

# HIV MONITORING QUARTERLY REPORT

#### FOR FRASER HEALTH

FOURTH QUARTER 2014

\* Please see foreword

















#### Foreword

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

The objectives of these reports are to:

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

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

For Indicator 5 (page 20) and 9 (page 29), 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.

<sup>\*</sup> For Indicator 1, new figures (Figure 1.6 and 1.7 on page 13) have been added to 2014 Q4 report for the following: HIV Test Episodes for Non-prenatal Females in Fraser Health by HSDA and HIV Test Episodes for Males in Fraser Health by HSDA.

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## Acknowledgements and Contributions



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



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

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

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

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

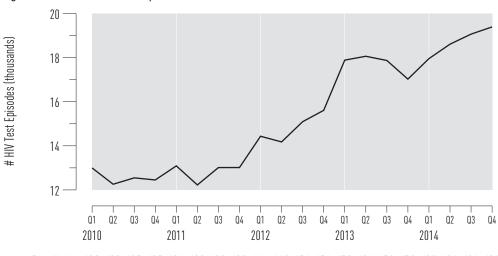
The goals of HA-led STOP-funded initiatives are to work toward achieving these goals. To these ends some outcome measures or indicators of progress have been drafted that should be considered in the design and implementation phases of these initiatives.

## **HIV Testing Episodes and Rates**

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

#### Indicator 1. HIV Testing Episodes

Figure 1.1 HIV Test Episodes for Fraser Health, 2010 Q1-2014 Q4



Fraser Health 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 18.1 17.9 17.0 18.0 18.6 19.1 19.4

Figure 1.2 HIV Test Episodes by Gender and Prenatal Status for Fraser Health, 2010 Q1–2014 Q4  $^{\rm 1}$ 

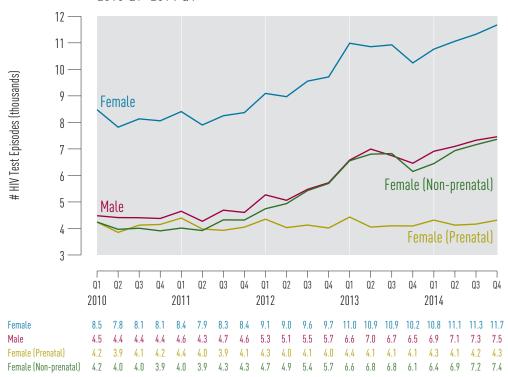
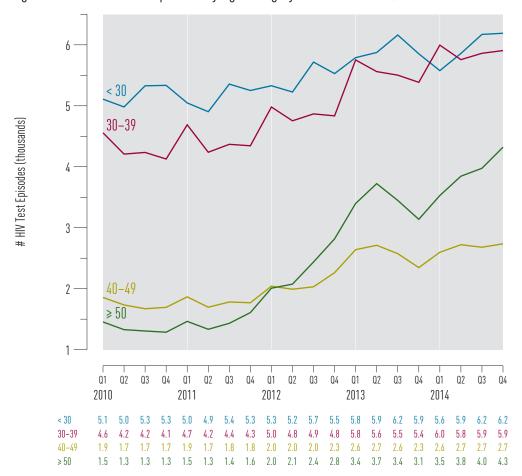


Figure 1.3 HIV Test Episodes by Age Category for Fraser Health, 2010 Q1–2014 Q4 1.2



Point-of-Care HIV Tests for Fraser Health, 2010 Q4-2014 Q4 Figure 1.4 500 # Point-of-Care HIV Tests 400 300 200 100 0 -Q4 Q2 Q3 Q4 Q2 Q3 Q4 Q2 Q3 Q1 Q2 Q3 Q1 Q1 Q1 2010 2011 2012 2013 2014 Fraser Health 12 121 31 158 296 187 182 302 254 426 377 238

#### Limitations:

- *i* Repeat tests in individuals who test using various identifiers may not be identified and these individuals may be counted more than once.
- ii In Fraser Health, POC testing data are available from March 2011 forward.
- Testing does not include point of care tests.

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

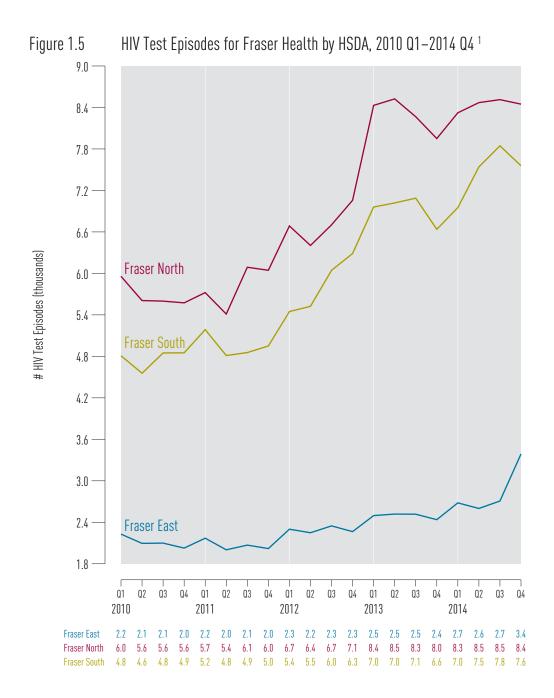
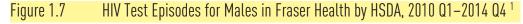
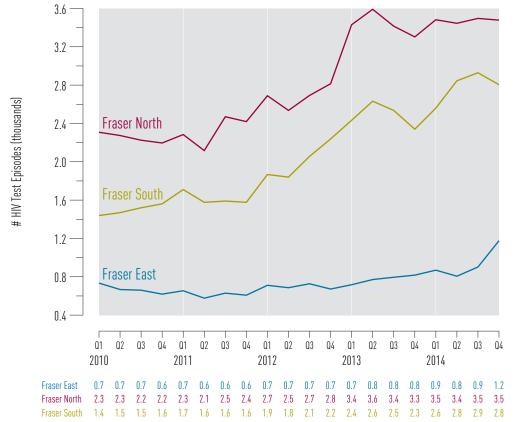


Figure 1.6 HIV Test Episodes for Non-prenatal Females in Fraser Health by HSDA, 2010 Q1–2014 Q4 <sup>1</sup>







