Association of highly active antiretroviral therapy coverage, population viral load, and yearly new HIV diagnoses in British Columbia, Canada: a population-based study

Patricia Daly, Perry Kendall

Summary

Background Results of cohort studies and mathematical models have suggested that increased coverage with highly active antiretroviral therapy (HAART) could reduce HIV transmission. We aimed to estimate the association between plasma HIV-1 viral load, HAART coverage, and number of new cases of HIV in the population of a Canadian province.

Methods We undertook a population-based study of HAART coverage and HIV transmission in British Columbia, Canada. Data for number of HIV tests done and new HIV diagnoses were obtained from the British Columbia Centre for Disease Control. Data for viral load, CD4 cell count, and HAART use were extracted from the British Columbia Centre for Excellence in HIV/AIDS population-based registries. We modelled trends of new HIV-positive tests and number of individuals on HAART using generalised additive models. Poisson log-linear regression models were used to estimate the association between new HIV diagnoses and viral load, year, and number of individuals on HAART.

Findings Between 1996 and 2009, the number of individuals actively receiving HAART increased from 837 to 5413 (547% increase; p=0.002), and the number of new HIV diagnoses fell from 702 to 338 per year (52% decrease; p=0.001). The overall correlation between number of individuals on HAART and number of individuals newly testing positive for HIV per year was −0.89 (p<0.0001). For every 100 additional individuals on HAART, the number of new HIV cases decreased by a factor of 0.97 (95% CI 0.96–0.98), and per 1 log10 decrease in viral load, the number of new HIV cases decreased by a factor of 0.86 (0.75–0.98).

Interpretation We have shown a strong population-level association between increasing HAART coverage, decreased viral load, and decreased number of new HIV diagnoses per year. Our results support the proposed secondary benefit of HAART used within existing medical guidelines to reduce HIV transmission.

Funding Ministry of Health Services and Ministry of Healthy Living and Sport, Province of British Columbia; US National Institute on Drug Abuse; US National Institutes of Health; Canadian Institutes of Health Research.

Introduction Despite recent progress, neither a cure nor a preventive vaccine against HIV-1 infection is likely to become available in the near future. Several preventive strategies have not been wholly effective for several reasons, including insufficient support, logistical difficulties, poor implementation, and underuse. As a result, the global effect of HIV/AIDS continues to grow. In 2008, an estimated 33.4 million people were living with HIV, there were 2.0 million AIDS-related deaths, and an alarming 2.7 million new HIV infections occurred. These developments have prompted the UN Joint Programme on HIV/AIDS to call for an urgent redoubling of efforts in the fight against HIV/AIDS.

HIV treatment has advanced remarkably since 1996, with the development and refinement of highly active antiretroviral therapy (HAART). HAART stops HIV replication on a sustained basis and, as a result, plasma HIV-1 RNA concentrations (henceforth viral load) typically become undetectable. This change allows for immune reconstitution to take place, leading to long-term disease remission and aversion of the otherwise fatal course. By 2006, at least 3 million years of life had been saved in the USA as a direct result of HAART within a decade. In high-income countries, life expectancy of HIV-positive individuals aged 20 years who were taking HAART was roughly two-thirds of that of the general population.

Interest has increased in the possible secondary effect of HAART—reduction of HIV transmission. The association between high plasma HIV-1 RNA concentration and high risk of HIV transmission has long been understood. In addition to decreasing plasma viral load to undetectable levels, HAART decreases viral load in other biological fluids, including semen and vaginal secretions. Although exceptions have been reported, from a public health perspective the association between viral load and other bodily fluids is quite strong, especially in the setting of long-term, sustained, and effective HAART. Strong proof of principle regarding the effect of
HAART on HIV transmission has been shown in studies of vertical transmission in resource-rich and resource-poor settings.21

New evidence suggests that HAART can decrease HIV transmission in other settings. Reductions in rates of HIV transmission of more than 90% have been reported in several cohort studies of heterosexual HIV-serodiscordant couples in whom the index partner was treated with HAART.24,25 Similarly, reduction in community viral load as a result of HAART was shown to be a key determinant of decreasing HIV incidence in a cohort of injecting drug users (IDUs) in Vancouver, Canada.26 Investigators have also documented this effect in retrospective population-based observational studies in Taiwan,27 the province of British Columbia, Canada,28 and the city of San Francisco, USA.29 However, the estimated effect of increased HAART coverage on HIV transmission varies greatly between mathematical models, with results ranging from elimination10,13 to potential worsening29 of the HIV epidemic. We therefore undertook this study with the aim of analysing at a population level the potential association between expansion of HAART coverage, viral load, and new HIV diagnoses per year in a Canadian province with free access to HIV care.

