

HIV MONITORING QUARTERLY REPORT

FOR FRASER 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.

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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. 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.



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

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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.

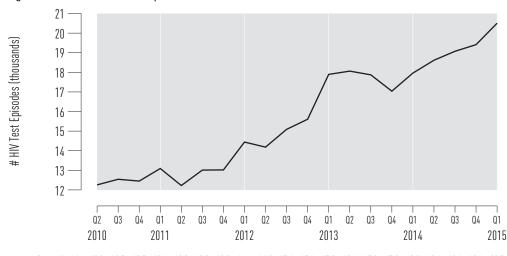
The goals of HA-led STOP-funded initiatives are to work toward achieving these goals. To these ends some outcome measures or indicators of progress have been drafted that should be considered in the design and implementation phases of these initiatives.

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



Fraser Health 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 20.5



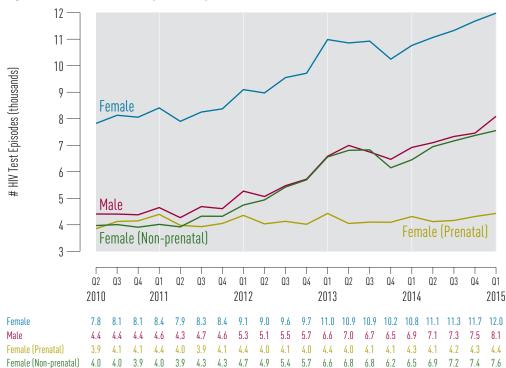
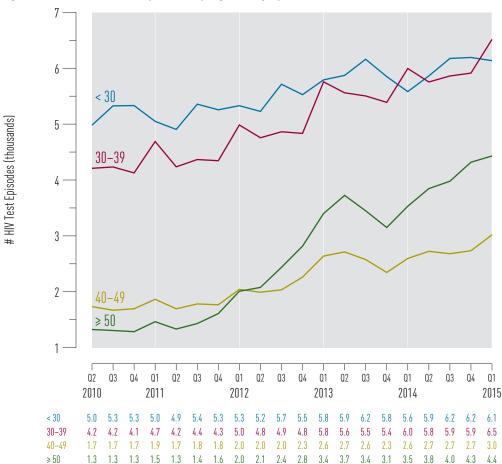
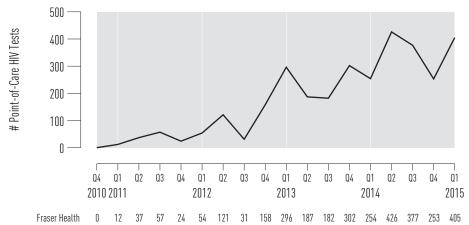


Figure 1.3 HIV Test Episodes by Age Category for Fraser Health 1,2







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 In Fraser Health, POC testing data are available from March 2011 forward.
- Testing does not include point of care tests.

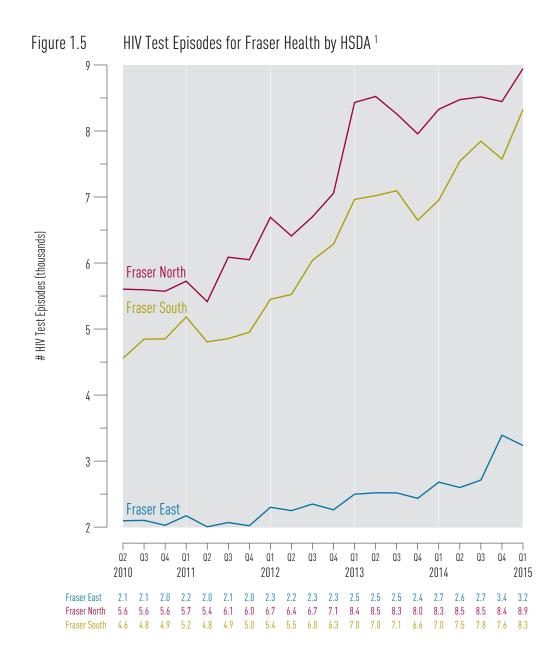


Figure 1.6 HIV Test Episodes for Non-prenatal Females in Fraser Health by HSDA ¹

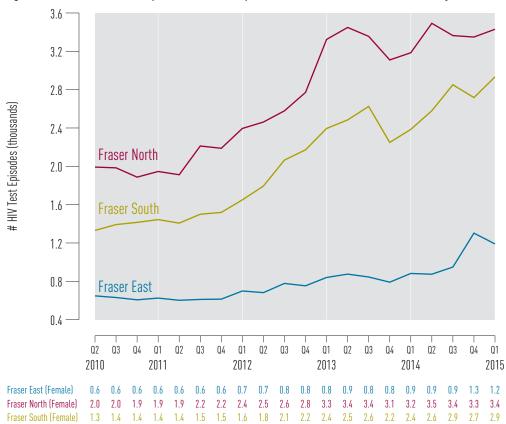
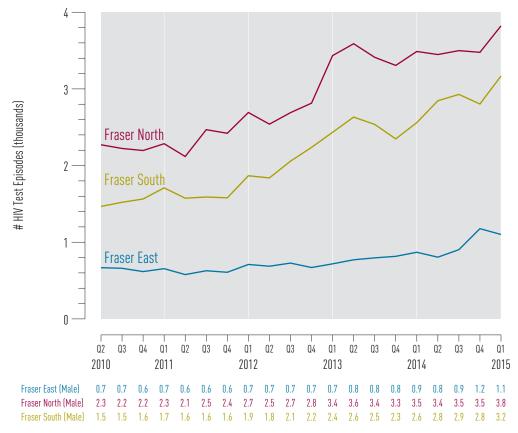


