Inhibition of Two Temporal Phases of HIV-1 Transfer from Primary Langerhans Cells to T Cells: The Role of Langerin

Najla Nasr, Joey Lai, Rachel A. Botting, Sarah K. Mercier, Andrew N. Harman, Min Kim, Stuart Turville, Rob J. Center, Teresa Domagala, Paul R. Gorry, Norman Olbourne and Anthony L. Cunningham

J Immunol 2014; 193:2554-2564; Prepublished online 28 July 2014;
doi: 10.4049/jimmunol.1400630
http://www.jimmunol.org/content/193/5/2554

Supplementary Material
http://www.jimmunol.org/content/suppl/2014/07/26/jimmunol.1400630.DCSupplemental

References
This article cites 60 articles, 30 of which you can access for free at: http://www.jimmunol.org/content/193/5/2554.full#ref-list-1

Why The JI? Submit online.
• Rapid Reviews! 30 days* from submission to initial decision
• No Triage! Every submission reviewed by practicing scientists
• Fast Publication! 4 weeks from acceptance to publication

Subscription
Information about subscribing to The Journal of Immunology is online at: http://jimmunol.org/subscription

Permissions
Submit copyright permission requests at: http://www.aai.org/About/Publications/JI/copyright.html

Email Alerts
Receive free email-alerts when new articles cite this article. Sign up at: http://jimmunol.org/alerts
Inhibition of Two Temporal Phases of HIV-1 Transfer from Primary Langerhans Cells to T Cells: The Role of Langerin

Najla Nasr,*† Joey Lai,*† Rachel A. Botting,*† Sarah K. Mercier,*† Andrew N. Harman,*† Min Kim,*† Stuart Turville,*† Rob J. Center,‡ Teresa Domagala,§ Paul R. Gorry,§ Norman Oborne,‖ and Anthony L. Cunningham*†

Epidermal Langerhans cells (eLCs) uniquely express the C-type lectin receptor langerin in addition to the HIV entry receptors CD4 and CCR5. They are among the first target cells to encounter HIV in the anogenital stratified squamous mucosa during sexual transmission. Previous reports on the mechanism of HIV transfer to T cells and the role of langerin have been contradictory. In this study, we examined HIV replication and langerin-mediated viral transfer by authentic immature eLCs and model Mutz-3 LCs. eLCs were productively infected with HIV, whereas Mutz-3 LCs were not susceptible because of a lack of CCR5 expression. Two successive phases of HIV viral transfer to T cells via cave/vesicular trafficking and de novo replication were observed with eLCs as previously described in monocyte-derived or blood dendritic cells, but only first phase transfer was observed with Mutz-3 LCs. Langerin was expressed as trimers after cross-linking on the cell surface of Mutz-3 LCs and in this form preferentially bound HIV envelope protein gp140 and whole HIV particles via the carbohydrate recognition domain (CRD). Both phases of HIV transfer from eLCs to T cells were inhibited when eLCs were pretreated with a mAb to langerin CRD or when HIV was pretreated with a soluble langerin trimeric extracellular domain or by a CRD homolog. However, the langerin homolog did not inhibit direct HIV infection of T cells. These two novel soluble langerin inhibitors could be developed to prevent HIV uptake, infection, and subsequent transfer to T cells during early stages of infection. The Journal of Immunology, 2014, 193: 2554–2564.

H

Received for publication March 10, 2014. Accepted for publication June 23, 2014.

The work was supported by Project Grants APP1028014 and 1050909 from the Australian National Health and Medical Research Council and by a grant from the Australian Centre for HIV and Hepatitis Virology.

The protein sequence data presented in this article have been submitted to the SwissProt database (http://www.uniprot.org) under accession number Q9UJ71.

Address correspondence and reprint requests to Prof. Anthony L. Cunningham at the current address: Centre for Virus Research, Westmead Millennium Institute, 176 Hawkesbury Road, PO Box 412, Westmead, NSW 2145, Australia; E-mail address: tony.cunningham@sydney.edu.au

The online version of this article contains supplemental material.

Abbreviations used in this article: CLR, C-type lectin receptor; CRD, carbohydrate recognition domain; DC, dendritic cell; DC-SIGN, DC-specific ICAM-3-grabbing nonintegrin; DSS, disuccinimidyl suberate; ECD, extracellular domain; eLC, epidermal Langerhans cell; hpi, hour postinfection; MDDC, monocyte-derived DC; MOI, multiplicity of infection; MR, mannose receptor; VCC, virus-containing compartment; VSV-G, G protein vesicular stomatitis virus.

