Proteome-wide analysis of HIV-specific naive and memory CD4+ T cells in unexposed blood donors

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The preexisting HIV-1–specific T cell repertoire must influence both the immunodominance of T cells after infection and immunogenicity of vaccines. We directly compared two methods for measuring the preexisting CD4+ T cell repertoire in healthy HIV-1–negative volunteers, the HLA–peptide tetramer enrichment and T cell library technique, and show high concordance (r = 0.989). Using the library technique, we examined whether naïve, central memory, and/or effector memory CD4+ T cells specific for overlapping peptides spanning the entire HIV-1 proteome were detectable in 10 HLA diverse, HIV-1–unexposed, seronegative donors. HIV-1–specific cells were detected in all donors at a mean of 55 cells/million naive cells and 38.9 and 34.1 cells/million in central and effector memory subsets. Remarkably, peptide mapping showed most epitopes recognized by naïve (88%) and memory (56%) CD4+ T cells had been previously reported in natural HIV-1 infection. Furthermore, 83% of epitopes identified in preexisting memory subsets shared epitope length matches (8–12 amino acids) with human microbiome proteins, suggestive of a possible cross-reactive mechanism. These results underline the power of a proteome-wide analysis of peptide recognition by human T cells for the identification of dominant antigens and provide a baseline for optimizing HIV-1–specific helper cell responses by vaccination.
performed a direct comparison of the T cell library (Geiger et al., 2009) and tetramer enrichment methods (Su et al., 2013) using varicella zoster virus (VZV) as a model antigen. We established CD4+ T cell libraries of 192 wells per subset from naive (CD45RA+CCR7+), central memory (CD45RA−CCR7+), and effector memory (CD45RA−CCR7−) subsets of five anonymous HLADRB1*1501+ blood donors. Lines were polyclonally expanded for 16–20 d before screening for proliferative response to two VZV epitopes (glycoprotein E [gE] and immediate early phosphoprotein 63 [IE63]), which are known to be restricted by and commonly detected in HLADRB1*1501 donors (Jones et al., 2007; Malavige et al., 2008). In parallel 25–70 × 10^6 CD4+ T cells from the same time point were screened using custom-made gE and IE63 HLADRB1*1501 tetramers. We show that the two techniques are highly comparable (Pearson’s correlation, r = 0.999), with no significant difference in the precursor frequencies obtained by each technique (paired Student’s t test, P = 0.1099; Fig. 1). These data support the T cell library technique for comprehensive analysis of naive and memory CD4+ T cell repertoires.

HIV-1–specific T cells in HIV-1–unexposed, seronegative donors

Naive, central memory, and effector memory CD4+ T cell subsets from 10 HLA diverse (Table S1), healthy, HIV-1 seronegative (HESN) donors were screened for HIV-1–specific responses using the T cell library method (Geiger et al., 2009). As above, 192 cell lines per subset were established at limiting dilution and polyclonally expanded. Each cell line was then screened for reactivity to peptides spanning the entire consensus clade C HIV-1 proteome, pooled by proteins Gag, Env, Pol, Nef, and a mix of Vpr, Vpu, Tat, Rev, and Vif (designated Nef/Acc). Responding cells were detected by 3H thymidine incorporation.

The existence of HIV-1–specific memory cells in seronegative donors was originally suggested by studies of highly exposed HIV-1 seronegative (HESN) donors. It has been shown that 25–61% of HESNs have demonstrable HIV-1–specific memory cells, probably primed by exposure to the virus. Surprisingly, HIV-1–specific CD4+ T cells were also detected in 24–44% of unexposed donors (Ritchie et al., 2011), although it was not clear whether the latter came from cross-reactive memory T cells or naive T cells primed in vitro. More recently, the existence of low frequency (1–10/ million) memory CD4+ T cells, specific for a known HIV-1 Gag epitope, was demonstrated by HLA DR4 tetramers in 50% of HIV-1 unexposed HLA DR4+ adults (Su et al., 2013), but it was not clear how generalizable the HIV-1 result was beyond the single epitope–HLA DR4 combination.

The present study first validates the library technique by direct comparison with the tetramer enrichment method for measuring precursor T cell frequencies. We then use the T cell library technique to provide the first proteome-wide analysis of the frequencies and specificities of preexposure HIV-1–specific naive and memory CD4+ T cells in a HLA diverse population of HIV-1 unexposed donors.

