

Increasing rates of earlier antiretroviral treatment associated with elevated rates of optimal virologic response among HIV-positive illicit drug users during a Treatment-as-Prevention initiative in Vancouver, Canada

M.-J. Milloy^{1,2}, T. Kerr^{1,3}, R. Hogg^{1,3}, S. Guillemi¹, J. Montaner^{1,2}, E. Wood^{1,2}

1. B.C. Centre for Excellence in HIV/AIDS; 2. Division of AIDS, Department of Medicine, University of British Columbia; 3. Faculty of Health Sciences, Simon Fraser University

Background

- Treatment-as-Prevention (TasP) initiative seek to promptly engage HIV-positive individuals in HIV/AIDS treatment and care in order to minimise HIV/AIDS-associated morbidity, mortality and viral transmission.
- Owing to individual-, social- and structural-level barriers, HIV-positive people who use illicit drugs (PWUD) typically experience sub-optimal rates of antiretroviral therapy (ART) access, adherence and viral suppression
- TasP-based campaigns have yet to be fully evaluated among PWUD
- Additionally, there are concerns that initiating ART earlier in the disease course (i.e., at higher CD4 cell counts) among asymptomatic individuals may lead poorer adherence and degraded virologic response
- Objectives:
 1. Characterize temporal trends in CD4 cell count at ART initiation
 2. Test rates of short-term virologic response by CD4 cell count at ART initiation

Methods

- Data were derived from the AIDS Care Cohort to evaluate Exposure to Survival Services (ACCESS), a long-running prospective cohort of community-recruited drug users in Vancouver, Canada, a setting of free and universal access to HIV care and other essential health services during a TasP-based ART scale-up effort
- Longitudinal cohort data were linked to comprehensive HIV clinical monitoring and ART dispensation records from the BC Centre for Excellence in HIV/AIDS.
- Individuals eligible for these analyses received their first dispensation of ART between December 1, 2005 and June 1, 2013
- First, we examined the study sample characteristics, stratified by year of ART initiation, and produced plots of mean CD4 cell count at initiation by year of initiation
- Second, we used multivariable linear regression to model the relationship between CD4 cell count at initiation (outcome) and year of initiation (primary explanatory variable) and other secondary explanatory variables
- Third, we plotted survival curves using Kaplan-Meier methods to visualize time to non-detectable HIV RNA viral load (VL) (i.e., < 50 c/mL) in the first 365 days following ART initiation, stratified by CD4 cell count at initiation (i.e., < 200 vs. ≥ 200 and ≤ 350 and > 350 cells/mL)
- Finally, we used multivariable Cox proportional hazards regression to model the effect of CD4 cell count on time to VL non-detectability

