



BRITISH COLUMBIA
CENTRE *for* EXCELLENCE
in HIV/AIDS

HIV MONITORING QUARTERLY REPORT **FOR INTERIOR HEALTH**

FOURTH QUARTER 2013



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



Foreword

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

The objectives of these reports are to:

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

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

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Acknowledgements and Contributions



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



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

Other Data Sources:

The above databases were supplemented with:

- (I) The BC Vital Statistics database which was used to calculate Indicator 5. The HIV Cascade of Care and Indicator 13. HIV-Related Mortality.
- (II) Linkage and preparation of the de-identified individual-level database used for calculating Indicator 5. The HIV Cascade of Care was facilitated by the British Columbia Ministry of Health.
- (III) The Statistics Canada database: BC and HIV-positive population counts were acquired through the statistics Canada website to calculate HIV-specific mortality rates for Indicator 13. HIV-Related Mortality.

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

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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 highly active antiretroviral therapy (HAART) initiation.

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

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

HIV Testing Episodes and Rates

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

Indicator 1. HIV Testing Episodes

Figure 1.1 HIV Test Episodes for Interior Health, 2009 Q1–2013 Q4

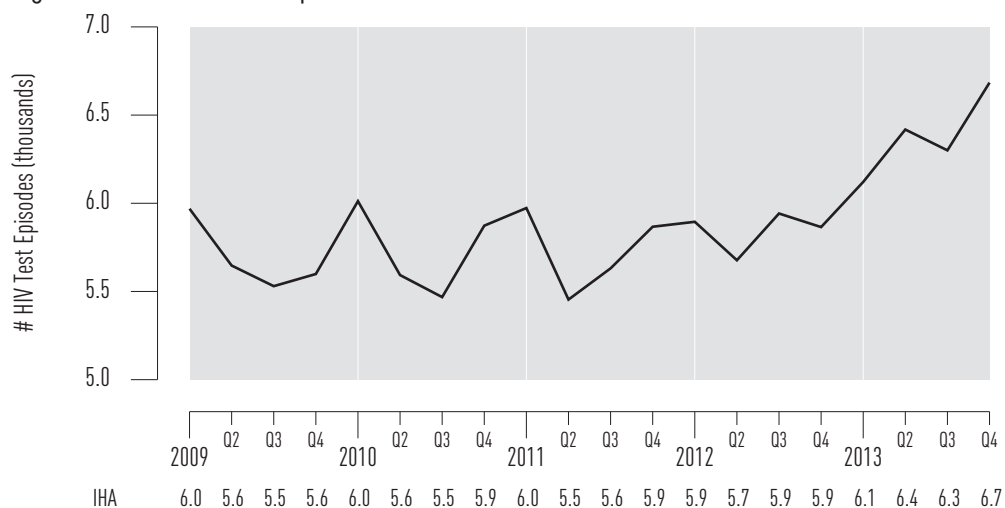
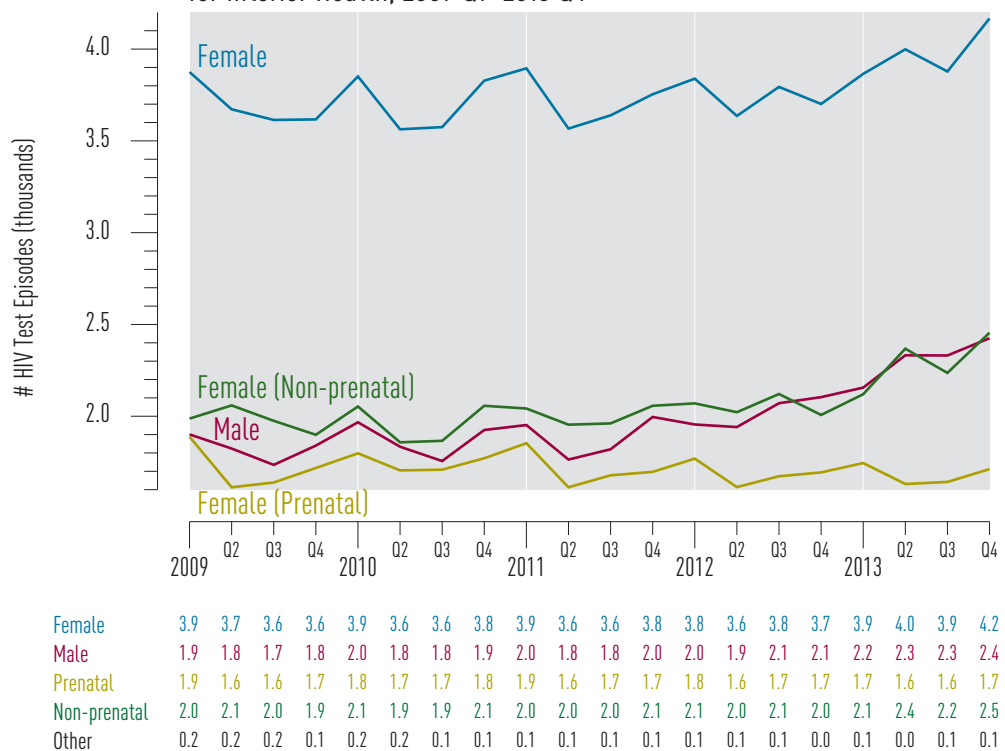


Figure 1.2 HIV Test Episodes by Gender and Prenatal Status for Interior Health, 2009 Q1–2013 Q4 ¹



¹ NB: Testing does not include point of care tests.

Figure 1.3 HIV Test Episodes by Age Category for Interior Health, 2009 Q1–2013 Q4 ^{1,2}

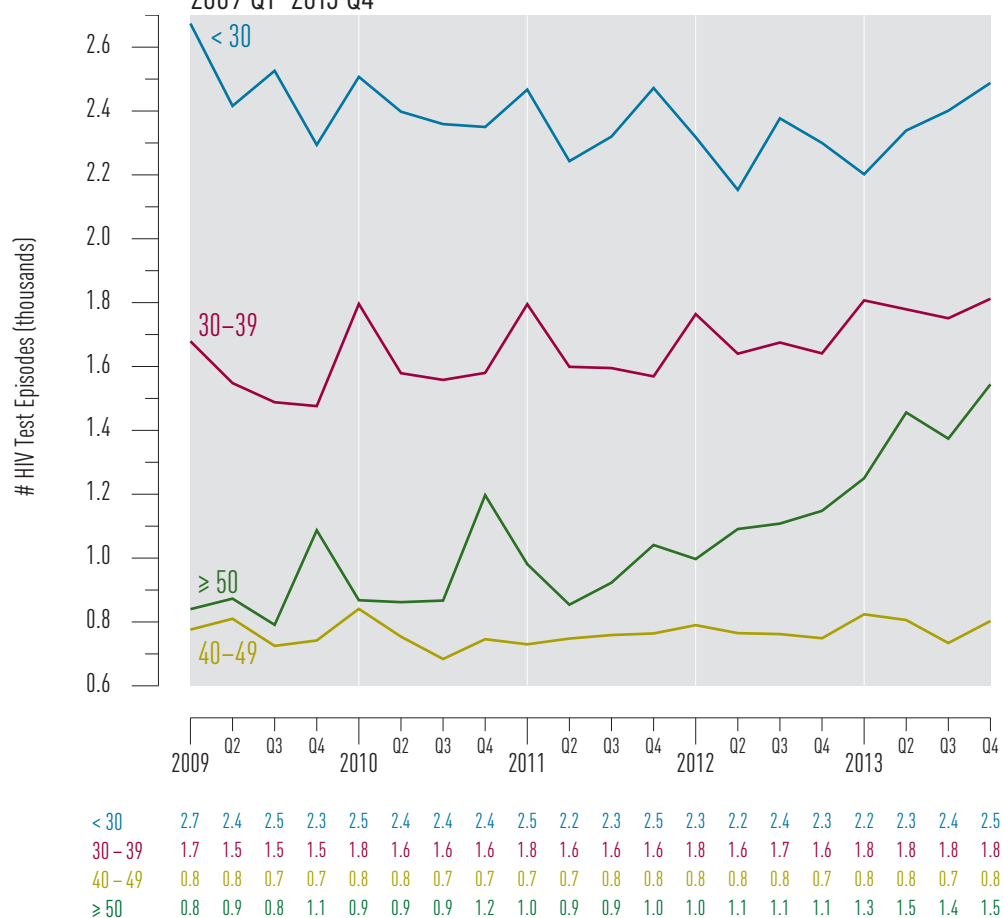
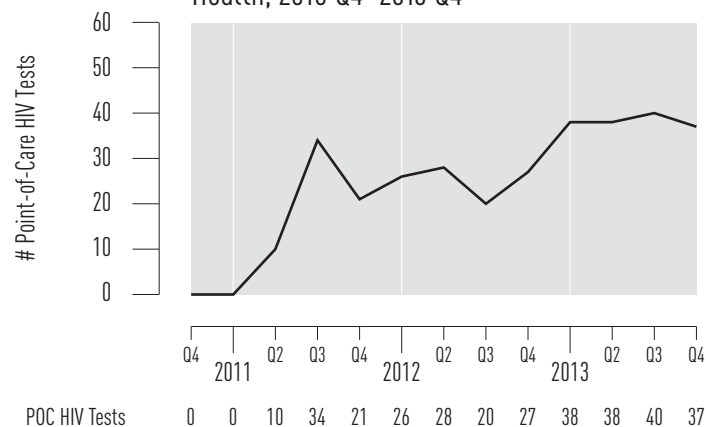


Figure 1.4 Point-of-Care HIV Tests for Interior Health, 2010 Q4–2013 Q4



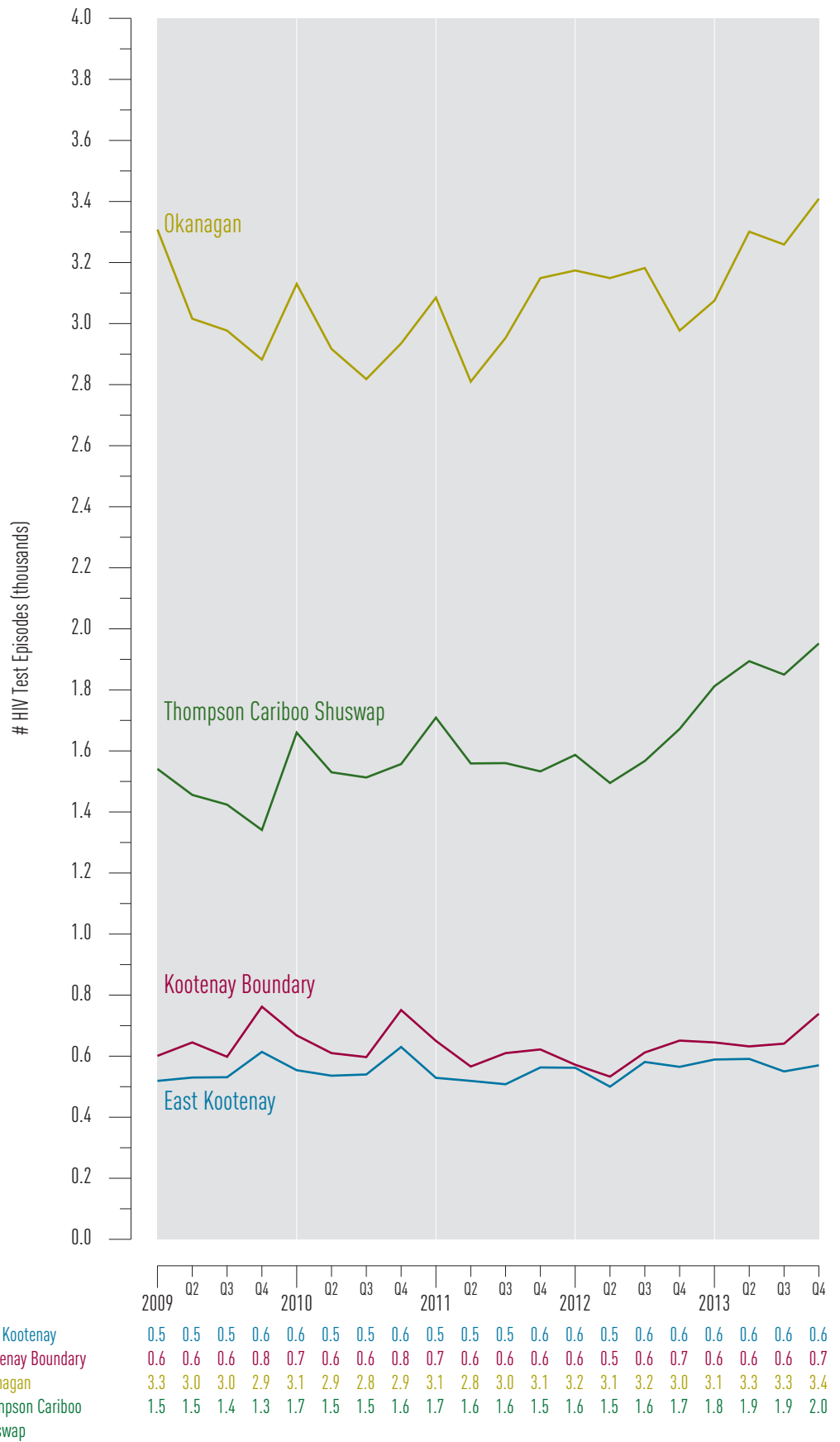
2 Data Source: The BC Public Health Microbiology and Reference Laboratory (BCPHMRL) courtesy of the BC Centre for Disease Control (BCCDC).