## Indicator 2. HIV Testing Rates

Figure 2.1 Rate of HIV Testing for Fraser Health and HSDAs, 2009–2014  $^{\rm 2}$ 

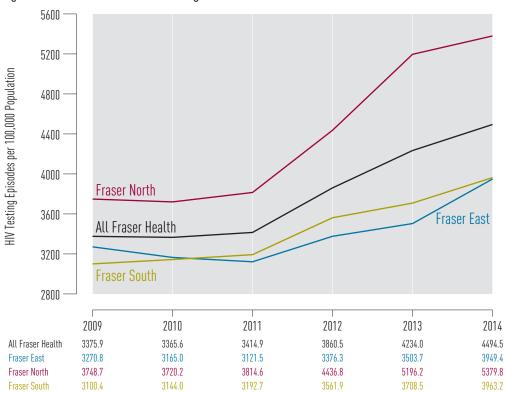
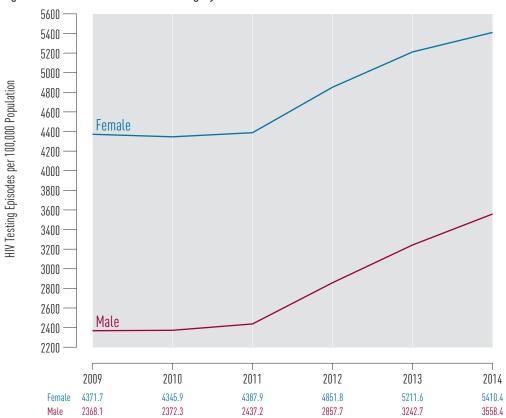
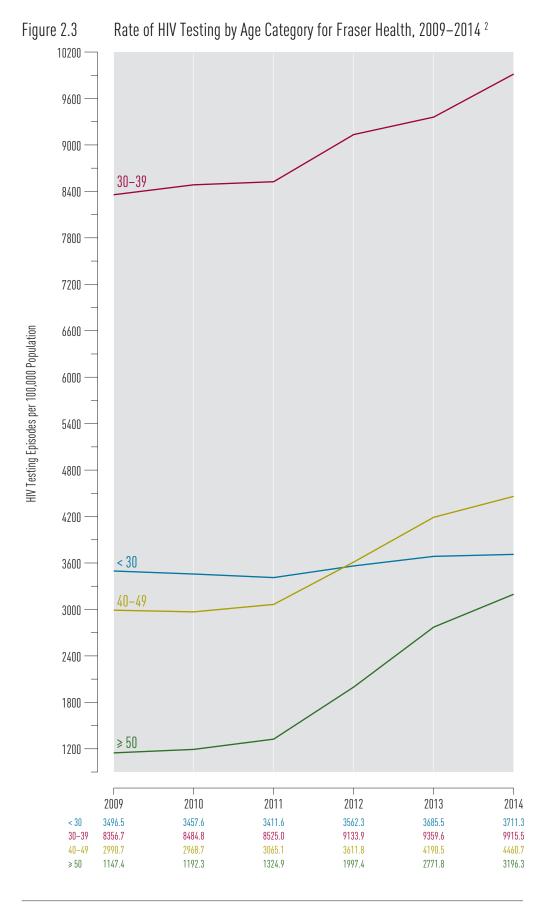


Figure 2.2 Rate of HIV Testing by Gender for Fraser Health, 2009–2014 <sup>2</sup>





<sup>2</sup> Testing does not include point of care tests.

## New HIV Diagnoses

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

#### Indicator 3. New HIV Diagnoses

Figure 3.1 New HIV Diagnoses for Fraser Health, 2010 Q1-2014 Q4 <sup>3</sup>

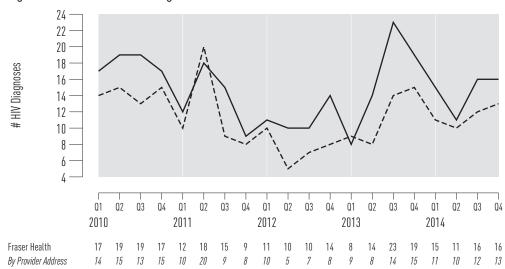


Figure 3.2 New HIV Diagnoses for Fraser Health by Gender, 2010 Q1–2014 Q4 <sup>3</sup>



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

Figure 3.3 New HIV Diagnoses for Fraser Health by Age Category, 2010 Q1-2014 Q4 $^{\rm 3}$ # HIV Diagnoses < 30 6 Q2 Q3 Q4 Q1 Q2 Q3 Q1 Q2 Q3 Q1 Q2 Q3 Q4 Q1 Q2 Q3 2010 2011 2012 2013 2014 < 30 3 5 3 5 5 4 30-39 2 3 3 1 3 2 3 8 6 5 5 4 6 40-49 5 5 5 3 3 ≥ 50 Figure 3.4 New HIV Diagnoses for Fraser Health by Exposure Category, 2010 Q1-2014 Q2 3,4 12 -HET 10 # HIV Diagnoses 8 MSM 6 IDU **Other** NIR/Unknown Q1 Q2 Q2 Q3 Q4 Q3 Q4 Q1 Q2 Q1 Q1 Q1 Q3 Q4 Q2 Q3 Q4 2010 2011 2012 2013 2014 10 MSM (men who have sex with men) 4 3 11 3 0 IDU (injection drug user) 2 4 2 0 0 2 10 HET (heterosexual) 8 4 4 6 Other (other exposure identified) 1 2 0 0 0 0 0 0 2 0 0 0 NIR/Unknown (no identified exposure) Figure 3.5 New HIV Diagnoses for Fraser Health by HSDA, 2010 Q1-2014 Q4 <sup>3</sup> 15 Fraser Fraser South 12 North # HIV Diagnoses 9 6 3 Fraser East Q1 Q3 Q2 Q4 Q1 Q2 Q3 Q4 Q2 Q3 Q2 Q3 Q4 Q1 Q2 Q3 Q1 Q4 Q1 2010 2011 2012 2013 2014 Fraser East by Provider Address 0 2 0 0 0 7 3 9 Fraser North 6 6 6 4 6 5 9 14 8 10 by Provider Address 5 2 5 3 6 4 11 Fraser South 0 by Provider Address

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

<sup>4</sup> MSM=men who have sex with men; IDU= injection drug user; HET=heterosexual. NIR=No identified risk/exposure.