Methods
Study design and participants
We used two separate unlinked databases in this population-based cohort study. Data for number of HIV tests done and new HIV diagnoses in the province of British Columbia, Canada, between 1996 and 2009, were obtained from the British Columbia Centre for Disease Control (BCCDC).10 BCCDC is the single provincial agency that centralises all HIV surveillance data for the province and has access to HIV testing data from the provincial public health reference laboratory, which does more than 90% of all HIV testing in British Columbia. Mandatory (nominal or non-nominal) HIV reporting legislation has been in place in the province since 2003. This includes specific establishment of whether a new HIV positive result is from an individual previously known to be infected with HIV, but who has moved to the area from elsewhere. For individuals aged 18 months or older, BCCDC uses a screening test (ELISA) to detect HIV antibodies, with HIV diagnosis confirmed on the basis of a reactive western blot or nucleic acid amplification test.

Data for viral load, CD4 cell count, and HAART use were extracted from the British Columbia Centre for Excellence in HIV/AIDS (BCCFE) population-based registries, which include all people on HAART or accessing viral load testing in the province. BCCFE is the single provincial agency that centrally distributes all antiretroviral drugs in the province, providing treatment free of charge to all residents who are infected with HIV. BCCFE maintains a set of independently generated HIV/AIDS management guidelines, which have remained consistent with the International AIDS Society-USA (IAS-USA) guidelines since 1996. BCCFE guidelines recommend that once an individual starts treatment, viral load and CD4 cell count be tested at regular intervals (eg, every 3–4 months) or after virological rebound. Capture of baseline and on-treatment data for viral load by BCCFE is 100%, because all viral load measurements in the province are done under the auspices of BCCFE by the virology laboratory at St Paul’s Hospital (Providence Health Care, Vancouver, BC, Canada).

Procedures
Table 1 shows the different HIV-1 RNA viral load assays that were used during the study period and their quantification ranges. For analytical purposes, we truncated data for viral load to the range 500–100 000 copies per mL, because these values were within the measurement range of all assays used during the study. We estimated population-level or community-level viral load using a conservative approach by recording the highest viral load for every individual in a specific year. To accommodate irregular frequency of viral-load measurements or missing values, the highest yearly

<table>
<thead>
<tr>
<th>Assay</th>
<th>Range of quantification (copies per mL)</th>
</tr>
</thead>
<tbody>
<tr>
<td>June 1, 1996 Roche HIV-1 monitor test; microplate format</td>
<td>500–1 000 000</td>
</tr>
<tr>
<td>April 18, 1997 Roche HIV-1 monitor test; microplate format</td>
<td>400–750 000</td>
</tr>
<tr>
<td>Oct 22, 1997 Roche HIV-1 monitor test version 1.5; included new primers to detect different HIV-1 subtypes</td>
<td>400–750 000</td>
</tr>
<tr>
<td>Dec 4, 1998 Roche COBAS HIV-1 monitor test version 1.5; first time automated amplification and detection were used, and the standard method was used for specimen preparation</td>
<td>400–750 000</td>
</tr>
<tr>
<td>April 1, 1999 Roche COBAS HIV-1 Ampliprep Amplicor Monitor ultrasensitive assay version 1.5; automated amplification and detection and the ultrasensitivity method was used for specimen preparation</td>
<td>50–75 000</td>
</tr>
<tr>
<td>March 8, 2000 Roche COBAS HIV-1 Ampliprep Amplicor Monitor ultrasensitive assay version 1.5; automated amplification and detection and the ultrasensitivity method detection range changed with time</td>
<td>50–100 000</td>
</tr>
<tr>
<td>Feb 1, 2008 (until present) Roche COBAS HIV-1 Ampliprep Taqman assay; method of automated amplification and detection changed</td>
<td>40–1 000 000</td>
</tr>
</tbody>
</table>