Limitations:

- 1 Repeat tests in individuals who test using various identifiers may not be identified and these individuals may be counted more than once.
- 2 POC testing data is available from the fourth quarter of 2010 and onwards.

Figure 1.5

HIV Test Episodes for Interior Health, 2009 Q1–2013 Q4



Indicator 2. HIV Testing Rates

Figure 2.1 Rate of HIV Testing for Interior Health and HSDAs, 2009–2013 ¹

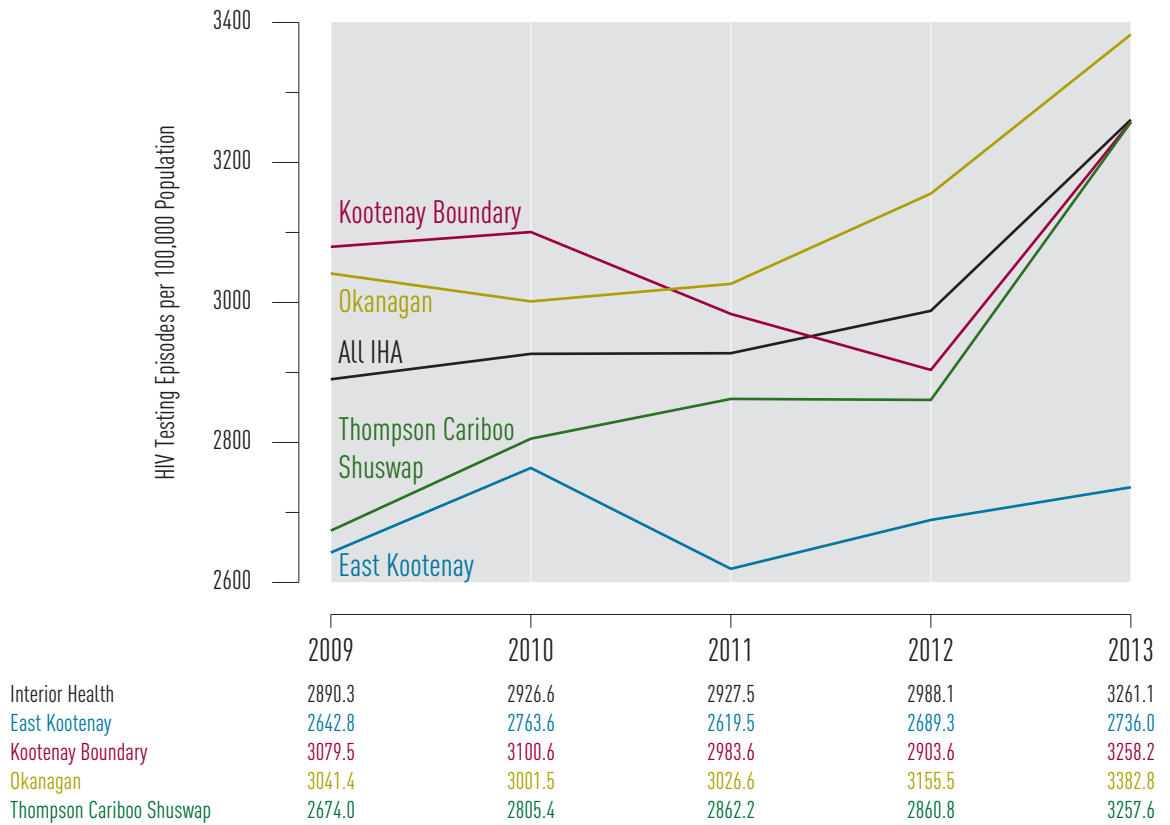


Figure 2.2 Rate of HIV Testing by Gender for Interior Health, 2009–2013 ¹

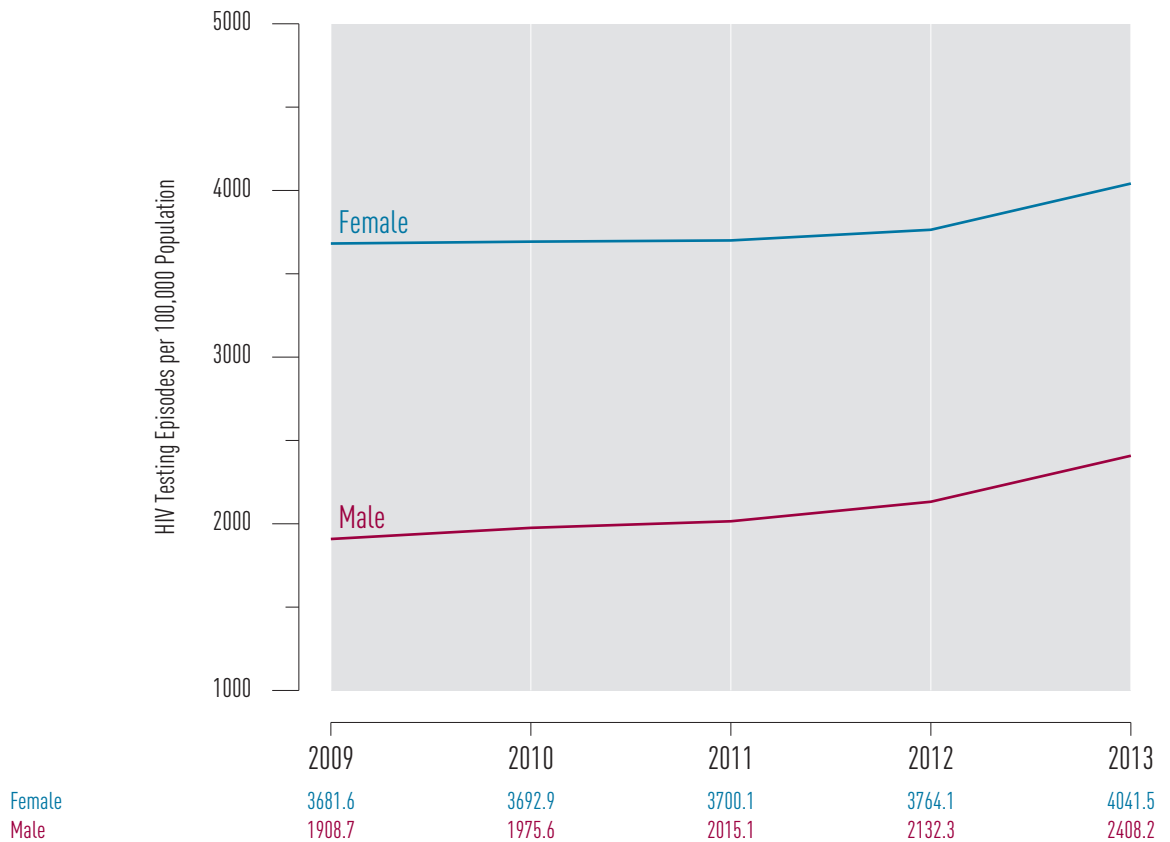
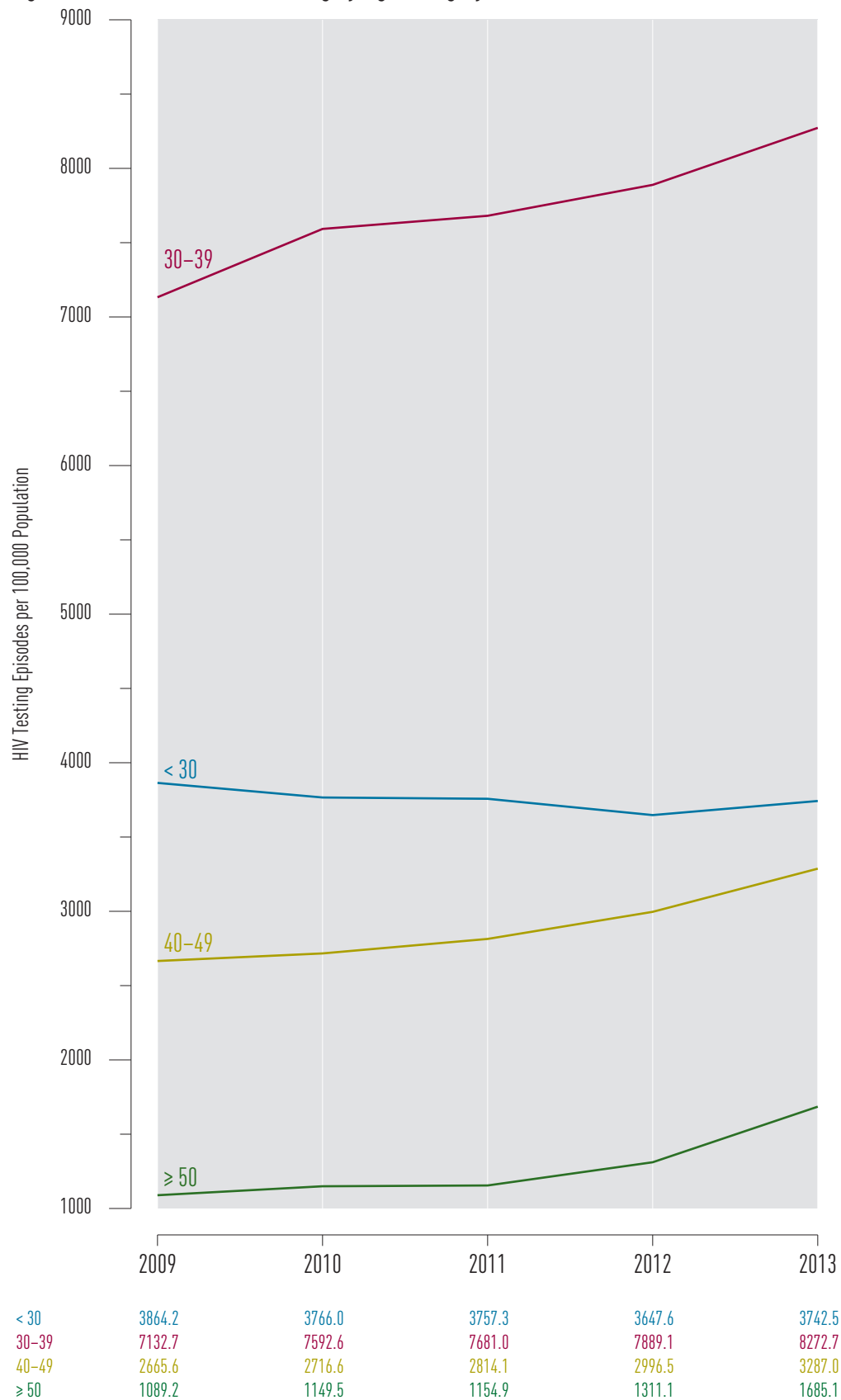


Figure 2.3 Rate of HIV Testing by Age Category for Interior Health, 2009–2013 ¹



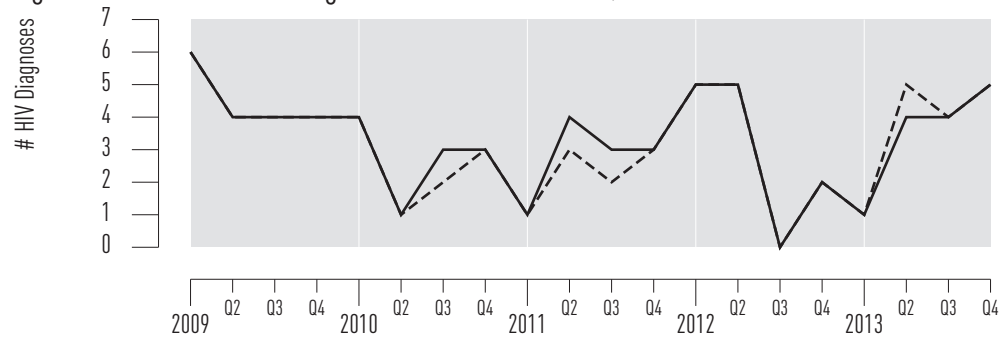
New HIV Diagnoses

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 Interior Health, 2009 Q1–2013 Q4 ³