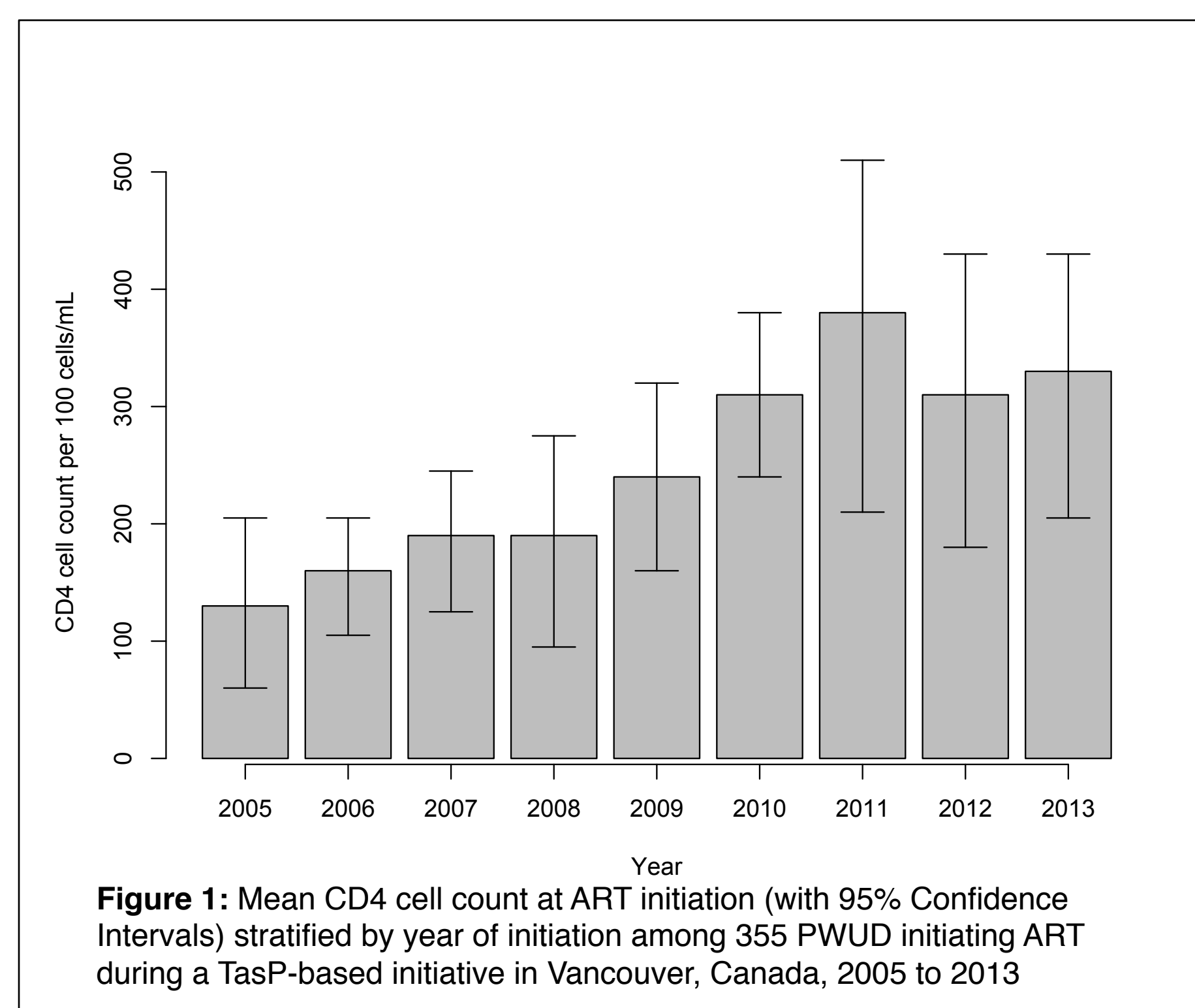


Figure 1: Mean CD4 cell count at ART initiation (with 95% Confidence Intervals) stratified by year of initiation among 355 PWUD initiating ART during a TasP-based initiative in Vancouver, Canada, 2005 to 2013

Results

- Between December 1, 2005 and June 1, 2013, 816 individuals were recruited; among these, 355 (42%) received their first dispensation of ART during the study period and were eligible for these analyses
- Among these individuals, 130 (37%) reported non-male gender and 200 (56%) reported Caucasian ancestry
- Mean CD4 cell count at initiation over the study period was 200 c/mL, increasing from 130 c/mL (2005) to 330 c/mL (2013); highest value was 380 c/mL in 2011
- In a multivariable linear regression model adjusted for gender and the experience of the prescribing physician, later year of initiation was associated with an increase of 29.5 (95% Confidence Interval [95% CI]: 21.0 – 37.9) CD4 cells at initiation
- In a multivariable Cox proportional hazards regression model adjusted for age and VL at ART initiation, year of initiation was positively associated with swifter rates of virologic response (Adjusted Hazards Ratio = 1.14 per 100 cells, 95% CI: 1.06 – 1.23)