Table 1: HIV-1 RNA viral load assays used in British Columbia, Canada, 1996–2009
value was carried forward until a new measurement was available, and patients were censored if they moved out of the province or died. We divided follow-up into three discrete time periods on the basis of data for HAART use in British Columbia, all of which were consistent with contemporary treatment guidelines. The first period, from 1996 to 1999, relates to the initial rollout of HAART. The second, from 2000 to 2003, relates to a steady-state phase of HAART use. The third, from 2004 to 2009, relates to the second HAART expansion. We estimated rates of HIV, infectious syphilis, genital gonorrhoea, and genital chlamydia by dividing the yearly number of new diagnosed cases of each disease in British Columbia by the yearly population of the province, and then multiplied by 100 000. Population estimates for the years studied were obtained from British Columbia Stats reports. We applied the HIV case rate for the first year of each study period to all subsequent years in that period to estimate the number of new HIV diagnoses that would have been expected had the case rate remained constant during each phase of the study. We then calculated rate ratios by dividing the number of reported cases by the expected number of cases, and calculated confidence intervals using Byar’s approximation methods.

CD4 cell counts were measured by flow cytometry (Beckman Coulter Inc, Mississauga, ON, Canada). Baseline CD4 cell counts measured before initiation of antiretroviral treatment were obtained from the BCCFE registry, which captures more than 80% of all tests of CD4 cell count done in the province. Physicians are requested to provide patients’ baseline information before initiation of antiretroviral therapy, including a pretreatment CD4 cell count. We tested trends in baseline CD4 cell count in each of the study periods using the Cochran-Armitage trend test for the proportion of measurements lower than 200 cells per μL.

We have shown previously that reduction in community viral load as a result of HAART was a key determinant of decreasing HIV incidence in a cohort of IDUs in the downtown eastside of Vancouver. Therefore, we repeated analyses in the present study after stratification by history of injecting drug use. Grouping of patients on the basis of injecting drug use relied mainly on self-reports and case-report forms completed by health-care providers and public health nurses, which were collected by the BCCDC. Additionally, data for history of injecting drug use were strengthened by linkages to all BCCFE-based cohort studies, including the Vancouver Injection Drug Users Study, the At-Risk Youth Study, the Scientific Evaluation of Supervised Injecting site, and MAKA cohorts.

The BCCFE received ethics approval from the University of British Columbia ethics review committee at the St Paul’s Hospital, Providence Health Care site (P05-123) to undertake this study. The programme also conforms to the province’s Freedom of Information and Protection of Privacy Act.

### Statistical analysis

We modelled trends of new HIV positive tests and number of individuals on HAART using generalised additive models. Our motivation for using such models was the potential to model the non-linear temporal effects without making strong assumptions about the parametric form of these trends. Poisson log-linear regression models were used to estimate the association between the outcome (new HIV positive tests per 100 population) and the covariates (viral load [log_{10}], transformed), year, and number of individuals on HAART). All models were used and implemented in R (version 2.8.1), p values are double-sided, and significance is at the 5% level. All p values are for trend, unless otherwise specified.

### Role of the funding source

The sponsors had no role in the design, data collection, data analysis, data interpretation, and writing or revisions of the report. The corresponding author had full access to all data in the study and had final responsibility to submit for publication.

### Results

Between 1996 and 2009, the number of individuals actively receiving HAART in British Columbia increased from 837 to 5413 (547%; p=0·002), and the number new HIV diagnoses fell from 702 to 338 cases per year (–52%; p=0·001). The overall correlation between number of individuals on HAART and number of new HIV diagnoses per year was –0·89 (p=0·0001).