Copyright © 2014 by The American Association of Immunologists, Inc. 0022-1767/14/$16.00 8. In the SIV–macaque model, foci of infected T cells underlying intact mucosa of the vagina and ectocervix have been observed by 48 h postinfection (hpi) (9), and some evidence for LC uptake also has been shown (10–12). LCs can either encounter HIV as the virus diffuses between the tight junctions of keratinocytes in an intact mucosa (4) or where there is a breach in the mucosa caused by genital ulcerative diseases such as HSV or epithelial abrasions that occur during intercourse. In foreskin/skin tissue explants (13) or in emigrant LCs infected in situ (14–16), epidermal LCs (eLCs) were permissive to de novo infection by CCR5 using HIV-1 viral strains (R5 HIV-1) at >24–72 hpi.

LCs characteristically express CD1a (17) and the HIV entry receptors CD4 and CCR5 (18) in their immature state, rendering them susceptible to infection primarily by R5 HIV-1 (2, 4, 12). They also uniquely express the HIV binding C-type lectin receptor (CLR) langerin (19) but not DC-specific ICAM-3-grabbing non-integrin (DC-SIGN) or mannose receptor (MR). Langerin has an extracellular domain consisting of a single carbohydrate recognition domain (CRD), a neck domain involved in oligomerization, a short transmembrane domain, and a cytoplasmic tail, which mediates endocytosis, vesicular trafficking, and cytoskeletal movement (20–22). After pathogen or Ag acquisition, LCs usually undergo a process of maturation characterized by upregulation of molecules associated with T cell activation (CD80, CD86, CD83, and CD40), Ag presentation (MHC class I and II) and migration (CCR7), and downregulation of molecules associated with Ag uptake (langerin) and epithelial adhesion (E-cadherin) (23, 24).

Monocyte-derived DCs (MDDCs) are usually used as a cell model to study the interaction of HIV with DCs because tissue DCs are difficult to isolate, require enzymatic digestion resulting in their maturation, and often yield low numbers. Unlike LCs, MDDCs express the CLR s DC-SIGN and MR at high levels. Viral transfer from MDDCs (25, 26) and primary blood myeloid DCs (27) to CD4+ T cells has been reported to occur in two successive phases,
corresponding to the two main routes of HIV entry into DCs. After binding to the concentrating CLR, the majority of HIV is either taken up into a neutral pH tetraspan CD81 compartment termed a virus-containing compartment (VCC) or into the endolysosomal system (30). The minority (<10%) is transferred from CLR to the HIV entry receptors CD4 and CCR5 (31). Virions that enter the endosomal system are degraded by acid–proteolytic digestion within 6–24 h. If the DC contacts CD4+ T cells in the first few hours after encountering the virus (probably in the mucosa), HIV residing in the VCC can then be transferred to T cells across an “infectious synapse” representing what is termed “first-phase” transfer (32). HIV entering via CD4/CCR5 undergoes de novo replication within 48 h (26), and the virus is then transferred to T cells across a viral synapse (33). This “second phase” of viral transfer occurs at a later stage than the viral transfer from VCCs and increases with time.

Although DC-mediated infection and viral transfer to T cells is well documented using MDDCs and blood myeloid DCs, whether two temporal phases of HIV transfer from eLCs to T cells occurs is still controversial (4, 8, 14, 34–36). Specifically, differences in cell models used (35), cell isolation methods (36), potencies of anti-langerin inhibitory Abs, and titers of HIV inocula used (36) have led to contradictory reports on the relative roles of langerin and CD4/CCR5 in HIV binding, entry, and replication in LCs.

In this study, we have examined whether primary human immature eLCs and a model LC derived from the Mutz-3 cell line (Mutz-3 LCs) can transfer HIV to CD4+ T cells in two successive phases, corresponding to the two temporal phases of HIV trafficking in MDDCs and myeloid DCs. Because oligomerization of CLR such as DC-SIGN (37–39) and MR (40, 41) has been reported to be essential for the binding of oligosaccharide ligands, we showed that native trimeric HIV envelope protein (HIVAD8 gp140) and HIVav particles bound strongly to langerin trimers on Mutz-3 LCs, suggesting that soluble langerin homologs may need to be trimeric for maximal inhibition. Thus, we examined inhibition of HIV binding to langerin on eLCs and subsequent first- and second-phase viral transfer to T cells with a langerin-specific mAb and by langerin homologs consisting of soluble trimeric full-length extracellular domain (ECD) or the monomeric truncated carbohydrate recognition domain (CRD). These inhibitors could provide a useful strategy for blocking entry of HIV into anogenital LCs during sexual transmission, particularly if combined with others such as CCR5 inhibitors, perhaps as a novel microbicide.