Figure 1.7 HIV Test Episodes for Males in Fraser Health by HSDA ¹



Indicator 2. HIV Testing Rates

Figure 2.1 Rate of HIV Testing for Fraser Health and HSDAs ²

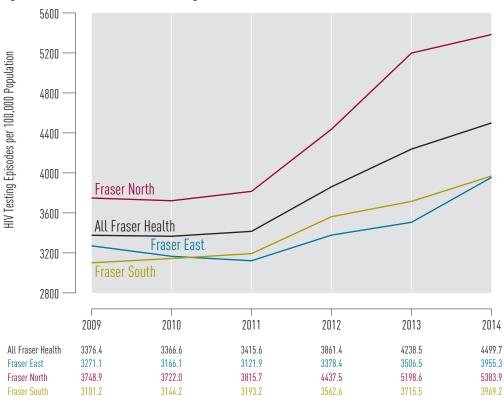
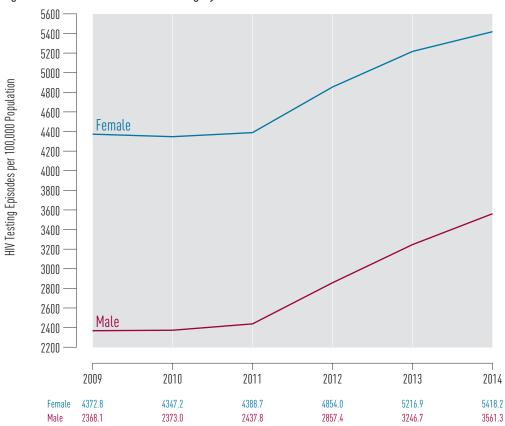
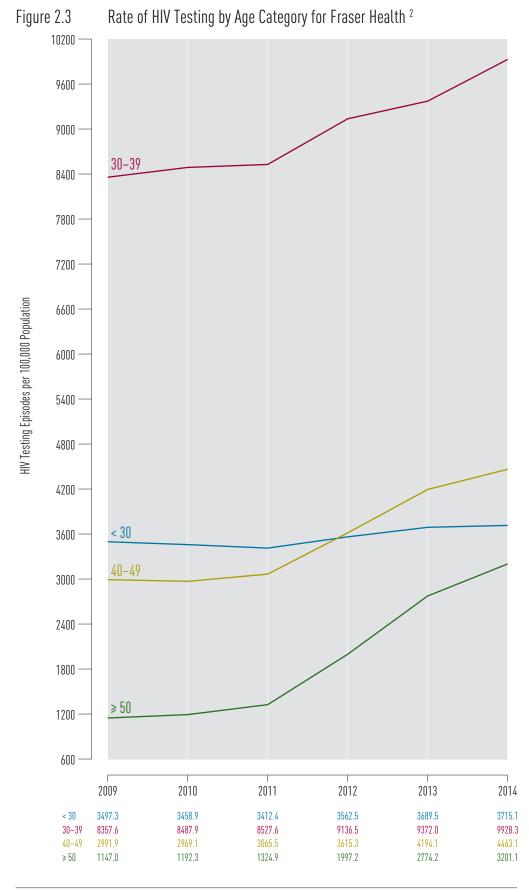


Figure 2.2 Rate of HIV Testing by Gender for Fraser Health ²





¹ 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 ³

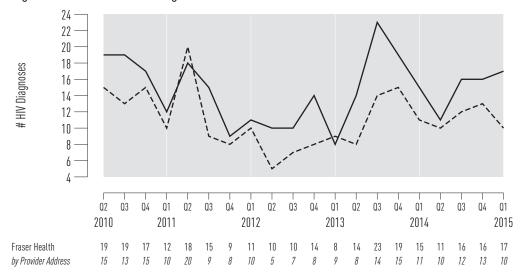
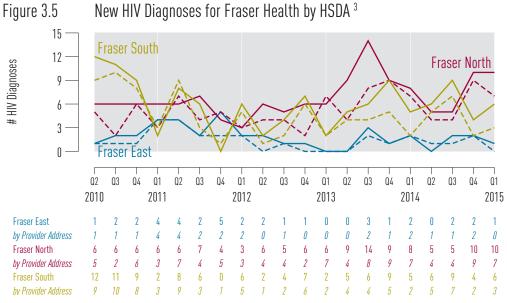


Figure 3.2 New HIV Diagnoses for Fraser Health by Gender ³



³ 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 ³ < 30 # HIV Diagnoses ≥ 50 Q2 Q4 Q2 Q4 Q2 Q3 Q1 Q2Q3 Q4 Q2 Q3 Q4 Q1 Q3 Q1 Q3 Q4 Q1 Q1 2010 2011 2012 2013 2014 2015 < 30 30-39 3 5 3 5 40-49 ≥ 50 Figure 3.4 New HIV Diagnoses for Fraser Health by Exposure Category 3,4 MSM 12 -10 # HIV Diagnoses 6 4 IDU 2 Other NIR/Unknown Q2 Q3 Q4 Q1 Q2 Q3 Q4 Q1 Q2 Q3 Q4 Q1 Q2 Q3 Q4 Q1 Q2 2010 2011 2012 2013 2014 MSM (men who have sex with men) 3 11 10 11 3 0 IDU (injection drug user) 4 0 0 0 10 3 HET (heterosexual) 4 9 0 0 0 0 Other (other exposure identified) 0 N 0 N NIR/Unknown (no identified exposure) 0 New HIV Diagnoses for Fraser Health by HSDA $^{\rm 3}$ Figure 3.5



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

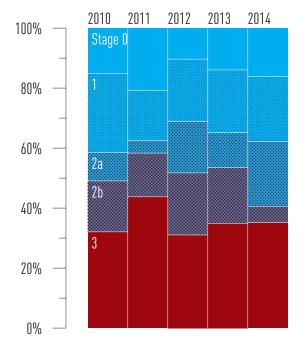
⁴ MSM=men who have sex with men; IDU= injection drug user; HET=heterosexual. NIR=No identified risk/exposure.