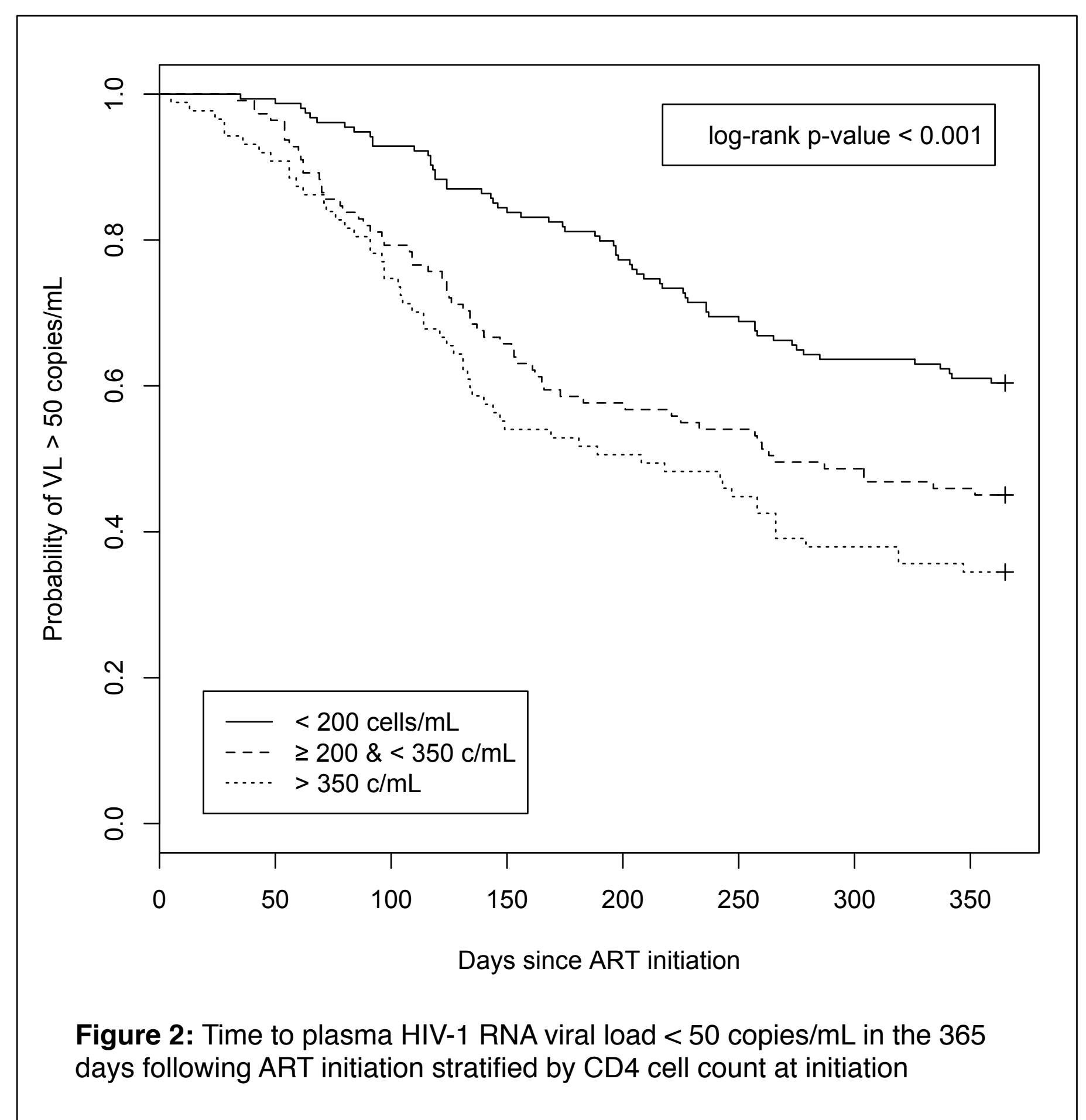


Figure 2: Time to plasma HIV-1 RNA viral load < 50 copies/mL in the 365 days following ART initiation stratified by CD4 cell count at initiation

Discussion

- We observed that individuals were initiating ART earlier in the disease course in a community-based TasP ART scale-up initiative
- Our findings stand in contrast to earlier studies among HIV+ illicit drug users, who typically do not initiate treatment or do so at lower CD4 cell counts
- In contrast to concerns that earlier initiation would lead to poorer treatment outcomes, we found that individuals initiating at higher CD4 cell counts exhibited swifter rates of optimal virologic response
- Our findings support the continued scale-up of ART among PWUD in order to reach 90-90-90 targets to eliminate substantial HIV/AIDS associated morbidity, mortality and viral transmission among this vulnerable population

Acknowledgements

We wish to thank the study participants for their contribution to the research, as well as current and past researchers and staff.

The study is supported by the US National Institutes of Health (R01-DA021525)

Dr. Milloy is supported in part by the United States National Institutes of Health (R01-DA021525)

This work was supported in part by a Tier 1 Canadian Research Chair in Inner-City Medicine awarded to Dr. Wood.

Dr. Montaner is supported by the British Columbia Ministry of Health and through and Avante-Garde Award (No. 1DP1DA026182) from the National Institute of Drug Abuse (NIDA), at the US National Institutes of Health (NIH). He has also received financial support from the International AIDS Society, United Nations AIDS Program, World Health Organization, National Institutes of Health Research-Office of AIDS Research, National Institute of Allergy & Infectious Diseases, the United States President's Emergency Plan for AIDS Relief (PEPFAR), UNICEF, the University of British Columbia, Simon Fraser University, Providence Health Care and Vancouver Coastal Health Authority