Decay of HIV DNA in the Reservoir and the Impact of Short Treatment Interruption in Kenyan Infants

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We compared change in HIV reservoir DNA following continued antiretroviral therapy (ART) vs short treatment interruption (TI) in early ART-treated Kenyan infants. While HIV DNA in the reservoir decayed with continued ART, HIV DNA levels were similar to pre-TI HIV DNA reservoir levels in most children after short TI.

Keywords. antiretroviral therapy; HIV DNA; infants; reservoir; treatment interruption.

During acute HIV infection, a reservoir of long-lived infected cells is established that persists during antiretroviral treatment (ART) and causes viral rebound upon ART cessation [1, 2]. Experimental nonhuman primate models show that this reservoir is generated within days of infection [3]. Initiating ART early in HIV infection may limit reservoir size, increase time to viral rebound upon treatment cessation, and increase likelihood of post-treatment viral control—but does not ensure remission [2, 4–6]. In case reports, HIV-infected infants with early ART had long periods of remission [7, 8], encouraging efforts to identify interventions that augment early ART to promote post-treatment control in pediatric populations. Evaluating these approaches requires analytical treatment interruption (TI). Thus, it is important to understand whether short TI leads to sustained increases in latently infected cells in the HIV reservoir.

Few studies have measured the impact of TI on HIV reservoir. In adults, TI has been associated with initially increased HIV DNA levels, which return to pre-TI levels after >6 months of ART resumption, suggesting that TI may not cause a lasting increase in HIV-infected cell reservoirs [6, 9]. In contrast, a preliminary report from the Children with HIV Early Antiretroviral Therapy trial (CHER) showed sustained increased HIV DNA levels in 17 infants 2–3 years following a median 11-month TI [10], and a smaller study showed similar results [11]. To understand the impact of shorter TI in infants, we quantified blood HIV DNA reservoir levels of Kenyan infants who were randomized to continued ART vs TI in the Optimizing Pediatric HIV-1 Treatment study (OPH).

METHODS

Study Population

OPH (NCT00428116) was a randomized controlled trial in which, following 24 months of continuous ART (median age at initiation, 5 months), children were randomized to continued ART or TI [12]. Blood was collected at 3-month intervals for CD4 measurements in real time and stored for HIV viral load (performed retrospectively). ART restart criteria were CD4% <20–25%, or >one-third decrease from peak CD4, more advanced World Health Organization stage, or weight-for-age decrease, as previously described [12]. After ART restart, there was no further interruption. Children were excluded from this substudy if HIV RNA was ≥1000 copies/mL during the 6 months prior to the HIV DNA measurements at 24 months post–ART initiation (time of randomization) and 42 months post–ART initiation (18 months following randomization; n = 27), or if samples were not available (n = 1). Seven of these children had virally suppressed samples available for HIV DNA measurement at 75 months post–ART initiation (51 months following randomization).

Laboratory Methods

Plasma HIV RNA was quantified using the Gen-Probe HIV-1 RNA assay (Gen Probe, San Diego, CA) with a limit of detection (LOD) of 2.18 log_{10} copies/mL. DNA was extracted from peripheral blood mononuclear cells (PBMCs) using QIAamp DNA (Qiagen, Valencia, CA). Cellular DNA was quantified using RPP30 ddPCR assay (Bio-Rad, Hercules, CA). HIV DNA was quantified in duplicate by in-house cross-subtype pol polymerase chain reaction (PCR) [13] modified for ddPCR, with LOD of 5 copies/10^6 cells determined using previously validated DNA from ACH2 cells that have a single HIV provirus per cell and HIV-negative genomic DNA controls (Supplementary Figure 1). If results were below the LOD or were >2-fold
discordant between duplicates, additional replicates were performed until >2.5e5 cells were tested. HIV DNA was normalized to RPP30 and HIV DNA copies/10⁶ PBMCs reported.

**Statistical Analysis**
Analysis was performed using R (version 3.4). HIV DNA fold change was compared between continued and TI arms using the Wilcoxon rank-sum test. Correlation between postrandomization peak HIV RNA and HIV DNA fold change was determined using Spearman’s rank-order correlation.

**RESULTS**

**Cohort Characteristics During Initial 24 Months of ART**
During the OPH study, children were treated with 24 months of ART (range, 23–28 months), after which 42 were randomized to TI (n = 21) or continued ART (n = 21). Fourteen children from OPH met criteria for this laboratory substudy (see “Methods”), 7 in each arm. At ART initiation, the median age of the 14 infants was 4.8 months (interquartile range [IQR], 4.4–7 months), median CD4% was 22.5% (IQR, 15%–25%), and the median viral load was 6.5 log₁₀ copies/mL (IQR, 5.7–6.9 log₁₀ copies/mL) (Supplementary Table 1). Initial ART was NNRTI-based for 10 infants and PI-based for 4 infants. Twenty-four copies/mL (IQR, 101–235 HIV DNA copies/10⁶ PBMCs) in the reservoir, and children in the TI arm had a median of 90 HIV DNA copies/10⁶ PBMCs (IQR, 54–485 DNA copies/10⁶ PBMCs; P = .9).