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 Fraser Health, 2010–2014 ⁵

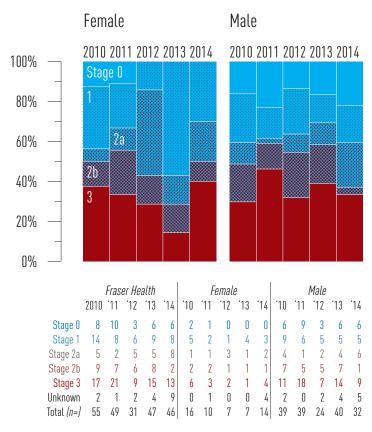


Indicator 4. Stage of HIV Infection at Diagnosis

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

Stage	Criteria													
0	previous	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	N anct2		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 Fraser Health, 2010–2014 ⁵



Data Source: BCCDC

Figure 4.3 Stage of HIV Infection at Diagnosis by Age Category for Fraser Health, 2010–2014 ⁵

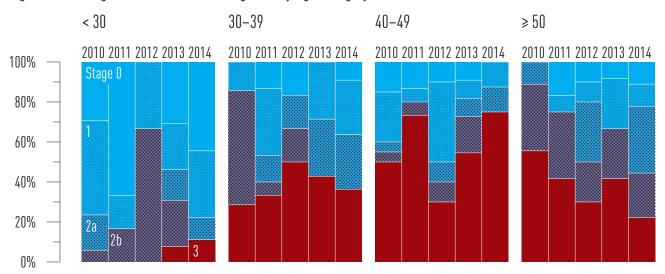
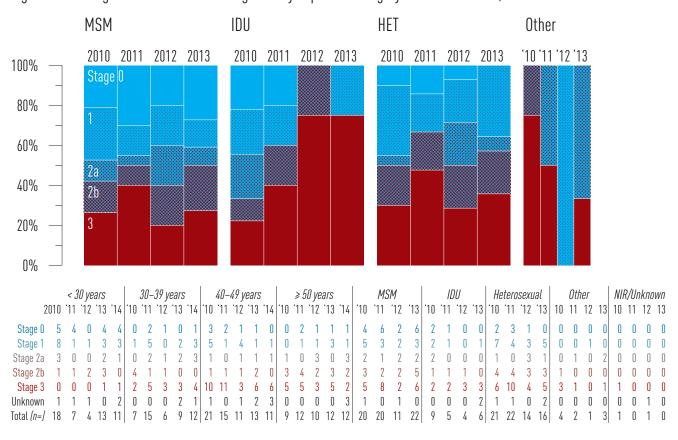


Figure 4.4 Stage of HIV Infection at Diagnosis by Exposure Category for Fraser Health, 2010–2013 5.6

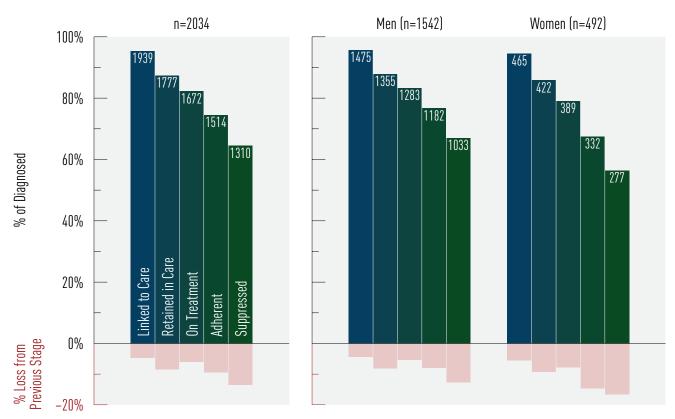


⁶ MSM=men who have sex with men; IDU= injection drug user; HET=heterosexual. NIR=No identified risk/exposure.

Indicator 5. HIV Cascade of Care

The success of seek, test, treat and retain (STTR) strategies like STOP is reliant on early diagnosis of HIV, linking newly diagnosed HIV-positive persons with ongoing care, retaining persons in HIV-care; initiating ART based on best evidenced practices and maintaining optimal ART adherence to ensure a suppressed viral load. These stages of HIV-care can be summarized as: 1. HIV diagnosis, 2. 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 Figure 5.2 Estimated Cascade of Care for Fraser Health Health, Year Ending 2015 Q1 7 Estimated Cascade of Care for Fraser Health by Gender, Year Ending 2015 Q1 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.

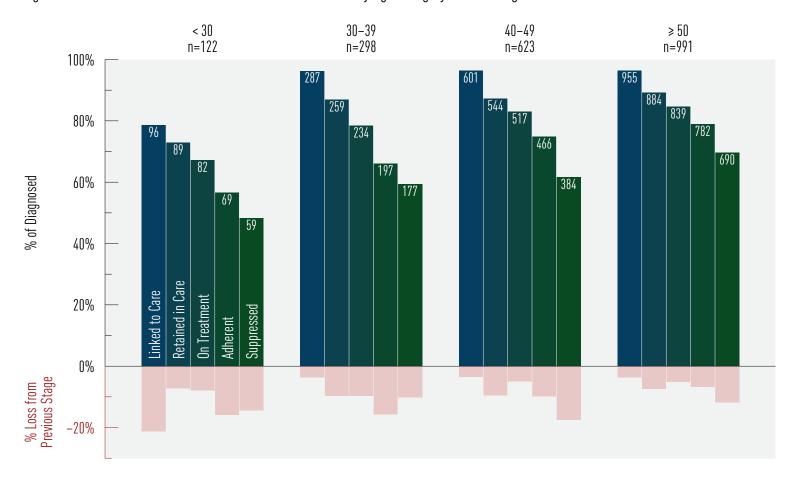
NB: Transgender has been assigned to their biological sex.