Materials and Methods

Ethics statement

Written and informed consent approved by the Western Sydney Local Health District human research ethics committee (Westmead Hospital, Sydney, NSW, Australia) was obtained from all patients undergoing breast reduction or abdominoplasty.

Isolation of immature skin LCs

Skin was collected immediately after surgery and was cleared of s.c. fat using a dermatome. Skin nets were generated using a skin graft mesh (Zimmer, Warsaw, IN) and were incubated at 4 °C in RPMI 1640 medium supplemented with 25 μg/ml gentamicin (Sigma-Aldrich) and 2.5 mg/ml disperse II (Roche Biochemicals). After overnight incubation, the epidermis and dermis were separated using forceps. Epidermal sheets were washed twice in PBS and incubated for 60 min at 37 °C in RPMI 1640 medium containing collagenase blend F (2.5 mg/ml) and DNase (50 μg/ml). Alternatively, trypsin (0.06%) was used for epidermal dissociation. Viability of eLCs was >90% in both collagenase and trypsin preparations. Isolated eLCs were stained for CD1a, CD45, CD3, and CD14. eLCs were sorted by gating on the CD3+CD4+CD1a+CD45− population (1–3 × 106 LC) on a FACSVantage fluorescence-activated cell sorter to a purity of >97%. eLCs were cultured in RPMI 1640 medium supplemented with 10% heat-inactivated human AB serum (RH10; Invitrogen) and 200 ng/ml GM-CSF (Invitrogen).

Generation of Mutz-3 LCs, MDDCs, and isolation of CD4+ T cells

Human CD34+ acute myeloid leukemia Mutz-3 cells provided by S.I.A.M. Santegoets (VU University Medical Center, Amsterdam, the Netherlands) were cultured and treated as previously reported by Santegoets et al. (42) to generate Mutz-3 LCs. MDDCs were generated from CD14+ monocytes using CD14 magnetic beads (Miltenyi Biotec) as described previously (26). CD4+ T cells were isolated from PBMCs using CD4 magnetic beads (Miltenyi Biotec). They were then activated with PHA (5 μg/ml; Sigma-Aldrich) for 48 h, washed, and cultured in 20 U/ml IL-2 (Roche Molecular Biochemicals, Indianapolis, IN).

Transfer of HIVav from MDDCs, eLCs, and Mutz-3 LCs to T cells and inhibition assays

A total of 2.5–5 × 106 MDDCs, Mutz-3 LCs, or eLCs were mock treated or inoculated with either the laboratory adapted R5 strain using HIVav, the laboratory adapted four times using strain NL43 or the primary strain NB2 (43) at multiplicities of infection (MOIs) ranging from 0.005 to 10 for 2 h at 37 °C to assess their infectivity and whether viral transfer to T cells is M01 dependent. They were then washed three times in PBS and plated in duplicate in 96-well plates. For the viral transfer inhibition assays, Mutz-3 LCs and eLCs were treated with either neutralizing mAb to langerin (HIV-anti-lang; 1.4 μM) purchased from Dentridics (clone 817G7) or the reverse transcriptase inhibitor AZT, obtained from National Institutes of Health AIDS Reagent Program, Division of AIDS, National Institute of Allergy and Infectious Diseases, National Institutes of Health (HIV+AZT; 50 μM), for 30 min at 4 °C or 37 °C, respectively, before HIVav inoculation (MOI of 0.5) for 2 h at 37 °C. In other settings, HIVav, pre-treated with soluble langerin ECD (aa 70–328) or CRD (aa 200–328) generated in a mammalian Chinese hamster ovary expression system (Apollo Life Sciences Pty, Sydney, NSW, Australia) for 1 h at 37 °C before addition to the cells at the final concentrations of 0.1 μM HIV+ECOD or HIV+CRD. Infected cells were washed three times in PBS and 1 × 106 PHA-activated CD4+ T cells or the CD4+ T cell line JLTR were added to mock, HIV-ECD, HIV-CRD, mAb to langerin, AZT, or HIV+C-RD-treated cells at the indicated time points. Half the medium was replenished every 48 h. HIV infection was determined by either staining with an intracellular p24 MAb (KC57-RD1; Beckman Coulter, Fullerton, CA) or by measuring the GFP fluorescence intensity of JLTR cells. Samples were run on a FACScan™-II flow cytometer (BD Biosciences) and FlowJo (Tree Star, Ashland, OR) was used for data analysis.

