Introduction

Elite controllers (EC) represent a group of individuals living with HIV-1 who naturally suppress viral replication to levels below 50–75 copies/mL for at least 12 months [1] without the use of antiretroviral therapy (ART) and rarely progress to AIDS. How these individuals control HIV-1 infection has been a major focus of research as it could provide valuable clues to guide research on vaccines and viral eradication. Multiple non-exclusive reasons have been put forward to explain the control of viremia in this subgroup of individuals. First, the infecting virus could be crippled [2,3], or secondly, the adaptive immune response could result in control of the infection [4–6]. Genetic determinants such as HLA [2,3], or secondly, the adaptive immune response could result in viral control [8] as broadly neutralising antibodies can be identified in a large subset of non-controllers and are less effective on viral control [8] as broadly neutralising antibodies can be identified in a large subset of non-controllers and are less effective on viral control [8].

Activated CD4 T cells from a controller-discordant couple, one partner being an EC and the other an HIV-1 progressor (PR), and healthy controls (HC) were isolated, activated and infected with VSV-G pseudotyped yellow fluorescent protein-encoding single-round HIV-1 virus (HIV-YFP). Viral reverse transcripts, 2-LTR circles and integrated proviral HIV-1 DNA were monitored by quantitative PCR (qPCR) and integration sites were analysed. We further measured LEDGF/p75 and p21 mRNA expression levels by qPCR.

Results: Infection of activated CD4 T cells with HIV-YFP was reduced in EC compared with the PR partner, and HC. Evaluation of viral DNA forms suggested a block after entry and during the early steps of HIV-1 reverse transcription in EC. The integration site distribution pattern in EC, PR and HC was similar. The p21 expression in CD4 T cells of EC was elevated compared with the PR or HC, in line with previous work.

Conclusions: Our study suggests a reduced permissiveness to HIV-1 infection of CD4 T cells from EC due to a block of HIV-1 replication after entry and before integration that might contribute to the EC phenotype in our patient.

Abbreviations: ART, antiretroviral therapy; EC, elite controllers; HIV, human immunodeficiency virus; PR, progressor; HC, healthy controls; CCR5Δ32, CCR5 deficiency; YFP, yellow fluorescent protein; qPCR, quantitative PCR; LEDGF/p75, Lamin-associated DNA binding protein p75.
(CCTCTTCTCTCTATTCCGACA), expected to generate a 190 bp amplicon in case of CCR5Δ32 deletion instead of a 222 bp amplicon in wild-type virus [14].

HLA-typing was performed for HLA B*27 and HLA B*57:01 using a commercially available assay.

HIV subtype and genotypic drug resistance were determined as described by Pineda-Peña et al. [15]. For the EC, a viral blip 2 years after the start of the study was analysed demonstrating an HIV-1 subtype A1 without resistance mutations in HIV-1 reverse transcriptase, protease or integrase. For the PR HIV-1 subtype A1 was identified. No baseline resistance information could be obtained, although, resistance to various treatment regimens developed over the years.

Peripheral blood mononuclear cells (PBMCs) from HIV-1 positive individuals and healthy controls (HC) were isolated in parallel using density gradient centrifugation and stored in liquid nitrogen. Defrosted PBMCs were enriched for CD4 T cells using sorting for CD4 expression with immunomagnetic beads (MACS, Miltenyi) or anti-human CD3.8 bispecific monoclonal antibody according to the manufacturer’s protocol. The latter reagent was obtained through the NIH AIDS Reagent Program, Division of AIDS, NIAID, National Institutes of Health (Anti-Human CD3/8 Bi-specific Monoclonal from Drs Johnson Wong and Galit Alter). Cells were activated and expanded using IL-2 (50 U/mL) and anti-CD3/CD28 antibodies in case bead-based sorting occurred prior to further experiments. CD4 expression was evaluated using flow cytometry with CD4-PE.

VSV-G pseudotyped yellow fluorescent protein-encoding single-round HIV-1 virus (HIV-YFP) was generated using pNL4-3 IRES-YFP and pseudotyped with vesicular stomatitis virus G (VSV-G) as described previously [16]. For transduction with HIV-YFP, cells were seeded at 30,000–500,000 per well in a 96–24-well plate, respectively, and a serial dilution of HIV-YFP was added and analysed for YFP expression using flow cytometry on day 2 and 7 post-transduction.

