Cumulative Viral Load and Virologic Decay Patterns after Antiretroviral Therapy in HIV-Infected Subjects Influence CD4 Recovery and AIDS

Vincent Marconi, Emory University
Greg Grandits, University of Minnesota
Jason F. Okulicz, San Antonio Military Medical Center
Glenn Wortmann, Walter Reed Army Medical Center
Anuradha Ganesan, National Naval Medical Center
Nancy Crum-Cianflone, Naval Medical Center San Diego
Michael Polis, National Institutes of Health
Michael Landrum, San Antonio Military Medical Center
Matthew J. Dolan, Wilford Hall United States Air Force Medical Center
Sunil K. Ahuja, University of Texas Health Science Center

Only first 10 authors above; see publication for full author list.

Journal Title: PLoS ONE
Volume: Volume 6, Number 5
Publisher: Public Library of Science | 2011-05-20, Pages e17956-e17956
Type of Work: Article | Final Publisher PDF
Publisher DOI: 10.1371/journal.pone.0017956
Permanent URL: http://pid.emory.edu/ark:/25593/f7jd7

Final published version:
http://www.plosone.org/article/info%3Adoi%2F10.1371%2Fjournal.pone.0017956

Copyright information:
This is an Open Access work distributed under the terms of the Creative Commons Universal : Public Domain Dedication License (http://creativecommons.org/publicdomain/zero/1.0/).

Accessed February 22, 2018 11:00 PM EST
Cumulative Viral Load and Virologic Decay Patterns after Antiretroviral Therapy in HIV-Infected Subjects Influence CD4 Recovery and AIDS

Vincent C. Marconi1,2*, Greg Grandits2,3, Jason F. Okulicz2,4, Glenn Wortmann2,5, Anuradha Ganesan2,6, Nancy Crum-Cianflone2,7, Michael Polis2,8, Michael Landrum2,4, Matthew J. Dolan9, Sunil K. Ahuja10,11,12, Brian Agan2, Hemant Kulkarni10,11,12, the Infectious Disease Clinical Research Program (IDCRP) HIV Working Group*

1 Division of Infectious Diseases, Emory University School of Medicine, Atlanta, Georgia, United States of America, 2 Infectious Disease Clinical Research Program, Uniformed Services University of the Health Sciences, Bethesda, Maryland, United States of America, 3 Division of Biostatistics, University of Minnesota, Minneapolis, Minnesota, United States of America, 4 Infectious Disease Service, San Antonio Military Medical Center, Brooke Army Medical Center, Fort Sam Houston, Texas, United States of America, 5 Infectious Disease Service, Walter Reed Army Medical Center, Washington, D.C., United States of America, 6 Infectious Disease Clinic, National Naval Medical Center, Bethesda, Maryland, United States of America, 7 Infectious Disease Clinic, Naval Medical Center San Diego, San Diego, California, United States of America, 8 National Institute for Allergy and Infectious Diseases, National Institutes of Health, Bethesda, Maryland, United States of America, 9 Henry M. Jackson Foundation, Wilford Hall United States Air Force Medical Center, Lackland Air Force Base, Texas, United States of America, 10 Veterans Administration Research Center for AIDS and HIV-1 Infection, South Texas Veterans Health Care System, San Antonio, Texas, United States of America, 11 Department of Medicine, University of Texas Health Science Center, San Antonio, Texas, United States of America, 12 Department of Microbiology and Immunology, and Biochemistry, University of Texas Health Science Center, San Antonio, Texas, United States of America

Abstract

Background: The impact of viral load (VL) decay and cumulative VL on CD4 recovery and AIDS after highly-active antiretroviral therapy (HAART) is unknown.

Methods and Findings: Three virologic kinetic parameters (first year and overall exponential VL decay constants, and first year VL slope) and cumulative VL during HAART were estimated for 2,278 patients who initiated HAART in the U.S. Military HIV Natural History Study. CD4 and VL trajectories were computed using linear and nonlinear Generalized Estimating Equations models. Multivariate Poisson and linear regression models were used to determine associations of VL parameters with CD4 recovery, adjusted for factors known to correlate with immune recovery. Cumulative VL higher than the sample median was independently associated with an increased risk of AIDS (relative risk 2.38, 95% confidence interval 1.56–3.62, p < 0.001). Among patients with VL suppression, first year VL decay and slope were independent predictors of early CD4 recovery (p = 0.001) and overall gain (p < 0.05). Despite VL suppression, those with slow decay during the first year of HAART as well as during the entire therapy period (overall), in general, gained less CD4 cells compared to the other subjects (133 vs. 195.4 cells/μL; p = 0.001) even after adjusting for potential confounders.