Measurement of gene expression by quantitative PCR

Total unamplified RNA was DNase I treated (Promega, Madison, WI) and reverse transcribed using oligo(dT) and superscript III (Invitrogen). The cDNA was subject to QPCR using CCR5 primers (Sigma-Aldrich), accession number NM-000579 and SYBR Green (Invitrogen) as described previously (24).

Affinity purification of langerin complexes and protein identification using mass spectrometry

Chemical cross-linking of cell surface molecules using disuccimimidyl suberate (DSS) (Pierce, Rockford, IL) has been described previously (37). Affinity beads were generated by coupling 100 μg IgG1 or anti-langerin (R&D Systems) to cyanogen bromide–activated Sepharose (Sigma-Aldrich), according to the manufacturer’s instructions. Non-specifically bound proteins were removed by incubating the lysate from 2 × 107 DSS-transferred Mutz-3 LCs with the IgG Sepharose beads for 2 h at 4°C. The lysate was then incubated with 200 μl anti-langerin Sepharose beads, and the beads were washed with 10 mM TBS containing 0.1% Triton X-100 (pH 8). Langerin complexes were eluted from the beads by boiling for 5 min in SDS buffer. Affinity purified proteins were identified by electrospray ionization mass spectrometry as described previously (40).

HIVAD8 gp140 and whole HIVav binding to Mutz-3 LCs and inhibition assay

For HIVav gp140 binding assays, mock or cross-linked cells were resuspended in binding buffer (RPMI 1640 medium, 1% BSA, and 10 mM HEPES [pH 7.5]) at 1 × 106 cells/50 μl. Three micrograms of biotinylated trimeric HIVAD8 gp140 was added to cells for 30 min at 4 °C as described previously (40). To inhibit gp140 binding, 5 mg/ml mannan, 10 mM EGTA,
and 1.4 μM anti-langerin (122D5/DCGM4) were added to the cells for 30 min at 4°C prior to gp140 addition. Cells were then washed three times with PBS, and bound ligands were incubated by labeling the cells with 1 μg/ml streptavidin-PE for 30 min for detection by flow cytometry. For HIV$_{nat}$ binding assays, 3 x 10$^6$ mock-treated and cross-linked Mutz-3 LCs were incubated with HIV$_{nat}$, ranging from 0.27 to 4.25 μg/ml p24 Ag for 2 h at 4°C. Cells were washed three times with PBS and then lysed in lysis buffer (10 mM HEPES, 140 mM NaCl, 1% Triton X-100, and 10 mM DTT with protease inhibitors). Total protein content was measured using a DC Protein Assay Kit (Bio-Rad, Hercules, CA), and p24 Ag was quantified by ELISA (Beckman Coulter, Brea, CA).

**Statistical analysis**

Differences between treatments were analyzed by Student t test and modified for unequal variance. A p value < 0.05 was considered statistically significant.

**Results**

**HIV$_{nat}$ infection of eLCs and two phase transfer of HIV$_{nat}$ to T cells**

Ex vivo immature eLCs were isolated from epidermis by enzymatic collagenase digestion and flow sorting to collect the CD3$^-$ CD14$^-$ CD45$^-$ population (Supplemental Fig. 1A). Because LCs are the only epidermal cells that coexpressed CD1a and langerin (Supplemental Fig. 1B), CD1a was used as a marker to select for eLCs, leaving the langerin epitope unbound and thus available to bind HIV immediately after sorting while eLCs are still in an immature state. To determine whether immature isolated eLCs can be infected with HIV, eLCs from five donors were inoculated with HIV$_{nat}$ at different MOIs for 24 and 72 h. The proportion of HIV p24 Ag-positive (Ag$^+$) eLCs was 3.6 ± 1.2%, 2 ± 0.6%, 0.8 ± 0.5%, and 0.3 ± 0.2% at MOI of 10, 5, 1, and 0.1, respectively, at 72 hpi (Fig. 1A, 1B). No intracellular p24 Ag$^+$ cells were detected at 24 hpi. Extracellular p24 Ag was detected by ELISA in supernatants of infected cells at 72 but not at 24 hpi. In addition, no intracellular p24 Ag$^+$ cells or extracellular virus were detected at 72 hpi when eLCs were pretreated with the antiretroviral reverse transcriptase inhibitor, zidovudine (AZT), for 1 h before HIV inoculation (Fig. 1B).

