

HIV MONITORING QUARTERLY REPORT

FOR ISLAND HEALTH

FOURTH 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 Health Service Delivery Areas (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 Island Health

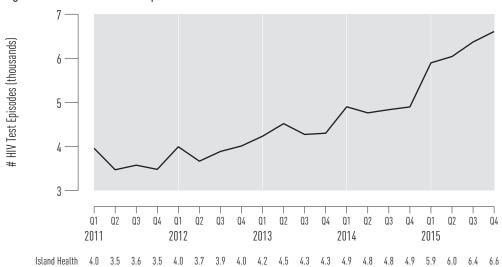


Figure 1.2 HIV Test Episodes by Gender for Island Health 1,2

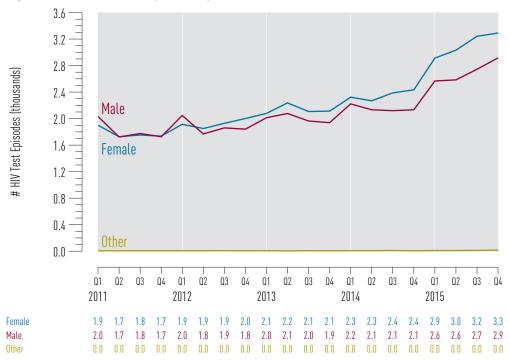


Figure 1.3 HIV Test Episodes by Age Category for Island Health 1,2

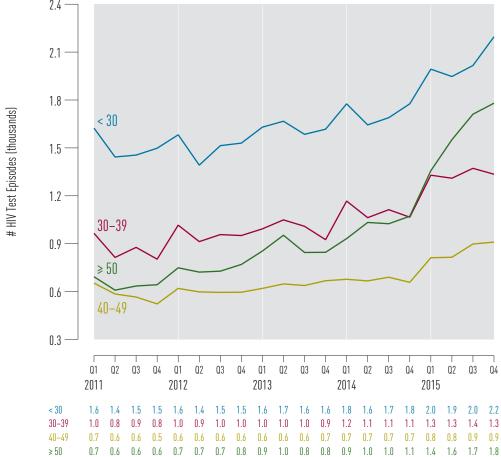
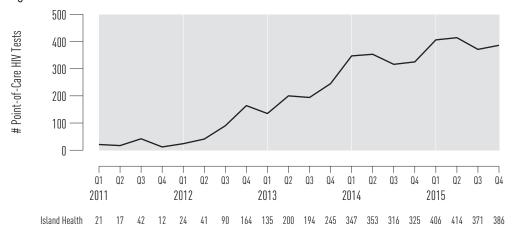


Figure 1.4 Point-of-Care HIV Tests for Island Health



Data Source: The BC Public Health Microbiology and Reference Laboratory (BCPHMRL) courtesy of the BC Centre for Disease Control (BCCDC). HIV screening tests conducted by the VIHA Laboratory are not included.

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.
- Testing does not include point of care tests.

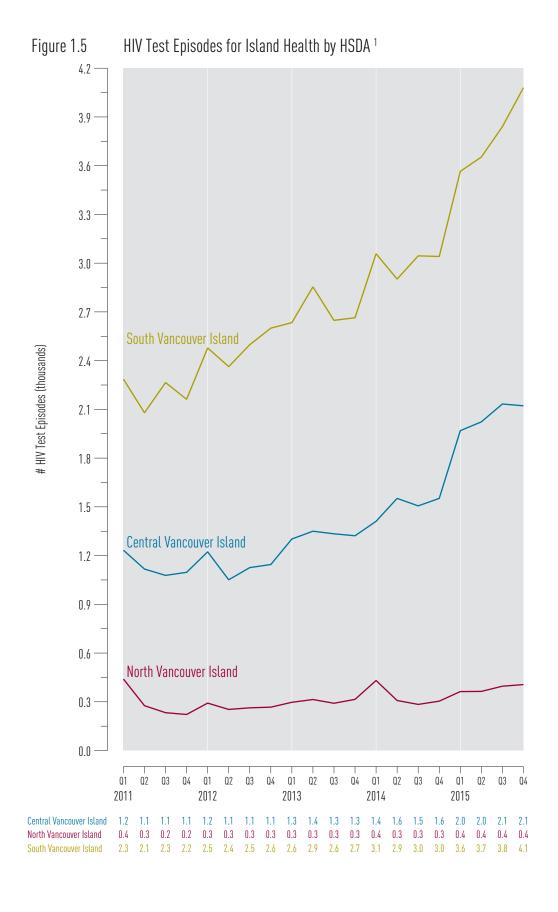
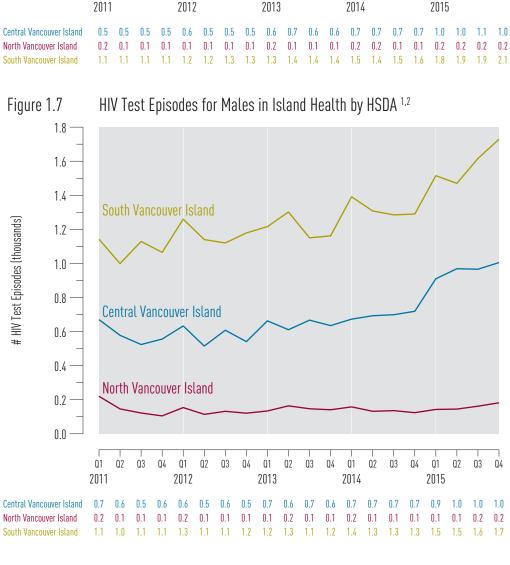
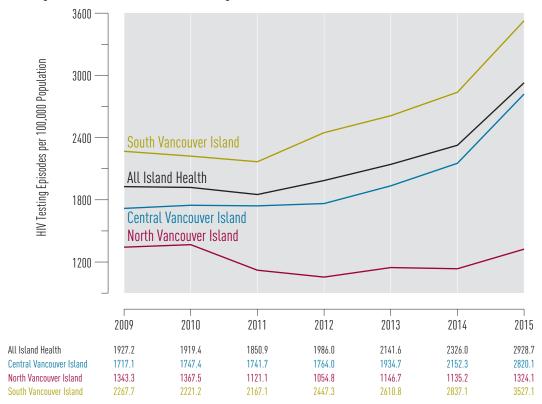