Conclusions: In a cohort with free access to healthcare, independent of established predictors of AIDS and CD4 recovery during HAART, cumulative VL and virologic decay patterns were associated with AIDS and distinct aspects of CD4 reconstitution.

Introduction

The initial goal of highly-active antiretroviral therapy (HAART) was to improve AIDS-free survival and attempt to mitigate the harmful effects of treatment. Immune reconstitution via CD4 recovery served as an intermediate marker for response to HAART because of its predictive capacity for AIDS events and death.[1,2,3] Thereafter, virologic suppression became the
primary target for therapy because it was shown to be an appropriate, early predictor of immunologic response and clinical outcomes.[4,5,6,7] Furthermore, it was demonstrated that incomplete suppression of viral replication allowed for the emergence of drug resistance and ultimately virologic failure.[8,9] These findings led to recommendations in the U.S. Department of Health and Human Services guidelines that patients should achieve complete virologic suppression (viral load [VL] <400 copies/mL by 24 weeks or <50 copies/mL by 48 weeks) and maintain suppression thereafter.[10]

Even among patients reaching these virologic targets, there are significant inter-individual differences in the recovery of CD4+ T cells and risk of clinical events, suggesting that other factors may relate to these outcomes.[11,12,13,14,15,16] Age at HAART initiation, pre-HAART VL and CD4 cell count, magnitude of and time to VL suppression all have been shown to influence CD4 recovery and clinical outcomes.[4,13,17,18,19,20,21,22,23,24,25,26] Although the relationship of virologic decay patterns with VL changes during HAART has been described,[23,27,28,29] the impact of these decay patterns on CD4 reconstitution and risk of subsequent clinical AIDS events has not been fully elucidated. Furthermore, it is also conceivable that the overall VL burden, represented as the cumulative VL during HAART, may also influence CD4 recovery and risk of AIDS events. Hence, we determined whether the patterns of virologic decay and the cumulative VL during HAART were associated with AIDS and CD4 recovery after HAART initiation independent of the currently recommended dichotomous measures of VL suppression[10] within a large, observational cohort with free access to medications and care, high rates of adherence, and low rates of injection drug use.[26,30] If virologic decay measures are independently associated with outcomes, this could provide some explanation as to why some individuals experience inadequate treatment response despite achieving virologic suppression. Additionally, cumulative viral load could serve as a sensitive marker for risk of AIDS after HAART beyond traditional measures.

Materials and Methods

Study Participants

The U.S. Military HIV Natural History Study (NHS) is a prospective multicenter observational study of HIV-infected active duty military personnel and other beneficiaries (spouses, dependents, and retired military personnel) from the Army, Navy/Marines and Air Force. Seroconverters (SC) were defined as patients having a documented HIV seronegative date prior to the first positive HIV date (see Table S1). The estimated date of seroconversion for SC was defined as the midpoint between the two dates. All CD4 count, VL, and other measurements were done as part of routine clinical care.[31] The clinically-approved methodology for this testing varied by site and over time. Prior ARV use referred to any antiretroviral therapy not meeting the NHS definition of HAART.[26] HAART initiation was the date when HAART was first prescribed.

Ethics Statement

Participants who provided written informed consent and initiated HAART through July 1, 2006 regardless of regimen continuation were included in the present study. The NHS and this substudy have been approved by each center’s Institutional Review Board and the Uniformed Services University of the Health Sciences Institutional Review Board.

Statistical Analysis

VL Parameters. A primary aim of this study was to capture and summarize the overall and early VL dynamics in such a manner as to permit their eventual use in clinical practice. In that regard, we made the following assumptions: i) by the time HAART is typically initiated for an individual in the NHS a natural steady state VL exists; ii) once potent HAART is initiated there is a rapid decline in the VL followed by a slower decline; and iii) such a typical pattern of VL can be explained on the basis of an exponential decay in the circulating VL. The definitions of the parameters used in this study are shown in Table S1, and the theoretical bases for the estimation of these parameters are further described in Note S1. The composite “virologic decay” refers to the application of an exponential decay equation which has been fitted to all viral loads available for an individual after the initiation of HAART. For a majority of participants in this cohort who have a high level of adherence, the virologic decay pattern corresponds to the concatenation of each “classical” (first, second, etc.) phase of decay for that individual. For some participants, their virologic decay does not follow these patterns due to suboptimal adherence, inadequate drug levels, drug resistance, and treatment interruption.

