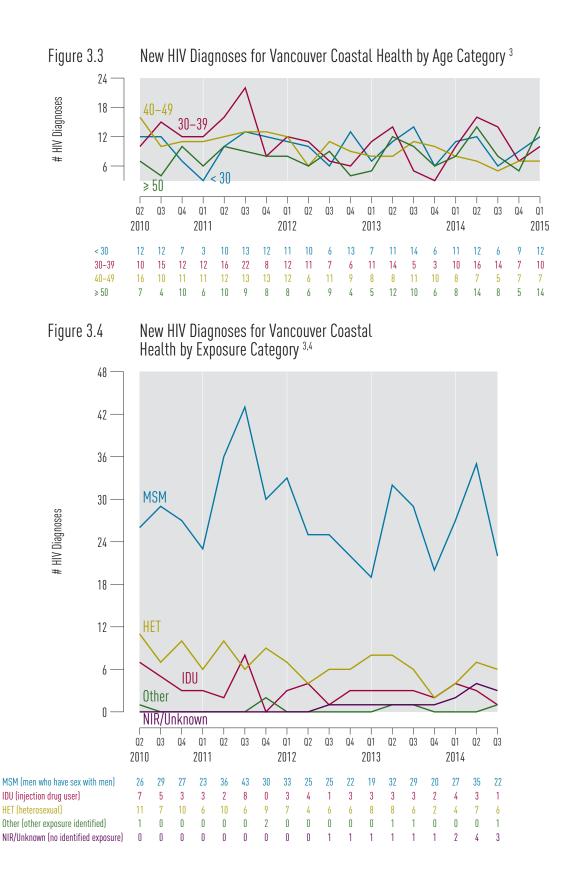
## New HIV Diagnoses

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.



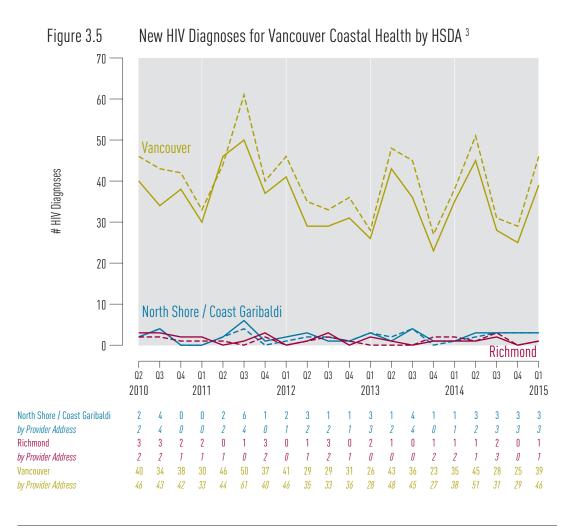


3 Data Source: BCCDC. "By Provider Address" is graphed as dashed line in same colour.



3 Data Source: всссос. "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.



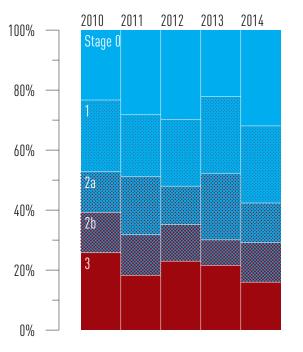
3 Data Source: BCCDC. "By Provider Address" is graphed as dashed line in same colour.

# Stage of HIV infection at diagnosis

Classification of stage of HIV infection, in the absence of information regarding recent testing history, is reliant on clinical information available at the time of diagnosis, including first CD4+ cell count, laboratory results suggestive of acute HIV infection, and clinical presentation with an AIDS-defining illness (Table 1). The benefits of Treatment as Prevention (TasP) are maximized when antiretroviral therapy (ART) is initiated at high CD4 cell counts. Accordingly, it is preferable that individuals newly diagnosed with HIV be in the early stages of HIV infection (stage o 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 Figure 4.2 Vancouver Coastal Health, 2010–2014 <sup>5</sup>

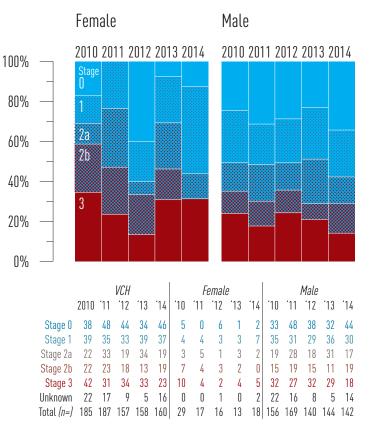


## Indicator 4. Stage of HIV Infection at Diagnosis

# Table 1Staging Classifications of Infection at Time<br/>of HIV Diagnosis Based on CDC HIV<br/>Surveillance Case Definitions

Stage	Criteria										
0	previous r	negativ	ria met for acute ve or indeterminat firmed positive H	e HIV i	test within 180						
1			CD4 ≥500								
2a			CD4 350-499	and	No AIDS case report						
2b	Stage O		CD4 200-349		ισμοιτ						
3	not met	and	( CD4 <200	or	AIDS case ) report						
Unknown	wn No available and No AIDS case CD4 report										

Stage of HIV Infection at Diagnosis by Gender for Vancouver Coastal Health, 2010–2014 <sup>5</sup>