Figure 1.6 HIV Test Episodes for Non-prenatal Females in Island Health by HSDA 1,2 2.2 -2.0 1.8 1.6 # HIV Test Episodes (thousands) South Vancouver Island 1.4 -1.2 -1.0 -0.8 -Central Vancouver Island 0.6 0.4 -North Vancouver Island 0.2 0.0 Q2 Q3 Q4 Q1 Q2 Q3 Q4 Q1 Q2 Q3 Q4 Q1 Q2 Q4 Q1 Q2 2012 2013 2014 2015 2011 0.6 0.7 0.7 0.7 0.5 0.5 0.5 0.6 0.5 0.5 0.5 0.6 0.7 0.6 0.7 1.0 Central Vancouver Island 0.1 0.2 0.1 0.1 0.2 North Vancouver Island 0.1 0.1 0.1 0.1 0.1 0.1 0.1 0.2 0.1 0.1 0.2 South Vancouver Island 1.1 1.1 1.1 1.2 1.2 1.3 1.3 1.3 1.4 1.4 1.4 1.5 1.4 1.5



Indicator 2. HIV Testing Rates

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







5700 -5400 -5100 -4800 -4500 -4200 -30-39 3900 -HIV Testing Episodes per 100,000 Population 3600 -3300 -3000 -2700 -< 30 2400 -40-49 2100 -1800 -1500 -1200 -900 ≥ 50 600 2009 2010 2011 2012 2013 2014 2015 2652.5 4322.3 2449.0 2805.5 3263.8 2490.5 2493.9 2381.6 < 30 30-39 3963.9 4042.8 3836.6 4231.4 4710.6 5533.4 2193.7 823.9 2365.6 954.0 **2597.0** 1095.7 2818.4 1253.4 2239.8 813.8 40-49 2216.7 3536.4 1875.7 830.6

Figure 2.3 Rate of HIV Testing by Age Category for Island Health $^{\rm 2}$

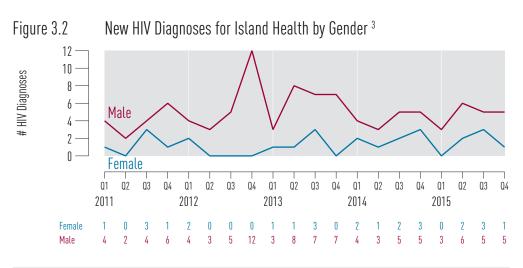
 $Testing\ does\ not\ include\ point\ of\ care\ tests.$

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 Island Health ³ 12 -10 # HIV Diagnoses 8 Q3 Q4 Q2 Q4 02 Q1 Q3 Q4 Q1 Q2 Q3 Q3 Q2 Q3 Q1 Q4 Q1 Q2 2013 2011 2012 2014 2015 Island Health 12 10 3 By Provider Address



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

Figure 3.3 New HIV Diagnoses for Island Health by Age Category ³ # HIV Diagnoses ≥ 50 30 - 39Q3 Q4 Q2 Q1 Q2 Q3 < 30 30-39

Figure 3.4 New HIV Diagnoses for Island Health by Exposure Category 3.4

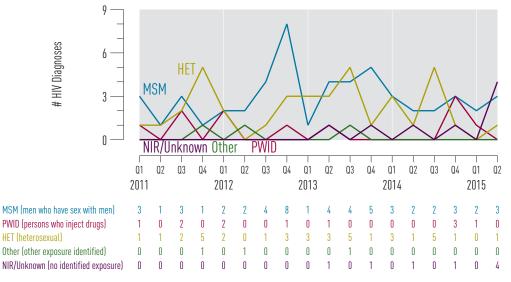
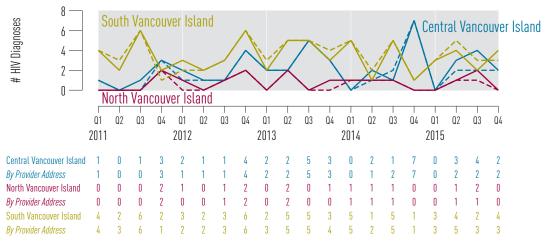


Figure 3.5 New HIV Diagnoses for Island Health by HSDA $^{\rm 3}$



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

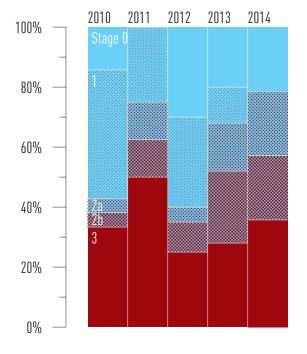
⁴ MSM=men who have sex with men; PWID=people who inject drugs; HET=heterosexual. NIR=No identified risk/exposure.