We computed four VL parameters at the level of each individual: (i–ii) exponential decay constant of VL change during entire duration of HAART (overall) and during the first year of HAART, respectively; (iii) VL slope during the first year of HAART obtained using linear Generalized Estimating Equations (GEE) models; and (iv) cumulative VL (Table S1). The VL parameters described above in i, ii, and iii are designated as VL kinetic parameters. Similarly, we computed the following four CD4 count parameters at the level of the individual: (i–ii) slope of the CD4 count change during and after the first two years of HAART; (iii) mean CD4 count after the first two years of HAART; and (iv) overall gain in CD4 counts (Table S1).

Cohort level analyses. The cohort-level analyses made use of all available CD4+ T cell content and VL data for all subjects to generate time-trend lines or curves using linear and non-linear GEE models, assuming an equal correlation structure. The time-trend curves derived by non-linear GEE modeling were refined further using spline smoothed curves with knots at the end of each year since HAART initiation. The resulting curves describe an overall or composite VL pattern for the cohort.

Association analyses. We estimated the parameters detailed in Table S1 for each individual. The association of these individual level parameters with the risk of AIDS (defined using 1993 clinical criteria[32] but did not include a CD4 count <200 cells/µL as an endpoint) was assessed by Poisson regression models, and with recovery of CD4 counts by linear regression models. In these models we accounted for the potential confounding due to VL suppression by HAART by including two covariates - achievement of overall or composite VL pattern for the cohort.

Results

Cohort-level VL and CD4 changes after HAART initiation

Characteristics of the 2278 participants who initiated HAART are in Table 1. The average follow up time after HAART for
participants was 5.63 years (SD 3.98). Cohort-level non-linear GEE modeling of VL from time-of-HAART initiation in all subjects revealed the following pattern: a precipitous decline in VL during the first year, a temporary rebound at ∼1.6 years post-HAART, followed by a relatively steady-state VL thereafter (Fig. 1A). The VL trajectory of subjects who developed AIDS versus those who remained AIDS-free differed significantly as a decline in VL after HAART initiation was not observed in patients who developed AIDS (Fig. 1B). In all subjects (Fig. 1C) and in those who attained VL suppression (Fig. 1D), VL trajectories differed according to the tertiles of the pre-HAART VL such that those who started with higher VLs (upper and middle tertiles of pre-HAART VL) displayed a sharper decline in VL than those subjects categorized to the lower pre-HAART VL tertile (Fig. 1C-D, Table S2).

The cohort-level trajectories in CD4 counts during HAART revealed two phases of CD4 count changes. In phase I, for all subjects initiating HAART there was a rapid increase in CD4 counts during the first two years, followed in phase II by a slower, sustained gain in CD4 cells (Fig. 1A). We stratified the cohort-level changes in CD4 count gains according to whether subjects attained VL suppression (Fig. 1E). This analysis revealed that during the first year of HAART, rapid and similar gains in CD4 counts (~200 cells on average) were observed in those who did (brown curve) or did not (black curve) attain VL suppression (Fig. 1E). However, in contrast to those who attained VL suppression, the initial gains in CD4 counts were not durable among those who did not achieve VL suppression (Fig. 1E).

### VL kinetic parameters and AIDS risk after HAART

The association of the three VL kinetic parameters and cumulative VL with risk of developing AIDS during HAART was evaluated in separate multivariate models adjusted for length of follow up. For these and the other analyses described later, we dichotomized subjects based on the VL parameters using the median value of the parameter as the cut-off. We included into each multivariate model additional covariates that have been shown to be predictive of immunologic recovery during HAART, including time to VL suppression (Table 2) [26,33,34,35,36,37,38].

The slope and exponential decay constant for VL during the first year of HAART were not predictive of AIDS, whereas a slower overall VL decay showed a statistical trend towards predicting AIDS during HAART, independent of the other covariates (RR 1.38, p = 0.058). A higher than average cumulative VL during HAART (RR = 2.38, 95% CI 1.56–3.62) was associated with the greatest risk of developing AIDS. These results were similar when the VL parameters were estimated excluding the VL measurements recorded after the AIDS event occurred (Table 2) or when the analyses were restricted to seroconverters only (Table S3).