HAART usage and new yearly HIV diagnoses showed three distinct phases during the study (figure 1). Between 1996 and 1999, we retrospectively noted a steep increase in number of individuals on HAART (258%; p=0·021), during the initial rollout of HAART in the province as a result of the 1996 IAS-USA guidelines. During this period, new HIV diagnoses per year decreased sharply (–40%; p=0·003). Between 2000 and 2003, HAART use increased slightly (9%; p=0·001) because of the balance between treatment interruptions and HAART initiations (new and returning patients). During this period, new HIV diagnoses per year also remained fairly stable (5%; p=0·954). Between 2004 and 2009, we prospectively recorded a second slow but steady increase in the number of individuals on HAART (51%; p=0·001), attributable to the emerging 2004 IAS-USA guidelines, which recommended against structured treatment interruptions. During this third period, new HIV diagnoses per year decreased substantially (–23%; p=0·0001). When data were stratified by history of injecting drug use, there was a 50% reduction in new diagnoses per year in individuals with a history of injecting drug use, from 159 cases in 1999, to 80 cases in 2009 (p=0·003). By contrast, the number of new positive tests remained stable (p=0·6229) in individuals with no history of injecting drug use.

We compared the number of new HIV diagnoses that would have been expected if the rate of new diagnoses...
had remained constant in each of the three study phases with the number of diagnoses reported. For 1996–2000, we noted that there were 30% (rate ratio 0·70, 95% CI 0·67–0·72) fewer new HIV diagnoses than were expected (figure 2). For 2001–03, we showed that the number of reported and expected new HIV diagnoses were almost the same—a decrease of 2% (0·98, 0·93–1·04). In the last period, 2004–09, there was a 17% (0·83, 0·80–0·87) reduction in new HIV diagnoses compared with expected values.

Figure 1: Number of active HAART participants and number of new HIV diagnoses per year in British Columbia, Canada, 1996–2009
p values are for trend and were obtained from the generalised additive model. Injecting drug user (IDU) refers to individuals who have ever injected illicit drugs. HAART=highly active antiretroviral therapy. BC=British Columbia. NA=not available.

Figure 2: Reported and expected number of new HIV diagnoses per year in British Columbia, Canada, during the three phases of the study, 1996–2009
p values refer to the total reported number of HIV diagnoses compared with the total expected number of HIV diagnoses at the end of each study phase.
strikingly during the study, rising from lower than 10% in 1996, to more than 50% in 2009 (p<0·0001). Additionally, between 2004 and 2009, the proportion of patients with no history of injecting drug use who had a viral load lower than 500 copies per mL increased by 36% (p<0·0002) and lower than 50 copies per mL by 42% (p=0·001). By contrast, the same proportions in patients with a history of injecting drug use increased by 82% (p=0·001) and 86% (p=0·002), respectively. Poisson regression modelling of the association between new HIV diagnoses, viral load, year, and number of individuals on HAART showed that for every 100 additional individuals on HAART, the estimated number of new HIV cases decreased by a factor of 0·97 (95% CI 0·96–0·98), and per 1 log10 decrease in viral load, the estimated number of new HIV cases decreased by a factor of 0·86 (0·75–0·98).

We considered whether the fall in new HIV diagnoses could have resulted from reduced HIV testing during periods of increased HAART use, but concluded that the total number of HIV tests done in the province had actually increased steadily during the study. The average number of HIV tests done was 137 585 per year between 1996 and 1999 (an average of 3·5% of the population), 139 464 per year between 2000 and 2003 (3·4%), and 168 924 per year between 2004 and 2008 (4·0%). Also of note, rates of infectious syphilis, genital gonorrhoea, and genital chlamydia increased steadily between 1996 and 2008. For infectious syphilis, the crude rate per 100 000 population increased from 0·5 in 1996 to 7·4 in 2008; for genital gonorrhoea, the crude rate per 100 000 population increased from 12·6 in 1996 to 31·3 in 2008; and for genital chlamydia, the crude rate per 100 000 population increased from 106·2 in 1996 to 239·3 in 2008. The rate of hepatitis C decreased from 158·2 per 100 000 population in 1996, to 87·0 per 100 000 population in 2003 (45% decrease; p<0·0001), and to 55·8 per 100 000 population in 2008 (36% decrease from 2003; p<0·0001).