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 Island 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 i	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. AIDO										
2a		and	CD4 350-499	and	No AIDS case report										
2b	Stage 0		CD4 200-349		ichnir										
3	not met		(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 Island Health, 2010-2014 ⁵

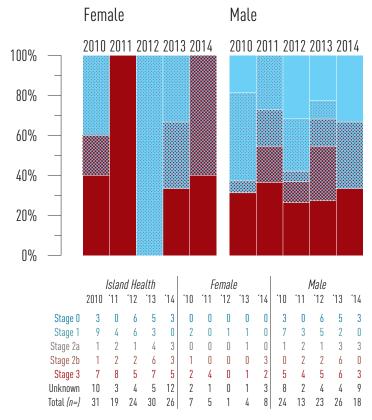


Figure 4.3 Stage of HIV Infection at Diagnosis by Age Category for Island Health, 2010–2014 ⁵

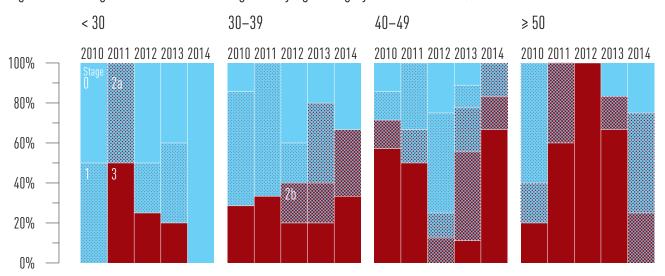
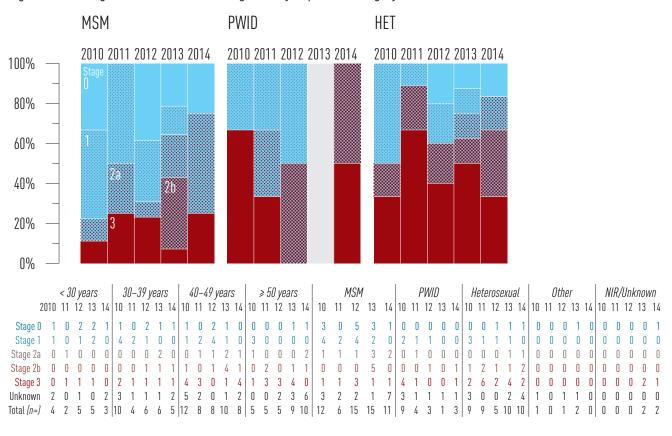


Figure 4.4 Stage of HIV Infection at Diagnosis by Exposure Category for Island Health, 2010–2014 5.6



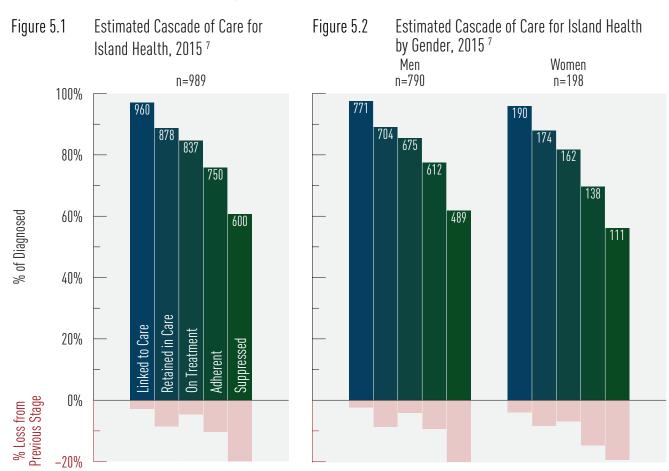
⁵ Data Source: BCCDC

⁶ MSM=men who have sex with men; PWID=people who inject drugs; HET=heterosexual. NIR=No identified risk/exposure.

HIV Cascade of Care

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 period 2015 Q1–2015 Q4 in Island Health and stratified by sex and age.



⁷ Data is for the period 2015 Q1–2015 Q4. Data Sources:

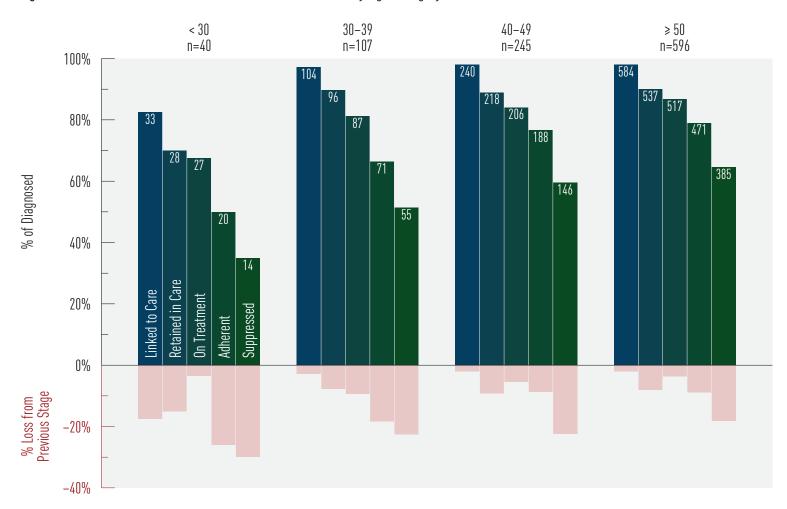
Limitations: HA assignment is based on the most recent HA of residence of the patient, if not available of the HIV-care provider. If the most recent HA of residence is not updated then the designated HA may be incorrect.

NB: Transgender have been assigned to their biological sex.

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 Island Health by Age Category, 2015 8



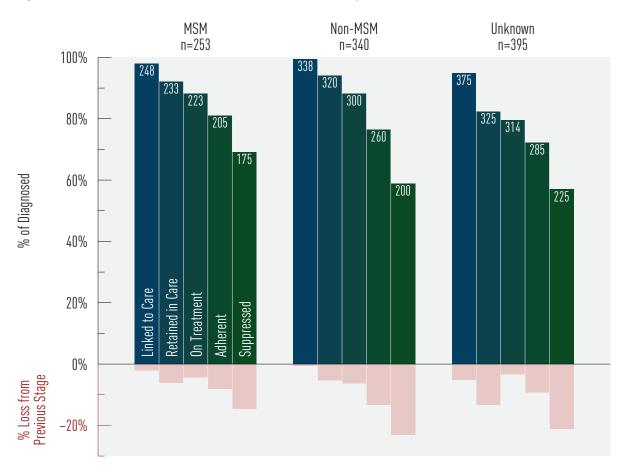
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Limitations: HA assignment is based on the most recent HA of residence of the patient, if not available of the HIV-care provider. If the most recent HA of residence is not updated then the designated HA may be incorrect.