We next assessed the ability of eLCs to transfer virus to the JLTR CD4 T cell line. JLTR cells were used as a target for viral transfer from eLCs because they encode the GFP flanked by the HIV-1 promoter, and therefore, they fluoresce green when infected with HIV. eLCs were infected with HIV$_{nat}$ (MOI of 1 and 0.1) for 2, 24, 48, and 96 hpi and then cocultured with JLTR cells for 4 d. The amount of transferred virus declined progressively between 2 and 48 hpi as shown by the reduction in the number of GFP$^+$ JLTR cells. This was followed by an increase in GFP$^+$ JLTR cells when trypsin at either 2 or 96 hpi. Viral transfer was observed when JLTRs were added at 2 hpi from both trypsin- and collagenase- isolated eLCs as shown by the number of GFP$^+$ JLTR cells. However, at 96 hpi, GFP$^+$ cells were detected in collagenase- but not trypsin-isolated eLCs after 4 d of coculture (Fig. 1G). Also, there was a progressive increase in HIV spread in JLTR cells cocultured in collagenase-treated eLCs up to 8 d but still no detectable GFP$^+$ cells in those cocultured with trypsin-treated eLCs (data not shown).

To exclude the fact that our findings were not influenced by fluctuation in viability and proliferation of JLTRs used as T cells in the viral transfer assay and which might affect viral replication, we assessed their proliferation using CFSE staining and their viability using DAPI at different time points after passage. There was no to little fluctuation in viability (90 ± 7%) over 6 d postpassage (Supplemental Fig. 2A). Similarly, over this 6-d culture period, the cells continued to proliferate uniformly as shown by the gradual decrease in the intensity of CFSE stain (Supplemental Fig. 2B). Furthermore, the HIV$_{nat}$ infection of JLTRs showed a uniform increase in the proportion of GFP$^+$ cells over time (Supplemental Fig. 2C).

**HIV$_{nat}$-infected Mutz-3 LCs only mediate first-phase viral transfer**

Model Mutz-3 LCs were produced after 10 d of cytokine treatment of the parental Mutz-3 cells as reported previously (42). Mutz-3 LCs expressed CD1a, langerin, and CD4 but not DC-SIGN, MR, and CCR5 (Fig. 2A). Thus, they resembled authentic eLCs apart from the undetectable cell surface CCR5 expression. This lack of CCR5 surface expression on Mutz-3 LCs was investigated and compared with precursor Mutz-3 cells and also eLCs. Prior to cytokine treatment, Mutz-3 expressed low levels of cell surface CCR5 (7 ± 1.5%) and moderate levels intracellularly (20 ± 2.7%) (Fig. 2B). After cytokine treatment to convert them into Mutz-3 LCs, CCR5 was detected at high levels intracellularly (69 ± 5%) but was not expressed on the cell surface. In contrast, eLCs from four donors variably expressed cell surface CCR5 (12 ± 6.3%) (two donors shown in Fig. 2B). In addition, CCR5 mRNA levels were slightly higher in Mutz-3 LCs than Mutz-3 cells, but both were markedly and significantly lower when compared with eLCs from three different donors (Fig. 2C). The lack of CCR5 expression by Mutz-3 LCs is partly due to their maturation state as measured by the proportion of cells expressing costimulatory molecules CD80 (80 ± 4.3%), 86 (30 ± 3.2%), and especially CD83 (25 ± 2.7%) (Fig. 2D, panel 1). In addition, Mutz-3 LCs could be further matured by the addition of maturation mix for 24–48 h as the expression of CD83 increased from 25 to 58%, whereas CD86 increased from 30 to 90% (Fig. 2D, panel 2). However, when freshly isolated eLCs (Fig. 2D, panel 3, CD83 7.3%) matured overnight (Fig. 2D, panel 4, CD83 64%), leading to a greater proportion of CD83 expression than Mutz-3 LCs (25%), CCR5 was still detectable on mature eLCs, indicating that the magnitude of CD83 expression is not an accurate predictor of CCR5 expression between cells. Thus, maturation of eLCs and Mutz-3 LCs downregulates CCR5, but the effect is greater on Mutz-3 LCs probably because of the lower expression levels on the parental Mutz-3 cell line.

When Mutz-3 LCs were inoculated with HIV$_{nat}$ at different MOIs (10, 4, 1, 0.5, and 0.005), they did not produce virus de novo as demonstrated by the fact that no intracellular p24 Ag was detected when compared with eLCs up to 96 hpi (Fig. 2E). In this LC model, HIV entry via fusion that usually leads to productive infection was impaired because Mutz-3 LCs did not express CCR5. However, Mutz-3 LCs infected at an MOI of 1 with the
G protein vesicular stomatitis virus (VSV-G) pseudotyped HIV$_{\text{Bal}}$, which enters cells independently of CD4/CCR5 for fusion and entry, showed a significant population of p24+ cells (24% ± 5%) by 96 hpi (Fig. 2F). Overall, these results show that Mutz-3 LCs express langerin but not CCR5 and therefore could not be infected de novo with HIV$_{\text{Bal}}$ because of failure of entry.