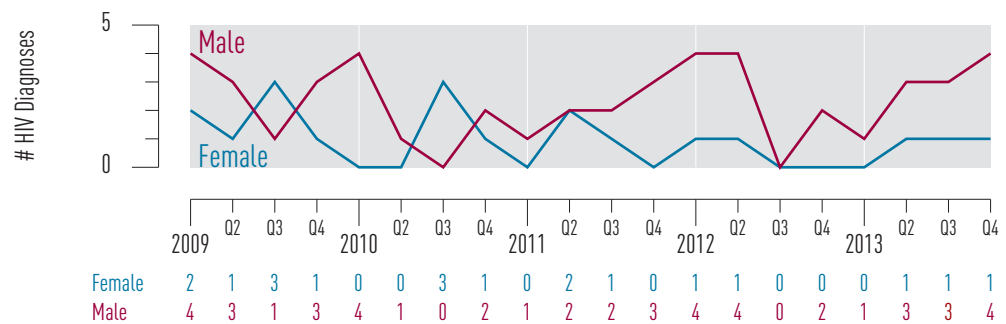


Interior Health By Client Residence
By Provider Address

6	4	4	4	4	1	3	3	1	4	3	3	5	5	0	2	1	4	4	5
6	4	4	4	4	1	2	3	1	3	2	3	5	5	0	2	1	5	4	5

Figure 3.2

New HIV Diagnoses for Interior Health by Gender, 2009 Q1–2013 Q4



Female
Male

2	1	3	1	0	0	3	1	0	2	1	0	1	1	0	0	0	1	1	1
4	3	1	3	4	1	0	2	1	2	2	3	4	4	0	2	1	3	3	4

Figure 3.3 New HIV Diagnoses for Interior Health by Age Category, 2009 Q1–2013 Q4

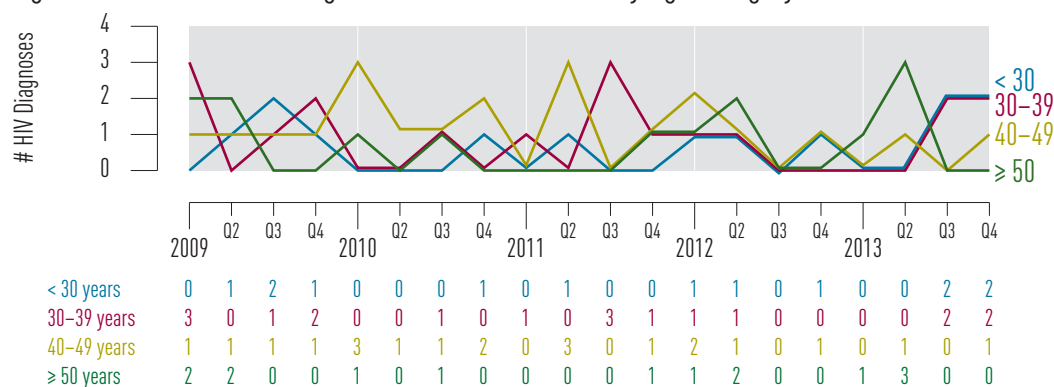


Figure 3.4 New HIV Diagnoses for Interior Health by Exposure Category, 2009 Q1–2013 Q2⁴

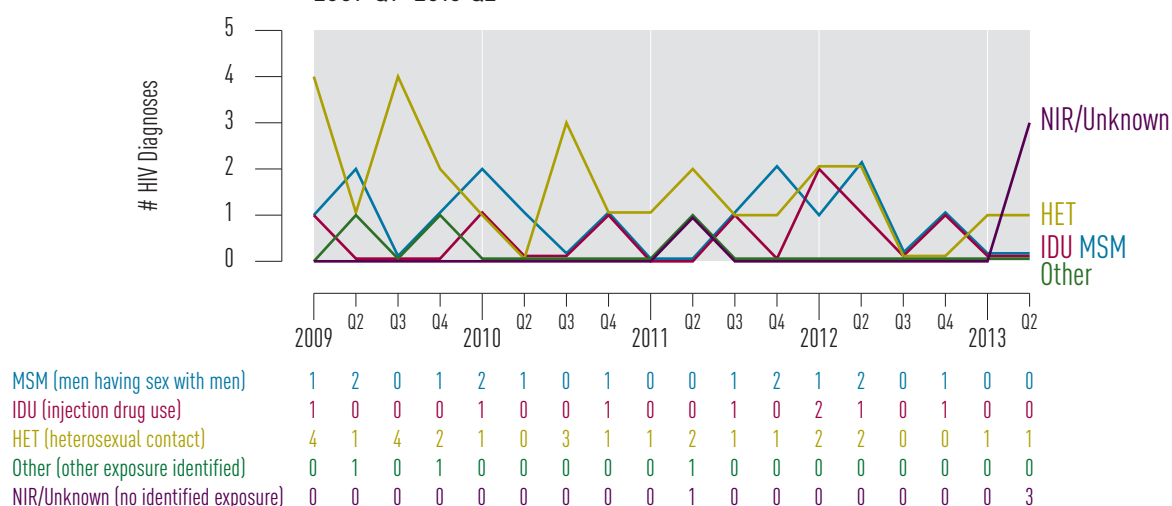
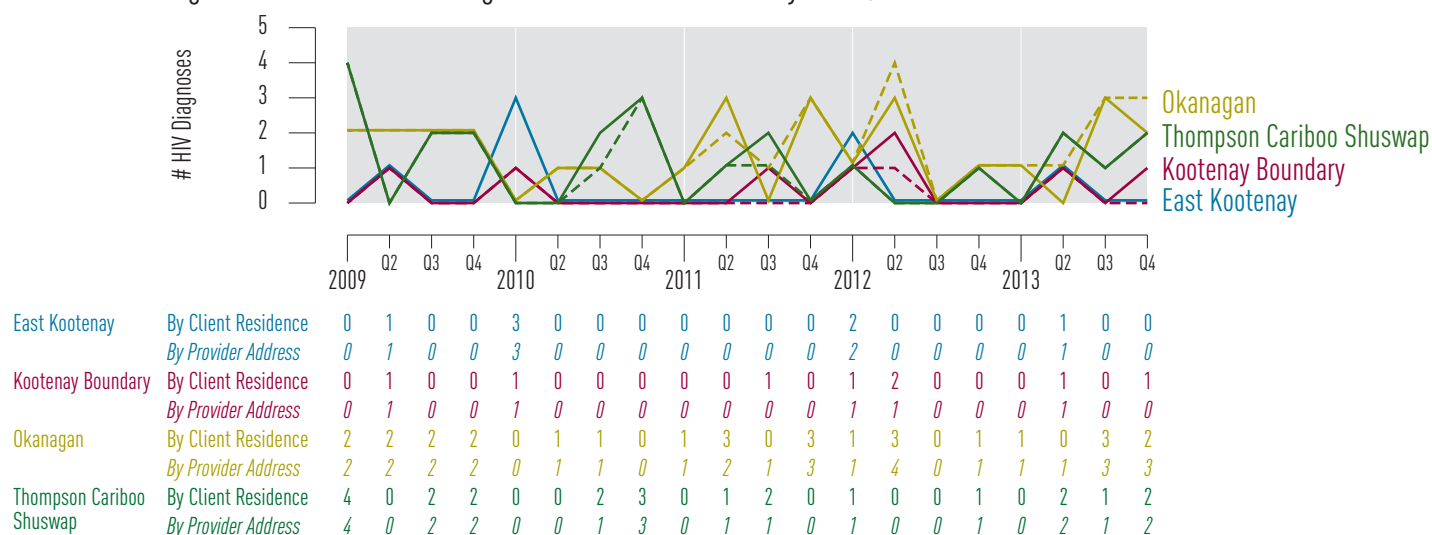


Figure 3.5 New HIV Diagnoses for Interior Health by HSDA, 2009 Q1–2013 Q4



⁴ "By Provider Address" is graphed as dashed line in same colour.

⁴ BCCDC: Data lags by 6 months.

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

Stage of HIV infection at diagnosis

Classification of stage of HIV infection, in the absence of information regarding recent testing history, is reliant on clinical information available at the time of diagnosis, including first CD4+ cell count, laboratory results suggestive of acute HIV infection, and clinical presentation with an AIDS-defining illness (Table 1). The benefits of Treatment as Prevention (TasP) are maximized when antiretroviral therapy (ART) is initiated at high CD4 cell counts. Accordingly, it is preferable that individuals newly diagnosed with HIV be in the early stages of HIV infection (stage 0 or 1) to allow for early ART initiation.

N.B. Interpretation of stage of HIV infection at diagnosis should proceed with caution. Early increases in diagnosis at late stage (i.e., low CD4 counts) may represent a “catching up” of previously missed long term infected individuals rather than a trend toward diagnosis at later stage of infection.

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

Figure 4.1 Stage of HIV Infection at Diagnosis for Interior Health, 2010–2013⁵

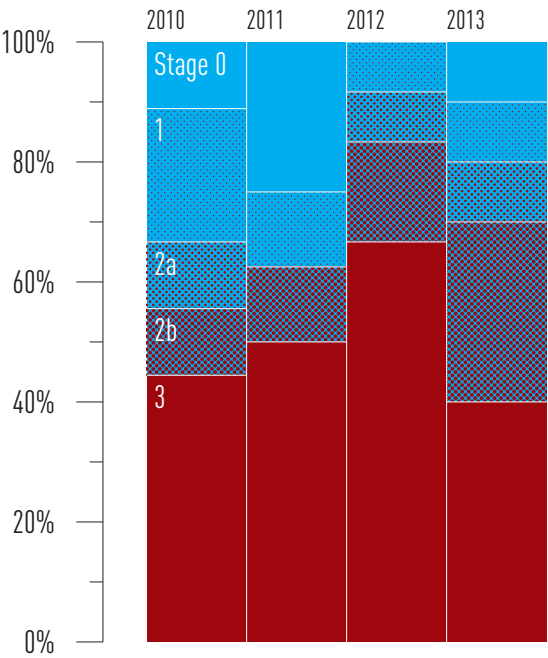
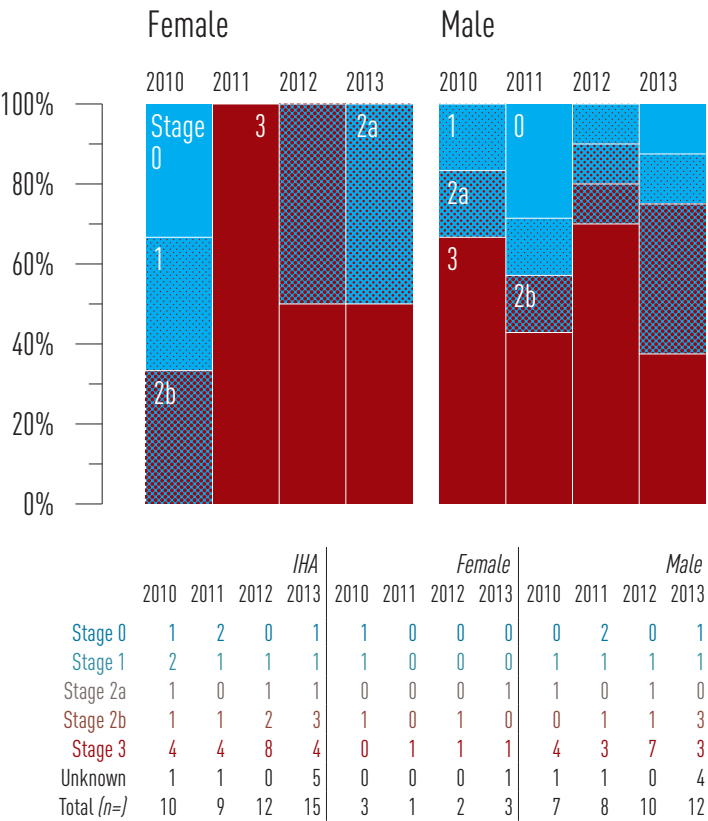


Figure 4.2 Stage of HIV Infection at Diagnosis by Gender for Interior Health, 2010–2013⁵



5 Data Source: BCCDC

Figure 4.3 Stage of HIV Infection at Diagnosis by Age Category for Interior Health, 2010–2013 ⁵