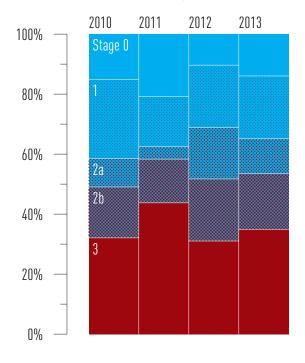
## Stage of HIV infection at diagnosis

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.

for Fraser Health, 2010–2013 <sup>5</sup>

Figure 4.1 Stage of HIV Infection at Diagnosis

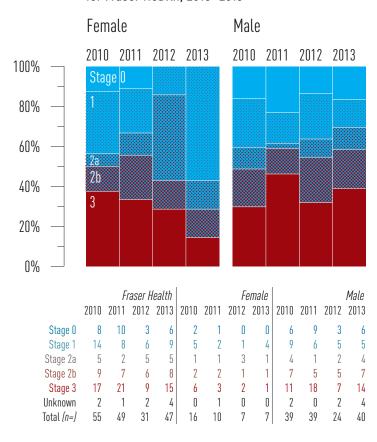


#### Indicator 4. Stage of HIV Infection at Diagnosis

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

Stage	Criteria													
0	previous	Laboratory criteria met for acute HIV infection, or previous negative or indeterminate HIV test within 180 days of first confirmed positive HIV test.												
1			CD4 ≥500		N. AIDC									
2a			CD4 350-499	and	No AIDS case report									
2b	Stage 0		CD4 200-349		торого									
3	not met	and	( CD4 <200	or	AIDS case report									
Unknown		,	No available CD4	and	No AIDS case report									

Figure 4.2 Stage of HIV Infection at Diagnosis by Gender for Fraser Health, 2010-2013 <sup>5</sup>



Data Source: BCCDC

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

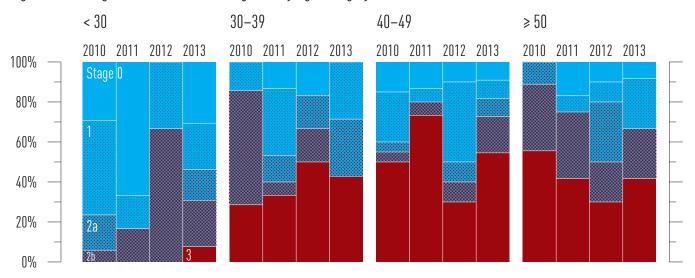
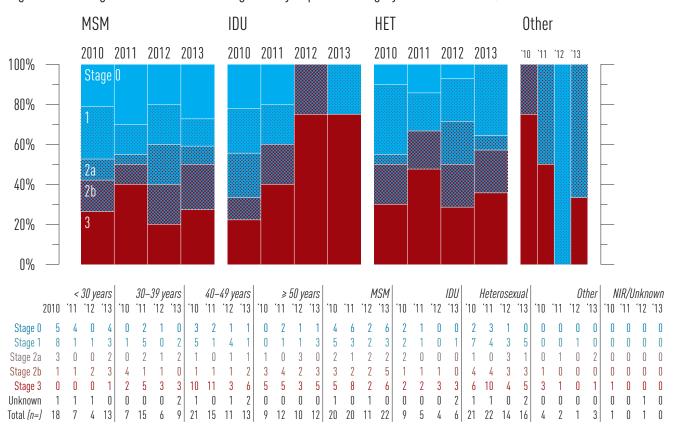


Figure 4.4 Stage of HIV Infection at Diagnosis by Exposure Category for Fraser Health, 2010–2013 5.6



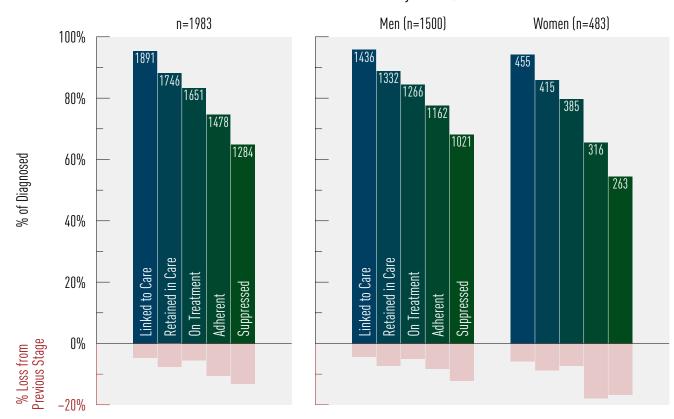
<sup>6</sup> MSM=men who have sex with men; IDU= injection drug user; HET=heterosexual. NIR=No identified risk/exposure.

#### 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. 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.

Figure 5.1 Estimated Cascade of Care for Fraser Health, 2014 7

Figure 5.2 Estimated Cascade of Care for Fraser Health by Gender, 2014 7



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.

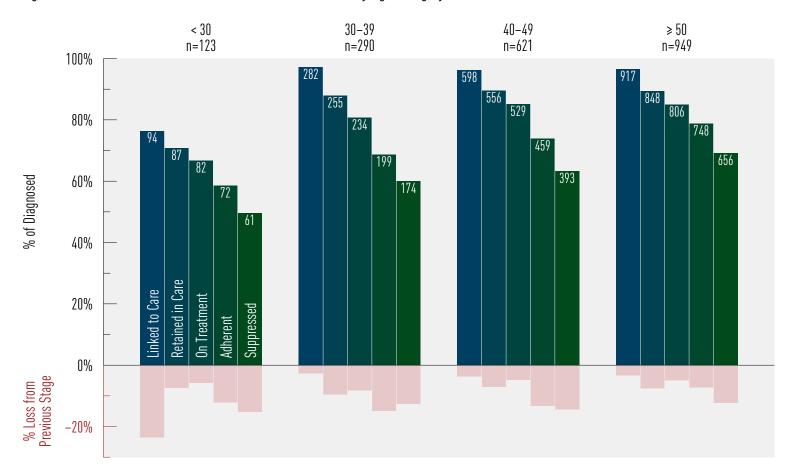
NB: Transgender has been assigned to their biological sex.