5 Data Source: BCCDC

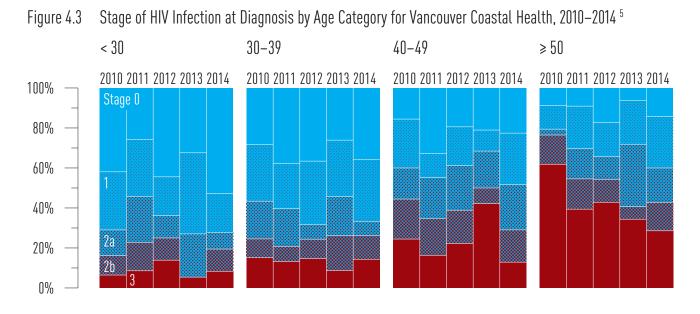
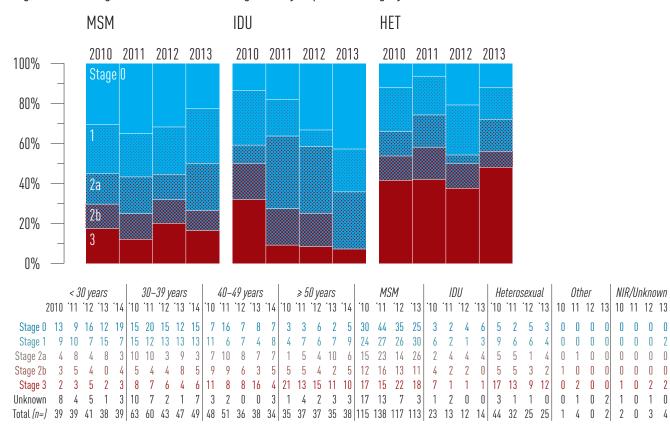


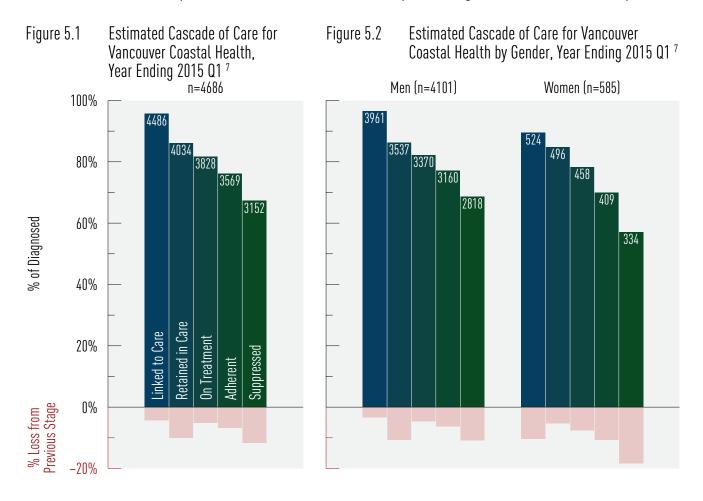
Figure 4.4 Stage of HIV Infection at Diagnosis by Exposure Category for Vancouver Coastal Health, 2010–2013<sup>5,6</sup>



<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. Linked to HIV care, 3. Retained in HIV care, 4. On ART, 5. Adherent to ART and 6. Achieving a suppressed VL; collectively, they are referred to as the cascade of care. Leakage between any of these stages of HIV-care means a reduction in the potential of ART as a benefit to the HIV-positive individual and as an HIV transmission prevention method on a population level. Thus, when interpreting trends in the cascade of care, we strive to see increases along each step of the cascade of care (i.e. reduced attrition) with the ultimate goal being 100% within each stage of the cascade. Monitoring the Cascade of Care provides a picture as to where deficiencies lie in the delivery and uptake of HIV-care. In this section we present the cascade of care for the year 2012 in BC overall and stratified by sex and age for each Health Authority.



7 Data is for the period 2014 Q2–2015 Q1.

- Data Sources:
- *i* British Columbia Centre for Excellence Drug Treatment Program (DTP) Database (ARV use, VL and CD4 count).

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

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

NB: Transgender has been assigned to their biological sex.

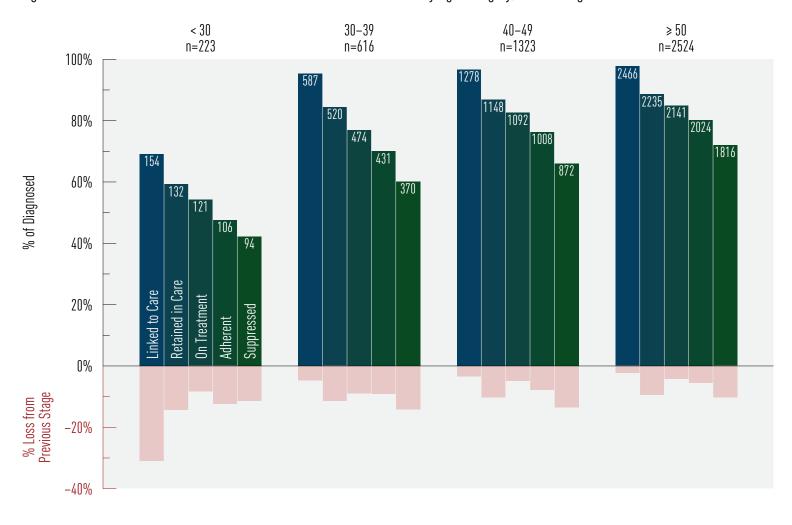


Figure 5.3 Estimated Cascade of Care for Vancouver Coastal Health by Age Category, Year Ending 2015 Q1 <sup>8</sup>

8 Data is for the period 2014 Q2–2015 Q1.

Data Sources:

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

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

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

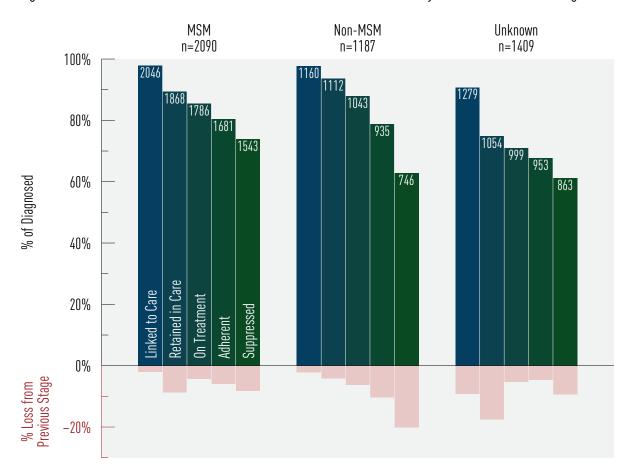


Figure 5.4 Estimated Cascade of Care for Vancouver Coastal Health by MSM Status, Year Ending 2015 Q1 <sup>9</sup>

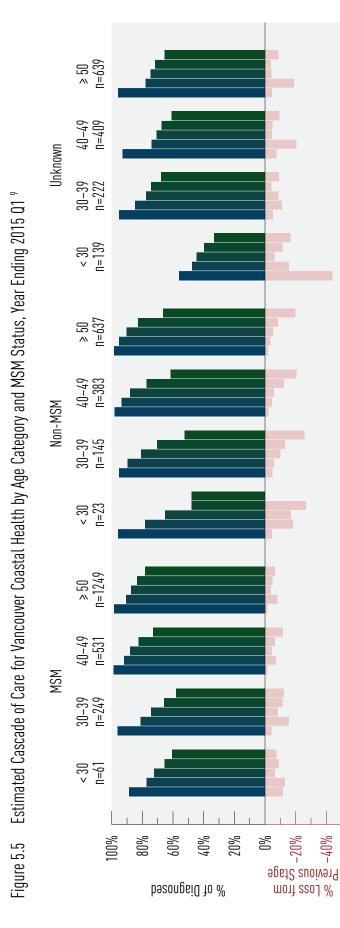
9 *Data is for the period 2014 Q2–2015 Q1.* 

#### Data Sources:

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

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

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



6

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)). ::1

Limitations: HA assignment is based on the most recent HA of residence of the patient, if not available of the HIV-care provider. If the most recent HA of residence is not updated then the designated HA may be incorrect. Recent updates to the DTP database have allowed for more comprehensive information on HIV risk group category. As a result, 2014 Q4 data may differ significantly from preceding reports in terms of total numbers ascribed to each risk group.

Data is for the period 2014 Q2–2015 Q1.

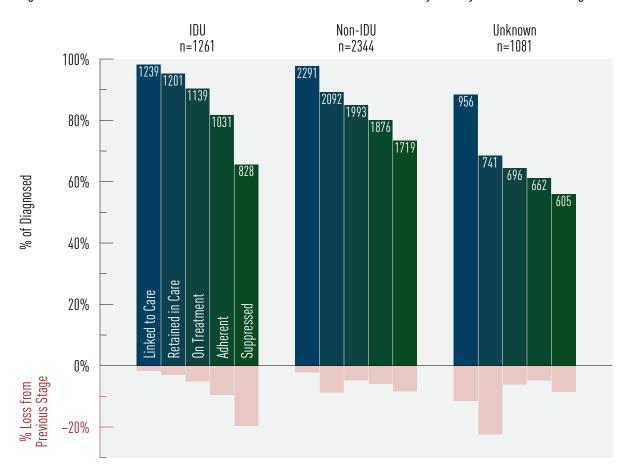


Figure 5.6 Estimated Cascade of Care for Vancouver Coastal Health by History of IDU, Year Ending 2015 Q1 <sup>9</sup>

9 *Data is for the period 2014 Q2–2015 Q1.* 

#### Data Sources:

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

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

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

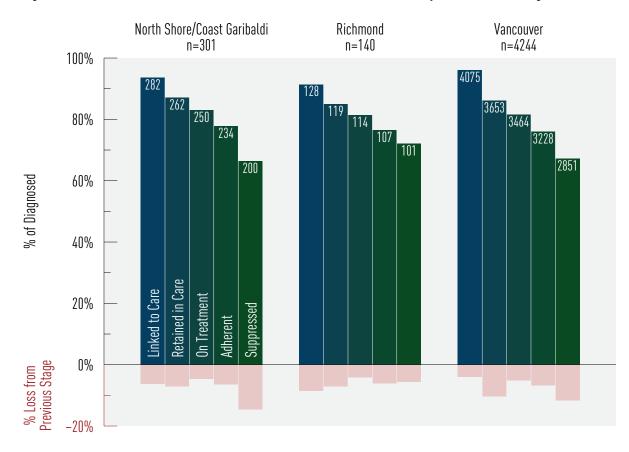


Figure 5.7 Estimated Cascade of Care for Vancouver Coastal Health by HSDA, Year Ending 2015 Q1 <sup>9</sup>

9 *Data is for the period 2014 Q2–2015 Q1.* 

Data Sources:

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

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

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

### 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/ $\mu$ L; 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)
0 (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* 