Quantitative PCR (qPCR) was performed as previously described for LEDGF/p75 [17] and p21 [7]. Late reverse transcripts (late RT), 2-LTR circles at different time points after transduction of activated CD4 T cells with VSV-G pseudotyped HIV-YFP were monitored by quantitative qPCR. Cell lysates, harvested 48 h after transduction, were used for detection of integrated proviral HIV-1 DNA using Alu-Gag PCR, as previously described [18].

LDL-receptor expression was evaluated using LDL-PE (mouse anti-human LDLR Clone 7 IgG-C7, Becton Dickinson) using flow cytometry.

For integration site analysis, cells were cultured 7 days post-transduction with HIV-YFP to minimise residual non-integrated DNA. Amplification and sequencing of proviral integration sites were performed using 454 pyrosequencing. Briefly, gDNA was digested using MseI and linkers were ligated. Proviral-host junctions were amplified by nested PCR using barcoded primers, which enabled pooling of PCR products. Following gel-purification, products were sequenced on the 454 GS-FLX. Authentic integration sites were aligned to the human genome (hg18) using BLAT requiring >98% sequence identity. Statistical methods are detailed in Berry et al [19]. Integration site counts were compared with matched random controls (MRCs) by Fisher’s exact test or by multiple regression models for integration intensity and a c-logit test for significance [19]. Analysis was carried out using R (www.r-project.org). Detailed analysis and bioinformatics were performed using Integration Site Pipeline and Database (INSIPID).

Results

The EC and PR were long-term partners, white and aged 67 years. They were diagnosed with HIV infection 18 years ago. They both had HIV-1 subtype A infection and were presumed to be infected with the same virus although viral sequences at the time of infection were unavailable. The EC had an undetectable HIV-1 viral load since diagnosis and a current CD4 T cell count of 710 cells/mm³. The EC was HLA B*27, B*57:01 and CCR5Δ32 negative. The PR partner had an undetectable HIV-1 viral load while on ART (ritonavir-boosted darunavir, zidovudine, lamivudine, tenofovir disoproxil and raltegravir). The PR’s CD4+ T cell count nadir was 113 cells/mm³ and current count was 540 cells/mm³, and was HLA B*27 and CCR5Δ32 negative, and HLA B*57:01 positive.

Ex vivo transduction of isolated and activated CD4 T cells with a VSV-G pseudotyped HIV-YFP resulted in a reduced transduction (about two-fold lower) in CD4 T cells from the EC compared to the PR partner (Figure 1a), and healthy controls (data not shown), suggesting an intrinsic block of HIV transduction in the EC at a post-entry step. Of note, LDL-receptor expression, a potential cause of reduced VSV-G-mediated entry [20], was similar in activated CD4 T cells from EC and PR (data not shown). Expression of PSIP1 mRNA encoding the LEDGF/p75 protein, a known cofactor for HIV integrase [21], was similar in EC, PR and HC (Figure 1b), but p21 expression was slightly elevated in the EC compared to the PR and HC, consistent with previous results [7] (Figure 1c).

To further dissect the replication block, we have determined viral intermediates at different time points post HIV-YFP transduction (Figure 1d). Already, at 12 hours post-transduction, late RT products in the EC cells were lower compared with PR and HC cells. This indicated an earlier replication block occurring around the RT step. As expected, the number of integrated copies was lower, although this was not due to an additional block at the integration step, since the reduction was similar in terms of late RT products, and the 2-LTR circles were not elevated compared to PR or HC cells or a raltegravir (RAL)-treated HC in whom a clear increase was observed (Figure 1d).