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

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, Year Ending 2015 Q1 8

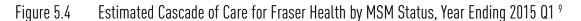


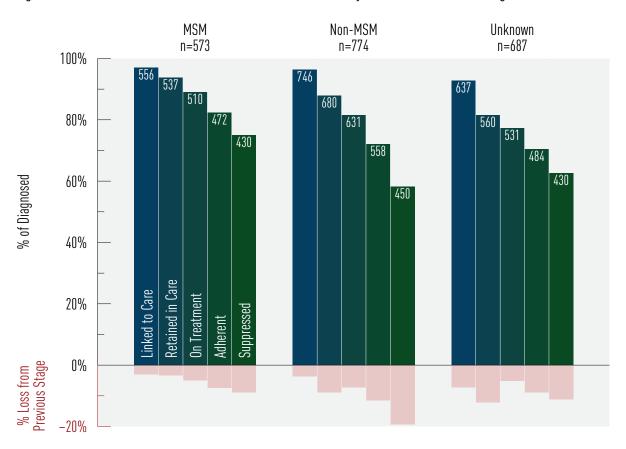
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Data Sources:

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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.

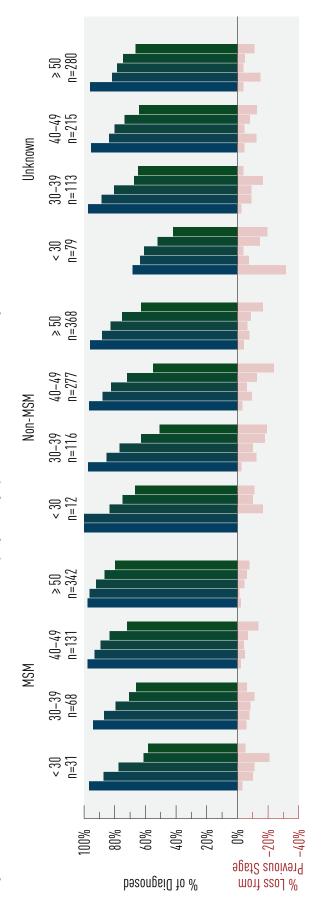
Recent updates to the DTP database have allowed for more comprehensive information on HIV risk group category. As a result, 2014 Q4 data may differ significantly from preceding reports in terms of total numbers ascribed to each risk group.

⁹ Data is for the period 2014 Q2-2015 Q1.

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 5.5 Estimated Cascade of Care for Fraser Health by Age Category and MSM Status, Year Ending 2015 Q1 ⁹

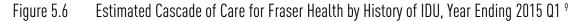


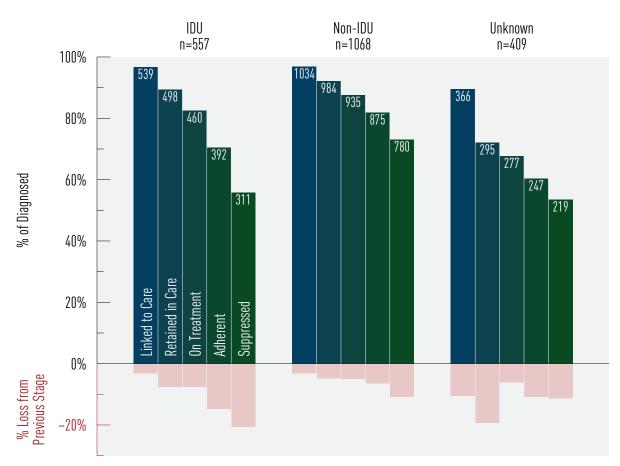
Data is for the period 2014 Q2-2015 Q1.

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.

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: 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.

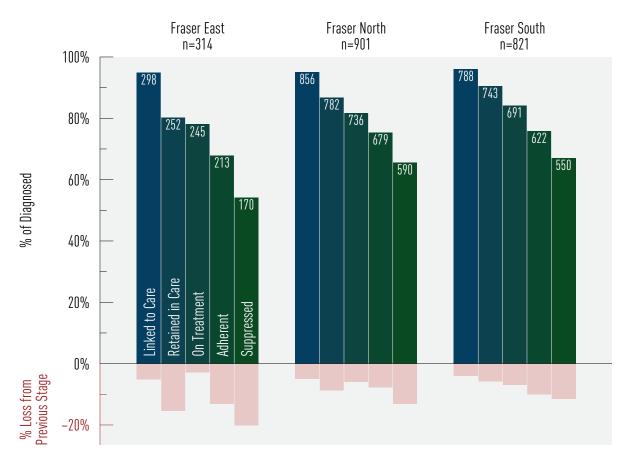
Recent updates to the DTP database have allowed for more comprehensive information on HIV risk group category. As a result, 2014 Q4 data may differ significantly from preceding reports in terms of total numbers ascribed to each risk group.

Data is for the period 2014 Q2-2015 Q1.

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.

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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)).