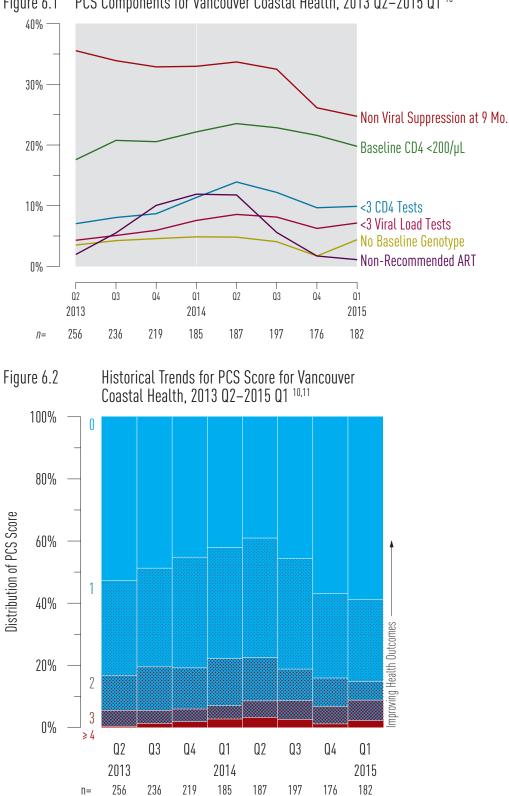


Figure 6.1 PCS Components for Vancouver Coastal Health, 2013 Q2-2015 Q1 <sup>10</sup>

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

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. 11 NB: A score of o is the best score and a score of 4 or more is the worst score.

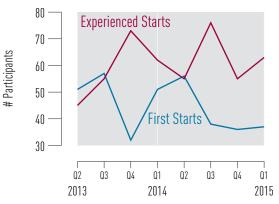
# Antiretroviral Uptake

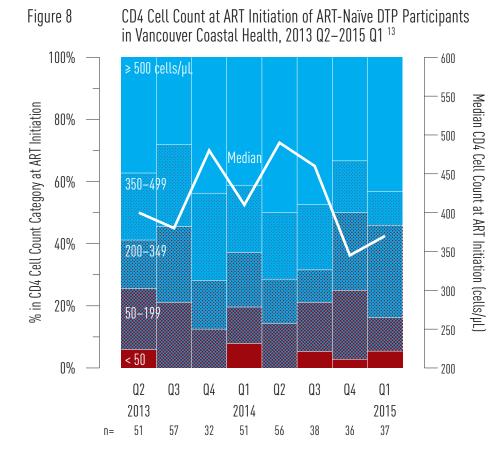
In this section we present trends in ART uptake, the number and proportion of new HIV treatment initiations and the number of active and inactive DTP participants. Trends in ART uptake should be interpreted under the consideration of changing BC HIV treatment guidelines. BC HIV treatment guidelines are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

## Indicator 8. CD4 Cell Count at ART Initiation

## Indicator 7. New Antiretroviral Therapy Starts in Vancouver Coastal Health

Figure 7 BC-CfE Drug Treatment Program Enrollment: New ART Participants in Vancouver Coastal Health, 2013 Q2–2015 Q1 <sup>12</sup>





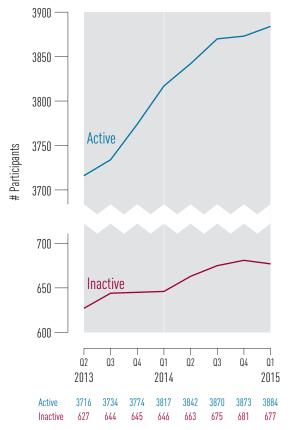
- 12 Data Source: Drug Treatment Program Database Limitation: DTP participants are designated to an HA based on most current residence provided by the participant.
- 13 Data Source: Drug Treatment Program Database Limitations: CD4 cell count data is approximately 80% complete.

## Indicator 9. Active and Inactive DTP Participants

Table 3. Distribution of People on ART for Vancouver Coastal Health, 2015 Q1 <sup>14</sup>

Age	< 30	148
	30-39	507
	40-49	1157
	≥ 50	2072
Gender	Male	3424
	Female	460
Exposure	MSM	1824
	IDU	1126
Total		3884





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

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

#### Definitions:

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

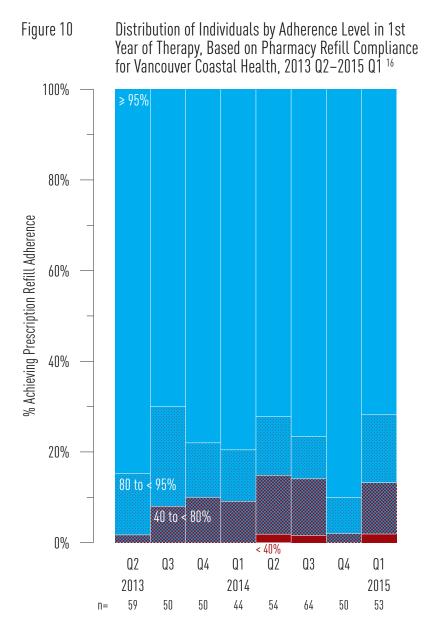
'Unknown/not stated' defined as being on treatment in the current quarter, and city of residence unknown

15 Active DTP participants: An individual who has had medication prescribed at least once in the preceding quarter. Inactive DTP participants: Persons no longer prescribed drugs through the HIV/AIDS Drug Treatment Program in the last quarter.

# Antiretroviral Adherence Level

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

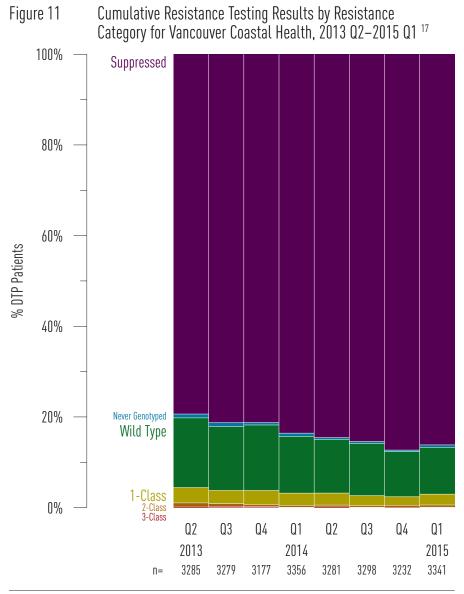




<sup>16</sup> Data Source: Drug Treatment Program Database Limitation: Prescription refill adherence is used as a proxy for patient adherence.