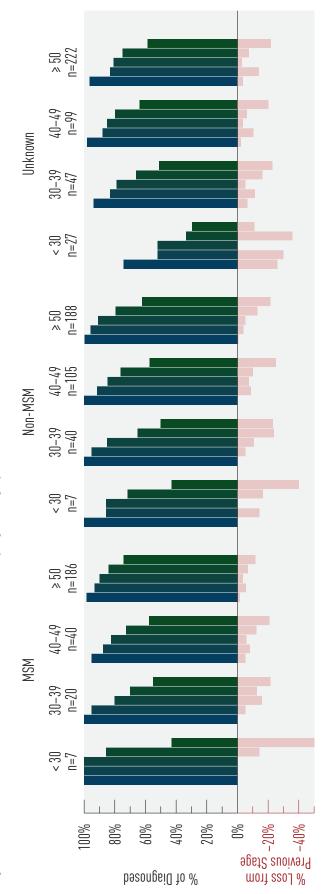
Recent updates to the DTP database have allowed for more comprehensive information on HIV risk group category. As a result, 2014 Q4 data may differ significantly from preceding reports in terms of total numbers ascribed to each risk group.

⁹ Data is for the period 2015 Q1-2015 Q4. 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)).

Estimated Cascade of Care for Island Health by Age Category and MSM Status, 2015 $^{\it 9}$ Figure 5.5



Data is for the period 2015 Q1-2015 Q4.

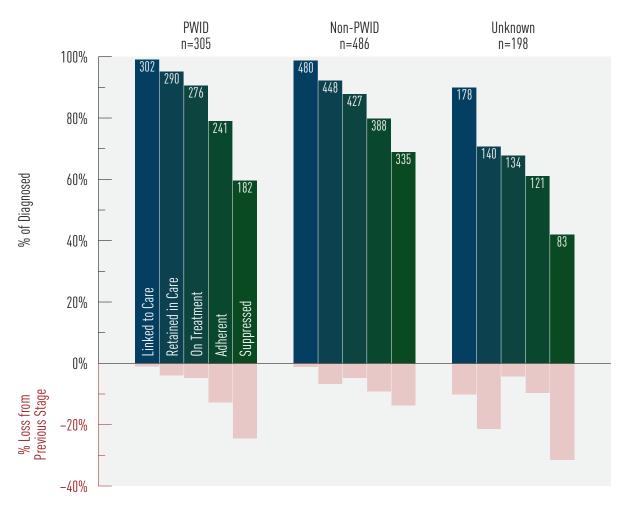
Data Sources:

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. 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.

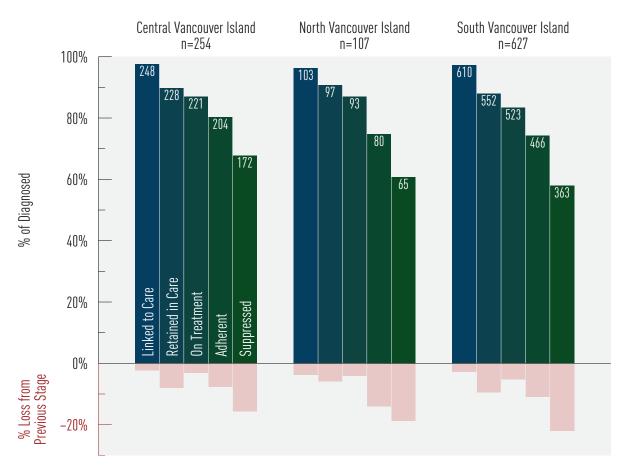
Recent updates to the DTP database have allowed for more comprehensive information on HIV risk group category. As a result, 2014 Q4 data may differ significantly from preceding reports in terms of total numbers ascribed to each risk group.

⁹ Data is for the period 2015 Q1-2015 Q4. 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.

⁹ Data is for the period 2015 Q1-2015 Q4. 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)).

Programmatic Compliance Score

Indicator 6. 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/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. 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 Island Health, 2014 Q1–2015 Q4 10

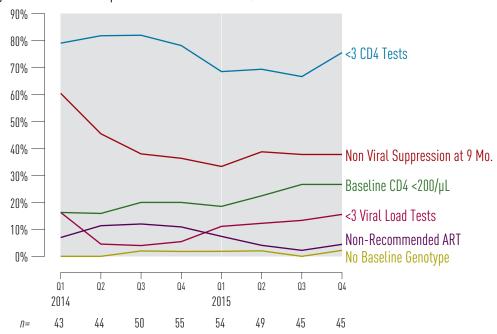
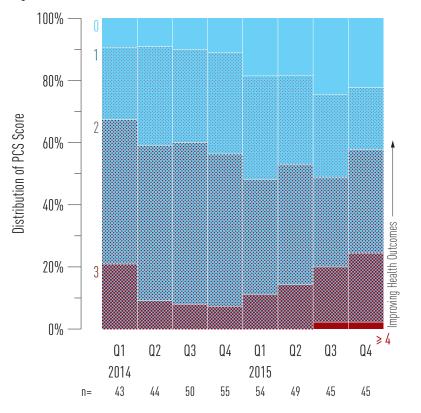


Figure 6.2 Historical Trends for PCS Score for Island Health, 2014 Q1-2015 Q4 10,11



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

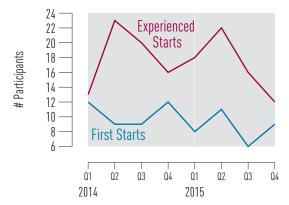
Each quarter's data is calculated as the sum of the 4 quarters leading up to it. e.g. 2013 Q1 is calculated from 2012 Q2 – 2013 Q1. NB: A score of 0 is the best score and a score of 4 or more is the worst score.