Eighteen months later, after children in the TI arm had resumed ART for >15 months, HIV DNA levels were median 50 HIV DNA copies/10⁶ PBMCs (IQR, 24–174 HIV DNA copies/10⁶ PBMCs) and 185 HIV DNA copies/10⁶ PBMCs (IQR, 92–280 HIV DNA copies/10⁶ PBMCs) in the continued and TI arms, respectively (P = .53). During the 18 months after randomization, 1 of 7 children in the continued arm had increased HIV DNA, 5 of 7 had decreased HIV DNA (range, 0.13–0.63), and 1 of 7 children had HIV DNA levels below detection at both time points (Figure 1B, Supplementary Table 1). In the TI arm, 3 of 7 children had increases in HIV DNA (range, 1.3–4.7), while 4 of 7 had decreased or unchanged HIV DNA levels (range, 0.26–1.04). The median HIV DNA declines during the 18 months after randomization were –5.69 HIV DNA copies/1e6 PBMCs/month in the continued arm and 0.18 in the interrupted arm.

The median HIV DNA fold changes were 0.32 (IQR, 0.22–0.58) in children randomized to continued ART and 1.04 (IQR, 0.72–1.64) in children randomized to TI (P = .14) (Figure 1C). Similar results were observed when we excluded the 2 children in the continued arm with postrandomization viremia: median HIV DNA fold changes were 0.32 (IQR, 0.23–0.46) vs 1.04 (IQR, 0.71–1.64) in the continued and TI arms, respectively (P = .04). In children with viremia during the 18 months after randomization, change in HIV DNA did not correlate with peak viremia (Spearman’s rho, 0.12; P = .77).

In a limited number of children with longer follow-up (2 in the TI arm and 5 in the continued arm), a virally suppressed sample was available from 51 months after randomization (75 months after initial ART). The median HIV DNA declines from 0 to 51 months following randomization were –1.96 HIV DNA copies/1e6 PBMCs/month in the continued arm and –0.37 in the interrupted arm. The median HIV DNA fold changes were 0.44 (IQR, 0.44–0.75) and 0.92 (IQR, 0.84–1.00) in children randomized to continued ART and TI, respectively (Supplementary Figure 2).

**DISCUSSION**
We measured HIV DNA in the reservoir in children that started ART during the first year of life and were randomized 2 years later to continue or interrupt ART. We compared HIV DNA levels at randomization and again 18 months later, after all children had resumed ART for >15 months and achieved viral suppression. HIV DNA decayed in children with continued viral suppression, while the median HIV DNA fold change after TI was 1.04, suggesting that TI lessens the decay that occurs on continued ART. Indeed, for 2 children in the TI arm with ~4 years of follow-up, HIV DNA reservoir size remained relatively unchanged over time (Supplementary Figure 2). The fact that most children in the TI arm had similar HIV DNA levels before and after TI suggests that the reservoir reseeding that occurred during TI was followed by decay after ART resumption. Mechanisms of reservoir seeding include new infections of cells that become quiescent and clonal proliferation of cells containing provirus [14, 15]. Our study could not
distinguish between these mechanisms as we did not characterize the reservoir composition or replication competence due to limited sample volume and cell viability. In addition, our cohort did not include infants treated with very early ART (within hours of birth), which may have different viral reservoir decay dynamics.

Our study did not quantify HIV DNA in tissue reservoirs and was limited by small sample size with few evaluated time points. However, our data add substantially to the 2 previous studies on changes in HIV DNA following TI in children [10, 11], and they support results from adult cohorts [6, 9]. Analysis of 15 adults in the SPARTAC trial with transient TI showed that HIV DNA returned to pre-TI levels after ≥6 months of ART [6]. Another TI study of 10 adults with very low HIV DNA levels in the reservoir prior to a median TI of 4 weeks observed similar results, with HIV DNA levels returning to
pre-TI values after treatment resumption [9]. These studies suggest that increases in HIV DNA can be minimized or reversed by rapid treatment resumption; however, larger studies with longer follow-up are needed.

Our findings add a new perspective that complements prior TI studies in perinatally infected infants, which focused on longer TI. The largest pediatric TI reservoir study to date included 17 infants treated earlier (<12 weeks at ART initiation) and with longer TI (median, 11 months) than in our study, and observed increased HIV DNA in the reservoir 26 months after treatment resumption [10]. Another study included only 3 infants with TI and showed 52.4-, 1.8-, and 12.2-fold increases in HIV DNA following TI of 0.75, 6.8, and 71 months, respectively [11]. Here we show that a short interruption of ~3 months does not appear to have sustained impact on HIV DNA levels in the reservoir in children; however, it may lessen the rate of decay provided by early continued ART. These data suggest that reseeding of the reservoir in pediatric HIV may be minimized with frequent viral load monitoring during short analytical TI.

Supplementary Data
Supplementary materials are available at Open Forum Infectious Diseases online. Consisting of data provided by the authors to benefit the reader, the posted materials are not copyedited and are the sole responsibility of the authors, so questions or comments should be addressed to the corresponding author.

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