RESULTS AND DISCUSSION

Comparison of the T cell library technique to tetramer enrichment

Before commencing a proteome-wide screen of the preexisting HIV-1–specific naive and memory CD4+ T cell repertoires, we first performed a direct comparison of the T cell library (Geiger et al., 2009) and tetramer enrichment methods (Su et al., 2013) using varicella zoster virus (VZV) as a model antigen. We established CD4+ T cell libraries of 192 wells per subset from naive (CD45RA+CCR7+), central memory (CD45RA−CCR7+), and effector memory (CD45RA−CCR7−) subsets of five anonymous HLADRB1*1501+ blood donors. Lines were polyclonally expanded for 16–20 d before screening for proliferative response to two VZV epitopes (glycoprotein E [gE] and immediate early phosphoprotein 63 [IE63]), which are known to be restricted by and commonly detected in HLADRB1*1501 donors (Jones et al., 2007; Malavige et al., 2008). In parallel 25–70 × 10^6 CD4+ T cells from the same time point were screened using custom-made gE and IE63 HLADRB1*1501 tetramers. We show that the two techniques are highly comparable (Pearson’s correlation, r = 0.999), with no significant difference in the precursor frequencies obtained by each technique (paired Student’s t test, P = 0.1099; Fig. 1). These data support the T cell library technique for comprehensive analysis of naive and memory CD4+ T cell repertoires.
In addition to the naive repertoire, HIV-1–specific T cells were also detected within the central memory (Fig. 2 B) and/or effector memory (Fig. 2 C) CD4+ T cell compartments of all donors. However, the observed frequencies were 15–20-fold lower (Fig. 3, B–D) than TT-specific memory CD4+ T cells (Fig. 3 D). The high circulating frequencies of TT-specific memory CD4+ T cells are consistent with previous reports (Geiger et al., 2009) and reflect successful prior immunization (Sallusto et al., 2010). In contrast, the low frequencies

Figure 2. HIV-1–specific responses were detected in the circulating naive and memory CD4+ T cell subsets of healthy, HIV-1 seronegative donors. For each of 10 donors, a mean of 187 cultured cell lines (each represented by a single dot) per naive (A), central memory (B), and effector memory (C) subsets were screened against pools of overlapping peptides spanning the entire HIV-1 proteome. Peptide pools were split according to protein with Nef and Accessory proteins, Vpr, Vpu, Tat, Rev, and Vif included as a single pool referred to as Nef/Acc. All data presented are expressed as the counts per minute, after subtraction of background, nonspecific proliferation (delta cpm). Positive responses are defined as SI >5 and >3,000 delta cpm with cutoff criteria represented by a horizontal line. Cell lines with background counts of ≥3,000 cpm were excluded from analysis. Proliferative responses to known recall antigen TT are shown for each donor.

28 to 129 cells/million (Fig. 3, A and D), comparable to precursor frequencies of tetanus toxoid (TT)–specific naive CD4+ cells (Fig. 3 D). Although the use of a single consensus virus sequence allowed a comprehensive assessment of preexisting specificity, it must underestimate the total response specific for this highly variable virus. However, because 70% of new HIV-1 infections are established by a single founder virus (Keele et al., 2008), the frequencies presented herein should realistically predict an individual’s CD4+ T cell response in acute infection.
of HIV-1–specific memory CD4+ T cells in unexposed donors suggest that these T cells arose because of rare cross-reactivities with non–HIV-1 antigens.

Considerable inter-donor variation in both the specificity and frequency of naive and memory HIV-1–specific CD4+ T cells was observed (Fig. 3, A–C), probably contributing to the great variation in adaptive immune responses seen after natural infection and vaccination. Overall, the HIV-1–specific memory T cell frequencies detected herein were ~10-fold higher than those recently reported using tetramer enrichment (Su et al., 2013). However, this is not a real discrepancy because we included peptides spanning the entire HIV-1 proteome, whereas they focused their analysis to a single known epitope; therefore, the findings are fully compatible.