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.

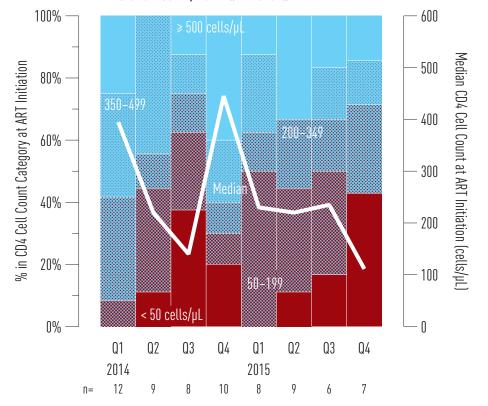
Indicator 7. New Antiretroviral Therapy Starts in Island Health

Figure 7 BC-CfE Drug Treatment Program Enrollment: New ART Participants in Island Health, 2014 Q1–2015 Q4 12



Indicator 8. CD4 Cell Count at ART Initiation

Figure 8 CD4 Cell Count at ART Initiation of ART-Naïve DTP Participants in Island Health, 2014 Q1–2015 Q4 ¹³



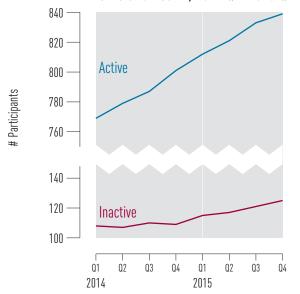
- Data Source: Drug Treatment
 Program Database
 Limitation: DTP participants
 are designated to an HA based
 on most current residence
 provided by the participant.
- 3 Data Source: Drug Treatment Program Database Limitations: CD4 cell count data is approximately 80% complete.

Indicator 9. Active and Inactive DTP Participants

Table 3. Distribution of People on ART for Island Health, 2015 Q4 14

Age	< 30	26
	30-39	94
	40-49	223
	≥ 50	496
Gender	Male	680
	Female	159
Exposure	MSM	228
	PWID	274
Total		839

Figure 9 Active and Inactive DTP Participants for Island Health, 2014 Q1-2015 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.

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.

Definition:

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

15 Active DTP participants: An individual who has had medication prescribed at least once in the preceding quarter.

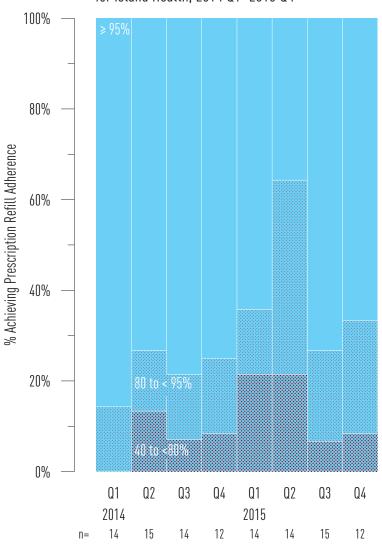
Inactive DTP participants: Persons no longer prescribed drugs through the HIV/AIDS Drug Treatment Program in the last quarter.

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 Island Health, 2014 Q1–2015 Q4 ¹⁶



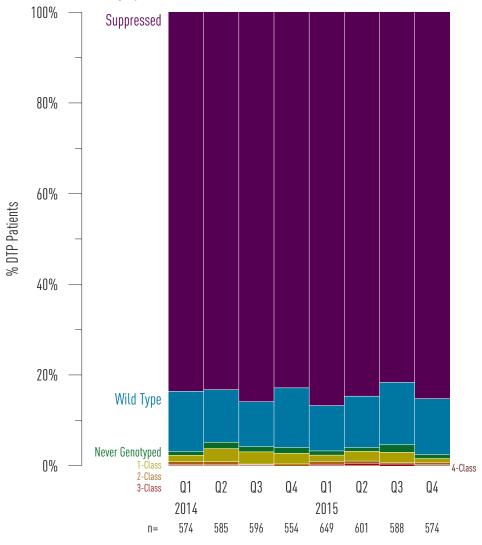
¹⁶ Data Source: Drug Treatment Program Database Limitation: Prescription refill adherence is used as a proxy for patient adherence.

Resistance Testing and Results

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, three, or four 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.





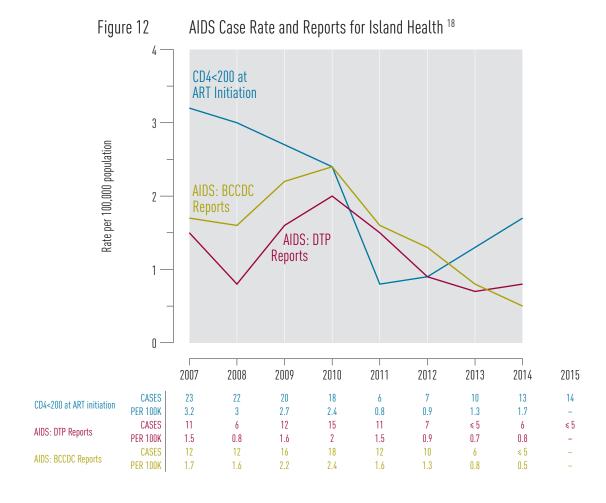
¹⁷ Data Source: Drug Treatment Program Database

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

AIDS-Defining Illness

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.