**Discussion**

Our results show a strong and significant association between increased HAART coverage, reduced community viral load, and decreased number of new HIV diagnoses per year in the population of a Canadian province. We had a unique opportunity to characterise the evolution of these variables during a 15-year period within a universal health-care environment with centralised and free access to HAART. During our study, the pattern of use of HAART changed strikingly on the basis of contemporary therapeutic guidelines. Three distinct phases were noted: two phases of steady expansion of HAART coverage, separated by a period of stable HAART use between 2000 and 2003. Community viral load and new HIV diagnoses...
per year decreased substantially during both phases of HAART expansion. By contrast, community viral load and new HIV diagnoses per year remained stable during the intervening steady period of HAART use. New HIV diagnoses fell by 30% in 1996–2000, remained fairly stable in 2001–03 (2% reduction), and decreased by 17% in 2004–09. Notably, rates of viral load suppression have increased steadily in British Columbia since 1996, as reported by Gill and colleagues.36 The results in whom median community viral load was the strongest driving force behind HIV seroconversion.26 The results are also consistent with those of Moupali Das-Douglas and colleagues,28 who described an association between decreasing community plasma viral load and new HIV diagnoses in San Francisco. Taken together, the reports confirm that expansion of HAART coverage within medical guidelines is associated with reductions in population or community plasma viral load. Reassuringly, this effect was also present in individuals with a history of injecting drug use.27 Furthermore, the results of our multivariable Poisson log-linear regression model are consistent with the previously proposed secondary preventive benefit of HAART used within existing therapeutic guidelines.

Since our report is based on an ecological study, our results cannot be taken as definitive proof of causality. Notably, however, our findings occurred against a background of an increasing number of HIV tests done each year in the province, suggesting that case-finding efforts did not decrease during the study. Furthermore, mandatory (nominal or non-nominal) reporting was implemented in the province in 2003, which improved follow-up and risk ascertainment. This change would have allowed for improved identification of individuals migrating to the province who were infected with HIV, and might have produced a small liberal bias (for the entire cohort after 2003 this subset represented fewer than two per 1 000 000 population). However, the liberal bias is likely to have been overcompensated for by a conservative bias resulting from the overall increase in number of new cases identified by the new strategy. A rise in cases associated with increased screening is consistent with

<table>
<thead>
<tr>
<th>Individuals who have never injected drugs</th>
<th>Individuals who have ever injected drugs</th>
</tr>
</thead>
<tbody>
<tr>
<td>n</td>
<td>Median HIV-1 RNA plasma concentration (copies per mL, IQR)</td>
</tr>
<tr>
<td>-----------------------------------------</td>
<td>-----------------------------------------</td>
</tr>
<tr>
<td>1996</td>
<td>2093</td>
</tr>
<tr>
<td>1997</td>
<td>2848</td>
</tr>
<tr>
<td>1998</td>
<td>3224</td>
</tr>
<tr>
<td>1999</td>
<td>3740</td>
</tr>
<tr>
<td>2000</td>
<td>4114</td>
</tr>
<tr>
<td>2001</td>
<td>4535</td>
</tr>
<tr>
<td>2002</td>
<td>4950</td>
</tr>
<tr>
<td>2003</td>
<td>5303</td>
</tr>
<tr>
<td>2004</td>
<td>5848</td>
</tr>
<tr>
<td>2005</td>
<td>6174</td>
</tr>
<tr>
<td>2006</td>
<td>6426</td>
</tr>
<tr>
<td>2007</td>
<td>6745</td>
</tr>
<tr>
<td>2008</td>
<td>7301</td>
</tr>
<tr>
<td>2009</td>
<td>8001</td>
</tr>
</tbody>
</table>

Data are n, median (IQR), or n (%).