In the absence of productive infection, the kinetics of HIV$_{\text{Bal}}$ transfer from Mutz-3 LCs to JLTR cells was assessed to investigate their suitability as a model to study eLC-mediated first-phase viral transfer to T cells. Mutz-3 LCs were exposed to HIV$_{\text{Bal}}$ at an MOI of 1 before JLTR cells were added to the culture at serial time points up to day 4. As shown in Fig. 3A, first-phase transfer was observed from Mutz-3 LCs to JLTR cells between 0 and 48 hpi, but the rate of viral transfer gradually decreased over time, probably as a result of endolysosomal degradation. In addition, when Mutz-3 LCs were inoculated with HIV$_{\text{Bal}}$ at different MOIs (0.5 ± 0.05), the amount of transferred virus correlated with the MOI used (Fig. 3B). Varying the MOI did not affect transfer kinetics, because a relative decline in the amount of virus transferred over the 96-h period was observed, irrespective of the MOI used. Furthermore, treatment of Mutz-3 LCs with AZT prior to HIV exposure did not affect their ability to transfer virus to JLTR cells (Fig. 3C), demonstrating that productive infection of these cells was not required for first-phase transfer.
was higher than those infected with cell-free supernatants at a similar MOI (Fig. 3D), indicating that uninfected HIV-exposed Mutz-3 LCs are effective vehicles for HIV transfer to primary CD4+ T cells. In summary, this indicates that the degree of virus transfer from Mutz-3 LCs was MOI dependent and that they were only able to engage in first-phase vesicular trafficking and transfer.
A

FIGURE 3. HIV<sub>BaL</sub> transfer from Mutz-3 LCs to T cells. (A) JLTR cells were added to Mutz-3 LCs at 2, 4, 6, 10, 20, 24, and 48 hpi, and the percentage of GFP<sup>+</sup> JLTR cells was determined 96 h postcoculture by flow cytometry. Data are representative of three experiments. The percentage of infected JLTRs is shown in the top right of each histogram. (B) Mutz-3 LCs were mock treated, infected with HIV<sub>BaL</sub>, or HIV<sub>BaL</sub> and pretreated with AZT (50 µM) for 30 min at 37°C before HIV inoculation (MOI of 0.1) for 2 h. JLTR cells were added 2 hpi to assess first-phase transfer by flow cytometry. A representative of three separate experiments is shown. (C) Mutz-3 LCs were mock treated, infected with HIV<sub>BaL</sub>, or HIV<sub>BaL</sub> and pretreated with AZT (50 µM) for 30 min at 37°C before HIV inoculation (MOI of 0.1) for 2 h. JLTR cells were added 2 hpi to assess first-phase transfer by flow cytometry. A representative of three separate experiments is shown. (D) CD4<sup>+</sup> T cells were either mock treated, infected with HIV<sub>BaL</sub>, or HIV<sub>BaL</sub> and pretreated with AZT (50 µM) for 30 min at 37°C before HIV inoculation (MOI of 0.1) for 2 h. JLTR cells were added 2 hpi to assess first-phase transfer by flow cytometry. A representative of three separate experiments is shown.

Kinetics of viral transfer in Mutz-3 LCs, MDDCs, and eLCs

We next compared the kinetics of HIV transfer to T cells from langerin-expressing Mutz-3 LCs and eLCs with that of MDDCs (expressing MR and DC-SIGN) to T cells. Both MDDCs and eLCs showed two phases of viral transfer to ILTR cells, whereas Mutz-3 LCs exhibited only first-phase transfer (Fig. 4). When the kinetics of viral transfer were compared between MDDCs and eLCs, we observed that the first-phase transfer by eLCs was extended up to 48 hpi compared with <24 hpi in MDDCs, indicating that eLCs had slower kinetics of first-phase transfer compared with MDDCs. Mutz-3 LCs, similar to eLCs, exhibited extended first-phase transfer only.