## 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.

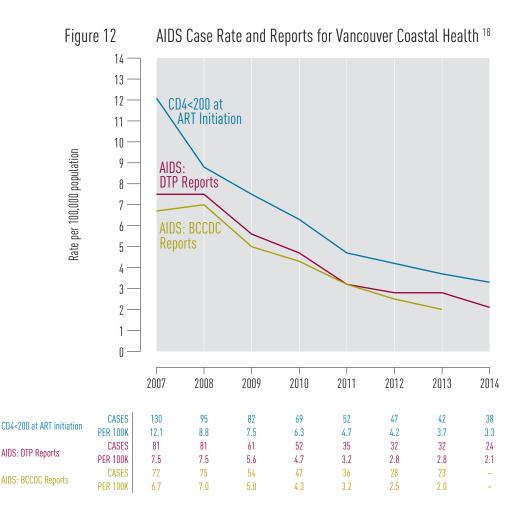


17 Data Source: Drug Treatment Program Database

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

## Indicator 12. AIDS-Defining Illness

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

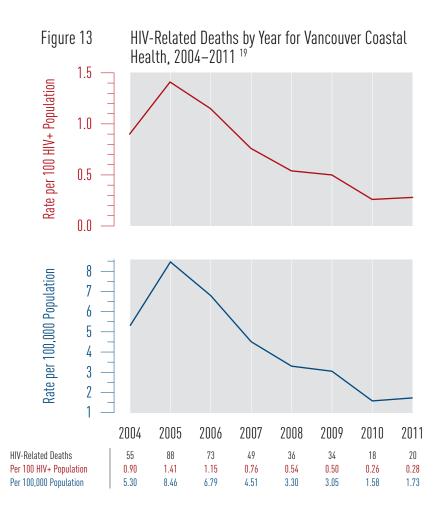


18 Data Source: DTP AIDS cases are obtained from the Drug Treatment Program Database; BCCDC AIDS cases are obtained from the BC-CDC; CD4<200 at ART initiation data came from the DTP database.

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/ $\mu$ L 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 HIVrelated mortality in British Columbia.



19 Data Source: BC Vital Statistics

Limitation:

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 Episodes (	1: Test (thousands)	2010 O2	Q3	04	2011 O1	Q2	Q3	Q4	2012 O1	2 02	Q3	Q4	2013 Q1	3 O2	Q3	Q4	2014 Q1	4 O2	03	Q4	2015 Q1
	Coastal Health	<u> </u>	<u> </u>	19.3				<u> </u>		<u> </u>	<u> </u>	<u> </u>	<u> </u>	<u> </u>	<u> </u>	<u> </u>	<u> </u>	<u> </u>	46.7	<u> </u>	<u> </u>
Gender	Female	9.7	10.0	9.8	10.6	9.7	10.7	11.6	13.6	13.5	14.7	14.8	17.2	18.3	17.7	17.2	19.7	21.6	23.8	23.3	25.8
	Male	8.2	8.2	8.2	8.7	8.0	9.0	9.6	11.3	11.3	12.4	12.6	14.3	15.3	14.4	14.2	16.8	18.3	20.4	19.9	21.8
	Other	0.4	0.4	0.4	0.4	0.4	0.4	0.4	0.3	0.2	0.2	0.3	0.2	0.2	0.2	0.1	0.2	0.2	0.2	0.1	0.2
Female (Pr	renatal)	3.1	3.4	3.5	3.7	3.4	3.5	3.7	4.1	3.7	3.6	3.6	3.6	3.5	3.7	3.7	3.9	3.8	3.7	3.7	4.0
Female (No	on-prenatal)	6.6	6.6	6.3	6.8	6.4	7.1	8.0	9.5	9.8	11.1	11.2	13.6	14.8	14.0	13.6	15.8	17.9	20.0	19.6	21.8
Age	< 30	6.0	6.3	6.1	6.1	5.9	6.7	6.8	7.3	7.2	8.1	8.0	8.5	9.0	9.2	9.1	9.6	9.9	11.1	10.8	11.0
	30-39	6.1	6.2	6.2	6.7	5.9	6.4	6.5	7.8	7.4	7.7	7.6	8.8	9.0	8.9	8.6	10.2	10.1	10.7	10.1	11.5
	40-49	2.9	2.8	2.8	3.1	2.8	3.0	3.2	3.9	3.8	3.9	4.1	5.2	5.5	5.0	4.9	5.7	6.0	6.5	6.3	7.0
	≥ 50	2.4	2.4	2.3	2.8	2.6	3.0	3.4	4.6	4.8	6.0	6.4	9.1	10.2	9.0	9.0	11.0	14.0	16.0	16.0	18.1
POC HIV		0.0	0.0	0.9	1.0	2.0	5.9	2.4	2.1	1.9	2.1	1.7	2.0	1.8	2.1	1.9	2.1	2.0	2.4	2.0	1.5
North Shor / Coast Ga		2.3	2.4	2.4	2.4	2.3	2.4	2.4	2.8	2.6	2.7	2.8	3.0	3.2	3.2	3.3	3.7	4.9	5.4	5.9	6.4
Female (	(Non-prenatal)	0.9	0.9	0.9	0.9	0.9	0.9	0.9	1.1	1.0	1.1	1.2	1.3	1.4	1.3	1.3	1.6	2.2	2.5	2.8	3.1
Male		0.9	0.8	0.9	0.9	0.9	0.9	0.9	1.0	1.0	1.0	1.1	1.2	1.3	1.3	1.3	1.5	2.1	2.2	2.5	2.7
Richmond		1.2	1.2	1.2	1.3	1.2	1.2	1.3	1.5	1.5	1.5	1.6	1.9	1.9	1.9	1.8	3.1	5.4	5.4	5.2	5.6
Female (	(Non-prenatal)	0.4	0.4	0.4	0.4	0.4	0.4	0.4	0.5	0.5	0.6	0.6	0.7	0.7	0.7	0.7	1.3	2.6	2.6	2.5	2.7
Male		0.4	0.4	0.4	0.4	0.4	0.4	0.5	0.5	0.5	0.5	0.6	0.7	0.7	0.7	0.7	1.2	2.2	2.3	2.2	2.4
Vancouver		14.8	15.1	15.7	17.0	16.6	22.5	20.2	22.9	22.9	25.2	25.0	28.7	30.6	29.3	28.4	32.1	31.8	36.0	34.3	37.3
Female (	Non-prenatal)	5.3	5.4	5.0	5.5	5.0	5.8	6.6	7.9	8.3	9.5	9.4	11.6	12.7	11.9	11.5	12.9	13.0	15.0	14.3	16.0
Male		6.9	6.9	6.9	7.4	6.8	7.8	8.2	9.8	9.9	10.9	11.0	12.4	13.3	12.4	12.3	14.1	14.1	15.9	15.1	16.7