HIV integration is not a random process and preferentially occurs in active transcription units [22]. The site of integration, therefore, can significantly influence transcription and hence viral replication [16,23–26]. Since a reduction in integrated HIV-1 copies might still be accompanied with an altered integration site selection, we have analysed the integration site distribution in EC, PR and HC samples. As the number of proviral DNA copies in CD4 T cells from EC or ART-treated individuals is estimated to be low, at around 10–1000 per million cells [12,27], ex vivo superinfection and analysis of these cells with laboratory viruses/HIV-derived vectors is deemed feasible. Therefore, we determined the integration site distribution pattern (Figure 1e). The genomic heat map demonstrated no clear difference between the integration site distribution pattern in EC, PR and HC (integration preference relative to random is compared to HC, indicated with ‘−’). Moreover, compared to the shift of integration outside RefSeq genes at knockdown or knockout of LEDGF/p75 or by using LEDGINs [25,28], the subtle differences are probably biologically irrelevant.

Discussion

EC individuals naturally control HIV replication in the absence of ART. Evidence suggests a block at the integration step in at least a subset of ECs [12,26]. The integration block in these individuals could be due to cellular restrictions affecting the integration step. Alternatively, the expression or functionality of cellular cofactors...
Figure 1. Analysis of ex vivo infection of a controller-discordant couple. (a) Percentage of YFP-positive CD4 T cells from EC and PR at day 2 and 7 after transduction with a serial dilution (1/10 and 1/20) of HIV-YFP; a representative experiment for multiple independent experiments (n=4), is shown. (b,c) LEDGF/p75 and p21 mRNA expression relative to beta-actin expression (mean, standard deviation) in stimulated CD4 T cells demonstrating only a mild increase in p21 in EC. Statistically significant differences using a one-way ANOVA are indicated with * (**: P = 0.03; ***: P = 0.0025). (d) Kinetics of viral intermediates (late reverse transcripts or Late RT, 2-LTR circles, integrated copies expressed relative to RNaseP) in CD4 T cells from EC, PR, or HC without efavirenz (EFV at 50xIC50) or raltegravir (RAL at 50xIC50) after transduction with HIV-YFP using qPCR demonstrating an early reduction in late RT products in EC compared with PR or HC. (e) For integration site analysis, cells were split and maintained for 7 days before determining the number of integrated copies and integration site analysis. Heat maps were developed to summarise relationships of proviral integration sites with genomic features using the receiver operating characteristic (ROC) area method [19]. The analysed genomic features are mentioned on the left of the corresponding row of the heat map. Tile colour indicates whether a chosen feature is favoured (red, enrichment compared with random) or disfavoured (blue, depletion compared with random) for integration for the respective data sets relative to their MRCs, as detailed in the coloured ROC area scale at the bottom of the panel. The different data sets used are indicated above the columns. The * denote significant differences of HIV integration compared to the LEDGF/p75 knockdown cell line for the respective features (*: P<0.05; ***: P<0.001, using Wald statistics referred to a chi-squared distribution), dashes overlay control tiles. The naming of the genomic features is described in Brady et al. [30]. TSS: transcription start site, EC: elite controller, HC: healthy controller, PR: progressor, YFP: yellow fluorescence protein.
involved in integration might be reduced in a subset of EC, resulting in less integrated HIV-1 provirus. In our EC case a reduction in integrated copies was observed, yet without an increase in 2-LTR circles. Together with a reduction in late RT products, this EC phenotype can be attributed to an early block in the replication cycle, pointing towards a heterogeneity within the EC population.

In line with previous work [7], we have also observed an upregulation of p21 mRNA in CD4 T cells from EC. This has been suggested to contribute to a reduced susceptibility to HIV-1, possibly through inhibiting viral reverse transcription and resulting in reduced 2-LTR cycles and integration [7]. P21 has the ability to inhibit the enzymatic activity of CDK9, a host protein essential for correct elongation of HIV-1 mRNA [29].

Most studies have aimed at examining viral, immunological or cellular biological differences between a group of EC and PR individuals. Multiple determinants underlie EC and considerable variations between experiments with primary cells could be observed. To overcome viral factors, we performed an in-depth study of the mechanisms of HIV-1 control in one EC who was presumed to have the same virus as the PR partner. This EC is characterised by an ex vivo block early after entry. Our work has certain limitations including the absence of evaluation of HIV-entry as well as virus–host adaptive immune responses. However, our findings warrant further validation in other controller-discordant couples and might inform the mechanisms of natural HIV-1 replication control.

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Declaration of interest

All authors declare no conflicts of interest.

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