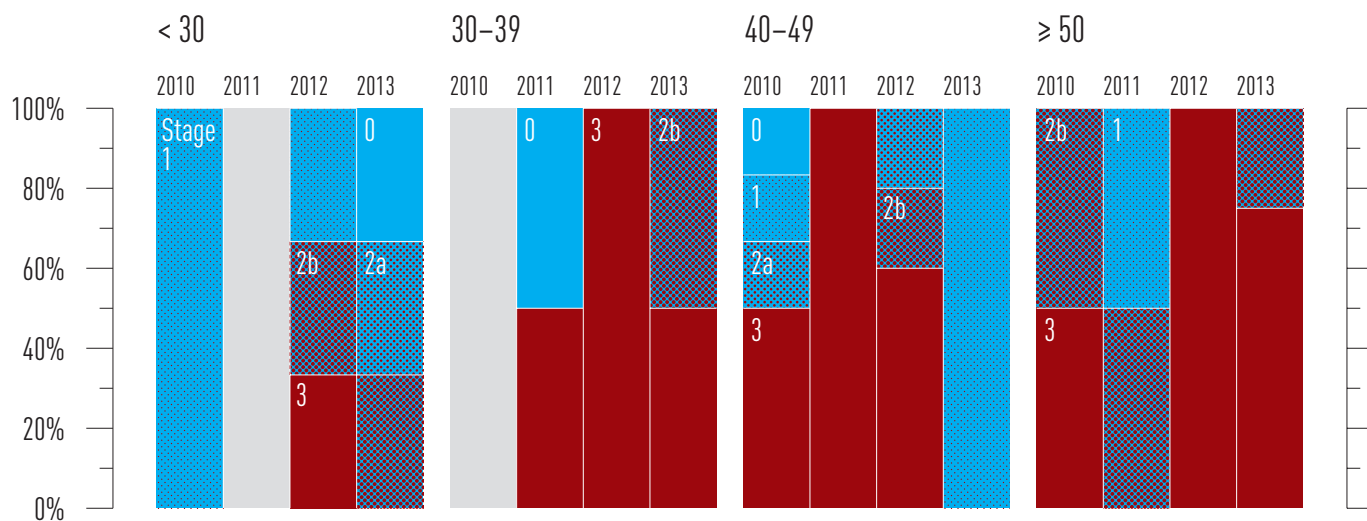
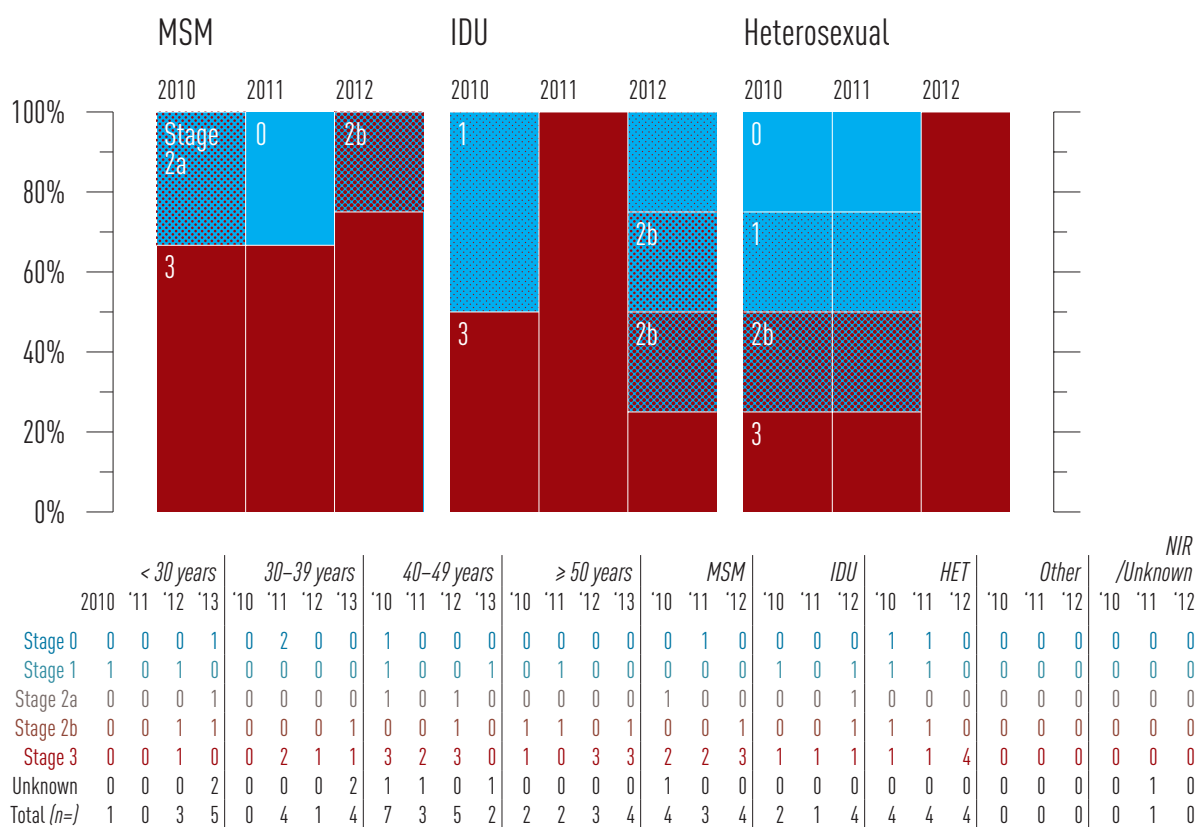


Figure 4.4 Stage of HIV Infection at Diagnosis by Exposure Category for Interior Health, 2010–2012 ^{5,6}



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

Figure 5.1 Estimated Cascade of Care for Interior Health, 2013 ⁷

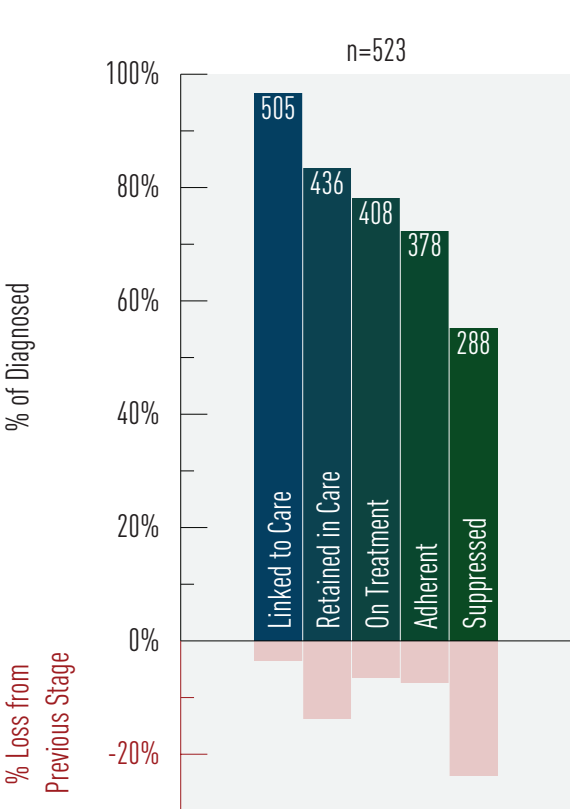
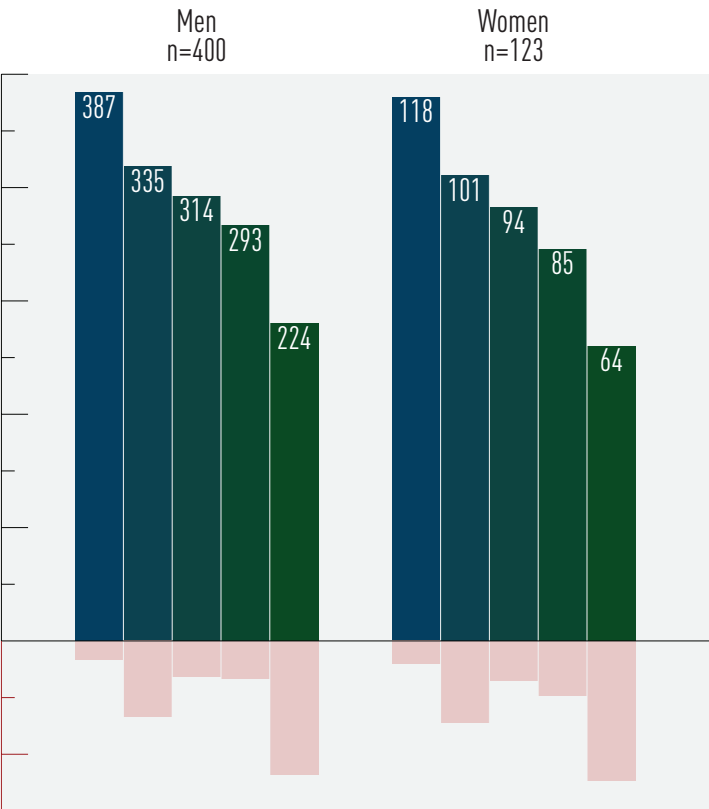


Figure 5.2 Estimated Cascade of Care for Interior Health by Gender, 2013 ⁸



7,8 Data is for the period 2013 Q1–2013 Q4.

Data Sources:

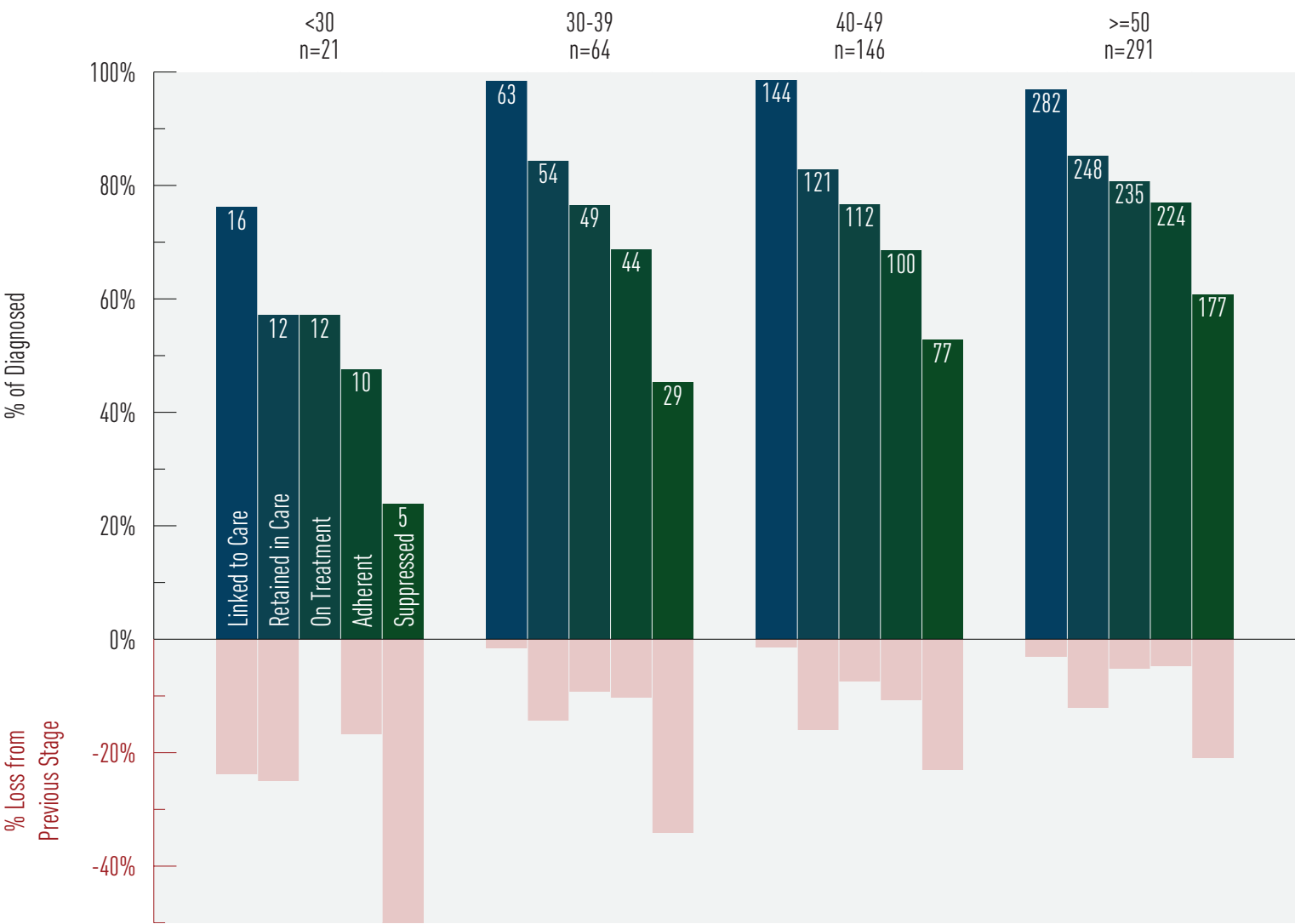
1 British Columbia Centre for Excellence Drug Treatment Program (DTP) Database (ARV use, VL and CD4 count).