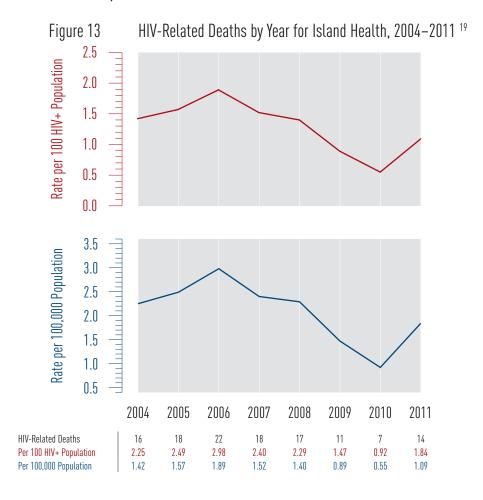
Data Source: DTP AIDS cases are obtained from the Drug Treatment Program Database; BCCDC AIDS cases are obtained from the BC-CDC; CD4<200 at ART initiation data came from the DTP database.

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

HIV-Related Mortality

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

	(thousands)	2011 Q1	Q2	Q3	Q4	2012 Q1	Q2	Q3	Q4	2013 Q1	Q2	Q3	Q4	2014 Q1	Q2	Q3	Q4	2015 Q1	Q2	Q3	Q4
Island Hea	lth	4.0	3.5	3.6	3.5	4.0	3.7	3.9	4.0	4.2	4.5	4.3	4.3	4.9	4.8	4.8	4.9	5.9	6.0	6.4	6.
Gender	Female	1.9	1.7	1.8	1.7	1.9	1.9	1.9	2.0	2.1	2.2	2.1	2.1	2.3	2.3	2.4	2.4	2.9	3.0	3.2	3.
	Male	2.0	1.7	1.8	1.7	2.0	1.8	1.9	1.8	2.0	2.1	2.0	1.9	2.2	2.1	2.1	2.1	2.6	2.6	2.7	2.
	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.
Age	< 30	1.6	1.4	1.5	1.5	1.6	1.4	1.5	1.5	1.6	1.7	1.6	1.6	1.8	1.6	1.7	1.8	2.0	1.9	2.0	2
	30-39	1.0	0.8	0.9	0.8	1.0	0.9	1.0	1.0	1.0	1.0	1.0	0.9	1.2	1.1	1.1	1.1	1.3	1.3	1.4	1
	40-49	0.7	0.6	0.6	0.5	0.6	0.6	0.6	0.6	0.6	0.6	0.6	0.7	0.7	0.7	0.7	0.7	0.8	0.8	0.9	0
	≥ 50	0.7	0.6	0.6	0.6	0.7	0.7	0.7	0.8	0.9	1.0	0.8	0.8	0.9	1.0	1.0	1.1	1.4	1.6	1.7	1
POC HIV (not in tho		21	17	42	12	24	41	90	164	135	200	194	245	347	353	316	325	406	414	371	38
Central Va	incouver Islan	d 1.2	1.1	1.1	1.1	1.2	1.1	1.1	1.1	1.3	1.4	1.3	1.3	1.4	1.6	1.5	1.6	2.0	2.0	2.1	2
Female		0.5	0.5	0.5	0.5	0.6	0.5	0.5	0.5	0.6	0.7	0.6	0.6	0.7	0.7	0.7	0.7	1.0	1.0	1.1	1
Male		0.7	0.6	0.5	0.6	0.6	0.5	0.6	0.5	0.7	0.6	0.7	0.6	0.7	0.7	0.7	0.7	0.9	1.0	1.0	1
North Van	couver Island	0.4	0.3	0.2	0.2	0.3	0.3	0.3	0.3	0.3	0.3	0.3	0.3	0.4	0.3	0.3	0.3	0.4	0.4	0.4	0
Female		0.2	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.2	0.1	0.1	0.1	0.2	0.2	0.1	0.1	0.2	0.2	0.2	0
Male		0.2	0.1	0.1	0.1	0.2	0.1	0.1	0.1	0.1	0.2	0.1	0.1	0.2	0.1	0.1	0.1	0.1	0.1	0.2	0
South Van	couver Island	2.3	2.1	2.3	2.2	2.5	2.4	2.5	2.6	2.6	2.9	2.6	2.7	3.1	2.9	3.0	3.0	3.6	3.7	3.8	4
Female		1.1	1.1	1.1	1.1	1.2	1.2	1.3	1.3	1.3	1.4	1.4	1.4	1.5	1.4	1.5	1.6	1.8	1.9	1.9	2
Male		1.1	1.0	1.1	1.1	1.3	1.1	1.1	1.2	1.2	1.3	1.1	1.2	1.4	1.3	1.3	1.3	1.5	1.5	1.6	1.
sland Hea Central Va	ncouver Islan	d	1927 1717		1919.4 1850.9 1747.4 1741.7				1986.0 1764.0			2141.6 2326.0 1934.7 2152.3			2928.7 2820.1						
	couver Island couver Island		1343 2267		136 222			121.1 167.1		1054.8 2447.3			6.7		1135.2		1324. 3527.				
Gender	Female		1911		187			785.8		2447. 1968.		261 217		2837.1 2377.8			3060				
Gender	Male				194							210					3000 2780.:				
A ~~			1926					911.6		1996.					264.1						
Age	< 30		2490		249			381.6	2449.0			265			305.5		3263. 5522				
	30-39		3963		404			336.6				432			10.6		5533. 2526				
	40-49		2216		223			193.7		2365.		259			318.4		3536.				
	≥ 50		830		2011	3.8	8	323.9 201	12	954.		109	5./	12	253.4 2014		1875.)15		
Indicator 3	3: New HIV I	Diagnoses				Q2 (Q3 Q		1 Q2	Q3)2 Q	3 Q4			Q3 (2 Q3	Ç
Island Hea		By Client		ence	5	2			5 3	5	12	4	9 1			4	7	8		8 8	
		By Provide			5	3	6	6 3	3 3	5	12	5	9 1			4	8	8		8 6	
Gender		Female			1	0			2 0	0	0	1		3 (1	2	3		2 3	
		Male			4	2			4 3	5	12	3		7 7		3	5	5		6 5	
Age		< 30			1	0			2 2	1	2	1		2 1		0	0	2		1 3	
o-		30–39			1	1			2 1	1	2	1		3 (0	4	1		2 4	
		40-49			3	1			2 0	1	5	1		4 4		2	1	3		3 1	
		≥ 50			0	0			0 0	2	3	1		1 2		2	2	2		2 0	
Exposure		MSM			3	1			2 2		8	1		4 5		2	2	3		3 -	
		PWID			1	0			2 0	0	1	0		0 0		0	0	3		0 -	
		1 1111			1	0	_	5	_ 0	U	1	0		5 (, 0	U	J	9		_	