Langerin forms trimers on the cell surface of Mutz-3 LCs

To determine the predominant oligomerization state of langerin on the cell surface to guide the use of inhibitors, Mutz-3 LCs were cross-linked with increased concentrations of DSS. DSS maintains protein–protein interactions between langerin molecules, thus preserving its oligomeric state on the cell surface. In the absence of DSS, the oligomeric langerin dissociated and resolved to a monomeric molecular mass of 40 kDa (Fig. 5A). However, langerin resolved to a dimeric molecular mass of 80 kDa when the cells were exposed to low DSS concentrations (15–120 µM). Langerin dimers gradually increased in abundance with increasing DSS concentration, and this increase also was associated with both the appearance of a trimeric band and a concomitant decrease in the abundance of the monomeric band. Langerin trimers continued to increase in abundance and remained the predominant oligomerization species at saturating DSS concentrations (1–2 mM), whereas no monomeric langerin and little to no dimeric langerin were detected with DSS concentrations >1 mM (Fig. 5A). To investigate the possibility that proteins other than langerin may be participating in the formation of langerin complexes, matrix-assisted laser desorption/ionization time-of-flight mass spectrometry analysis was performed on peptides isolated from trypsin-digested affinity-purified monomeric and trimeric langerin from DSS<sup>+</sup> Mutz-3 LCs. Table I lists the peptides identified from gel slices corresponding to monomeric and trimeric langerin. A total of eight langerin-derived peptides were identified in the monomeric band (expectation value = 5.7 × 10<sup>−3</sup>) and six were identified in the trimeric band (expectation value = 0.023), confirming the presence of langerin in these bands. The absence of peptides from other potentially interacting proteins suggests that langerin mainly exists as homotrimers on cell membranes of Mutz-3 LCs.

Purified HIV<sub>AD8</sub> gp140 and HIV<sub>BaL</sub> bind to langerin trimers on the surface of Mutz-3 LCs via the CRDs in a Ca<sup>2+</sup>-dependent manner

To determine the effect of oligomerization of cell surface langerin on its ability to bind trimeric HIV<sub>AD8</sub> gp140, langerin expressed on the surface of Mutz-3 LCs was cross-linked with saturating concentrations of DSS prior to the addition of HIV<sub>AD8</sub> gp140 or HIV<sub>BaL</sub>. The DSS concentration that was chosen (1 mM) was based on the results shown in Fig. 5A, which showed maximal detection of langerin trimmer concomitant with minimal monomeric langerin detection in the absence of cytotoxicity. DSS<sup>+</sup> cells displayed an enhanced gp140 binding level, with a 2-fold increase in gp140 binding compared with that in DSS<sup>−</sup> cells, as detected by flow cytometry (Fig. 5B). Similarly, a 2-fold enhancement in the binding of HIV<sub>BaL</sub> viral particles was demonstrated on DSS<sup>+</sup> cells by p24 ELISA over a range of input virus (Fig. 5C). The gp140–langerin interaction was further characterized by using various inhibitors to specifically block gp140 binding to langerin on DSS<sup>−</sup> and DSS<sup>+</sup> cells. The inhibitors used included the following: 1) yeast derived mannan, which like gp140, is highly glycosylated and expresses carbohydrate structures recognized by the CRD of langerin (45); 2) EGTA, which inhibits Ca<sup>2+</sup>-dependent binding; and 3) a mAb to langerin specific for the CRD (clones 122D5/DCGM4 and 817G7). In DSS<sup>−</sup> cells, these inhibitors reduced total gp140 binding by ~50% (Fig. 5B). However, in DSS<sup>+</sup> cells, mannan, EGTA, and anti-langerin reduced gp140 binding to baseline levels (95% inhibition), suggesting that gp140 binds to langerin in both a carbohydrate and Ca<sup>2+</sup>-dependent manner.
Second-phase transfer was inhibited by 30% showed a 66% reduction in second-phase transfer from immature eLCs to JLTR cells. Higher levels of inhibition of first-phase transfer and a 79% reduction in second-phase transfer when JLTR cells were added 96 h after eLC inoculation. HIV-CRD (0.1 µM) did not affect their ability to stimulate T cell proliferation (Supplemental Fig. 3).