<sup>7</sup> Data is for the period 2014 Q1–2014 Q4.

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)).

Figure 5.3 Estimated Cascade of Care for Fraser Health by Age Category, 2014 8



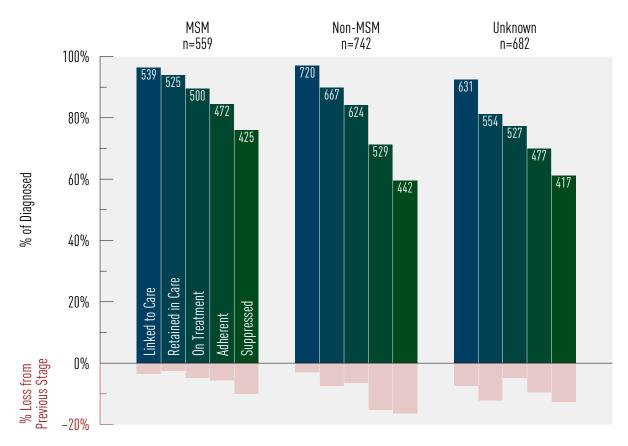
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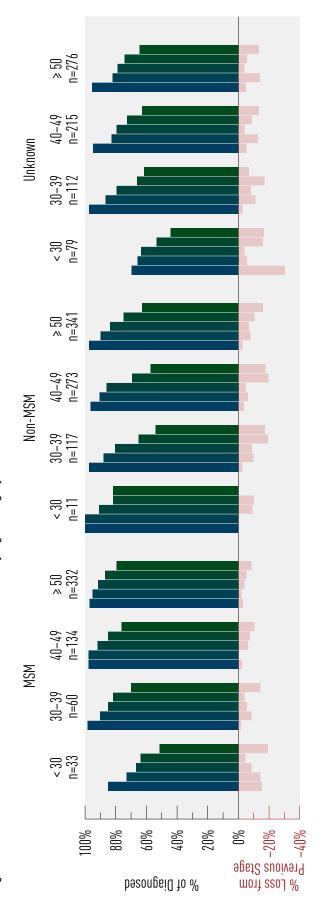
Limitations: HA assignment is based on the most recent HA of residence of the patient, if not available of the HIV-care provider. If the most recent HA of residence is not updated then the designated HA may be incorrect.

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.

<sup>9</sup> Data is for the period 2014 Q1–2014 Q4. Data Sources:

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



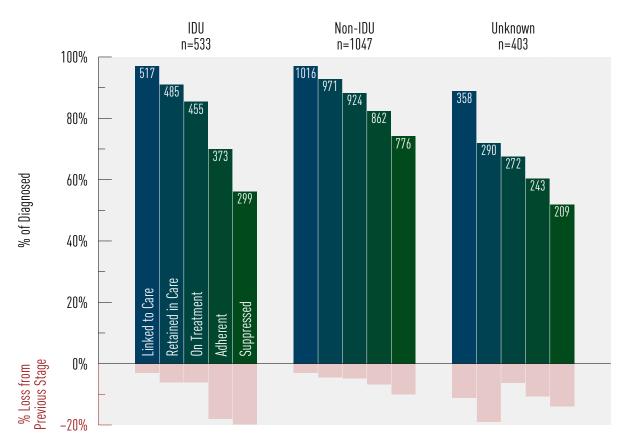
Data is for the period 2014 Q1-2014 Q4. Data Sources:

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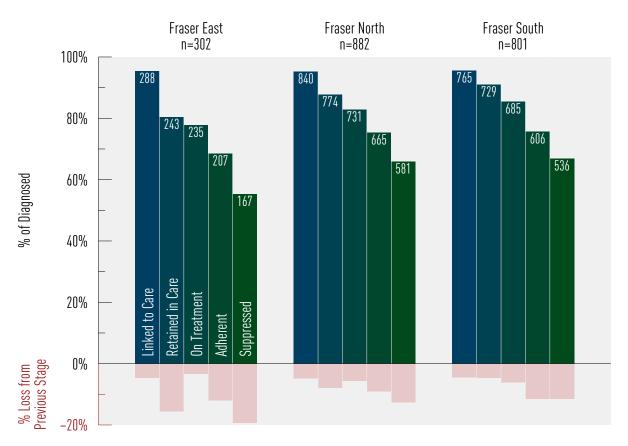
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i British Columbia Centre for Excellence Drug Treatment Program (DTP) Database (ARV use, VL and CD4 count).

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Data Sources:

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

## Indicator 6. The Programmatic Compliance Score (PCS)

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

- having <3 CD4 cell count tests in the first year after starting antiretroviral therapy (ART);
- 2. having <3 plasma viral load (VL) tests in the first year after starting ART;
- not having drug resistance testing done prior to starting ART;
- 4. starting on a non-recommended ART regimen;
- 5. starting therapy with CD4<200 cells/μL; and
- 6. not achieving viral suppression within 9 months since ART initiation.

In this section we provide PCs scores and their components over time for the province of BC. A decline to 0%, (i.e., all individuals having a score of o) is the eventual goal.

Table 2. The Probability of Mortality, Immunologic Failure and Virologic Failure based on the Programmatic Compliance Score

Programmatic Compliance Score	Mortality Risk Ratio (95% Confidence Interval)	Immunologic Failure Risk Ratio (95% CI)	Virologic Failure Risk Ratio (95% CI)
O (Best score)	1 (-)	1 (-)	1 (-)
1	3.81 (1.73-8.42)	1.39 (1.04–1.85)	1.32 (1.05–1.67)
2	7.97 (3.70–17.18)	2.17 (1.54-3.04)	1.86 (1.46–2.38)
3	11.51 (5.28-25.08)	2.93 (1.89-4.54)	2.98 (2.16-4.11)
4 or more (Worst score)	22.37 (10.46–47.84)	9.71 (5.72–16.47)	3.80 (2.52–5.73)

Reference: Lima VD, Le A, Nosyk B, Barrios R, Yip B, et al. (2012) Development and Validation of a Composite Programmatic Assessment Tool for HIV Therapy. PLoS ONE 7(11): e47859. doi:10.1371/journal.pone.0047859