After normalization of the dataset for viral protein size, HIV-1 Gag and Env proteins were the immunodominant targets of the preexposure naive and memory CD4+ T cell subsets (Fig. 3 E), similar to the known patterns of immunodominance in natural HIV-1 infection (Kaufmann et al., 2004; Ranasinghe et al., 2012). Consistent with the low frequency of Pol-specific responses observed after infection (Kaufmann et al., 2004; Ranasinghe et al., 2012), relatively minimal numbers of mapped epitopes were detected within the Pol protein (Fig. 4). The LANL database was screened to determine whether mapped epitopes had previously been detected in natural HIV-1 infection. Epitopes that had previously been detected in HIV-1 infection are represented with a solid line, whereas those that had not previously been characterized are shown with a dashed line. The number of mapped epitopes that had previously been reported in natural HIV-1 infection as compared with the total number of mapped epitopes for each protein is presented in B. To minimize bias, the number of amino acids covered by epitopes identified in this study was normalized to the total number of amino acids in HIV-1 reference strain HBX2 covered by previously identified or as yet unknown CD4+ T cell epitopes and compared between memory and naive subsets using a Fisher’s exact test (C). Significance (*) was defined as P = 0.0028.
few naive or memory CD4+ T cells specific for Pol were detected in the preexposure repertoire. Low expression of Pol protein in infected cells may account for the lack of Pol-specific responses after infection but cannot explain why these CD4+ T cells are rare in the naive repertoire. Because Pol is a relatively conserved protein, it is possible that T cells with this specificity are more likely to be deleted during thymic selection due to cross-reactivities with endogenous retrovirus sequences.

Peptide specificity and avidity of preexisting HIV-1–specific T cells
We next mapped the fine specificity of 68 preexisting naive, central memory, and effector memory HIV-1–specific CD4+ T cells in a total 68 T cell lines (Fig. 4 and Table S2). Epitopes were found to span the entire HIV-1 proteome (Fig. 4) with the majority of positive cell lines (76%) showing a single, unique peptide specificity, consistent with previous observations that preexisting memory CD4+ T cell responses are largely monoclonal (Geiger et al., 2009; Su et al., 2013).

The functional avidity of a subset of naive (n = 7; Fig. 5 A), central memory (n = 11; Fig. 5 B), and effector memory (n = 7; Fig. 5 C) CD4+ T cell lines was assessed by peptide titration. Overall, the avidities were comparable to those found in T cells responding to natural HIV-1 infection, with both high and low avidity responses (EC50, 3.125–0.001577 µM) detected, but no significant difference between subsets.
The immunogens that primed the preexisting HIV-1–specific memory CD4+ T cells detected within seronegative donors are unknown and likely to come from many sources. Sequence identity searches (Altschul et al., 1990; Edgar, 2010) performed using the reactive HIV-1 peptide sequences identified potential epitope-length (8–12 aa) subsequence matches to a variety of human (Tables S4 B and S5 B) and human microbiome proteins (Tables S4 and S5). Epitope matches ranged from 1 to 230 microbial sequences, with 83% of all HIV-1 epitopes mapped to the CD4+ memory subsets shown to have epitope length (8–12 aa) matches to human microbiome proteins (Tables S4 A and S5 A). These data suggest that microbial proteins could have contributed to T cell priming. Our list of potential cross-reactive epitopes (Tables S4 and S5) is unlikely to be exhaustive because the degree of sequence identity required to activate cross-reactive T cell responses is unpredictable, and highly divergent epitopes can elicit cross-reactive responses in mice (Birnbaum et al., 2014). Moreover, database searches are limited to sequenced organisms and gene prediction algorithms which may miss cryptic epitopes that could be processed and presented during the course of infection (Ho and Green, 2006).

Known immunoprevalent epitopes are strongly represented in the preexposure repertoire

We next compared the mapped epitopes identified in the preexisting repertoire against HIV-1 epitope data stored in the Los Alamos National Laboratory (LANL) database (LANL-Immunology-Database, http://www.hiv.lanl.gov/content/immunology). We found that 70% of epitopes detected had previously been reported in natural HIV-1 infection (Fig. 4 A and Table S3). Many of these epitopes are presented by multiple Class II HLA types, making them immunoprevalent across the population (Table S3). Indeed, 10% of the mapped epitopes detected within this study were recognized by more than one donor (Fig. 4 and Table S2), independent of HLA type (Table S1). These data imply that T cell precursor frequency and specificity in both the naive and memory subsets before infection could play a large part in determining what is immunodominant after infection.