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

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

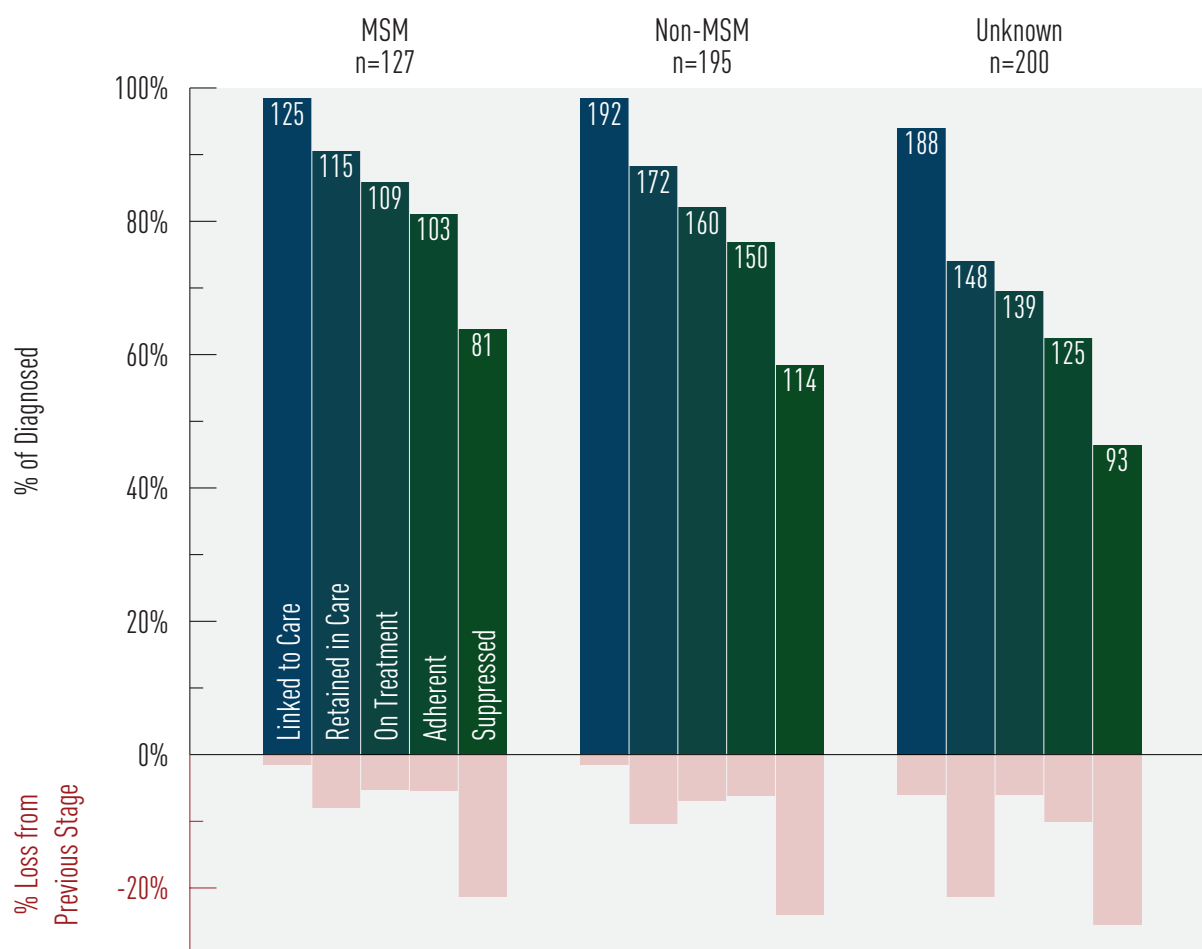
NB: Transgender has been assigned to their biological sex.

Figure 5.3 Estimated Cascade of Care for Interior Health by Age Category, 2013 ⁹



⁹ Data is for the period 2013 Q1–2013 Q4.
Data Sources:
1 British Columbia Centre for Excellence Drug Treatment Program (DTP) Database (ARV use, VL and CD4 count).
2 Administrative data (ex. MSP billings; hospitalization data from the Discharge Abstract Database (DAD)).
Limitations: HA assignment is based on the most recent HA of residence of the patient, if not available of the HIV-care provider.
If the most recent HA of residence is not updated then the designated HA may be incorrect.

Figure 5.4 Estimated Cascade of Care for Interior Health by MSM Status, 2013 ¹⁰



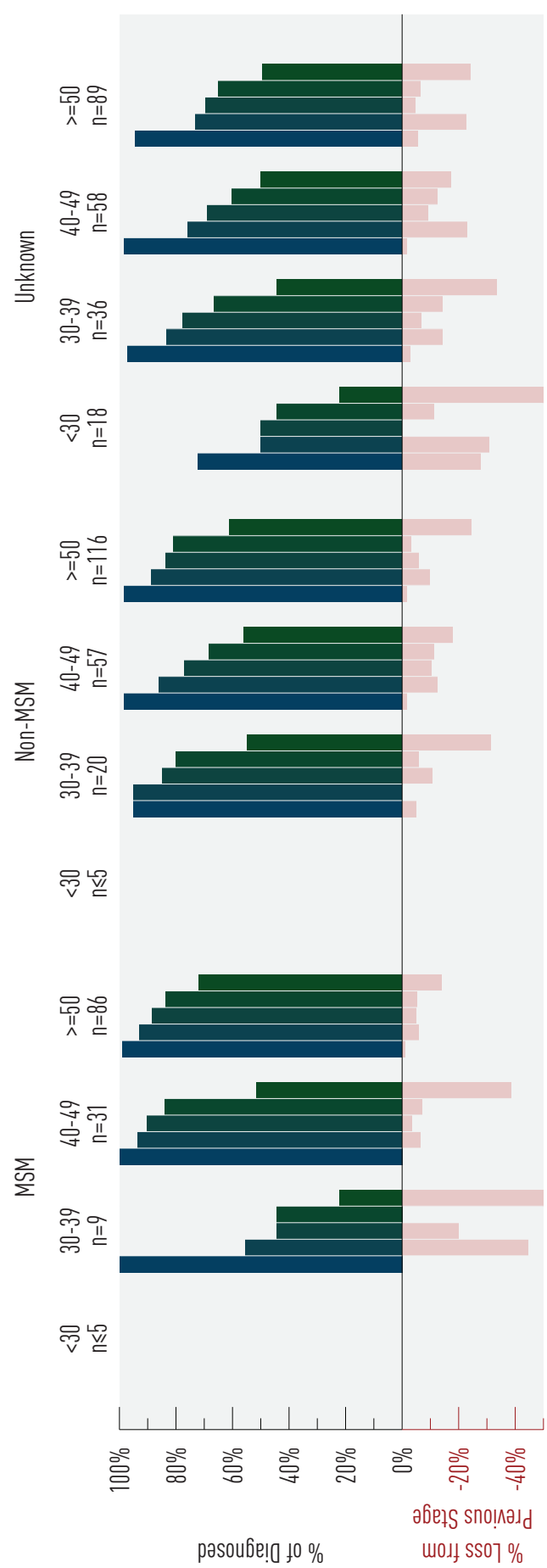
¹⁰ Data is for the period 2013 Q1–2013 Q4.

Data Sources:

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

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

Figure 5.5 Estimated Cascade of Care for Interior Health by Age Category and MSM Status, 2013 ¹¹



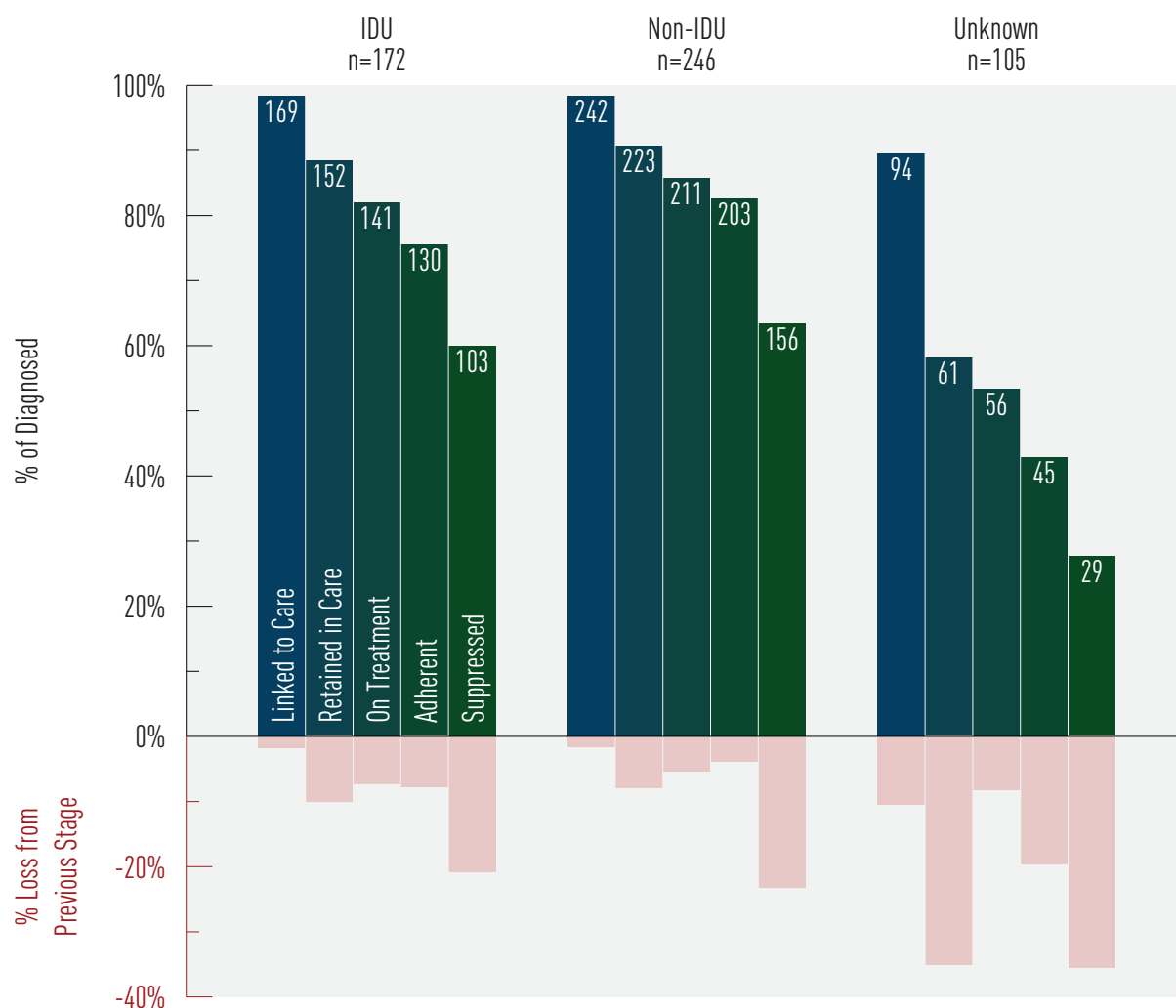
¹¹ Data is for the period 2013 Q1–2013 Q4.

Data Sources:

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

Limitations: HA assignment is based on the most recent HA of residence of the patient, if not available of the HIV-care provider. If the most recent HA of residence is not updated then the designated HA may be incorrect. Where n ≤ 5, data has been withheld for concerns of statistical significance as well as privacy. Authorized parties may contact the British Columbia Centre for Excellence in HIV/AIDS to obtain this information.