Figure 6.1 PCS Components for Fraser Health, 2013 Q1-2014 Q4  $^{10}$ 

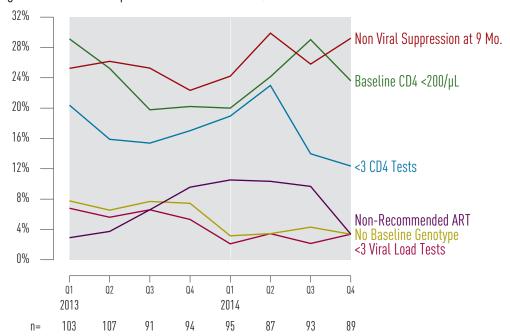
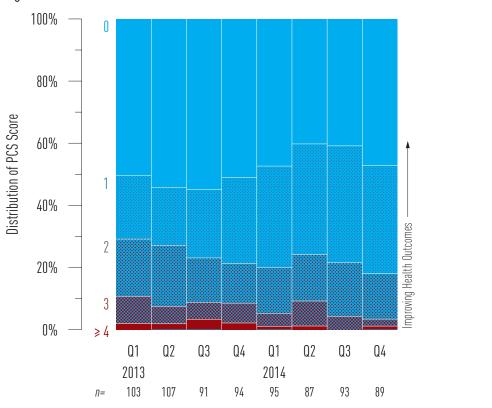


Figure 6.2 Historical Trends for PCS Score for Fraser Health, 2013 Q1-2014 Q4 10,11



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

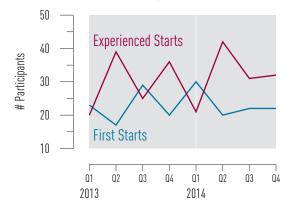
Each quarter's data is calculated as the sum of the 4 quarters leading up to it. e.g. 2013 Q1 is calculated from 2012 Q2 – 2013 Q1. NB: A score of o is the best score and a score of 4 or more is the worst score.

## Antiretroviral Uptake

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

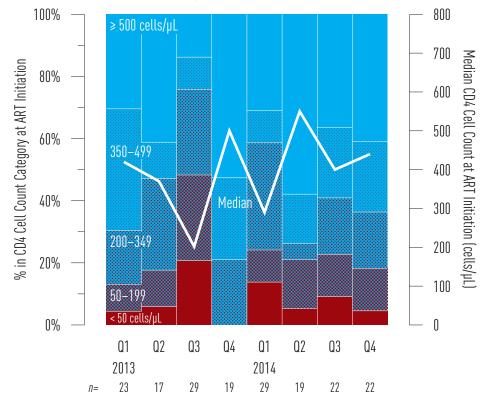
## Indicator 7. New Antiretroviral Therapy Starts in Fraser Health

Figure 7 BC-CfE Drug Treatment Program
Enrollment: New ART Participants in
Fraser Health, 2013 Q1–2014 Q4 12



#### Indicator 8. CD4 Cell Count at ART Initiation

Figure 8 CD4 Cell Count at ART Initiation of ART-Naïve DTP Participants in Fraser Health, 2013 Q1–2014 Q4 <sup>13</sup>



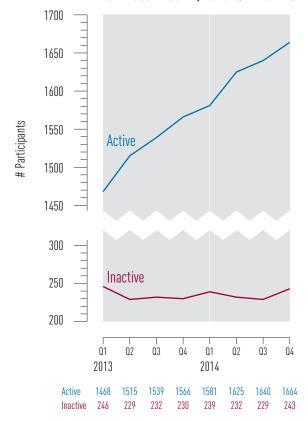
- 2 Data Source: Drug Treatment Program Database Limitation: DTP participants are designated to an HA based on most current residence provided by the participant.
- Data Source: Drug Treatment Program Database Limitations: CD4 cell count data is approximately 80% complete.

## Indicator 9. Active and Inactive DTP Participants

Table 3. Distribution of People on ART for Fraser Health, 2014 Q4  $^{14}$ 

Age	< 30	89
	30-39	250
	40-49	542
	≥ 50	783
Gender	Male	1282
	Female	382
Exposure	MSM	519
	IDU	441
Total		1664

Figure 9 Active and Inactive DTP Participants for Fraser Health, 2013 Q1-2014 Q4 15



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

The recent update in DTP database allows improved classification of some individuals in the risk groups who were previously identified as unknown. 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: are those who are prescribed one or more drugs in the last six months.

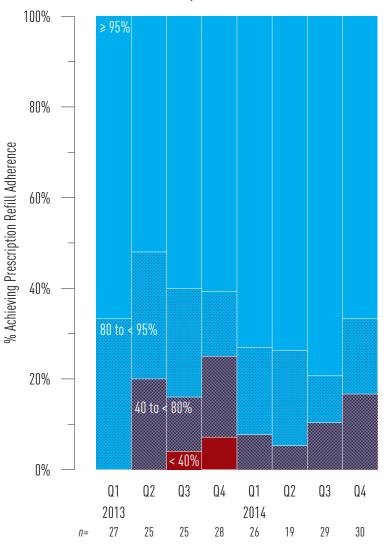
Inactive DTP Participants: Persons no longer prescribed drugs through the HIV/AIDS Drug Treatment Program in the last quarter.

#### Antiretroviral Adherence Level

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

#### Indicator 10. Antiretroviral Adherence

Figure 10 Distribution of Individuals by Adherence Level in 1st Year of Therapy, Based on Pharmacy Refill Compliance for Fraser Health, 2013 Q1–2014 Q4 <sup>16</sup>



<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.