In separate analyses, where attainment of VL suppression during HAART was replaced with VL suppression by 6 or 12 months, overall VL decay constant was a significant independent predictor of AIDS in patients who attained VL suppression at 6 (RR = 1.63, p = 0.007, 95% CI = 1.14–2.38) and 12 (RR = 1.43, p = 0.055, 95% CI = 0.99–2.05) months, whereas in these models, the VL slope or exponential VL decay during the first year were not predictive of AIDS. The cumulative VL remained highly predictive of AIDS risk in those who attained VL suppression during 6 (RR = 1.96, p = 0.004, 95% CI = 1.24–3.13) and 12 (RR = 2.33, p = 0.001, 95% CI = 1.44–3.80) months of HAART. Collectively, these data indicated that a slow overall VL exponential decay and a high cumulative VL during HAART increased AIDS risk after initiation of HAART.

### VL parameters and CD4 Recovery

We next determined whether the VL kinetic and other parameters that were included in the models to assess AIDS risk during HAART also associated with the rate of CD4 gain (Table 3). We found that the VL parameters predicted different aspects of CD4 count recovery even after accounting for factors that we found to be highly predictive of AIDS risk, including prior

### Table 1. Characteristics of subjects on HAART studied.

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Median (IQR) or Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age at HAART (y)</td>
<td>34.27 (29.15–39.61)</td>
</tr>
<tr>
<td>Female gender</td>
<td>188 (8.3%)</td>
</tr>
<tr>
<td>Ethnicity</td>
<td></td>
</tr>
<tr>
<td>European Americans</td>
<td>1006 (44.2%)</td>
</tr>
<tr>
<td>African Americans</td>
<td>1003 (44.0%)</td>
</tr>
<tr>
<td>Hispanic Americans</td>
<td>186 (8.2%)</td>
</tr>
<tr>
<td>Others</td>
<td>83 (3.6%)</td>
</tr>
<tr>
<td>Baseline CD4 (cells/μl)</td>
<td>466 (330–637)</td>
</tr>
<tr>
<td>Nadir CD4 (cells/μl)</td>
<td>278 (167–378)</td>
</tr>
<tr>
<td>Time from nadir CD4 to HAART initiation (y)</td>
<td>0.26 (0.03–1.34)</td>
</tr>
<tr>
<td>Baseline VL (log_{10} copies/ml)</td>
<td>43.8 (3.75–4.88)</td>
</tr>
<tr>
<td>Pre HAART VL (log_{10} copies/ml)</td>
<td>43.5 (3.76–4.85)</td>
</tr>
<tr>
<td>Overall VL decay constant (x10^{-2})</td>
<td>2.57 (1.19–7.06)</td>
</tr>
<tr>
<td>VL decay constant during year one of HAART (x10^{-2})</td>
<td>5.22 (7.8–55.2)</td>
</tr>
<tr>
<td>VL slope (log_{10} copies/ml/month) during year one of HAART</td>
<td>0.16 (0.26–1.42)</td>
</tr>
<tr>
<td>Cumulative VL (log_{10} copies*months/ml)</td>
<td>16.31 (7.14–24.94)</td>
</tr>
<tr>
<td>Average time to HAART initiation (y)</td>
<td>3.60 (0.46–7.88)</td>
</tr>
<tr>
<td>Late HAART era</td>
<td>1579 (69.3%)</td>
</tr>
<tr>
<td>Prior use of ARV</td>
<td>1087 (47.7%)</td>
</tr>
<tr>
<td>AIDS before HAART initiation</td>
<td>139 (6.1%)</td>
</tr>
<tr>
<td>Duration of follow-up on HAART (y)</td>
<td>5.63</td>
</tr>
<tr>
<td>VL measurements per individual per yearb</td>
<td>3.07</td>
</tr>
<tr>
<td>CD4 measurements per individual per yearb</td>
<td>2.65</td>
</tr>
<tr>
<td>AIDS after HAART (%)</td>
<td>12.27%</td>
</tr>
<tr>
<td>VL suppression</td>
<td>1925 (84.5%)</td>
</tr>
<tr>
<td>First twelve months</td>
<td>1113 (64.6%)</td>
</tr>
<tr>
<td>First six months</td>
<td>1178 (65.8%)</td>
</tr>
<tr>
<td>First three months</td>
<td>837 (64.7%)</td>
</tr>
<tr>
<td>CD4 slope in first 2 years after HAART</td>
<td>56.6 (18.0–128.8)</td>
</tr>
<tr>
<td>Mean CD4 count 2 years after HAART (cells/μl)</td>
<td>3.55 (25.3–34.2)</td>
</tr>
</tbody>
</table>