Figure 5.6 Estimated Cascade of Care for Interior Health by History of IDU, 2013¹²



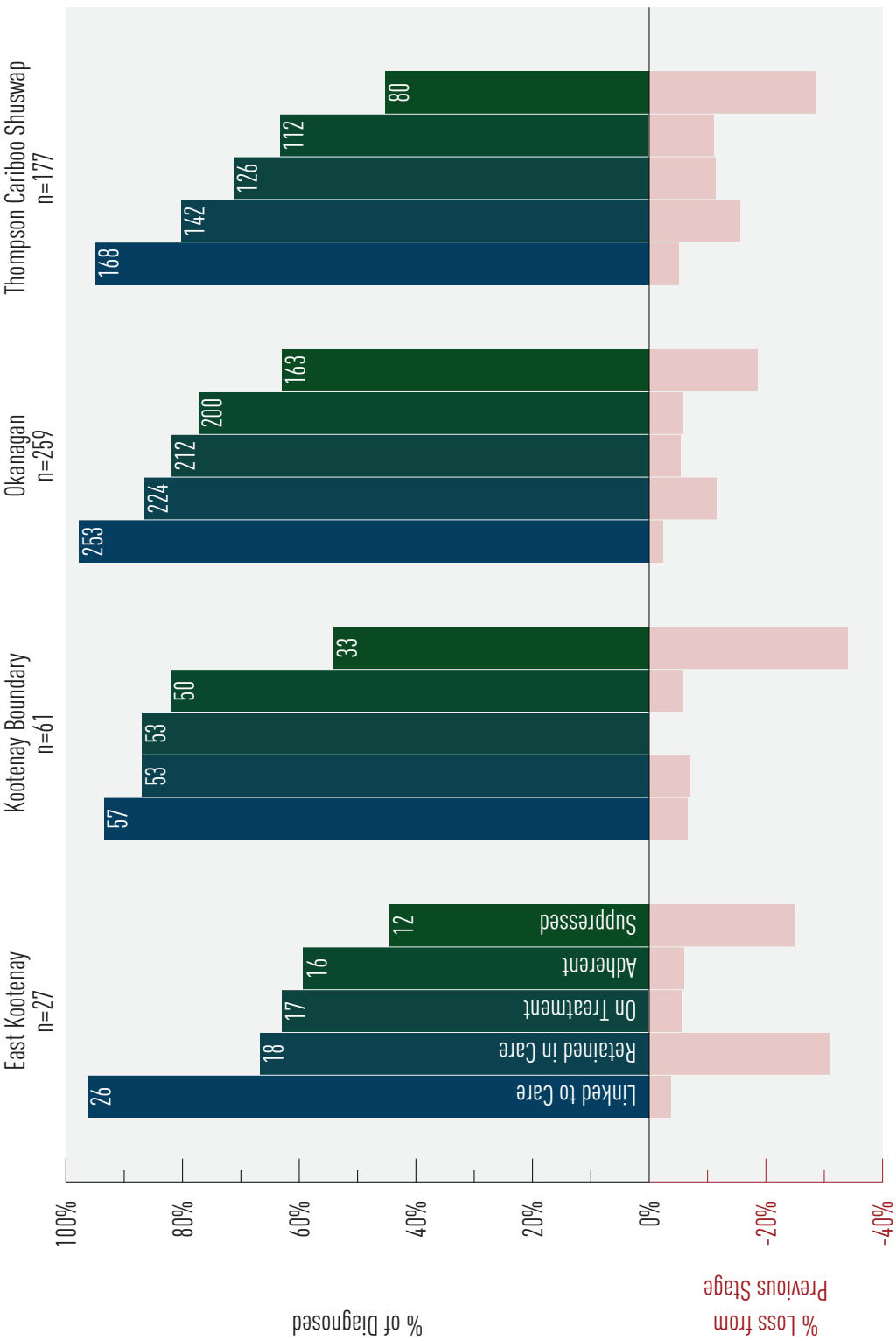
¹² Data is for the period 2013 Q1–2013 Q4.

Data Sources:

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

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

Figure 5.7 Estimated Cascade of Care for Interior Health by HSDA, 2013¹³



¹³ Data is for the period 2013 Q1-2013 Q4.

Data Sources:

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

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

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 0–6 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 $PCS \geq 4$ 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 0. 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:

1. 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/ μ L; and
6. not achieving viral suppression within 9 months since ART initiation.

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

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

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

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

Figure 6.1 PCS Components for Interior Health, 2011–2013 ¹⁴

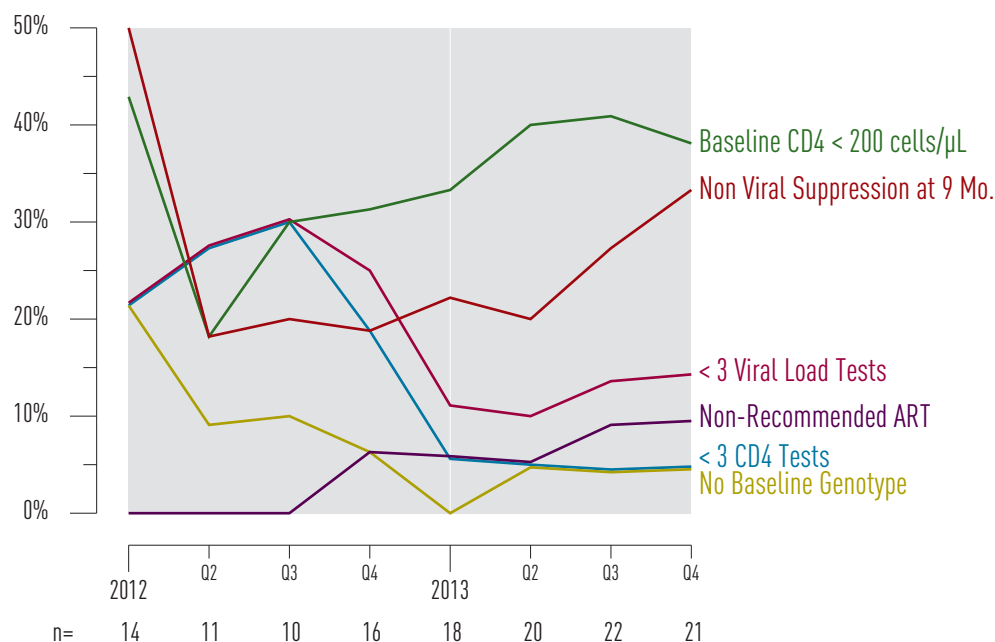
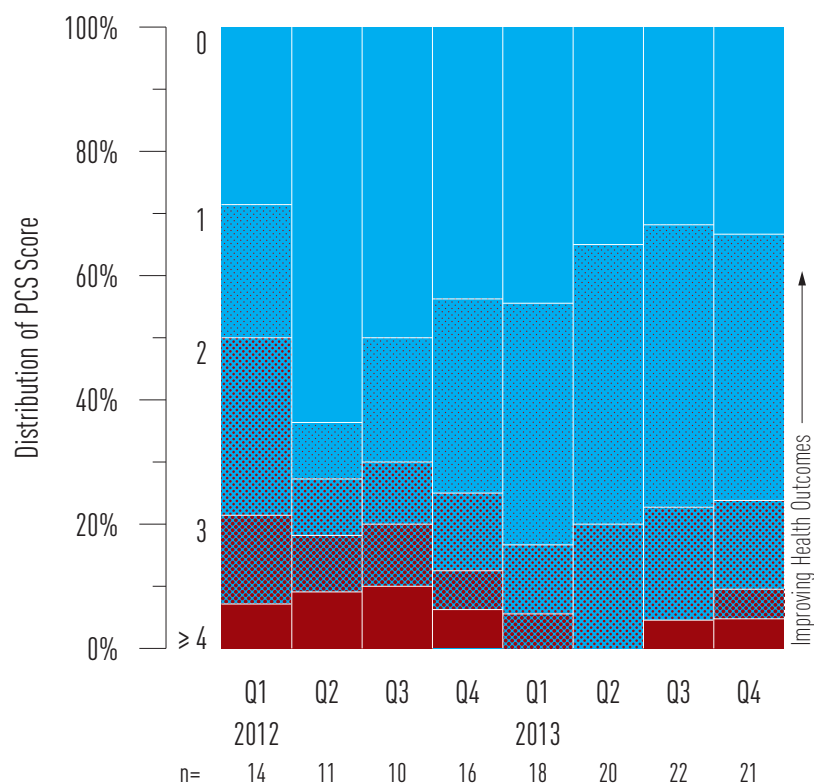


Figure 6.2 Historical Trends for PCS Score for Interior Health, 2011 Q1–2013 Q4 ¹⁵



NB: A score of 0 is the best score and a score of 4 or more is the worst score.

¹⁴ Data Source: British Columbia Centre for Excellence Drug Treatment Program (DTP) Database.
Limitations: CD4 cell count capture is approximately 80%.

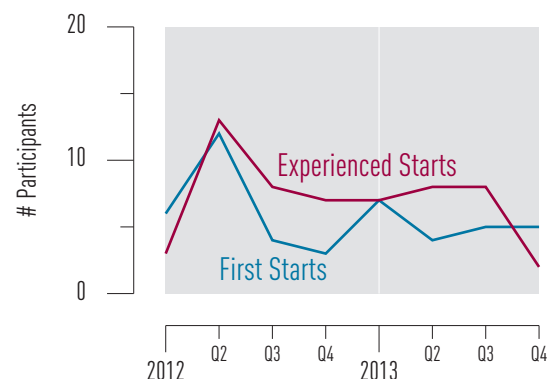
¹⁵ Data Source: British Columbia Centre for Excellence Drug Treatment Program (DTP) Database.
Each quarter's data is calculated as the sum of the 4 quarters leading up to it. e.g. 2012 Q1 is calculated from 2011 Q2 – 2012 Q1.

Antiretroviral Uptake

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

Indicator 7. New Antiretroviral Therapy Starts in Interior Health

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



Indicator 8. CD4 Cell Count at ART Initiation

Figure 8 CD4 Cell Count at ART Initiation of ART-Naïve DTP Participants in Interior Health, 2012 Q1–2013 Q4 ¹⁷

The majority of cells in this figure have $n \leq 5$, which is considered statistically insignificant as well as a possible risk to patient privacy. For this reason, this figure has been omitted. Authorized parties may contact the British Columbia Centre for Excellence in HIV/AIDS to obtain this information.

¹⁶ Data Source: Drug Treatment Program Database

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

¹⁷ Data Source: Drug Treatment Program Database

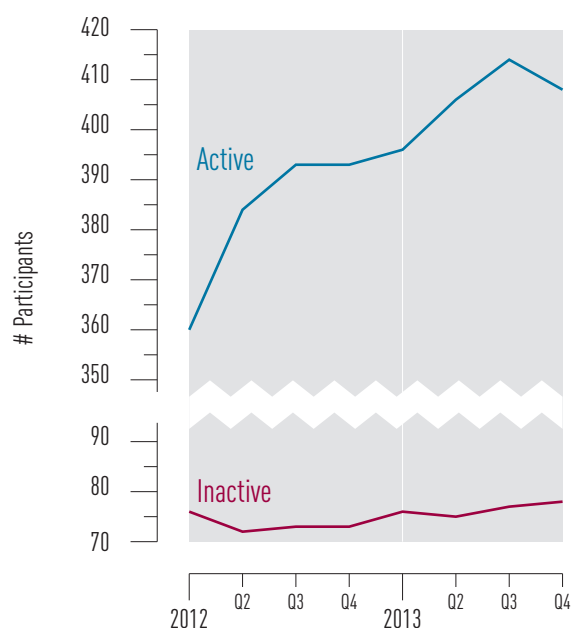
Limitations: CD4 cell count data is approximately 80% complete.

Indicator 9. Active and Inactive DTP Participants

Table 3. Distribution of People on ART for Interior Health, 2013 Q4 ¹⁶

Age	< 30	10
	30–39	51
	40–49	111
	≥ 50	236
Gender	Male	315
	Female	93
Exposure	MSM	109
	IDU	143
Total		408

Figure 9 Active and Inactive DTP Participants in Interior Health, 2012 Q1–2013 Q4 ¹⁹



¹⁸ Data Source: Drug Treatment Program Database

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

Definitions:

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

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

¹⁹ Active DTP participants: are those who are prescribed one or more drugs in the last six months.

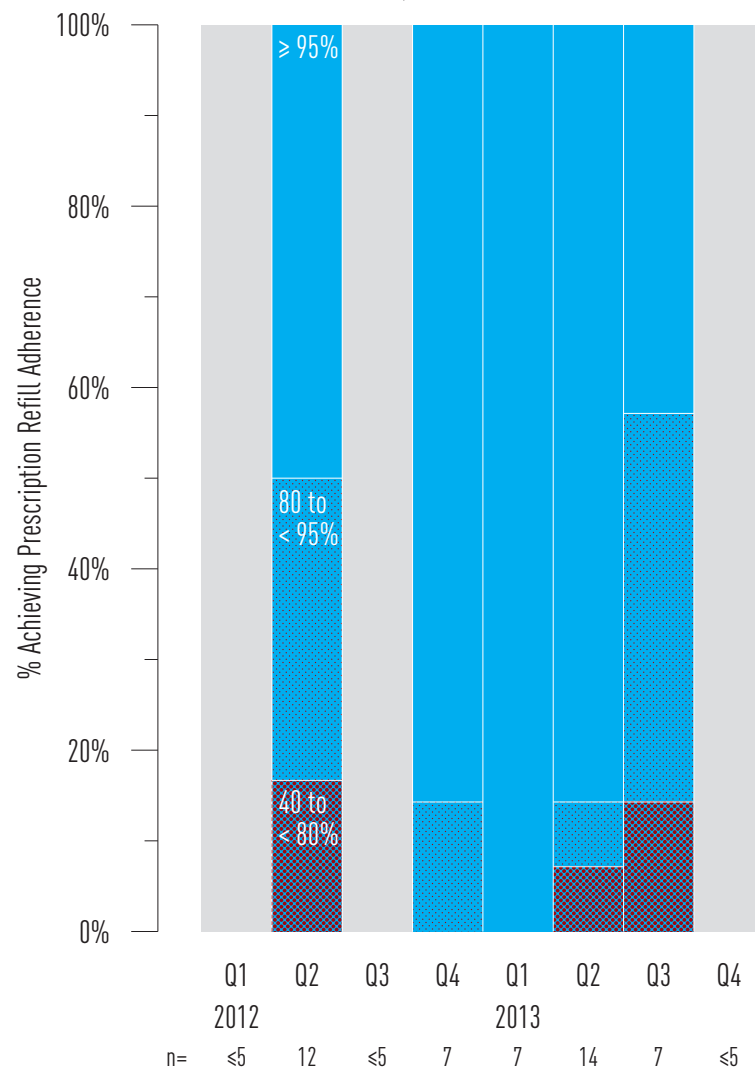
Inactive DTP Participants: Persons no longer prescribed drugs through the HIV/AIDS Drug Treatment Program in the last quarter.