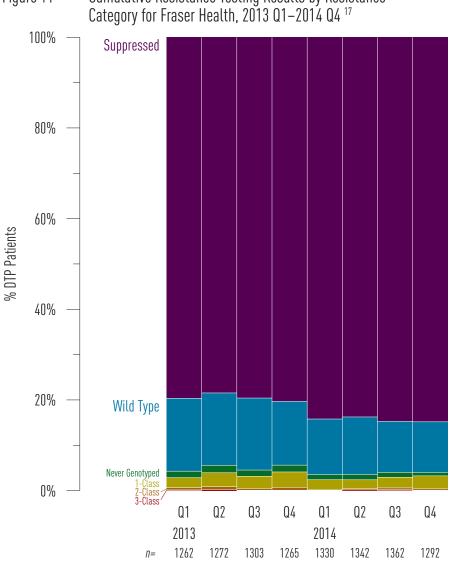


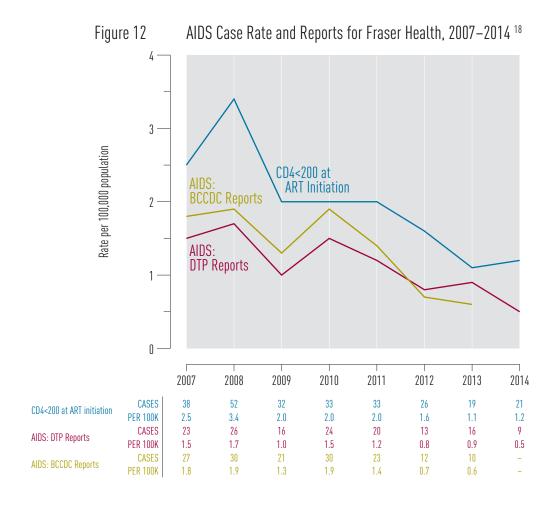
Figure 11 Cumulative Resistance Testing Results by Resistance

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

## Indicator 12. AIDS-Defining Illness

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

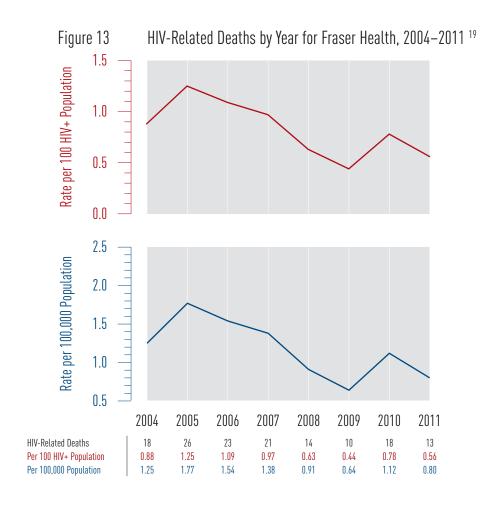


Data Source: DTP AIDS cases are obtained from the Drug Treatment Program Database; BCCDC AIDS cases are obtained from the BCCDC; CD4<200 at ART initiation data came from the DTP database.

Limitation: AIDs case reporting was investigated using 3 definitions: First, using AIDs cases reported in AIDs case report forms from the DTP; Second, using AIDs cases reported via the BCCDC and third, using a CD4 cell count of <200 cells/µL at time of ART initiation using DTP data. AIDs case reporting is passive in BC, thus; AIDs case reporting is not well captured. The DTP sends out AIDs reporting forms to physicians annually. The BCCDC uses DTP AIDs case reports as well as physician AIDs case reports made directly to the BCCDC. Interpreting AIDs case reports should be done with these limitations in mind. AIDs data is updated annually as very few AIDs cases reports are reported in general and trends would be difficult to notice if reported quarterly.

## Indicator 13. HIV-Related Mortality

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



#### Limitation:

<sup>19</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 Episodes	r 1: Test s (thousands)	2010 Q1	O2	О3	04	2011 O1	Q2	O3	Q4	2012 Q1	Q2	Q3	Q4	2013 Q1	Q2	Q3	Q4	2014 Q1	1 O2	Q3	Q4
Fraser He	ealth	13.0	12.3	12.5	12.5	13.1	12.2	13.0	_	14.4	14.2	15.1	15.6		_	17.9	_	_	18.6	_	
Gender	Female	8.5	7.8	8.1	8.1	8.4	7.9	8.3	8.4	9.1	9.0	9.6	9.7	11.0	10.9	10.9	10.2	10.8	11.1	11.3	11.7
	Male	4.5	4.4	4.4	4.4	4.6	4.3	4.7	4.6	5.3	5.1	5.5	5.7	6.6	7.0	6.7	6.5	6.9	7.1	7.3	7.5
	Other	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Female (1	Prenatal)	4.2	3.9	4.1	4.2	4.4	4.0	3.9	4.1	4.3	4.0	4.1	4.0	4.4	4.1	4.1	4.1	4.3	4.1	4.2	4.3
Female (1	Non-prenatal)	4.2	4.0	4.0	3.9	4.0	3.9	4.3	4.3	4.7	4.9	5.4	5.7	6.6	6.8	6.8	6.1	6.4	6.9	7.2	7.4
Age	< 30	5.1	5.0	5.3	5.3	5.0	4.9	5.4	5.3	5.3	5.2	5.7	5.5	5.8	5.9	6.2	5.9	5.6	5.9	6.2	6.2
	30-39	4.6	4.2	4.2	4.1	4.7	4.2	4.4	4.3	5.0	4.8	4.9	4.8	5.8	5.6	5.5	5.4	6.0	5.8	5.9	5.9
	40-49	1.9	1.7	1.7	1.7	1.9	1.7	1.8	1.8	2.0	2.0	2.0	2.3	2.6	2.7	2.6	2.3	2.6	2.7	2.7	2.7
	≥ 50	1.5	1.3	1.3	1.3	1.5	1.3	1.4	1.6	2.0	2.1	2.4	2.8	3.4	3.7	3.4	3.1	3.5	3.8	4.0	4.3
POC HIV	V Tests (not in tho	usands	)		0	12	37	57	24	54	121	31	158	296	187	182	302	254	426	377	238
Fraser Ea	ıst	2.2	2.1	2.1	2.0	2.2	2.0	2.1	2.0	2.3	2.2	2.3	2.3	2.5	2.5	2.5	2.4	2.7	2.6	2.7	3.4
Female	e (Non-prenatal)	0.7	0.6	0.6	0.6	0.6	0.6	0.6	0.6	0.7	0.7	0.8	0.8	0.8	0.9	0.8	0.8	0.9	0.9	0.9	1.3
Male		0.7	0.7	0.7	0.6	0.7	0.6	0.6	0.6	0.7	0.7	0.7	0.7	0.7	0.8	0.8	0.8	0.9	0.8	0.9	1.2
Fraser No	orth	6.0	5.6	5.6	5.6	5.7	5.4	6.1	6.0	6.7	6.4	6.7	7.1	8.4	8.5	8.3	8.0	8.3	8.5	8.5	8.4
Female	e (Non-prenatal)	2.1	2.0	2.0	1.9	1.9	1.9	2.2	2.2	2.4	2.5	2.6	2.8	3.3	3.4	3.4	3.1	3.2	3.5	3.4	3.3
Male		2.3	2.3	2.2	2.2	2.3	2.1	2.5	2.4	2.7	2.5	2.7	2.8	3.4	3.6	3.4	3.3	3.5	3.4	3.5	3.5
Fraser So	outh	4.8	4.6	4.8	4.9	5.2	4.8	4.9	5.0	5.4	5.5	6.0	6.3	7.0	7.0	7.1	6.6	7.0	7.5	7.8	7.6
Female	e (Non-prenatal)	1.5	1.3	1.4	1.4	1.4	1.4	1.5	1.5	1.7	1.8	2.1	2.2	2.4	2.5	2.6	2.2	2.4	2.6	2.9	2.7
Male		1.4	1.5	1.5	1.6	1.7	1.6	1.6	1.6	1.9	1.8	2.1	2.2	2.4	2.6	2.5	2.3	2.6	2.8	2.9	2.8