*Values represent the mean.

doi:10.1371/journal.pone.0017956.t001
The history of AIDS, nadir CD4, age at HAART initiation and time to VL suppression. The overall decay rate constant was not predictive of rate of CD4 gain in the first 2 years, but was significantly associated with the rate of CD4 gain after two years of HAART, the mean CD4 count two years after HAART, and the overall gain in CD4 cells (Table 3). The decay constant and VL slope in the first year of HAART were mostly predictive of the rate of CD4 cell gain during the first two years and the overall gain in CD4 cells (Table 3). By contrast, the cumulative VL was only predictive of rate of CD4 gains after 2 years and not the overall gain in the CD4 count (Table 3).

The aforementioned data suggested that a slower VL decay during the first year of HAART is associated with both a reduced rate of CD4 gain in the first two years of HAART and overall gain in CD4 cells (Table 3). By contrast, a slower overall VL decay is more predictive of a reduced rate of CD4 gain after 2 years of HAART, lower mean gains in CD4 counts after 2 years of HAART as well as a reduced overall gain in CD4 cells (Table 3). On the basis of these results, we posited that VL suppressors who had a slow VL decay in the first year of HAART and the entire therapy course (overall) would fare the worst with respect to CD4 recovery. To test this, we categorized VL suppressors into two
groups: those with a slow decay in the first year and slow overall decay were categorized into one group, whereas the remainder (rapid/rapid, rapid/slow, slow/rapid decay in the first year and overall decay, respectively) were grouped together because they had very similar CD4 count trajectories (data not shown). Subjects categorized to the slow early/slow overall decay group were similar to other subjects with respect to age at HAART initiation, ethnicity and nadir CD4 (all p values >0.2).

Notably, VL suppressors categorized to the slow/slow decay category had a significantly muted CD4 cell gains during HAART compared with all other subjects (Fig. 1F). Concordantly, VL suppressors categorized to the slow/slow decay category had a lower rate of CD4 recovery in the first two years (54.6 vs. 80.2 cells/µL/yr, p = 0.002) and after two years (>14.8 vs. 16.5 cells/µL/yr, p = 0.002) of HAART, a lower mean CD4 count after two years (564.6 vs. 614.8 cells/µL, p = 0.005) and a lower absolute CD4 gain (153 vs. 195.4 cells/µL, p = 0.001). We also conducted these analyses for subjects who achieved VL suppression within 6 and 12 months and found highly concordant results (data not shown).

**Discussion**

Not all patients on HAART display robust CD4 cell gains, despite VL suppression.[11,12,13,14,15,16] This has been attributed previously to factors such as pre-HAART VL and nadir CD4, age at HAART initiation, and depth of and time to VL suppression. In this study, we modeled the VL decay and cumulative VL and applied these relatively unique parameters to a large well-characterized cohort in order to determine whether these factors were associated with AIDS risk and CD4 recovery during HAART independent of currently recommended benchmarks of VL suppression at 6 and 12 months.[10] In the participants that we evaluated, the initiation of HAART was associated with a predictable decline in VL that was concomitantly associated with an increase in CD4 counts. Subjects who did or did not achieve VL suppression both experienced, on average, a gain of 200 CD4 cells/µL during the first year of HAART. However, in contrast to those who attained VL suppression, these gains were not sustainable among non-VL suppressors. Notwithstanding the importance of attaining VL suppression or minimizing the time to VL suppression, our data show that in addition to these endpoints, both a slow early (first year of HAART) and slow overall (during entire treatment period) VL decay were independently associated with both a slower rate of and lower absolute CD4 cell gain during HAART. Furthermore, a slower overall VL decay in those who attained VL suppression within 6 and 12 months of HAART initiation, and a higher cumulative VL during HAART were each independent predictors of increased AIDS risk during HAART. These findings suggest that the patterns of VL decay are important factors in addition to VL suppression for CD4 reconstitution and risk of AIDS during HAART.