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 Interior Health, 2012 Q1–2013 Q4 ²⁰



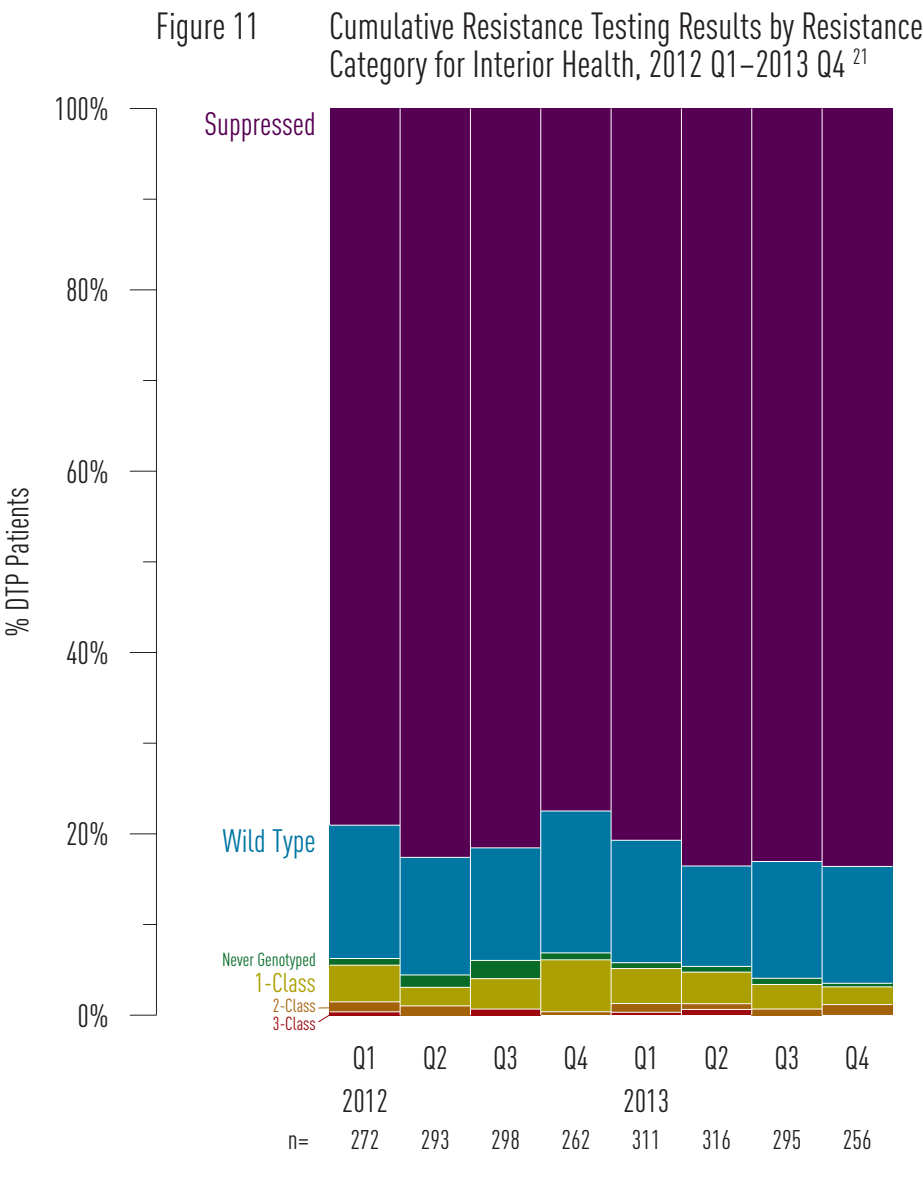
²⁰ Data Source: Drug Treatment Program Database

Limitation: Prescription refill adherence is used as a proxy for patient adherence.

Where $n \leq 5$, data has been withheld for concerns of statistical significance as well as privacy. Authorized parties may contact the British Columbia Centre for Excellence in HIV/AIDS to obtain this information.

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.



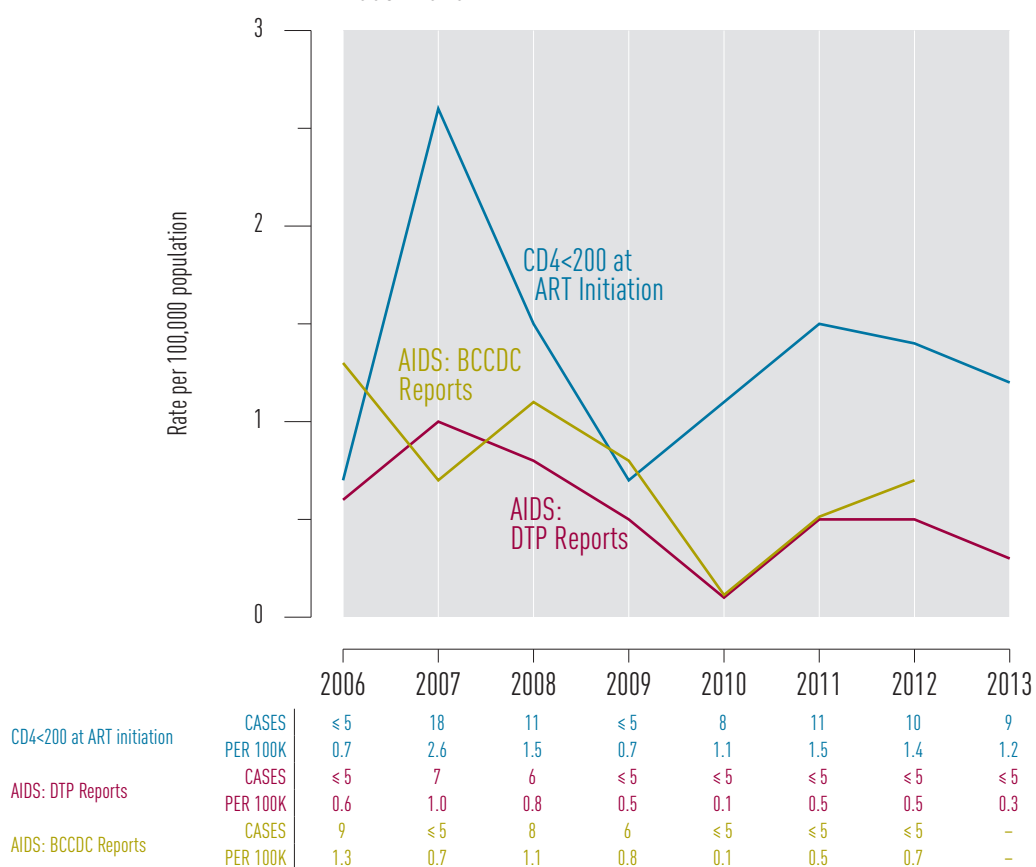
²¹ Data Source: Drug Treatment Program Database

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

Indicator 12. AIDS-Defining Illness

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

Figure 12 AIDS Case Rate and Reports for Interior Health, 2006–2013²²



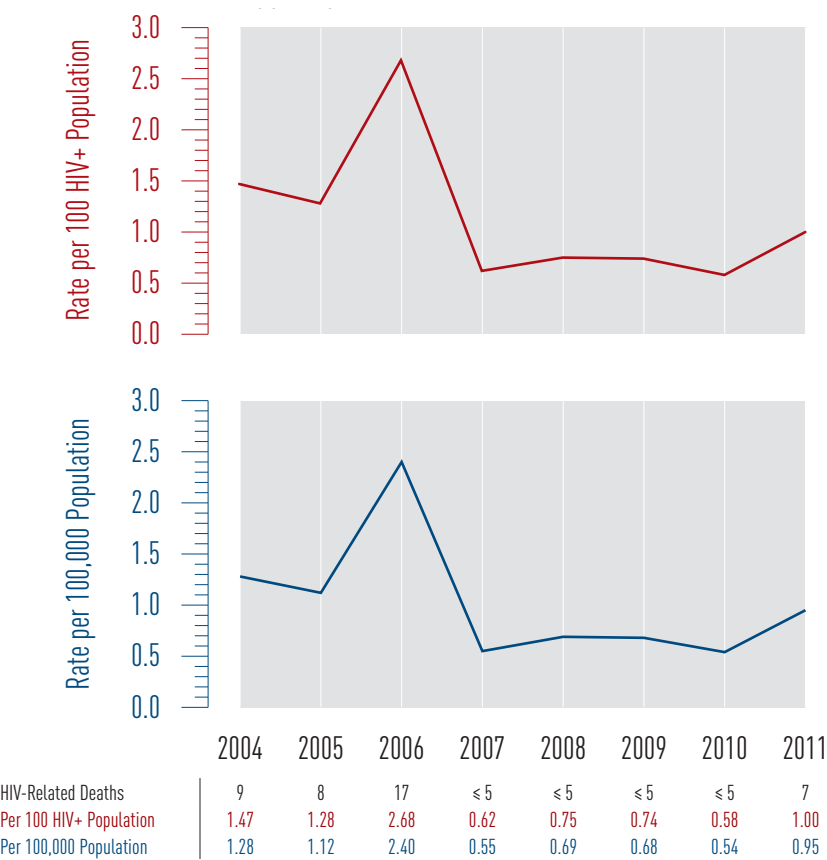
²² Data Source: Drug Treatment Program Database

Limitation: AIDS case reporting was investigated using 2 definitions: First, using AIDS cases reported in AIDS case report forms from the DTP, and second, using a CD4 cell count of <200 cells/μL at time of ART initiation using DTP data. AIDS case reporting is passive in BC, thus; AIDS case reporting is not well captured. The DTP sends out AIDS reporting forms to physicians annually. Interpreting AIDS case reports should be done with these limitations in mind. AIDS data is updated annually as very few AIDS cases reports are reported in general and trends would be difficult to notice if reported quarterly.

Indicator 13. HIV-Related Mortality

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

Figure 13 HIV-Related Deaths by Year for Interior Health, 2004–2011 ²³



²³ Data Source: BC Vital Statistics

Limitation:

1. DTP participants are designated to an HA based on most current residence provided by the participant.
2. Mortality data is updated annually.
3. The most recent available data was used.

Appendices

Indicator 1: Test Episodes (thousands)		2009				2010				2011				2012				2013			
		Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4
Interior Health		6.0	5.6	5.5	5.6	6.0	5.6	5.5	5.9	6.0	5.5	5.6	5.9	5.9	5.7	5.9	5.9	6.1	6.4	6.3	6.7
Gender	Female	3.9	3.7	3.6	3.6	3.9	3.6	3.6	3.8	3.9	3.6	3.6	3.8	3.8	3.6	3.8	3.7	3.9	4.0	3.9	4.2
	Male	1.9	1.8	1.7	1.8	2.0	1.8	1.8	1.9	2.0	1.8	1.8	2.0	2.0	1.9	2.1	2.1	2.2	2.3	2.3	2.4
	Other	0.2	0.2	0.2	0.1	0.2	0.2	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.0	0.1	0.0	0.1	0.1
Female (Prenatal)		1.9	1.6	1.6	1.7	1.8	1.7	1.7	1.8	1.9	1.6	1.7	1.7	1.8	1.6	1.7	1.7	1.7	1.6	1.6	1.7
Female (Non-prenatal)		2.0	2.1	2.0	1.9	2.1	1.9	1.9	2.1	2.0	2.0	2.0	2.1	2.1	2.0	2.1	2.0	2.1	2.4	2.2	2.5
Age	< 30	2.7	2.4	2.5	2.3	2.5	2.4	2.4	2.4	2.5	2.2	2.3	2.5	2.3	2.2	2.4	2.3	2.2	2.3	2.4	2.5
	30–39	1.7	1.5	1.5	1.5	1.8	1.6	1.6	1.6	1.8	1.6	1.6	1.6	1.8	1.6	1.7	1.6	1.8	1.8	1.8	1.8
	40–49	0.8	0.8	0.7	0.7	0.8	0.8	0.7	0.7	0.7	0.7	0.8	0.8	0.8	0.8	0.8	0.7	0.8	0.8	0.7	0.8
	≥ 50	0.8	0.9	0.8	1.1	0.9	0.9	0.9	1.2	1.0	0.9	0.9	1.0	1.0	1.1	1.1	1.1	1.3	1.5	1.4	1.5
POC HIV Tests (not in thousands)									0	0	10	34	21	26	28	20	27	38	38	40	37
East Kootenay		0.5	0.5	0.5	0.6	0.6	0.5	0.5	0.6	0.5	0.5	0.5	0.6	0.6	0.5	0.6	0.6	0.6	0.6	0.6	0.6
Kootenay Boundary		0.6	0.6	0.6	0.8	0.7	0.6	0.6	0.8	0.7	0.6	0.6	0.6	0.6	0.5	0.6	0.7	0.6	0.6	0.6	0.7
Okanagan		3.3	3.0	3.0	2.9	3.1	2.9	2.8	2.9	3.1	2.8	3.0	3.1	3.2	3.1	3.2	3.0	3.1	3.3	3.3	3.4
Thompson Cariboo Shuswap		1.5	1.5	1.4	1.3	1.7	1.5	1.5	1.6	1.7	1.6	1.6	1.5	1.6	1.5	1.6	1.7	1.8	1.9	1.9	2.0