Analysis of the mapped HIV-1 epitopes according to CD4+ T cell subset (Fig. 4, B and C) showed that independent of donor HLA or antigen sensitivity, 88% of the epitopes recognized by preexisting naive CD4+ T cells had previously been detected in natural infection, whereas a significantly lower proportion (56%) was found for epitopes recognized by memory CD4+ T cells (p = 0.0028). HIV-1 preferentially infects HIV-1–specific memory CD4+ T cells (Douek et al., 2002), and central memory CD4+ T cells in particular are rapidly depleted during acute infection leaving very few during chronic infection (Younes et al., 2003). Thus, preexisting memory HIV-1–specific T cells could be depleted during the acute stages of infection, allowing naive T cells to expand preferentially. Alternatively, the higher proportion of novel epitopes (42%) identified in the preexisting memory CD4+ T cell repertoire may help to promote diversity of the T cell response in natural infection and could be of clinical benefit to host (Rosenberg et al., 1997).

Detection of preexisting, cross-reactive memory CD4+ T cells in unexposed donors is not restricted to HIV-1

Finally, we asked whether detection of cross-reactive memory CD4+ T cells in unexposed donors is unique to HIV-1. We screened five of the same healthy donors for proliferative responses to peptides spanning the entire envelope protein of the Zaire reference strain of Ebola virus. Because Ebola infection is associated with an extremely high mortality rate (WHO, 1978), our donors cannot have been previously exposed. In addition to naive Ebola Env-specific cells (mean 25.26 specific cells/million; Fig. S1 A), Ebola-specific memory CD4+ T cells were detected in 4/5 donors tested (mean 142 specific cells/million; Fig. S1, B and C). The frequencies of Ebola Env-specific T cells were modestly higher than those observed for HIV-1 (Fig. S1 D) but still 10-fold lower in the memory subsets than TT-specific memory cells (Fig. S1 D and Fig. 3 B). These observations demonstrate that the findings on preexposure CD4+ repertoires reported herein are not limited to HIV-1.

Conclusion

The present dataset provides the first comprehensive, systematic, and quantitative analysis of the preexposure HIV-1–specific CD4+ T cell repertoire in HLA diverse seronegative donors. Using peptides spanning the entire HIV-1 proteome, we show that both specificity and avidity of the preexposure HIV-1–specific CD4+ T cell repertoire has considerable overlap with those of CD4+ T cells detected after natural HIV-1 infection. Furthermore, we suggest that some preexisting memory HIV-1–specific T cells may have been primed by microbial organisms present within the human microbiome. These data help explain immunodominance and immunoprevalence in natural HIV-1 infection and the variability in human immune responses to infection and vaccines.

MATERIALS AND METHODS

Study participants and approval. Leukapheresis samples were obtained from a total of 15 anonymous HIV-1 seronegative, healthy individuals recruited by the Basel Swiss Red Cross Blood Centre and National Blood Service (Bristol, UK). Informed, written consent was obtained from all donors and human primary cell protocols were approved by the Federal Office of Public Health (N. A000197/2 to F. Sallusto).

HLA typing. DNA was extracted using 5 PRIME Achieve Pure DNA kit (Prima Scientific) as per manufacturer’s recommendations. HLA typing (Weatherall Institute of Molecular Medicine, Oxford, England, UK) was performed using the sequence-specific primer method adapted from Bunce (2003), which uses allele–specific primer combination in PCR amplification to provide absolute HLA resolution to two digits and high probability resolution to four digits. HLA types for all donors are shown in Table S1.