The pre-HAART VL predicted the subsequent rate of decay during the first year of HAART. Since most patients were able to achieve suppression by 6–12 months, it is not surprising that the rate of decay would be greater for patients with higher initial VLs. This may also suggest that patients with higher initial VLs have a larger proportion of actively replicating, productively-infected cells that are more susceptible to HAART. This is consistent with previous studies that examined decay for patients receiving HAART.[39] It was also intriguing that the cohort-level analyses also revealed that after the initial precipitous decline in VL there was a transient rebound, regardless of initial VL (Fig. 1C). This is due to a combination of individual profiles including a proportion of patients experiencing virologic rebound with subsequent resuppression, a small percentage experiencing rebound and not achieving resuppression, and some patients experiencing blips. It is unclear if this temporary, population-level rebound represents a specific temporal relationship with average time to medication fatigue and/or the development of virologic drug resistance as nearly half of treated patients experience a change in therapy around this time both as reported in this cohort[26] and elsewhere.[40] Even in the absence of complete rebound from poor adherence or drug resistance, periods of increased replication can occur due to pharmacologic changes or altered drug activity in a particular compartment.[29] Modeling data from structured treatment interruption trials have shown that parametric resonance, such as that seen in our study, can occur even in the
absence of drug resistance and complete virologic rebound.[41] This phenomenon can be seen when a system undergoing small oscillations over time (such as during the dynamic equilibrium of viral load setpoint) undergoes a significant dampening (HAART oscillations over time (such as during the dynamic equilibrium of This phenomenon can be seen when a system undergoing small oscillations over time (such as during the dynamic equilibrium of viral load setpoint) undergoes a significant dampening (HAART oscillations over time (such as during the dynamic equilibrium of the overall VL decay pattern. These findings demonstrated that VL suppressors could be stratified into two categories such that those with both a slow early and overall VL decay (slow early/slow overall decay) will achieve CD4 recovery, but the gain in CD4 cells would be significantly muted relative to the absence of drug resistance and complete virologic rebound.[41] This phenomenon can be seen when a system undergoing small oscillations over time (such as during the dynamic equilibrium of viral load setpoint) undergoes a significant dampening (HAART initiation) and then experiences brief periods of external perturbation (brief treatment interruptions).

Although the importance of early virologic suppression and virologic failure on CD4 recovery has been well-described,[21,22,24,42,43,44] much less is known about the impact of rate of decay or detectable VL after initial suppression of VL on CD4 recovery.[25,38,46,47,48] In this study, we incorporated several of these elements within the first parameter (overall VL decay). This parameter provides information on the early trajectory as well as the durability of the VL response after the initial decay. We also evaluated cumulative VL because it could be argued that it is the overall exposure to virus that influences CD4 recovery and AIDS.[38,49,50,51] We found that cumulative VL was a stronger predictor of AIDS risk than CD4 recovery after HAART initiation. Additionally, VL decay or slope within the first year of HAART was not predictive of AIDS, whereas the overall VL decay predicted AIDS even among patients who attained VL suppression during 6 and 12 months of HAART. Thus, it is striking that the risk of AIDS is not impacted by the initial rapid phase of virologic decay, but rather by longitudinal assessments such as the overall decay or cumulative VL. These findings suggest that risk of AIDS during HAART is more sensitive to the VL over time rather than events that occur during the first year of HAART as has been suggested previously.[32,53,54,55] In contrast to this study which examined the impact after HAART, Cole et al. recently found that cumulative VL predicted AIDS or death in absence of HAART independent of known risk factors in the Multicenter AIDS Cohort Study.[56] As the number of serious non-AIDS events during HAART increases relative to the number of AIDS events over time, it will be important to determine the association of overall virologic decay and cumulative VL with serious non-AIDS events as has been demonstrated with cancer[57,58] and renal impairment.[59] This data would also suggest that perhaps the cumulative VL even prior to HAART could be associated with clinical events during HAART, supporting the notion that earlier diagnosis and treatment would further reduce the number of these adverse outcomes.

Table 3. Association of VL parameters with CD4 recovery after HAART initiation in subjects who did not develop AIDS.