#### Indicator 2: Rate of HIV Testing per 100,000

		2009	2010	2011	2012	2013	2014
Vancouver	Coastal Health	5242.3	5259.0	5554.7	7325.4	9221.7	11508.4
North Shor	e / Coast Garibaldi	3500.8	3540.2	3650.3	4182.7	4760.5	7044.0
Richmond		2590.2	2697.6	2794.2	3410.2	3955.7	9186.0
Vancouver		6816.6	6802.0	7233.2	9880.8	12734.2	14133.2
Gender	Female	5547.5	5534.5	5877.9	7828.9	10036.9	12342.1
	Male	4603.9	4677.7	4913.5	6628.6	8275.5	10525.4
Age	< 30	5052.7	5037.2	5234.6	6304.7	7442.6	8602.9
	30-39	11235.3	11585.5	11909.5	14080.8	17109.6	19295.4
	40-49	5017.8	4961.8	5233.1	6863.1	9707.7	11681.3
	≥ 50	1846.5	1913.2	2197.5	4206.9	7289.8	10769.4

			)		2011				2012	2			2013				2014	-		2	2015
Indicator 3: New HIV	Diagnoses	Q2	Q3	Q4	Q1																
Vancouver Coastal	By Client Residence	45	41	40	32	48	57	41	43	33	33	32	31	45	40	25	37	49	33	28	43
Health	By Provider Address	50	49	43	34	47	65	42	47	38	37	38	31	50	49	29	41	54	37	32	50
Gender	Female	8	5	3	3	7	2	2	5	4	3	2	4	4	1	1	3	4	3	4	6
	Male	37	36	37	29	41	55	39	38	29	30	30	27	41	39	24	33	44	30	24	36
Age	< 30	12	12	7	3	10	13	12	11	10	6	13	7	11	14	6	11	12	6	9	12
	30-39	10	15	12	12	16	22	8	12	11	7	6	11	14	5	3	10	16	14	7	10
	40-49	16	10	11	11	12	13	13	12	6	11	9	8	8	11	10	8	7	5	7	7
	≥ 50	7	4	10	6	10	9	8	8	6	9	4	5	12	10	6	8	14	8	5	14
Exposure	MSM	26	29	27	23	36	43	30	33	25	25	22	19	32	29	20	27	35	22	_	-
	IDU	7	5	3	3	2	8	0	3	4	1	3	3	3	3	2	4	3	1	_	-
	HET	11	7	10	6	10	6	9	7	4	6	6	8	8	6	2	4	7	6	_	-
	Other	1	0	0	0	0	0	2	0	0	0	0	0	1	1	0	0	0	1	_	-
	NIR/Unknown	0	0	0	0	0	0	0	0	0	1	1	1	1	1	1	2	4	3	-	-