Indicator 2: Rate of HIV Testing per 100,000

		2009	2010	2011	2012	2013
Interior Health		2890.3	2926.6	2927.5	2988.1	3261.1
East Kootenay		2642.8	2763.6	2619.5	2689.3	2736.0
Kootenay Boundary		3079.5	3100.6	2983.6	2903.6	3258.2
Okanagan		3041.4	3001.5	3026.6	3155.5	3382.8
Thompson Cariboo Shuswap		2674.0	2805.4	2862.2	2860.8	3257.6
Gender	Female	3681.6	3692.9	3700.1	3764.1	4041.5
	Male	1908.7	1975.6	2015.1	2132.3	2408.2
Age	< 30	3864.2	3766.0	3757.3	3647.6	3742.5
	30–39	7132.7	7592.6	7681.0	7889.1	8272.7
	40–49	2665.6	2716.6	2814.1	2996.5	3287.0
	≥ 50	1089.2	1149.5	1154.9	1311.1	1685.1

Indicator 3: New HIV Diagnoses		2009				2010				2011				2012				2013			
		Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4
Interior Health	By Client Residence	6	4	4	4	4	1	3	3	1	4	3	3	5	5	0	2	1	4	4	5
	By Provider Address	6	4	4	4	4	1	2	3	1	3	2	3	5	5	0	2	1	5	4	5
Gender	Female	2	1	3	1	0	0	3	1	0	2	1	0	1	1	0	0	0	1	1	1
	Male	4	3	1	3	4	1	0	2	1	2	2	3	4	4	0	2	1	3	3	4
Age	< 30	0	1	2	1	0	0	0	1	0	1	0	0	1	1	0	1	0	0	2	2
	30–39	3	0	1	2	0	0	1	0	1	0	3	1	1	1	0	0	0	0	2	2
	40–49	1	1	1	1	3	1	1	2	0	3	0	1	2	1	0	1	0	1	0	1
	≥ 50	2	2	0	0	1	0	1	0	0	0	0	1	1	2	0	0	1	3	0	0
Exposure	MSM	1	2	0	1	2	1	0	1	0	0	1	2	1	2	0	1	0	0	–	–
	IDU	1	0	0	0	1	0	0	1	0	0	1	0	2	1	0	1	0	0	–	–
	HET	4	1	4	2	1	0	3	1	1	2	1	1	2	2	0	0	1	1	–	–
	Other	0	1	0	1	0	0	0	0	0	1	0	0	0	0	0	0	0	0	–	–
	NIR/Unknown	0	0	0	0	0	0	0	0	0	1	0	0	0	0	0	0	0	3	–	–
East Kootenay	By Client Residence	0	1	0	0	3	0	0	0	0	0	0	0	2	0	0	0	0	1	0	0
	By Provider Address	0	1	0	0	3	0	0	0	0	0	0	0	2	0	0	0	0	1	0	0
Kootenay Boundary	By Client Residence	0	1	0	0	1	0	0	0	0	1	0	1	2	0	0	0	1	0	1	1
	By Provider Address	0	1	0	0	1	0	0	0	0	0	0	1	1	0	0	0	1	0	0	0
Okanagan	By Client Residence	2	2	2	2	0	1	1	0	1	3	0	3	1	3	0	1	1	0	3	2
	By Provider Address	2	2	2	2	0	1	1	0	1	2	1	3	1	4	0	1	1	1	3	3
Thompson Cariboo Shuswap	By Client Residence	4	0	2	2	0	0	2	3	0	1	2	0	1	0	0	1	0	2	1	2
	By Provider Address	4	0	2	2	0	0	1	3	0	1	1	0	1	0	0	1	0	2	1	2

Indicator 4: Stage of HIV Infection at Baseline

	IHA				Female				Male				< 30 years				30–39 years				40–49 years			
	'10	'11	'12	'13	'10	'11	'12	'13	'10	'11	'12	'13	'10	'11	'12	'13	'10	'11	'12	'13	'10	'11	'12	'13
Stage 0	1	2	0	1	1	0	0	0	0	2	0	1	0	0	0	1	0	2	0	0	1	0	0	0
Stage 1	2	1	1	1	1	0	0	0	1	1	1	1	1	0	1	0	0	0	0	0	1	0	0	1
Stage 2a	1	0	1	1	0	0	0	1	1	0	1	0	0	0	0	1	0	0	0	0	1	0	1	0
Stage 2b	1	1	2	3	1	0	1	0	0	1	1	3	0	0	1	1	0	0	0	1	0	0	1	0
Stage 3	4	4	8	4	0	1	1	1	4	3	7	3	0	0	1	0	0	2	1	1	3	2	3	0
Unknown	1	1	0	5	0	0	0	1	1	1	0	4	0	0	0	2	0	0	0	2	1	1	0	1
Total	10	9	12	15	3	1	2	3	7	8	10	12	1	0	3	5	0	4	1	4	7	3	5	2

	≥ 50 years				MSM			IDU			Heterosexual			Other Exposure			NIR/Unknown		
	'10	'11	'12	'13	2010	2011	2012	2010	2011	2012	2010	2011	2012	2010	2011	2012	2010	2011	2012
Stage 0	0	0	0	0	0	1	0	0	0	0	1	1	0	0	0	0	0	0	0
Stage 1	0	1	0	0	0	0	0	1	0	1	1	1	0	0	0	0	0	0	0
Stage 2a	0	0	0	0	1	0	0	0	0	1	0	0	0	0	0	0	0	0	0
Stage 2b	1	1	0	1	0	0	1	0	0	1	1	1	0	0	0	0	0	0	0
Stage 3	1	0	3	3	2	2	3	1	1	1	1	1	4	0	0	0	0	0	0
Unknown	0	0	0	0	1	0	0	0	0	0	0	0	0	0	0	0	0	1	0
Total	2	2	3	4	4	3	4	2	1	4	4	4	4	0	0	0	0	1	0

Indicator 5: HIV Cascade of Care

		DIAGNOSED	LINKED	RETAINED	ON ART	ADHERENT	SUPPRESSED
Interior Health		523	505	436	408	378	288
Age Category	< 30	21	16	12	12	10	5
	30–39	64	63	54	49	44	29
	40–49	146	144	121	112	100	77
	≥ 50	291	282	248	235	224	177
Age Category and MSM Status	MSM	< 30	≤ 5	≤ 5	≤ 5	≤ 5	≤ 5
		30–39	9	9	5	4	2
		40–49	31	31	29	28	16
		≥ 50	86	85	80	76	62
	Non-MSM	< 30	≤ 5	≤ 5	≤ 5	≤ 5	≤ 5
		30–39	20	19	19	17	11
		40–49	57	56	49	44	32
		≥ 50	116	114	103	97	71
	Unknown	< 30	18	13	9	9	4
		30–39	36	35	30	28	16
		40–49	58	57	44	40	29
		≥ 50	89	84	65	62	44
Gender	Male	400	387	335	314	293	224
	Female	123	118	101	94	85	64
Injection Drug Use	IDU	172	169	152	141	130	103
	Non-IDU	246	242	223	211	203	156
	Unknown	105	94	61	56	45	29
MSM Status	MSM	127	125	115	109	103	81
	Non-MSM	195	192	172	160	150	114
	Unknown	200	188	148	139	125	93
Health Authority	East Kootenay	27	26	18	17	16	12
	Kootenay Boundary	61	57	53	53	50	33
	Okanagan	259	253	224	212	200	163
	Thompson Cariboo Shuswap	177	168	142	126	112	80

Indicator 6: Programmatic Compliance Score (PCS)

	2012				2013			
	Q1	Q2	Q3	Q4	Q1	Q2	Q3	Q4
< 3 CD4 Tests	21.4%	27.3%	30.0%	18.8%	5.6%	5.0%	4.5%	4.8%
< 3 Viral Load Tests	21.4%	27.3%	30.0%	25.0%	11.1%	10.0%	13.6%	14.3%
No Baseline Genotype	21.4%	9.1%	10.0%	6.3%	0.0%	5.0%	4.5%	4.8%
Baseline CD4 < 200 cells/μL	42.9%	18.2%	30.0%	31.3%	33.3%	40.0%	40.9%	38.1%
Non-Recommended ART	0.0%	0.0%	0.0%	6.3%	5.6%	5.0%	9.1%	9.5%
Non Viral suppression at 9 Mo.	50.0%	18.2%	20.0%	18.8%	22.2%	20.0%	27.3%	33.3%
PCS Score: 0	4	7	5	7	8	7	7	7
PCS Score: 1	3	1	2	5	7	9	10	9
PCS Score: 2	4	1	1	2	2	4	4	3
PCS Score: 3	2	1	1	1	1	0	0	1
PCS Score: 4 or more	1	1	1	1	0	0	1	1
Total (n=)	14	11	10	16	18	20	22	21

Indicator 7: New DTP ARV Participants

First Starts	6	12	4	3	7	4	5	5
Experienced Starts	3	13	8	7	7	8	8	2

Indicator 8: CD4 Cell Count at ART Initiation for ARV-Naïve DTP Participants

CD4 ≥ 500	1	1	–	–	1	–	–	–
CD4 350–499	0	2	–	–	2	–	–	–
CD4 200–349	2	5	–	–	2	–	–	–
CD4 50–199	2	2	–	–	2	–	–	–
CD4 < 50	1	2	–	–	0	–	–	–
<i>CD4 Median (cells/μL)</i>	<i>195</i>	<i>220</i>	<i>–</i>	<i>–</i>	<i>310</i>	<i>–</i>	<i>–</i>	<i>–</i>
Total (n=)	6	12	≤ 5	≤ 5	7	≤ 5	≤ 5	≤ 5

Indicator 9: Active and Inactive DTP Participants

Active DTP Participants	360	384	393	393	396	406	414	408
Inactive DTP Participants	76	72	73	73	76	75	77	78

Indicator 10: Antiretroviral Adherence

≥ 95%	–	6	–	6	7	12	3	–
80% to < 95%	–	4	–	1	0	1	3	–
40% to < 80%	–	2	–	0	0	1	1	–
< 40%	–	0	–	0	0	0	0	–
Total (n=)	≤ 5	12	≤ 5	7	7	14	7	≤ 5

Indicator 11: Resistance Testing and Results

Suppressed	215	242	243	203	251	264	245	214
Wild Type	40	38	37	41	42	35	38	33
Never Genotyped	2	4	6	2	2	2	2	1
1-Class	11	6	10	15	12	11	8	5
2-Class	3	3	0	1	3	2	2	3
3-Class	1	0	2	0	1	2	0	0
Total (n=)	272	293	298	262	311	316	295	256

Indicator 12: AIDS-Defining Illness

		2006	2007	2008	2009	2010	2011	2012	2013
CD4 < 200 at	Cases	≤ 5	18	11	≤ 5	8	11	10	9
ART initiation	<i>Rate per 100,000</i>	<i>0.7</i>	<i>2.6</i>	<i>1.5</i>	<i>0.7</i>	<i>1.1</i>	<i>1.5</i>	<i>1.4</i>	<i>1.2</i>
AIDS Cases	Cases	≤ 5	7	6	≤ 5	≤ 5	≤ 5	≤ 5	≤ 5
(DTP Reports)	<i>Rate per 100,000</i>	<i>0.6</i>	<i>1.0</i>	<i>0.8</i>	<i>0.5</i>	<i>0.1</i>	<i>0.5</i>	<i>0.5</i>	<i>0.3</i>
AIDS Cases	Cases	9	≤ 5	8	6	≤ 5	≤ 5	5	–
(BCCDC Reports)	<i>Rate per 100,000</i>	<i>1.3</i>	<i>0.7</i>	<i>1.1</i>	<i>0.8</i>	<i>0.1</i>	<i>0.5</i>	<i>0.7</i>	<i>–</i>

Indicator 13: HIV-Related Mortality

	2004	2005	2006	2007	2008	2009	2010	2011
Interior Health	9	8	17	≤ 5	≤ 5	≤ 5	≤ 5	7
Per 100 HIV+ Population	1.47	1.28	2.68	0.62	0.75	0.74	0.58	1.00
Per 100,000 Population	1.28	1.12	2.40	0.55	0.69	0.68	0.54	0.95