		2009	2010	2011	2012	2013	2014
Fraser Hea	lth	3375.9	3365.6	3414.9	3860.5	4234.0	4494.5
Fraser East		3270.8	3165.0	3121.5	3376.3	3503.7	3949.4
Fraser Nor	th	3748.7	3720.2	3814.6	4436.8	5196.2	5379.8
Fraser Sout	th	3100.4	3144.0	3192.7	3561.9	3708.5	3963.2
Gender	Female	4371.7	4345.9	4387.9	4851.8	5211.6	5410.4
	Male	2368.1	2372.3	2437.2	2857.7	3242.7	3558.4
Age	< 30	3496.5	3457.6	3411.6	3562.3	3685.5	3711.3
	30-39	8356.7	8484.8	8525.0	9133.9	9359.6	9915.5
	40-49	2990.7	2968.7	3065.1	3611.8	4190.5	4460.7
	≥ 50	1147.4	1192.3	1324.9	1997.4	2771.8	3196.3

	2010		2011				2012	2		2013					2014						
Indicator 3: New HIV	V Diagnoses	Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4
Fraser Health	By Client Residence	17	19	19	17	12	18	15	9	11	10	10	14	8	14	23	19	15	11	16	16
	By Provider Address	14	15	13	15	10	20	9	8	10	5	7	8	9	8	14	15	11	10	12	13
Gender	Female	6	8	3	5	1	5	4	1	0	3	2	3	1	3	2	3	6	4	6	1
	Male	11	11	16	12	11	13	11	8	11	7	8	11	7	11	21	16	9	7	10	15
Age	< 30	3	5	9	1	1	3	3	3	3	1	1	2	1	3	5	6	5	1	4	4
	30-39	6	2	2	6	3	9	3	1	1	3	2	2	3	4	8	6	5	3	5	4
	40-49	6	6	5	5	2	5	4	4	3	2	5	4	2	4	6	4	3	3	1	3
	≥ 50	2	6	3	5	6	1	5	1	4	4	2	6	2	3	4	3	2	4	6	5
Exposure	MSM	4	3	11	4	7	5	8	4	6	5	6	5	6	9	10	11	5	3	_	_
	IDU	2	4	3	2	1	4	0	0	4	0	0	1	1	1	2	5	2	1	_	_
	HET	9	10	3	8	3	7	7	5	1	4	4	6	1	3	9	3	7	6	_	_
	Other	2	0	1	1	1	2	0	0	0	0	0	1	0	1	2	0	0	0	_	_
	NIR/Unknown	0	2	1	2	0	0	0	0	0	1	0	1	0	0	0	0	1	1	-	-
	HET Other	9	10		8	1	7 2	7	5	1 0	4	4	1 6 1 1		1	9	3	7	_	- - -	

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Praser East			•		er Ada		4	1	1	1	4	4	2 2		0	1	0	0	0	2	1	2	1	1	2
Fraser North			•		Resid		7	6	6	6	6	6	7 4		6	5	6	6		14	9	8	5	5	10
Traser North			•		er Ada		5	5	2	6	3	7	4 5			4	2	7	4	8	9	7	4	4	9
Fraser South			,		Resid		8		11	9	2	8	6 (		2	4	7	2	5	6	9	5	6	9	4
Traser South			•		er Ada		5	9	10	8	3	9	3		1	2	6	2	4	4	5	2	5	7	2
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Stage 0	8	10	3	6	2	1	0	0	6	9	3	6	5	4	0	4	0	2	1	$\frac{15}{0}$		3	2	1	1
Stage 1	14	8	6	9	5	2	1	4	9	6	5	5	8	1	1	3	1	5	0	2		5	1	4	1
Stage 2a	5	2	5	5	1	1	3	1	4	1	2	4	3	0	0	2	0	2	1	2		1	0	1	1
Stage 2b	9	7	6	8	2	2	1	1	7	5	5	7	1	1	2	3	4	1	1	0		1	1	1	2
Stage 3	17	21	9	15	6	3	2	1	11	18	7	14	0	0	0	1	2	5	3	3	10	) 1	1	3	6
Unknown	2	1	2	4	0	1	0	0	2	0	2	4	1	1	1	0	0	0	0	2		1	0	1	2
Total	55	49	31	47	16	10	7	7	39	39	24	40	18	7	4	13	7	15	6	9	2	1 1	15	11	13
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Stage 0	0	2	1	1	4	6	2	6	2	1	0	0	2	3	1	0	0	0	0	0	_	)	0	0	0
Stage 1	0	1	1	3	5	3	2	3	2	1	0	1	7	4	3	5	0	0	1	0		)	0	0	0
Stage 2a	1	0	3	0	2	1	2	2	2	0	0	0	1	0	3	1	0	1	0	2		)	0	0	0
Stage 2b	3	4	2	3	3	2	2	5	1	1		0	4	4	3	3	1	0	0	0		)	0	0	0
Stage 3	5	5	3	5	5	8	2	6	2	2	3	3	6	10	4	5	3	1	0	1		1	0	0	0
Unknown	0	0	0	0	1	0	1	0	0	0	0	2	1	1	0	2	0	0	0	0		)	0	1	0
Total	9	12	10	12	20	20	11	22	9	5	4	6	21	22	14	16	4	2	1	3		1	0	1	0
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Fraser Health								198			189			174				51		]	1478			12	284
Age Category								123				94			37			82			72				61
		-39						290			28			25				34			199				174
		-49						62			59			55				29			459				393
	≥ !							949			91			84				06			748			6	556
Age Category	M	SM			30			3.				28			24			22			21				17
and MSM Status					-39			60				59			54			51			49				42
Status					-49			134			13			13				23			114				102
	<b>N.T.</b>		03.4	≥.				332			32			31				04			288			- 2	264
	No	on-M	SM	< 3				11				1			.1			10			9				9
					-39 -49			117			11			10				94			76				63
					50			27. 34.			26 33			24 30				35 85			189 255				156 214
	T T.	ıknov	1792		30			79				55			52			50			42				35
	OI	IKIIOV	WII		-39			112			10				,z )7			89			74				69
					-49			21:			20			17				71			156			1	135
				≥.				270			26			22				17			205				178
Gender	M:	ale			30			1500			143			133				66		1	1162				021
Condo		male						48.			45			41				85		,	316				263
Injection	ID							53:			51			48				:55			373				299
Drug Use		on-ID	U					104			101			97				24			862				776
Ü		ıknov						40.			35			29				72			243				209
MSM Status		SM	_					559			53			52				00			472				425
		on-M	SM					742			72			66				24			529				142
		ıknov						682			63			55				27			477				417
Health		aser I						302			28			24				35			207				167
Authority			North	ı				882			84			77				31			665				581
-			South					80			76			72				85			606				536