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Richmond				1			idenc		-		2	0	1		-	0	1 3	3 0	2	-	0		1	1	1	2	0	1
							ddres		2 2	1	1	1	0		2	0	1 2	2 1	0	0	0		2	2	1	3	0	1
Vancouver				'			idenc		) 34	38	30	46	50	3	74	1 2	9 29	31	26	43	36	23	3 3	35 4	45	28	25	39
				By F	rovi	der A	ddres	s 40	5 43	42	33	44	61	4	0 4	6 3	5 33	3 36	28	48	45	22	7 3	38 5	51	31	29	46
Indicator 4:	Stag	ge of	f HI	V In	fect	ion a	t Bas	eline																				
		1	VCE					nale				ale					years			30-3					40-4			
	<b>'</b> 10	'11	'12	'13	'14	<b>'</b> 10	<b>'</b> 11 <b>'</b> 1	2 '13	'14	<b>'</b> 10	ʻ11 ʻ	12'1	3 '1	4 '	10 '	11'1	2 '13	3 '14	<b>'</b> 10	ʻ11 ʻ	12 '	13	'14	'10	<u>'11</u>	'12	'13	'14
Stage 0	38	48	44	34	46	5	0	6 1	2	33	48	38 3	32 4	4	13	9 1	6 12	2 19	15	20	15	12	15	7	16	7	8	7
Stage 1	39	35	33	39	37	4	4	3 3	7	35	31	29 3	36 3	0	9	10	7 15	5 7	15	12	13	13	13	11	6	7	4	8
Stage 2a	22	33	19	34	19	3	5	1 3	2	19	28	18 3	31 1	7	4	8	4 8	3 3	10	10	3	9	3	7	10	8	7	7
Stage 2b	22	23	18	13	19	7	4	3 2	2 0	15	19	15 1	11 1	9	3	5	4 (	) 4	5	4	4	8	5	9	9	6	3	5
Stage 3	42	31	34	33	23	10	4	2 4	5	32	27	32 2	29 1	8	2	3	5 2	2 3	8	7	6	4	6	11	8	8	16	4
Unknown	22	17	9	5	16	0	0	1 (	) 2	22	16	8	5 1	4	8	4	5	13	10	7	2	1	7	3	2	0	0	3
Total	185	187	157	158	160	29	17	16 13	8 18	1561	1691	40 14	44 14	2	39	<b>39</b> 4	1 38	3 39	63	60	43	47	49	48	51	36	38	34
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Stage 0	3	3	6	2	5	30	44	35	25	3	2	4	1	6	5	2	5	3	0	0	)	0	0	0		0	0	0
Stage 1	4	7	6	7	9	24	27	26	30	6	2	: 1	L	3	9	6	6	4	0	0	)	0	0	0		0	0	2
Stage 2a	1	5	4	10	6	15	23	14	26	2	4	. 4	1	4	5	5	1	4	0	1	L	0	0	0		0	0	0
Stage 2b	5	5	4	2	5	12	16	13	11	4	2	2	2	0	5	5	3	2	1	C	)	0	0	0		0	0	0
Stage 3	21	13	15	11	10	17	15	22	18	7	1	1	L	1	17	13	9	12	0	2	2	0	0	1		0	2	2
Unknown	1	4	2	3	3	17	13	7	3	1	2	(	)	0	3	1	1	0	0	1	L	0	2	1		0	1	0
Total	35	37	37	35	38	115	138	117	113	23	13	12	2 1	4	44	32	25	25	1	4	ł	0	2	2	(	0	3	4
Tu di sets u F				J	60.																							
Indicator 5					r Ca	re		DIAG	NOSE			LINI			RE	TAIN				ART		ADI			S	SUPP	PRESS	
Vancouver				ICN					468				486				)34		ć	3828				569			3	152
Age Catego	ory									23			154				32			121				106				94
		30-	39						6	16			587			5	520			474				431				370
		40-	49						132	23		1	278			11	48		1	092			1	008			8	872

Age Category	< 30		223	154	132	121	106	94
	30-39		616	587	520	474	431	370
	40-49		1323	1278	1148	1092	1008	872
	≥ 50		2524	2466	2235	2141	2024	1816
Age Category	MSM	< 30	61	54	47	44	40	37
and MSM		30-39	249	239	202	185	164	144
Status		40-49	531	524	487	466	437	387
		≥ 50	1249	1229	1132	1091	1040	975
	Non-MSM	< 30	23	22	18	15	11	11
		30-39	145	138	130	117	102	76
		40-49	383	375	358	337	296	236
		≥ 50	637	626	605	574	526	423
	Unknown	< 30	139	78	66	62	55	46
		30-39	222	211	188	172	165	150
		40-49	409	379	302	289	275	249
		≥ 50	639	611	497	476	458	418
Gender	Male		4101	3961	3537	3370	3160	2818
	Female		585	524	496	458	409	334