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	Mortality Risk Ratio (95% Confidence Interval)	Immunologic Failure Risk Ratio (95% CI)	Virologic Failure Risk Ratio
Compliance Score	(95% Confidence interval)	Katio (40% CI)	(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 Fraser Health, 2013 Q2-2015 Q1 10

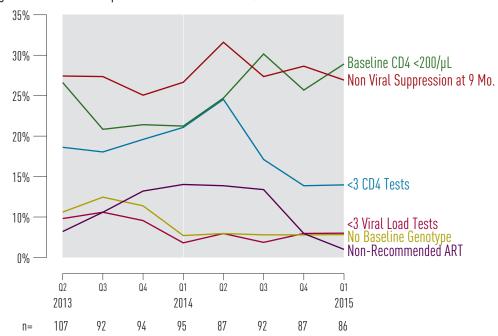
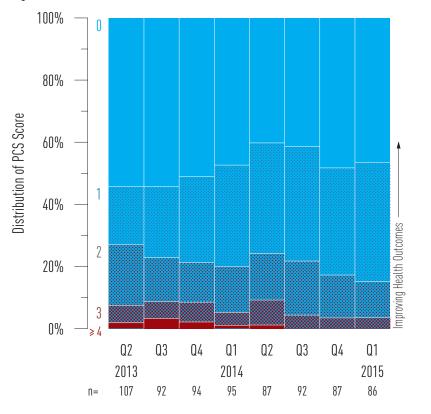


Figure 6.2 Historical Trends for PCS Score for Fraser Health, 2013 Q2-2015 Q1 10,11



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

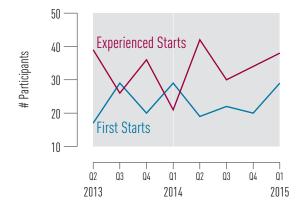
¹¹ Each quarter's data is calculated as the sum of the 4 quarters leading up to it. e.g. 2013 Q1 is calculated from 2012 Q2 – 2013 Q1. NB: A score of o is the best score and a score of 4 or more is the worst score.

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.

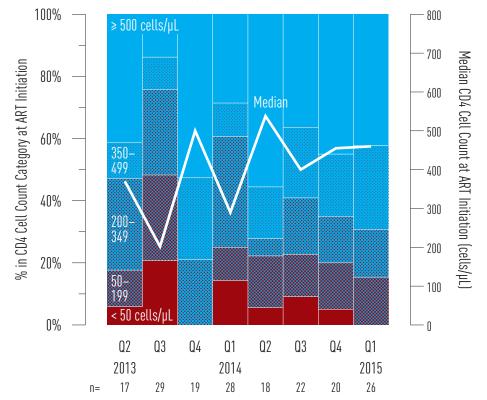
Indicator 7. New Antiretroviral Therapy Starts in Fraser Health

Figure 7 BC-CfE Drug Treatment Program Enrollment: New ART Participants in Fraser Health, 2013 Q2-2015 Q1 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 Q2—2015 Q1 ¹³



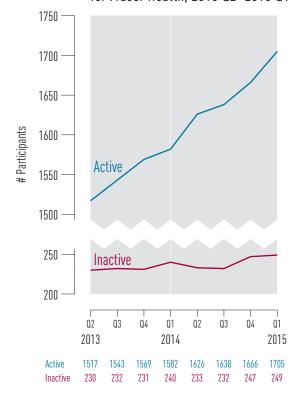
- Data Source: Drug Treatment Program Database
 Limitation: DTP participants are designated to an HA based on most current residence provided by the participant.
- Data Source: Drug Treatment Program Database Limitations: CD4 cell count data is approximately 80% complete.

Indicator 9. Active and Inactive DTP Participants

Table 3. Distribution of People on ART for Fraser Health, 2015 Q1 14

Age	< 30	89
	30-39	259
	40-49	543
	≥ 50	814
Gender	Male	1309
	Female	396
Exposure	MSM	521
	IDU	461
Total		1705

Figure 9 Active and Inactive DTP Participants for Fraser Health, 2013 Q2-2015 Q1 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.

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.

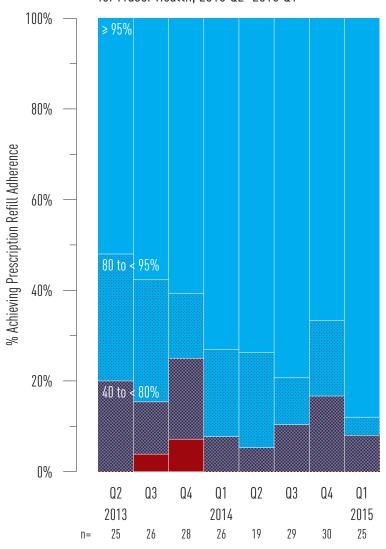
Inactive DTP participants: Persons no longer prescribed drugs through the HIV/AIDS Drug Treatment Program in the last quarter.

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.

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 Q2–2015 Q1 ¹⁶



¹⁶ 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.

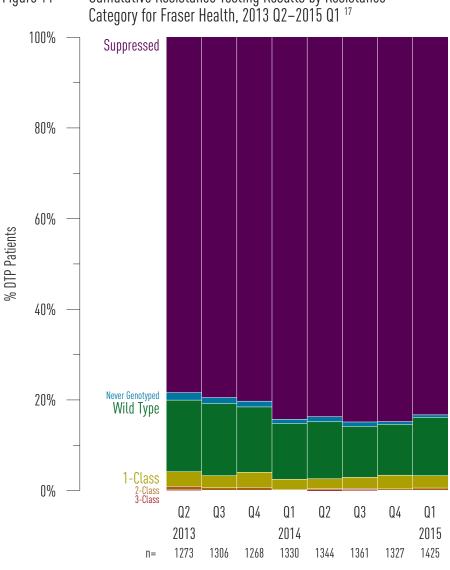


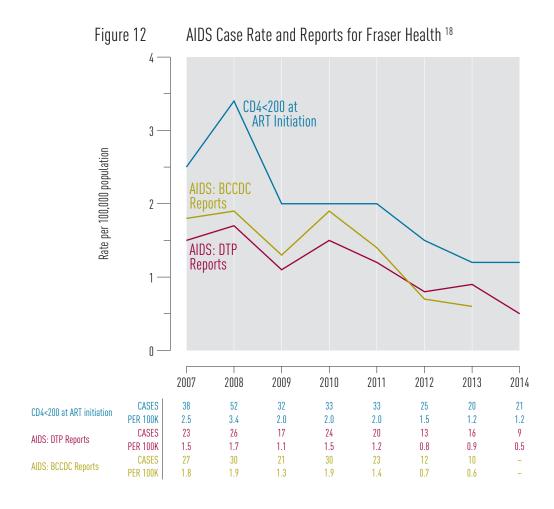
Figure 11 Cumulative Resistance Testing Results by Resistance

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

¹⁷ 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.