<table>
<thead>
<tr>
<th>Outcomes and Adjustment</th>
<th>Overall decay constant</th>
<th>Decay constant in first year</th>
<th>Slope in first year</th>
<th>Cumulative VL</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Coeff (SE)</td>
<td>p</td>
<td>Coeff (SE)</td>
<td>p</td>
</tr>
<tr>
<td>Model 1: Rate of CD4 gain in first 2 years (cells/μl/year)</td>
<td>All subjects</td>
<td>6.98 (9.29)</td>
<td>0.453</td>
<td>32.01 (10.43)</td>
</tr>
<tr>
<td></td>
<td>VL suppressors</td>
<td>3.32 (9.29)</td>
<td>0.721</td>
<td>33.22 (10.42)</td>
</tr>
<tr>
<td></td>
<td>Seroconverters</td>
<td>1.12 (10.54)</td>
<td>0.915</td>
<td>33.09 (11.67)</td>
</tr>
<tr>
<td>Model 2: Rate of CD4 gain after 2 years (cells/μl/year)</td>
<td>All subjects</td>
<td>40.59 (8.46)</td>
<td>&lt;0.001</td>
<td>-5.29 (9.86)</td>
</tr>
<tr>
<td></td>
<td>VL suppressors</td>
<td>41.31 (8.61)</td>
<td>&lt;0.001</td>
<td>-4.63 (10.02)</td>
</tr>
<tr>
<td></td>
<td>Seroconverters</td>
<td>43.59 (9.87)</td>
<td>&lt;0.001</td>
<td>-1.81 (11.34)</td>
</tr>
<tr>
<td>Model 3: Mean CD4 count 2 years after HAART (cells/μl)</td>
<td>All subjects</td>
<td>40.21 (14.40)</td>
<td>0.004</td>
<td>33.67 (16.00)</td>
</tr>
<tr>
<td></td>
<td>VL suppressors</td>
<td>24.97 (14.10)</td>
<td>0.077</td>
<td>23.87 (16.02)</td>
</tr>
<tr>
<td></td>
<td>Seroconverters</td>
<td>32.08 (15.66)</td>
<td>0.041</td>
<td>34.36 (17.58)</td>
</tr>
<tr>
<td>Model 4: Overall gain of CD4 cells (cells/μl)</td>
<td>All subjects</td>
<td>53.62 (17.45)</td>
<td>0.002</td>
<td>41.77 (18.80)</td>
</tr>
<tr>
<td></td>
<td>VL suppressors</td>
<td>51.06 (17.49)</td>
<td>0.004</td>
<td>40.35 (18.96)</td>
</tr>
<tr>
<td></td>
<td>Seroconverters</td>
<td>43.78 (19.40)</td>
<td>0.025</td>
<td>47.42 (20.54)</td>
</tr>
</tbody>
</table>

The results are from multivariate linear regression models, and shown are the linear regression coefficients and their standard errors along with significance values. Each model set has three models for the indicated subjects. Each model is adjusted for covariates that in previous analyses were shown to associate with risk of AIDS and were age at HAART initiation, gender, time from entry into cohort to HAART initiation, African American ethnicity, previous receipt of ARV, AIDS prior to HAART, pre-HAART VL, time to VL suppression and late HAART era (after 2000; Marconi et al.).

doi:10.1371/journal.pone.0017956.t003

Viral Load Decay Modeling
all other subjects (Fig. 1F). Because studies have identified polymorphisms that track the durability of CD4 recovery, it will be important to evaluate whether the patterns of VL decay are in part related to such host factors.\[11\]

The association of the extent of CD4 recovery was strongest with the overall VL decay and not the first year VL decay or the cumulative VL, suggests that these VL parameters may be capturing different aspects of VL changes during HAART (early trajectory and maintenance of suppression). The overall decay provides information throughout the duration of treatment and is not limited to one year of information. Hence, it is probable that the decay pattern occurring after virologic suppression (third phase of VL decay)[60] indexed to the decay patterns that occur immediately after HAART initiation[23] together contribute to the ability of a patient to experience durable immune reconstitution. It remains unclear if the latter phases of immune reconstitution are affected by “blips” or primarily by more substantial viral rebound.[61,62]

In contrast to the overall decay pattern, the cumulative VL is a coarse measure of overall VL burden (total virus exposure) during HAART and does not account for VL decay patterns. For example, a patient who suppresses early but has late rebound might have a comparable cumulative VL to that of a patient with predominantly late virologic suppression. This may partly explain why this parameter as computed may not associate strongly with CD4 recovery. However, another explanation hinges on the use of detectable VLs to compute this parameter. Certainly, patients with complete or repetitive virologic rebounds may experience a loss of CD4 recovery; however, the vast majority of patients in this cohort achieved suppression within the first year and the rate of rebound was low.[26] Therefore, at the frequency of available measurements, the cumulative VL may not capture some of the intermittent or ongoing low-level viremia during HAART which may represent actual viral replication in the setting of periodic HAART interruption. Hence, it is conceivable that computation of the cumulative VL using more frequent measurements and/or single copy assays that assess VL below the detectable threshold of commercial assays might reveal that the cumulative VL is a more sensitive marker of not only AIDS risk but also CD4 recovery.

