Trends of non-HIV chronic comorbidities among HIV-positive individuals on highly active antiretroviral therapy in British Columbia from 2000-2009

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Background

- Since the advent of highly active antiretroviral therapy (HAART) in 1996, successes in reducing HIV and AIDS related morbidity and mortality have led to emerging issues of aging with HIV.
- Due to the complex medical history of HIV positive patients, they are at higher risk for a number of chronic conditions that are typically associated with aging in HIV negative individuals.
- We aim to characterize ten year follow-up of chronic disease incidence among HIV positive BC residents during a period of time within the HAART era.

Table 1: Baseline Cohort Characteristics (n=8620)								
Characteristic	Percent	n						
Age		8620						
≤35	29.4%							
36-50	55.2%							
>50	15.3%							
Sex		8619						
Male	83.2%							
Female	16.8%							
Ethnicity		5249						
First Nation	20.6%							
Asian	5.3%							
Black	3.6%							
White	75.0%							
Hispanic	3.2%							
HIV Risk Group								
Known MSM	49.1%	6096						
Known Hetero Risk	30.0%	6096						
Known Blood Risk	4.6%	6096						
Known PWID	43.8%	6956						
Death during study period	23.6%							
Median CD4 cell count (cells/µL; IQR)	240 (130-370)	8448						
Median pVL (copies/mL; IQR)	80000 (20900-110000)	6270						
Weighted CCI categories		7259						
No morbidity (CCI = 0)	67.4%							
Low morbidity (CCI = 1 to 5)	2.8%							
High morbidity (CCI ≥ 6)	29.8%							
MSM: Men who have sex with men; PWID: People who inject drugs; pVL: plasma								
HIV-1 RNA viral load; IQR: inter-quartile range; CCI: Charlson Comorbidity Index								

Methods

Study Population

 The individuals included in this analysis were identified as HIV-positive BC residents aged ≥ 19 and initiating HAART between April 1st, 1996 to March 31st, 2010 from the Comparative Outcomes And Service utilization Trends study (COAST) dataset.

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Study Design

- We conducted a retrospective population-based cohort study of ageassociated chronic disease incidence from HIV positive individuals.
- We assessed incidence rates of six chronic diseases (diabetes mellitus (DM), hypertension (HTN), cardiovascular disease (CVD), chronic obstructive pulmonary disease / asthma (COPD), chronic liver disease (CLD) and chronic kidney disease (CKD)) in the study population; incidence rates were determined for each disease over a period of ten years, from January 1, 2000 to December 31, 2009.
- For each analysis, prevalent cases were identified pre-baseline during a 4-year washout period (April 1st, 1996 to December 31st, 1999) and were excluded from the analyses.

Statistical Analysis

- Poisson's log-linear regression analysis was used to measure trends in





Figure 2. Adjusted Incidence Rate (per 1000 person year) over time

incidence rates for each disease-specific analysis.

 Adjusted incidence rates were determined by controlling for variables: age at baseline, sex, baseline weighted Charlson Comorbidity Index (CCI), and baseline CD4 cell count and PVL (log-10).

Results

- The study cohort (n=8620) was predominantly white (75%, based on known ethnicity) male (83%) with a median CD4 cell count of 240 cells/ µL and PVL of 80,000 copies/mL at HAART initiation.
- Adjusted incidence rate trends per 1000 person-years of DM, HTN and CKD significantly increased over a 10-year span (p<0.001 for each disease).
- Conversely, adjusted incidence rate trend per 1000 person-years of CLD significantly decreased over a 10-year span (p<0.001). Incidence rates of CLD were also much greater than other chronic diseases at each time point.
- Incidence rate patterns for CVD and COPD/asthma did not significantly change over the study period (Figure 1).

Table 2: Crude and adjusted incidence rates (per 1000 person year) of 6 chronic
diseases from 2000-2009 and associated p-value for Poission log-linear regression
test for trend

	2000	2001	2002	2003	2004	2005	2006	2007	2008	2009	p-value
Crude Incidence Rates (per 1000 person year)											
DM	5.86	5.14	6.3	8.13	6.87	6.4	7.1	11.8	11.8	8.69	<0.001
CVD	12.7	9.6	10.4	10.6	13.4	13.7	12.4	12.7	11.2	12.4	0.508
HTN	10.1	5.75	9.3	8.75	9.73	9.9	14.7	15.8	21.6	14.7	<0.001
COPD	12.7	9.01	9.83	8.97	9.18	10.4	10	10.3	11.2	11.6	0.634
CLD	34.7	59.8	46	39.9	52.8	43.3	42	29.1	24.3	22.3	<0.001
CKD	4.68	1.87	2.58	5.21	8.6	8.31	6.46	8.31	8.45	7.47	<0.001

Adjusted Incidence Rates (per 1000 person year)

adjusted for age, sex, baseline weighted CCI, CD4 and PVL(log-10)



DM: diabetes mellitus; HTN: hypertension; CVD: cardiovascular disease; COPD: chronic obstructive pulmonary disease / asthma; CLD: chronic liver disease; CKD: chronic kidney disease * p<.001

DM	2.6	2.92	4.84	6.11	6.58	6.34	5.73	9.59	10.2	6.75	<0.001
CVD	7.64	6.62	5.74	9.1	12.3	9.23	7.6	9.77	11.8	9.49	0.055
HTN	5.21	5.11	5.76	5.2	6.11	5.92	10.4	12.6	17.6	9.76	<0.001
COPD	15.7	12.4	11	11.6	13.6	14.2	13.6	12.6	16.5	15.9	0.315
CLD	47.7	95.2	63.4	55.5	71.9	50.4	58.6	42	32.5	32.1	<0.001
CKD	5.6	1.94	1.86	3.5	7.5	7.56	7.23	6.59	9.57	8.93	<0.001

DM: diabetes mellitus; HTN: hypertension; CVD: cardiovascular disease; COPD: chronic obstructive pulmonary disease / asthma; CLD: chronic liver disease; CKD: chronic kidney disease; CCI: Charlson Comorbidity Index; PVL: plasma HIV-1 RNA viral load; 95% CI: 95% confidence intervals

Conclusions

- We observed marked population-level increases in incidence rates for DM, HTN and CKD and no changes for CVD and COPD/asthma among HIV-positive individuals on HAART over the ten-year period.
- Incidence rates of CLD were much greater than the five other chronic diseases at each time point and also had a decreasing trend over the ten-year period.
- Understanding how these chronic conditions affects future disability and death among the aging HIV-positive population is an important area of further research.



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