HET Other

NIR/Unknown

								2	2011				20	12			2	2013				2014	4			20	015			
Indicator 3	: Ne	w HI	V D	iag	nose	es (co	nt'd	1)	Q1	Q2	Q3	Q ²	4 Q	1 Q	2 (Q3 (Q4	Q1	Q2	Q3	Q4	Q1	Q2	Q3	3 Q) 4 (Q1 (Q2	Q3	Q4
Central Var Island	ıcou	ver		•		t Res			1 1	0	1			2 1	1	1	4	2 2	2 2	5 5	3	0	2			7 7	0	3 2	4 2	2 2
				,		der A				0					1	1			2											
North Vanc Island	ouv	er		•		t Res			0	0	0			1	0	1	2	0		0	1	1	1			0	0	1	2	0
				-		der A			0	0	0			0	0	1	2	0	2	0	0	1	1		1	0	0	1	1	0
South Vanc Island	ouve	er		•		it Res der A			4 4	2 3	6			3 2	2 2	3	6 6	2 3	5 5	5 5	3 4	5 5	1 2		5	1 1	3 3	4 5	2	4
Indicator 4:	Sta	ge of	ніч	V In	fect	ion a	t Ba	selir	ıe																					
		Island						emal					Male					0 ye				30-3					40-			1
	'10	'11	'12	'13	'14	'10	' 11	'12	'13	' 14	'10	' 11	'12	'13	'14	'10	' 11	'12	'13	' 14	'10	'11 '	12	'13 '	'14	'10	'11	'12	'13	'14
Stage 0	3	0	6	5	3	0	0	0	0	0	3	0	6	5	3	1	0	2	2	1	1	0	2	1	1	1	0	2	1	0
Stage 1	9	4	6	3	0	2	0	1	1	0	7	3	5	2	0	1	0	1	2	0	4	2	1	0	0	1	2	4	1	0
Stage 2a	1	2	1	4	3	0	0	0	1	0	1	2	1	3	3	0	1	0	0	0	0	0	0	2	0	0	1	1	2	1
Stage 2b	1	2	2	6	3	1	0	0	0	3	0	2	2	6	0	0	0	0	0	0	0	0	1	1	1	1	0	1	4	1
Stage 3	7	8	5	7	5	2	4	0	1	2	5	4	5	6	3	0	1	1	1	0	2	1	1	1	1	4	3	0	1	4
Unknown	10	3	4	5	12	2	1	0	1	3	8	2	4	4	9	2	0	1	0	2	3	1	1	1	2	5	2	0	1	2
Total	31	19	24	30	26	7	5	1	4	8	24	13	23	26	18	4	2	5	5	3	10	4	6	6	5	12	8	8	10	8
		≥ 50	0 ye	ars			N	иsм				P	WID)]	Hete	rose	xual		Ot	her I	Ехро	osure	9	N	IIR/I	Unk	nowi	n
	'10	' 11			' 14	'10	' 11	'12	'13	' 14	'10	' 11	'12	'13	' 14	'10	' 11	'12	' 13	'14				'13 '		'10	' 11	'12	'13	' 14
Stage 0	0	0	0	1	1	3	0	5	3	1	0	0	0	0	0	0	0	1	1	1	0	0	0	1	0	0	0	0	0	1
Stage 1	3	0	0	0	0	4	2	4	2	0	2	1	1	0	0	3	1	1	1	0	0	0	0	0	0	0	0	0	0	0
Stage 2a	1	0	0	0	2	1	1	1	3	2	0	1	0	0	0	0	0	0	1	1	0	0	0	0	0	0	0	0	0	0
Stage 2b	0	2	0	1	1	0	0	0	5	0	0	0	1	0	1	1	2	1	1	2	0	0	0	0	0	0	0	0	0	0
Stage 3	1	3	3	4	0	1	1	3	1	1	4	1	0	0	1	2	6	2	4	2	0	0	0	0	0	0	0	0	2	1
Unknown	0	0	2	3	6	3	2	2	1	7	3	1	1	1	1	3	0	0	2	4	1	0	1	1	0	0	0	0	0	0
Total	5	5	5	9	10	12	6	15	15	11	9	4	3	1	3	9	9	5	10	10	1	0	1	2	0	0	0	0	2	2
Indicator 5		V Ca	sca	de o	f Ca	re		DI	AGN	IOSE	D		LI	NKE	D	F	RETA	INEI)		ON .	ART		ADI	IERI	ENT		SUPF	RESS	ED
Island Hea	lth									98	9			96	0			878	3			837				750			6	500
Age Catego	ory	< 30								4	0:			3.	3			28	3			27				20				14
		30-3	39							10	7			10	4			96	5			87				71				55
		40-4	19							24	5			24	0			218	3			206				188			1	146
		≥ 50								59	6			58	4			537	7			517				471			3	385
Age Catego	ory	MSN	M		<	< 30					7				7			7	7			7				6				3
and MSM					3	30-39	9			2	0.			2	0			19)			16				14				11
Status					4	10-49	9			4	0			3	8			35	5			33				29				23
					2	≥ 50				18	6			18	3			173	3			167				156			1	38
		Non	-MS	SM	<	< 30					7				7			6	5			6				5				3
						30-39	9			4	0:			4	0			38	3			34				26				20
						10-49				10				10				96				89				80				60
						≥ 50				18				18				180				171				149				17
		Unk	now	'n		< 30					7			2				14				14				9				8
						30-39)				7			4				39				37				31				24
						10-49					9			9				87				84				79				63
						≥ 50				22				21				184				179				166				30
Gender		Male	9							79				77				704				675				612				189
Guidei		Fem								19				19				174				162				138				11
Injection		PW								30				30:				290				276				241				82
Drug Use		Non		ΔID						48				48				448				427				388				335
2148 000		Unk								19																				83
MSM Statu	10	MSN		11						25				178 248				140 233				134 223				121205				75
IVIOIVI Statu	13	Non		21/1						34				33								300				260				
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TT. 1/1		Unk				+	1.	1		39				37				325				314				285				225
Health		Cen						1		25				248				228				221				204				.72
Authority		Nor								10				10:				97				93				80				65
		Sout	n Va	anco	ouve	r Isla	ınd			62	/			610	U			552	2			523			4	466			3	363