Indicator 6: <b>Programmat</b>	•	e (PCS)							
	2013 Q1	Q2	Q3	Q4		2014 Q1	Q2	Q3	Q4
< 3 CD4 Tests	20.4%	15.9%	15.4%	17.0%	18	3.9%	23.0%	14.0%	12.4%
< 3 Viral Load Tests	6.8%	5.6%	6.6%	5.3%		2.1%	3.4%	2.2%	3.4%
No Baseline Genotype	7.8%	6.5%	7.7%	7.4%	3	3.2%	3.4%	4.3%	3.4%
Baseline CD4 < 200 cells/		25.2%	19.8%	20.2%		0.0%	24.1%	29.0%	23.6%
Non-Recommended ART	•	3.7%	6.6%	9.6%		).5%	10.3%	9.7%	3.4%
Non Viral suppression at		26.2%	25.3%	22.3%		1.2%	29.9%	25.8%	29.2%
PCS Score: 0	52	58	50	48	2,	45	35	38	42
PCS Score: 1	21	20	20	26		31	31	35	31
PCS Score: 2	19	21	13	12		14	13	16	13
PCS Score: 3	9					4		4	2
		6	5	6			7		
PCS Score: 4 or more	2	2	3	2		1	1	0	1
Total (n=)	103	107	91	94		95	87	93	89
Indicator 7: New DTP AF	RV Participants								
First Starts	23	17	29	20		30	20	22	22
Experienced Starts	20	39	25	36		21	42	31	32
Indicator 8: CD4 Cell Co	unt at ART Initiatio	n for ADV	Vaïve DTD I	Particinante					
CD4 ≥ 500	7	7	Naive DIP 1	10		9	11	8	9
CD4 2500 CD4 350-499	9	2	3	5		3	3	8 5	5
CD4 200-349	4	5	8	4		10	1	4	4
CD4 50–199	2	2	8	0		3	3	3	3
CD4 < 50	1	1	6	0		4	1	2	1
CD4 Median (cells/μL)	420	370	202	500		290	550	400	440
Total (n=)	23	17	29	19		29	19	22	22
Indicator 9: Active and In	nactive DTP Particit	pants							
Active DTP Participants	1468	1515	1539	1566	1	581	1625	1640	1664
Inactive DTP Participants		229	232	230		239	232	229	243
inactive B 11 Turticipums	, 210	22)	202	250		200	232	227	213
Indicator 10: Antiretrovi	ral Adherence								
≥ 95%	18	13	15	17		19	14	23	20
80% to < 95%	9	7	6	4		5	4	3	5
40% to < 80%	0	5	3	5		2	1	3	5
< 40%	0	0	1	2		0	0	0	0
Total (n=)	27	25	25	28		26	19	29	30
T 1: 4 11 D : 4 D	r (* 15 16								
Indicator 11: Resistance		000	1027	1016	1	120	1124	1154	1000
Suppressed	1005	998	1037	1016	1	120	1124	1154	1096
Wild Type	203	203	207	178		163	170	153	144
Never Genotyped	18	21	18	19		15	16	15	10
1-Class	28	39	35	43		30	26	32	37
2-Class	6	8	5	8		1	4	5	4
3-Class Total (n=)	2 1262	3 1272	1 1303	1 1265	1	330	2 1342	3 1362	1 1292
Total (II—)	1202	12/2	1303	1203	•	.550	1312	1302	1272
Indicator 12: AIDS-Defin		2007	2008	2009	2010	2011	2012	2013	2014
	Cases	38	52	32	33	33	26	19	21
	Rate per 100,000	2.5	3.4	2.0	2.0	2.0	1.6	1.1	1.2
	Cases	23	26	16	24	20	13	16	9
	Rate per 100,000	1.5	1.7	1.0	1.5	1.2	0.8	0.9	0.5
	Cases	27	30	21	30	23	12	10	-
(BCCDC Reports)	Rate per 100,000	1.8	1.9	1.3	1.9	1.4	0.7	0.6	_
Indicator 13: HIV-Related	d Mortality	2004	2005	2006	2007	2008	2009	2010	2011
Fraser Health		18	26	23	21	14	10	18	13
Per 100 HIV+ Population		0.88	1.25	1.09	0.97	0.63	0.44	0.78	0.56
Per 100,000 Population		1.25	1.77	1.54	1.38	0.91	0.64	1.12	0.80
- 11 100,000 Topulation		1.20	2.77	1.01	1.00	0.71	0.0 f	1.12	0.00