We investigated a large number of prospectively evaluated subjects who have equal access to healthcare and high rates of adherence to HAART.[26,30] This afforded an excellent opportunity to observe the impact of virologic parameters on CD4 recovery in a setting outside of a clinical trial, making these results more generalizable to the HIV-infected population at large. There are some limitations of this study. This study did not attempt to dissect the components of the VL decay[24,63] and determine what baseline and subsequent factors contribute to these components. For example, different regimens, the existence of drug resistance, variable pharmacokinetics and adherence patterns can result in different rates for the first and second phases of VL decay, respectively.[24,28] To this end, we used HAART era as a covariate in the multivariate model to adjust for regimen potency and prior single or dual ART. Furthermore, although rates of adherence in this cohort[26] are high, the rationale of these analyses was not to understand the impact of adherence on the rate of decay but instead how the decay patterns alone influence subsequent clinical/immunologic outcomes regardless of the level of adherence which in a clinical setting can often be unreliable. We also acknowledge that we studied a total of 52 multivariate models (shown in Tables 2 and 3) and at a global type I error rate of 0.05, 2–3 observed associations are likely to be erroneous. Given the fact, however, that we observed a total of 23 associations to be significant at 0.05 type I error rate, our study results are unlikely to have been influenced by false positive associations due to multiple testing. Finally, although the impact of drug resistance and pharmacokinetic interactions was not examined in this study, prior ARV use was used as a surrogate marker of baseline resistance in the multivariate models.

In summary, our findings underscore that the early and overall patterns of VL decay among VL suppressed patients is an independent determinant of CD4 recovery. In addition, the cumulative VL is a determinant of AIDS risk during HAART. Thus, inter-individual differences in VL decay patterns may partly explain the wide variability in CD4 recovery even among those individuals achieving VL suppression within the recommended timeframe. These results also suggest that regimens that produce the most rapid virologic decay and durable suppression could lead to better clinical/immunologic responses. These parameters could be further developed to enhance clinical trial assessment of ARV regimens and assist clinicians with identifying patients at risk for adverse events beyond standard indicators.

Supporting Information

Table S1 Definitions for various parameters used in this study.
(DOCX)

Table S2 Modeling of VL kinetics based on tertiles of pre-HAART VL.
(DOCX)

Table S3 Association of VL parameters with risk of AIDS development after initiation of HAART among seroconverters.
(DOCX)

Note S1 Statistical concepts in VL parameter estimation.
(DOCX)

Acknowledgments

The authors would like to thank our patients for their enormous contributions over the years and the IDC RP HIV Working Group including: Susan Banks, Mary Bavaro, MD, Helen Chiu, MD, Cathy Decker, MD, Connor Eggleston, Tomas Ferguson, MD, Heather Hairson, Clif Hawkes, MD, Arthur Johnson, MD, Erica Johnson, MD, Alan Lifson, MD, MPH, Grace Macalino, PhD, Jason Maguire, MD, Scott Merritt, Robert O’Connell, MD, Sheila Peel, PhD, John Powers, MD, Rosanne Resner, MD, Edmund Tramoni, MD, Timothy Whitman, MD, and Michael Zapor, MD. The content of this publication is the sole responsibility of the authors and does not necessarily reflect the views or policies of the NIH or the Department of Health and Human Services, the DoD or the Departments of the Army, Navy or Air Force. Mention of trade names, commercial products, or organizations does not imply endorsement by the U.S. Government. This work is original and has not been published elsewhere. Portions were presented at the 17th Conference on Retroviruses and Opportunistic Infections, February 2010, San Francisco, California (Abstract #306).

Author Contributions

Conceived and designed the experiments: VCM HK SKA JFO MD. Analyzed the data: HK VCM GG JFO. Contributed reagents/materials/analysis tools: HK SKA BA MP. Wrote the paper: VCM HK SKA GG. Gathered clinical data: VCM JFO GW AG NCC ML MD BA IDC RP-HWG. Analysis interpretation and manuscript review: VCM GG JFO GW AG NCC MP ML SKA BA HK.
References