Table 2: Distribution of the highest HIV-1 RNA plasma concentrations for all individuals in British Columbia, Canada who ever had an HIV-1 RNA level in a plasma test, 1996–2009
findings from the US Centers for Disease Control and Prevention, which showed a 15% increase in HIV diagnoses between 2004 and 2007 in 34 US states, partly attributable to enhanced testing efforts in these states.36

Rates of sexually transmitted infections increased during the last 15 years of our study, which implies that our findings cannot be accounted for by decreasing sexual HIV risk behaviour. Rates of hepatitis C decreased during the study, however, the timing of this decrease was clearly out of synchrony with the changes in new HIV diagnoses in IDUs. Furthermore, hepatitis C only became reportable in British Columbia in 1993, and as a result, a large proportion of the cases newly identified in the ensuing years were prevalent rather than incident cases. Therefore, the fall in yearly hepatitis C diagnoses, from 1996 to 2003 in particular, probably represents a reduction in the pool of prevalent cases identified rather than a steep decrease in new cases in IDUs. This effect is less likely to have been a contributor in later years of the study.

Notably, our results were internally reproducible, since we recorded decreases in new HIV diagnoses per year during two distinct periods of HAART expansion that were separated by a stable period of HAART use. The stable period was characterised on a prospective basis37 and was entirely consistent with the effect predicted by our mathematical models.38,39 Furthermore, the reductions in plasma viral load that were documented during HAART expansion provide a plausible mechanistic pathway to account for the association. Ample supportive evidence exists regarding the preventive effect of HAART in plasma viral load that were documented during the ensuing years were prevalent rather than incident cases. Therefore, the fall in yearly hepatitis C diagnoses, from 1996 to 2003 in particular, probably represents a reduction in the pool of prevalent cases identified rather than a steep decrease in new cases in IDUs. This effect is less likely to have been a contributor in later years of the study.

Thus, taken together, the available evidence strongly suggests that community viral load is a key driving force of new HIV diagnoses and can be successfully modulated through effective expansion of HAART coverage within medical guidelines. Our results provide a strong rationale for re-examination of the HIV prevention and treatment dichotomy, as has been strongly advocated by the UN Joint Programme on HIV/AIDS as part of a comprehensive combination prevention strategy.40 Furthermore, our results should serve to re-energise the G8’s universal access pledge as a means to curb the effect of AIDS and the growth of the HIV pandemic.

Contributors
JSGM, VDL, RB, EW, TK, KS, PRH, and RSH contributed to conception and design of the study. Data collection and linkages were undertaken by VDL, RB, EW, TK, KS, PRH, RSH, PD, and PK; Data analysis was done by VDL and BY. JSGM, VDL, RB, EW, TK, KS, PRH, RSH, PD, and PK interpreted the results. JSGM and VDL drafted the report. JSGM, VDL, RB, EW, TK, KS, PRH, RSH, PD, and PK contributed to editing and review of the report. JSGM secured financial support.

Conflicts of interest
JSGM has received funding from Merck, Gilead and ViV Healthcare to support research into Treatment as Prevention, consultancy fees from Merck, and speakers’ fees from Clinical Care Options. PRH holds the GlaxoSmithKline/Canadian Institutes of Health Research Chair in Clinical Virology. RSH has received a research grant from Merck and a conference travel grant from GlaxoSmithKline. PRH has received honoraria from Viiv, Virco, Quest, and Merck, travel grants to attend conferences from Viiv and Quest, research grants from Abbott, Viiv, Pfizer, and Merck, and speakers’ fees from Viiv. VDL, TK, EW, KS, RB, PD, FK, and BY declare that they have no conflicts of interest.

Acknowledgments
JSGM is supported by the Ministry of Health Services and the Ministry of Healthy Living and Sport, from the Province of British Columbia; through a Knowledge Translation Award from the Canadian Institutes of Health Research (CIHR); and through an Avant-Garde Award (No IDP1A026182-01) from the National Institute on Drug Abuse, at the US National Institutes of Health. TK is supported by CIHR and the Michael Smith Foundation for Health Research for Research Excellence. VDL is supported through Fellowship Awards by CIHR and Michael Smith Foundation for Health Research. The study was further supported by the US National Institutes of Health (RO1DA021525) and CIHR (MOP-79297, RAA-79918).

Merck, Gilead and ViV Healthcare support research into Treatment as Prevention. We thank Kelly Hsu for her assistance with preparation of the report, and Tom Patterson and Steffanie Strathdee for their helpful comments and advice.

References

www.thelancet.com Published online July 18, 2010 DOI:10.1016/S0140-6736(10)60936-1