Antigen preparation. Synthetic peptides were synthesized by Sigma-Aldrich, and/or the Medical Research Council Human Immunology Unit, WIMM (Oxford, UK), at 18mers overlapping by 10 aa. The Zaire Ebola Reference Strain (Zaire-strain Mayinga-76; FASTA ID VGP_EBOZM, UNIPROT Q05320; UniProt Consortium, 2012) was used to design peptides
Tetramer enrichment protocol. Untouched CD4+ T cells were isolated from PBMCs using magnetic microbeads (Miltenyi Biotec). Tetramer staining and enrichment was performed as described previously (Su et al., 2013). In brief, cells were incubated for 30 min with live/dead Aqua marker (Invitrogen), washed, and then labeled with either gE or IE63 tetramers at room temperature for 45 min (14 µg/ml). Surface markers AF700-labeled anti-CD3 (UCHT3; BD), FITC-labeled anti-CD4 (SK3; BD), Pacific blue-labeled anti-CD56 (B159; BD), anti-CD14 (M5E2; BD), and anti-CD8 (SK1; BD) were incubated at room temperature for 15 min. Before tetramer enrichment, 1/10th staining volume was removed and added to TruCount tubes (BD) to give an absolute count of the starting number of CD4+ naive and memory T cells. The remaining staining volume was enriched for tetramer-positive cells using anti-PE-microbeads (Miltenyi Biotec) and added to a separate TruCount tube. Samples were acquired using an LSR Fortessa (BD), and the frequency of tetramer-positive cells determined by dividing the absolute counts of tetramer positive cells by the starting number of CD4+ naive/memory T cells.

Cell purification and sorting for T cell library. CD14+ monocytes and CD4+ T cells were isolated from PBMCs by positive selection with anti-body-coated microbeads (Miltenyi Biotec). CD14+ monocytes were immediately cryopreserved and stored in liquid nitrogen until required for use as antigen-presenting cells in subsequent stimulation assays. CD4+ T cell subsets were cell sorted to 99% purity on a FACSAria (BD) after staining with FITC-labeled anti-CD45RA (ALB11; Beckman Coulter), allophycocyanin-labeled anti-CD4 (SK3; BD), and anti-CCR7 (150503; R&D Systems), followed by staining with biotinylated anti-IgG2a (SouthernBiotech) and streptavidin–Pacific blue (Invitrogen). PE-cyanine 7 (PECy7)–labeled anti-CD56 (B159; BD), anti-CD14 (M5E2; BD), and anti-CD8 (SK1; BD) were incubated at room temperature for 15 min. Before tetramer enrichment, 1/10th staining volume was removed and added to TruCount tubes (BD) to give an absolute count of the starting number of CD4+ naive and memory T cells. The remaining staining volume was enriched for tetramer-positive cells using anti-PE-microbeads (Miltenyi Biotec) and added to a separate TruCount tube. Samples were acquired using an LSR Fortessa (BD), and the frequency of tetramer-positive cells determined by dividing the absolute counts of tetramer positive cells by the starting number of CD4+ naive/memory T cells.

T cell libraries. The medium used throughout was RPMI 1640 supplemented with 2 mM glutamine, 1% (vol/vol) nonessential amino acids, 1% (vol/vol) sodium pyruvate, 50 U/mL penicillin, 50 µg/mL streptomycin, and 5% human serum (Swiss Red Cross). Cell-sorted naive (CD45RA+CCR7+), central memory (CD45RA+CCR7+), and effector memory (CD45RA-CCR7+) CD4+ T cell populations were seeded at 2,000 (naive) and 1,000 (memory) cells. Cell lines were polyclonally expanded in the presence of 2.5 × 105 irradiated (45 Gy) allogeneic feeder cells, 500 IU/mL IL-2, and 1 µg/mL PHA (Thermo Fisher Scientific). Throughout the culture period, cells were supplemented with exogenous complete media containing 500 IU/mL IL-2 and progressively transferred from 96- to 24-well tissue culture plates. The full protocol has previously been published by Geiger et al. (2009).

Stimulation assays. After a 16–20-d culture period, T cells lines were counted and 2.5 × 10^6 T cells/line removed. T cells were washed 4× in 180 µL PBS and rested for a minimum of 4 h at 37°C 5% CO2. Autologous CD14+ monocytes (2.5 × 10^9/well) were pulsed for 2 h with appropriate peptide pools, or control antigen before co-culture with T cells. Proliferation was measured on day 4 after 16 h incubation with 1 µCi/ml [3H] Thymidine (GE Healthcare). A stringent positivity criteria, defined as a stimulation index of >5 and a delta value (cpm in response to antigen-pulsed monocytes – cpm in response to unpulsed monocytes) of >3 × 10^3 cpm was adopted, based upon observations made across multiple negative and positive samples assessed by T cell library technique and represented the 99th percentile of delta cpm obtained from unstimulated samples (Geiger et al., 2009). Overall, <3% of lines were excluded because of nonspecific proliferation. Positive cultures identified in the first screening assay were subsequently reanalyzed using a three dimensional matrix mapping approach (Roederer and Koup, 2003) to identify epitope specificity. Precursor frequencies were calculated based on the number of negative wells according to the Poisson distribution and expressed per million cells (Lefkovits and Waldmann, 1979). EC50 values were determined from interpolated dose–response curves using Prism (version 5.00 for Windows; GraphPad Software).