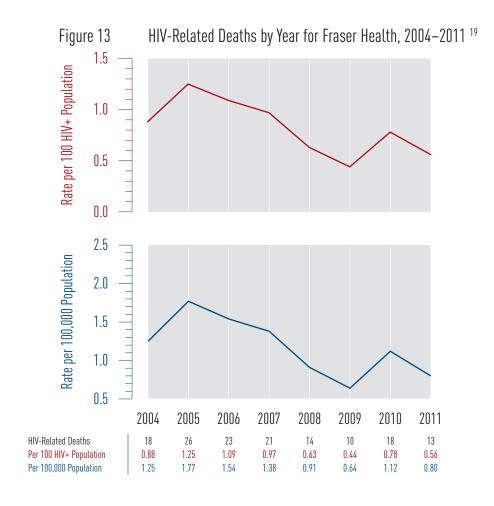


Data Source: DTP AIDS cases are obtained from the Drug Treatment Program Database; BCCDC AIDS cases are obtained from the BCCDC; CD4<200 at ART initiation data came from the DTP database.

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:

¹⁹ Data Source: BC Vital Statistics

^{1.} DTP participants are designated to an HA based on most current residence provided by the participant.

^{2.} Mortality data is updated annually.

^{3.} The most recent available data was used.

Appendices

Indicator 1 Episodes (1		201 Q2		Q4	2011 Q1	Q2	Q3	Q4	2012 Q1	2 Q2	Q3	Q4	2013 Q1	3 Q2	Q3	Q4	2014 Q1		Ç	23	Q4	2015 Q1
Fraser Heal	th	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 1	9.4	20.5
Gender	Female	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 1	1.7	12.0
	Male	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	8.1
	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 (Pre	enatal)	3.9	4.1	4.1	4.4	4.0	3.9	4.1	4.4	4.0	4.1	4.0	4.4	4.0	4.1	4.1	4.3	4.1	. 4	.2	4.3	4.4
Female (No	n-prenatal)	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	6.2	6.5	6.9	7	.2	7.4	7.6
Age	< 30	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	5.8	5.6	5.9	6	.2	6.2	6.1
	30-39	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	3 5	.9	5.9	6.5
	40-49	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	7 2	.7	2.7	3.0
	≥ 50	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	3 4	.0	4.3	4.4
POC HIV T				0	12	37	57	24	54	121	31	158	296	187	182	302	254	426	5 37	77 :	253	405
(not in thou	,																					
Fraser East		2.1		2.0	2.2	2.0	2.1	2.0		2.2	2.3			2.5	2.5	2.4				7	3.4	3.2
	Non-prenatal			0.6	0.6	0.6	0.6			0.7	0.8	0.8		0.9	0.8	0.8				1.9	1.3	1.2
Male	•	0.7		0.6	0.7	0.6	0.6			0.7	0.7	0.7		0.8	0.8	0.8				1.9	1.2	1.1
Fraser Nort		5.6		5.6	5.7	5.4	6.1	6.0		6.4	6.7	7.1	8.4	8.5	8.3	8.0				5.5	8.4	8.9
	Non-prenata			1.9	1.9	1.9	2.2			2.5	2.6			3.4	3.4	3.1	3.2			5.4	3.3	3.4
Male	1	2.3		2.2	2.3	2.1	2.5			2.5	2.7	2.8		3.6	3.4	3.3	3.5			5.5	3.5	3.8
Fraser Sout		4.6			5.2	4.8	4.9			5.5	6.0		7.0	7.0	7.1	6.6	7.0	7.5			7.6	8.3
	Non-prenatal	1.3		1.4	1.4	1.4	1.5	1.5	1.6	1.8	2.1	2.2	2.4	2.5	2.6	2.2	2.4			.9	2.7	2.9
Male		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	3.2
Indicator 2	: Rate of HI	V Testing	per 1	00,00	0																	
			20)9	2	010		2011		201	.2	2	013		2014							
Fraser Heal	th		3376	.4	336	6.6	3	415.6		3861.	.4	423	88.5	44	199.7							
Fraser East			3271	.1	316	6.1	3	121.9		3378.	.4	350	6.5	39	955.3							
Fraser Nort	h		3748	.9	372	22.0	3	815.7		4437.	.5	519	8.6	53	383.9							
Fraser South	h		3101	.2	314	4.2	3	193.2		3562.	.6	371	5.5	39	969.2							
Gender	Female		4372	8	434	7.2	4	388.7		4854.	.0	521	6.9	54	118.2							
	Male		2368	.1	237	3.0	2	437.8		2857.	.4	324	6.7	35	561.3							
Age	< 30		3497	.3	345	8.9	3	412.4		3562.	.5	368	39.5	37	715.1							
	30-39		8357	.6	848	37.9	8	527.6		9136.	.5	937	2.0	99	928.3							
	40-49		2991	.9	296	9.1	3	065.5		3615.	.3	419	4.1	44	163.1							
	≥ 50		1147	.0	119	2.3	1	324.9		1997.	.2	277	4.2	32	201.1							
					2010		20)11			2012			201	3			2014				2015
Indicator 3	: New HIV I	Diagnose	s		Q2	Q3 (2 Q3			Q2 (Q3 Q			Q3			Q2	Q3	Q4	
Fraser Heal		By Client		lence	19			12 1			11			4 8			19		11	16	16	
		By Provid			15		15	10 2			10	5	7	8 9	9 8	14	15	11	10	12	13	10
Gender		Female			8	3	5		5 4	1	0	3	2	3	1 3	2	3	6	4	6	1	6
		Male			11	16	12	11 1	3 11	8	11	7	8 1	1 7	7 11	21	16	9	7	10	15	11
Age		< 30			5	9	1	1	3 3	3	3	1	1	2	1 3	5	6	5	1	4	4	2
		30-39			2	2	6	3	9 3	1	1	3	2	2 3	3 4	8	6	5	3	5	4	5
		40-49			6	5	5	2	5 4	4	3	2	5	4 2	2 4	6	4	3	3	1	3	2
		≥ 50			6	3	5	6	1 5	1	4	4	2	6 2	2 3	4	3	2	4	6	5	8
Exposure		MSM			3	11	4	7	5 8	4	6	5	6	5 (5 9	10	11	5	3	7	-	_
		IDU			4	3	2	1	4 0	0	4	0	0	1	1 1	2	5	2	1	0	_	_
		HET			10	3	8	3	7 7	5	1	4	4	6	1 3	9	3	7	6	6	_	_
		Other			0	1	1	1	2 0	0	0	0	0	1 () 1	2	0	0	0	0	_	_
	NIR/Unk	nown		2	1	2	0	0 0	0	0	1	0	1 (0 (0	0	1	1	3	_	_	