Indicator 6: Programma	tic Compliance Scor	re (PCS)								
	2014 Q1	Q2		Q3	Q4	2015 Q1	Q	2	Q3	Q4
< 3 CD4 Tests	79.1%	81.8%	8	2.0%	78.2%	68.5%	69.49	6	66.7%	75.6%
< 3 Viral Load Tests	16.3%	4.5%		4.0%	5.5%	11.1%	12.29	6	13.3%	15.6%
No Baseline Genotype	0.0%	0.0%		2.0%	1.8%	1.9%	2.09	6	0.0%	2.2%
Baseline CD4 < 200 cells	s/μL 16.3%	15.9%	2	0.0%	20.0%	18.5%	22.49	6	26.7%	26.7%
Non-Recommended AR	T 7.0%	11.4%	1	2.0%	10.9%	7.4%	4.19	6	2.2%	4.4%
Non Viral suppression at	t 9 Mo. 60.5%	45.5%	3	8.0%	36.4%	33.3%	38.89	6	37.8%	37.8%
PCS Score: 0	4	4		5	6	10		9	11	10
PCS Score: 1	10	14		15	18	18	1		12	9
PCS Score: 2	20	22		26	27	20	1		13	15
PCS Score: 3	9	4		4	4	6		7	8	10
PCS Score: 4 or more	0	0		0	0	0		0	1	10
Total (n=)	43	44		50	55	54	4		45	45
Indicator 7: New DTP A										
First Starts	12	9		9	12	8	1	1	6	9
Experienced Starts	13	23		20	16	18	2		16	12
•	ADEL									
Indicator 8: CD4 Cell Co $CD4 \ge 500$	ount at ART Initiatio	on for ARV 0	-Naïve	DTP Parti	cipants 4	1		3	1	1
CD4 350-499	3 4	4		1	2	2		0	1	1
	_									
CD4 200-349	4	1		1	1	1		2	1	2
CD4 50-199	1	3		2	1	4		3	2	C
CD4 < 50	0	1		3	2	0		1	1	3
CD4 Median (cells/μL)	395	220		140	445	230	22		235	230
Total (n=)	12	9		8	10	8		9	6	7
Indicator 9: Active and I	nactive DTP Partici	pants								
Active DTP Participants	769	779		787	801	812	82	1	833	839
Inactive DTP Participant	ts 108	107		110	109	115	11	7	121	125
Indicator 10: Antiretrov	iral Adherence									
≥ 95%	12	11		11	9	9		5	11	8
80% to < 95%	2	2		2	2	2			3	
								6		3
40% to < 80%	0	2		1	1	3		3	1	1
< 40% Total (n=)	0 14	0 15		0 14	0 12	0 14	1-	0	0 15	12
		13		14	12	14	1.	1	13	12
Indicator 11: Resistance										
Suppressed	480	487		512	459	563	50		480	490
Wild Type	76	68		59	73	65	6		81	71
Never Genotyped	5	8		7	7	6		5	10	5
1-Class	8	17		16	13	9	1	3	13	5
2-Class	4	4		1	2	4		3	2	3
3-Class	1	1		1	0	2		3	2	C
Total (n=)	574	585		596	554	649	60	1	588	575
Indicator 12: AIDS-Defi	ning Illness	2007	2008	2009	2010	2011	2012	2013	2014	2015
	Cases	23	22	20	18	6	7	10	13	14
ART initiation	Rate per 100,000	3.2	3.0	2.7	2.4	0.8	0.9	1.3	1.7	_
	Cases	11	6	12	15	11	7	≤ 5	6	≤ 5
	Rate per 100,000	1.5	0.8	1.6	2.0	1.5	0.9	0.7	0.8	_
•	Cases	12	12	16	18	12	10	6	≤ 5	_
	Rate per 100,000	1.7	1.6	2.2	2.4	1.6	1.3	0.8	0.5	-
Indicator 12 IIIV D 1	ad Mantalit	2004	2005	2006	2007	2000	2000	2010	2011	
Indicator 13: HIV-Relate	eu Mortality	2004	2005	2006	2007	2008	2009	2010	2011	
Island Health		16	18	22	18	17	11	7	14	
Per 100 HIV+ Population	n	2.25	2.49	2.98	2.40	2.29	1.47	0.92	1.84	
Per 100,000 Population		1.42	1.57	1.89	1.52	1.40	0.89	0.55	1.09	