Epitope analysis. All mapped epitopes were screened against the LANL HIV-1 Molecular Immunology database (LANL-Immunology-Database) to determine whether they had previously been reported in natural HIV-1. HIV-1–specific epitopes mapped in the memory subsets were screened using the National Centre for Biotechnology and Information (NCBI) basic local alignment search tool (BLAST) search tool (Abecul et al., 1998) for epitope-length sequence matches to human pro tease and microbiome sequences. Short-sequence optimizations were used for BLAST, as the goal was detection of sequence similarity rather than of bona fide homology. Additional human proteome sequence data were downloaded from UniProt (UniProt Consortium, 2012), whereas further human microbiome sequences ( shotgun sequences derived from body location samples) were obtained from the Human Microbiome Project (HMP) clustered gene indices catalog (http://www.hmpdacc.org/HMGC/). The UniProt and HMP sequences were processed with UBLAST (Edgar, 2010) using highly non-stringent criteria [id, 0.01; evalue, 10,000; maxexacts, 500,000] to generate matches. Both NCBI-BLAST and UBLAST matches were then filtered on the number of amino acid identities and similarities (Tables S4 and S5). For NCBI, BLAST–included epitopes showed a minimum of 8 out of 9 matched amino acids with the highest sequence identities showing 12 of 12 matched amino acids. Sequences identified with UBLAST search algorithms had a minimum of 8 of 8 matched amino acids with the highest levels of sequence identity observed with 12 of 12 amino acids matched.

Statistics. Epitope maps obtained from the LANL immunology database show 60% of the HIV-1 proteome is covered by previously defined CD4+ T cell epitopes (LANL–Immunology-Database). Because of this greater proportion of sequence with previously known epitopes, we normalized the number of amino acids covered by epitopes identified in this study to the number of amino acids from the HBX2 reference strain of HIV-1 which contain known CD4+ T cell epitopes (1,875 aa) or which contained no epitopes (1,272 aa). Using a Fisher’s exact test, the percentage of HIV-1 proteome covered by previously reported epitopes from the naive and memory CD4+ T cell subsets could be compared. Significance was defined as P ≤ 0.005.

Pearson’s correlation was used to compare the precursor frequencies obtained using the T cell library technique and the tetramer enrichment protocol. Because the dataset included zero values, a log (x + 1) transformation was applied to all data points.

Online supplemental material. Fig. S1 shows data from the naive, central and effector memory subsets of five healthy leukapheresis donors who were screened for proliferative responses to a pool of peptides spanning the envelope protein of the Zaire reference strain of Ebola virus. Table S1 shows the HLA class I and II typing for all 10 donors studied within the context of this manuscript. Table S2 shows the amino acid sequence and delta cpm values for the 68 HIV-1–specific T cell responses detected in the naive, central memory, and effector memory CD4+ T cell subsets of five healthy, HIV-1
seronegative donors. Table S3 shows the results of a Los Alamos database search to determine whether any of the mapped epitopes detected in the preserocephyl repertoires had previously been reported in natural infection, and lists the citations for instances where the epitope had previously been reported. Table S4 displays the sequence identity matches of HIV-1 epitopes in the memory CD4+ T cell repertoire to the human microbiome, human sequences, and all GenBank/EMBL sequences excepting those classified as lentiviral, synthetic, or other. Table SS shows the results of screening mapped HIV-1-specific epitopes in the memory CD4+ T cell subsets of HIV-1 unexposed uninfected donors against predicted genes from the HMP, shotgun sequences and human sequences from Uniprot. Online supplemental material is available at http://www.jem.org/cgi/content/full/jem.20130555/DC1.

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