T 1:		**				,	.31		2010		04	201		02		2012		02	04	2013		02	04	201		2 0	2 04	2015
Indicator 3: New HIV Diagnoses (cont'd) Fraser East By Client Residence									$\frac{Q^2}{1}$	Q3 2				Q3 2	Q4 5	Q1 2	2	Q3 1	1	Q1 0	Q2 0	3	1	2		2 Q 0	3 Q4 2 2	
Fraser East				,		n Ke ider 1			1	1	1	4		2	2	2	0	1	0	0	0	2	1	2			2 2 1 2	
Fraser Nort	h			-					6	6				7	4	3	6	5	6	6	9	14	9	8			5 10	10
Fraser North By Client Residence By Provider Address						5	2				4	5	3	4	4	2	7	4	8	9				4 9				
Fraser Sout	h			-		nt Re			12		9			6	0	6	2	4	7	2	5	6	9				9 4	
				•		ider 1			9		8			3	1	5	1	2	6	2	4	4	5				7 2	
Indicator 4:	Sta	ge of		-					ne.																			
mulcator 4.		Frase										1./	Iale				20 ***			2	0.20		••		4	0 40		
						'10		emal '12		' 14	'10			3 '14	1 '10		30 y€ '12		' 14		0-39			، ا			year: 12 '13	
Stage 0	8		3				1	0	0	0	6	9		6 6					4	0	2		0	1	3	2	1 1	
Stage 1	14	8	6				2	1	4	3	9	6		5 5					3	1	5		2	3	5	1	4 1	
Stage 2a	5	2	5	5	8	1	1	3	1	2	4	1	2	4 6	5 3	3 (0 (2	1	0	2	1	2	3	1	0	1 1	. 1
Stage 2b	9	7	6	8	2	2	2	1	1	1	7	5	5	7 1	. 1	l 1	2	3	0	4	1	1	0	0	1	1	1 2	2 0
Stage 3	17	21	9	15	13	6	3	2	1	4	11	18		4 9) (0 (1	1	2	5	3	3	4	10	11	3 6	6
Unknown	2	1	2			1	1	0	0	4	2	0	_	4 5			_	-	2	0	0	0	2	1	1	0	1 2	
Total	55	49	31	47	46	16	10	7	7	14	39	39	24 4	0 32	2 18	3 7	7 4	13	11	7	15	6	9 1	12	21	15	11 13	3 11
	410	≥ 5			(1.4	(10		ASM		112	'10		DU 112	(1.2				exual			her E						nknov	
C+ 0		<u>'11</u>				_			12	'13	'10	'11			_			¹ 12	'13	'10	'11			_	10	<u>'11</u>	'12	<u>'13</u>
Stage 0 Stage 1	0	2	1 1	1 3		5		6 3	2	6	2 2	1 1				2 7	3 4	1 3	0 5	0	0	0		0	0	0	0	0
Stage 1 Stage 2a	1	0	3					<i>3</i>	2	2	2	(1	0	3	1	0	1	0		2	0	0	0	0
Stage 2b	3	4	2					2	2	5	1	1				4	4	3	3	1	0	0		0	0	0	0	0
Stage 3	5	5	3	5				8	2	6	2	2					10	4	5	3	1	0		1	1	0	0	0
Unknown	0	0	0	0	3	1		0	1	0	0	C	0	2	:	1	1	0	2	0	0	0		0	0	0	1	0
Total	9	12	10	12	12	20	2	0	11	22	9	5	4	6	2	1	22	14	16	4	2	1		3	1	0	1	0
Indicator 5	: HI	V Ca	isca	de c	of Ca	ıre		D	IAGI	NOSE	D		LINE	ŒD		RET	AINE	D		ON A	ART	Α	DH	EREI	NT	SU	PPRES	SSED
Fraser Heal	lth									203	34		19	939			177	7		1	672			15	14			310
Age Catego	ory	< 30)							12	22			96			8	9			82				69			59
		30-	39							29	8		2	287			25	9		:	234			1	97			177
		40-	49							62	23		(501			54	4			517			4	66			384
		≥ 50)							99	1		9	955			88	4		:	839			7	82			690
Age Catego	ory	MSI	M			< 30				3	31			30			2	7			24				19			18
and MSM						30-3	9			6	8			64			5	9			54				48			45
Status						40-4	9			13	31			128			12	2			117			1	09			94
						≥ 50				34	2		3	334			33	0			315			2	96			273
		Nor	ı-M	SM		< 30					.2			12			1				10				9			8
						30-3				11				113			9	9			89				73			59
						40-4	9			27				268			24				228				99			152
						≥ 50				36			3	353			32				304				77			231
		Unk	cnov	wn		< 30					9			54				0			48				41			33
						30-3				11				110			10				91				76			73
						40-4	9			21				205			18				172				58			138
		_			:	≥ 50				28				269			22				220				09			186
Gender		Mal								154				175			135				283			11]	.033
		Fem								49				165			42				389				32			277
Injection		IDU								55				539			49				460				92			311
		Nor								106)34			98				935				75			780
3 (O) (O		Unk		wn						40				366			29				277				47			219
MSM Status		MSI		03.6						57				556			53				510				72			430
		Non								77				746			68				531				58			450
TT141		Unk								68				537			56				531				84			430
Health Authority		Fras			L					31				298			25				245				13			170
Aumority		Fras								90				356			78 74				736				79 22			590
		Fras	er S	out	11					82	1		7	788			74	3		(591			6.	22			550