Indicator 5: F	HIV Cascade of Care	DI	AGNOSED	LINKED	RETAINED	0	N ART	ADHERENT	SUPPRESSED
Injection	IDU		1261	1239	1201		1139	1031	828
Drug Use	Non-IDU		2344	2291	2092		1993	1876	1719
	Unknown		1081	956	741		696	662	605
MSM Status	MSM		2090	2046	1868		1786	1681	1543
	Non-MSM		1187	1160	1112		1043	935	746
	Unknown		1409	1279	1054		999	953	863
Health Authority	North Shore / Coast Garibaldi		301	282	262		250	234	200
	Richmond		140	128	119		114	107	101
	Vancouver		4244	4075	3653		3464	3228	2851
Indicator 6: P	Programmatic Comp	liance Scor	e (PCS)						
	0	2013			2014				2015
		Q2	Q3	Q4	Q1	Q2	Q3		
< 3 CD4 Tests		7.0%	8.1%	8.7%	11.4%	13.9%	12.2%		9.9%
< 3 Viral Load		4.3%	5.1%	5.9%	7.6%	8.6%	8.1%		7.1%
No Baseline C		3.5%	4.2%	4.6%	4.9%	4.8%	4.1%		4.4%
	4 < 200 cells/μL	17.6%	20.8%	20.5%	22.2%	23.5%	22.8%		19.8%
Non-Recomn		2.0%	5.5%	10.0%	11.9%	11.8%	5.6%		1.1%
_	ppression at 9 Mo.	35.5%	33.9%	32.9%	33.0%	33.7%	32.5%		24.7%
PCS Score: 0		135	115	99	78	73	90		107
PCS Score: 1		78	75	78	66	72	70	48	48
PCS Score: 2		29	33	29	28	26	20		11
PCS Score: 3		13	10	9	8	10	12		12
PCS Score: 4	or more	1	3	4	5	6	5	2	4
Total (n=)		256	236	219	185	187	197	176	182
	New DTP ARV Partic	<u>^</u>							
First Starts		51	57	32	51	56	38		37
Experienced S	Starts	45	55	73	62	55	76	55	63
Indicator 8: C	CD4 Cell Count at A	RT Initiatio	on for ARV-N	aïve DTP Par	ticipants				
CD4 ≥ 500		19	16	14	21	28	18	12	16
CD4 350-499		11	15	9	11	12	8	6	4
CD4 200-349	9	8	14	5	9	8	4	. 9	11
CD4 50-199		10	12	4	6	8	e	8	4
CD4 < 50		10 3	12 0	4 0	6 4	8 0	e 2		
CD4 < 50 CD4 Median	(cells/µL)	3 400		0 480	4 410		2 460	1 345	2
CD4 < 50	(cells/µL)	3	0	0	4	0	2	1 345	2 <i>370</i>
CD4 < 50 CD4 Median Total (n=) Indicator 9: A	Active and Inactive D	3 400 51	0 <i>380</i> 57	0 480	4 410	0 490	2 460	1 345	2 370
CD4 < 50 CD4 Median Total (n=)	Active and Inactive D	3 400 51	0 <i>380</i> 57	0 480	4 410	0 490	2 460	2 1 345 36	2 370 37
CD4 < 50 CD4 Median Total (n=) Indicator 9: A	Active and Inactive D Participants	3 400 51 0 <b>TP Partici</b>	0 380 57 pants	0 480 32	4 410 51	0 490 56	2 460 38	2 1 345 36 3873	2 370 37 3884
CD4 < 50 CD4 Median Total (n=) Indicator 9: A Active DTP P Inactive DTP	Active and Inactive D Participants	3 400 51 7TP Partici 3716 627	0 380 57 pants 3734	0 480 32 3774	4 410 51 3817	0 490 56 3842	2 460 38 3870	2 1 345 36 3873	2 <i>370</i> 37 3884
CD4 < 50 CD4 Median Total (n=) Indicator 9: A Active DTP P Inactive DTP	Active and Inactive D Participants Participants	3 400 51 7TP Partici 3716 627	0 380 57 pants 3734	0 480 32 3774	4 410 51 3817	0 490 56 3842	2 460 38 3870	1 345 36 3873 681	2 370 37 3884 677
CD4 < 50 CD4 Median Total (n=) Indicator 9: A Active DTP P Inactive DTP Indicator 10:	Active and Inactive D Participants Participants Antiretroviral Adhe	3 400 51 7TP Partici 3716 627 rence	0 380 57 pants 3734 644	0 480 32 3774 645	4 410 51 3817 646	0 490 56 3842 663	2 460 38 3870 675	2 1 345 36 3873 681	2 370 37 3884 677 3883
CD4 < 50 <i>CD4 Median</i> Total (n=) Indicator 9: A Active DTP P Inactive DTP Indicator 10: ≥ 95%	Active and Inactive D Participants Participants Antiretroviral Adhe	3 400 51 TP Partici 3716 627 rence 50	0 380 57 pants 3734 644 35	0 480 32 3774 645 39	4 410 51 3817 646 35	0 490 56 3842 663 39	2 460 38 3870 675 49	1 345 36 3873 681 9 45 4	2 370 37 3884 677
CD4 < 50 CD4 Median Total (n=) Indicator 9: A Active DTP P Inactive DTP Indicator 10: $$ $\geq 95\%$ 80% to < 95%	Active and Inactive D Participants Participants Antiretroviral Adhe	3 400 51 7TP Partici 3716 627 rence 50 8	0 380 57 pants 3734 644 35 11	0 480 32 3774 645 39 6	4 410 51 3817 646 35 5	0 490 56 3842 663 39 7	2 460 38 3870 675 49 6	1 345 36 3873 681 45 4 4 1	2 370 37 3884 677 388 8

#### Indicator 11: Resistance Testing and Results

	2013 Q2	Q3	Q4	2014 Q1		Q2	Q3	Q4	2015 Q1
Suppressed	2607	2665	2581	2805	2	2774	2817	2822	2879
Wild Type	505	462	459	420		388	380	322	345
Never Genotyped	28	28	17	24		14	14	11	18
1-Class	111	96	98	91		87	74	61	80
2-Class	30	19	15	14		14	11	15	16
3-Class	4	9	7	2		4	2	1	3
Total (n=)	3285	3279	3177	3356	3	281	3298	3232	3341
Indicator 12: AIDS-I	Defining Illness	2007	2008	2009	2010	2011	2012	2013	2014
CD4 < 200 at	Cases	130	95	82	69	52	47	42	38
ART initiation	Rate per 100,000	12.1	8.8	7.5	6.3	4.7	4.2	3.7	3.3
AIDS Cases	Cases	81	81	61	52	35	32	32	24
(DTP Reports)	Rate per 100,000	7.5	7.5	5.6	4.7	3.2	2.8	2.8	2.1
AIDS Cases	Cases	72	75	54	47	36	28	23	-
(BCCDC Reports)	Rate per 100,000	6.7	7.0	5.0	4.3	3.2	2.5	2.0	-
Indicator 13: HIV-Re	lated Mortality	2004	2005	2006	2007	2008	2009	2010	2011
Vancouver Coastal H		55	88	73	49	36	34	18	20
Per 100 HIV+ Popula	ation	0.90	1.41	1.15	0.76	0.54	0.50	0.26	0.28
Per 100,000 Populatio	on	5.30	8.46	6.79	4.51	3.30	3.05	1.58	1.73