Indicator 6: Programmat	tic Compliance Scor	e (PCS)							
	2013	O2	04	2014			O2	04	2015
< 3 CD4 Tests	Q2	Q3 15.2%	Q4 17.0%	Q1 18.9%	Q2		Q3	Q4	Q1
< 3 CD4 Tests	15.9% 5.6%	6.5%	5.3%	2.1%	23.0%		4.1% 2.2%	10.3% 3.4%	10.5%
					3.4%				3.5%
No Baseline Genotype	6.5%	8.7%	7.4%	3.2%	3.4%		3.3%	3.4%	3.5%
Baseline CD4 < 200 cells/	•	18.5%	19.1%	18.9%	23.0%		9.3%	24.1%	27.9%
Non-Recommended ART		6.5%	9.6%	10.5%	10.3%		9.8%	3.4%	1.2%
Non Viral suppression at		26.1%	23.4%	25.3%	31.0%		6.1%	27.6%	25.6%
PCS Score: 0	58	50	48	45	35		38	42	40
PCS Score: 1	20	21	26	31	31		34	30	33
PCS Score: 2	21	13	12	14	13		16	12	10
PCS Score: 3	6	5	6	4	7		4	3	3
PCS Score: 4 or more	2	3	2	1	1		0	0	0
Total (n=)	107	92	94	95	87		92	87	86
Indicator 7: New DTP Al	RV Participants								
First Starts	17	29	20	29	19		22	20	29
Experienced Starts	39	26	36	21	42		30	34	38
Indicator 8: CD4 Cell Co	ount at ART Initiatio	n for ARV-!	Naïve DTP I	Particinants					
CD4 ≥ 500	7	4	10	8	10		8	9	11
CD4 2 500 CD4 350-499	2	3	5	3	3		5	4	7
CD4 200-349	5	8	4	10	1		4	3	4
CD4 200-349 CD4 50-199	2	8	0	3	3		3	3	4
CD4 50-199 CD4 < 50	1	6	0	4	1		2	1	0
	370	202	500	290	538		400	455	460
CD4 Median (cells/μL) Total (n=)	370 17	202 29	19	290	18		22	20	26
Total (II–)	17	29	19	20	10		22	20	20
Indicator 9: Active and In	nactive DTP Particip	ants							
Active DTP Participants	1517	1543	1569	1582	1626		1638	1666	1705
Inactive DTP Participants	s 230	232	231	240	233		232	247	249
Indicator 10: Antiretrovi	ral Adherence								
≥ 95%	13	15	17	19	14		23	20	22
80% to < 95%	7	7	4	5	4		3	5	1
40% to < 80%	5	3	5	2	1		3	5	2
< 40%	0	1	2	0	0		0	0	0
Total (n=)	25	26	28	26	19		29	30	25
Indicator 11: Resistance	Tooting and Dooulto								
	998	1020	1010	1121	1125		1155	1125	1107
Suppressed	201	1038 208	1018 183	1121 164	1125 169		1155 153	1125 149	1187 183
Wild Type Never Genotyped	201	208 17	163	164	159		133		8
1-Class	41	35	42	30	29		33	8 40	39
2-Class	10	7	8	2	4		4	40	7
3-Class	2	1	1	1	2		3	1	1
Total (n=)	1273	1306	1268	1330	1344		1361	1327	1425
7 1	. 711	•••	•	•	2010		2012	2012	2011
Indicator 12: AIDS-Defin		2007	2008	2009	2010	2011	2012	2013	2014
	Cases Rate per 100,000	38 2.5	52 3.4	32 2.0	33 2.0	33 2.0	25	20	21
	Kate per 100,000 Cases	2.5	3.4 26	2.0 17	2.0	2.0	1.5 13	1.2 16	1.2 9
	Rate per 100,000	23 1.5	26 1.7	1.1	1.5	1.2	0.8	0.9	0.5
=	Cases	27	30	21	30	23	12	10	0.5
	Rate per 100,000	1.8	30 1.9	1.3	1.9	23 1.4	0.7	0.6	_
•	•								
Indicator 13: HIV-Relate	d Mortality	2004	2005	2006	2007	2008	2009	2010	2011
Fraser Health		18	26	23	21	14	10	18	13
Per 100 HIV+ Population	1	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