To determine whether soluble langerin can act as a potential inhibitor of HIV transfer, HIV-BaL that was pretreated with either ECD or CRD was then added to eLCs or Mutz-3 LCs with final concentrations of 0.1 and/or 0.4 µM. The concentrations selected were based on surface plasmon resonance binding data of gp120 to CRD and ECD (data not shown), which is similar to recently published findings (46), and took into account our cytotoxicity studies and available amounts of reagent. eLCs or Mutz-3 LCs pretreated with AZT (50 µM) before HIV inoculation were used as positive controls for HIV transfer. HIV-ECD (0.1 µM) showed inhibitory activity against both first- and second-phase transfer (Fig. 6C), with a peak reduction of 55-12% in first-phase transfer of HIV from eLCs to JLTR cells when the latter cells were added at 2 h after HIV inoculation of eLCs and a 66-12.5% reduction in second-phase transfer when JLTR cells were added 96 h after eLC inoculation. HIV-CRD (0.1 µM) also effectively inhibited both phases of viral transfer, with a 46 ± 10% reduction in first-phase transfer and a 79 ± 7% reduction in second-phase transfer of virus to JLTR cells. Higher levels of inhibition of first-phase (56 ± 6%) and second-phase (85 ± 4%) transfer were observed at higher CRD concentrations (0.4 µM; data not shown). Also, we assessed whether the CRD-specific mAb (clone 817G7), which has been shown to strongly inhibit oligomeric langerin-gp140 interactions (Fig. 4B), can inhibit viral transfer from eLCs to T cells. eLCs pretreated with the mAb to langerin (1.4 µM) showed a 66 ± 5% reduction in first-phase viral transfer (Fig. 6C). Second-phase transfer was inhibited by 30 ± 10% when the mAb was added once to eLCs 30 min before and during the 2-h infection. However, when the mAb was also maintained postinfection and replenished every 48 h, there was an 82 ± 10% reduction in HIV transfer from immature eLCs to JLTR cells but sustained presence of the mAb in the medium was required to inhibit second-phase transfer.

Furthermore, to further verify the inhibitory effect of the ECD and anti-langerin on first-phase transfer, we treated Mutz-3 LCs with HIV-BaL that was preincubated with langerin ECD (0.1 µM) and showed that transfer from Mutz-3 LCs to PHA-activated CD4+ T cells was inhibited by 65 ± 7%. In addition, when Mutz-3 LCs were pretreated with either the anti-langerin mAb or AZT, there was 55 ± 8% reduction in first-phase transfer to CD4+ T cells (Fig. 6D), whereas AZT had no effect on such transfer. As a control, we examined the effects of langerin ECD on direct HIV transfer to T cells. eLCs or Mutz-3 LCs were mock or HIVBaL treated at an MOI of 1 for 2 h at 37˚C, followed by three washes in PBS and addition of JLTR at 2, 24, 48, and 96 hpi. The mean data of GFP+ JLTR cells from three experiments is shown with SE (± SE) bars.

**FIGURE 4.** Dynamics of viral transfer from MDDCs, Mutz-3 LCs, and eLCs. MDDCs, Mutz-3 LCs, and eLCs were mock or HIVBaL treated at an MOI of 1 for 2 h at 37˚C, followed by three washes in PBS and addition of JLTR at 2, 24, 48, and 96 hpi. The mean data of GFP+ JLTR cells from three experiments is shown with SE (± SE) bars.

**FIGURE 5.** Langerin oligomerisation and binding to HIV gp140 and HIV-BaL. (A) Cell surface langerin on Mutz-3 LCs was cross-linked with increasing concentrations of DSS (15, 30, 60, 120, 1000, and 2000 µM DSS), and langerin monomer, dimer, and trimer formation were detected by Western blot analysis. A representative Western blot from three independent experiments is shown. (B) Binding of gp140 to DSS- and DSS+ Mutz-3 LCs in the absence or presence of binding inhibitors (mannan, EGTA, and mAbs to the CRD of langerin; clones 122D5/DCGM4 and 817G7) is represented as the geometric mean of streptavidin PE fluorescence bound gp140 (n = 3, **p < 0.02 for all inhibitors on DSS−, ***p < 0.001 for all three inhibitors on DSS+). (C) Mutz-3 LCs were incubated with HIV-BaL at p24 concentrations ranging between 0.27 and 4.25 µg/ml for 2 h at 4˚C. The amount of virions bound to the surface of DSS− and DSS+ Mutz-3 LCs was quantified by p24 ELISA. Values are represented as the mean captured p24 Ag from bound virions ± SE (n = 3, *p < 0.05).
infection of JLTR cells and found that there was no difference in
the proportion of infected JLTR cells whether they were inoculated
with either HIV or HIV preincubated with langerin ECD (Fig. 6E).

Discussion

In this study, authentic immature primary eLCs were infected with
CCR5 using laboratory (HIVBal) and primary (NB2) strains at
different MOIs ranging from 0.05 to 10. Two successive phases of
HIV transfer from eLCs to T cells were observed at all MOIs:
first-phase transfer mediated by the CLR langerin and a second
phase of transfer followed de novo replication mediated through
CCR5 and CD4 and facilitated by HIV binding to langerin. These
two phases of viral transfer from eLCs to T cells were similar to
those previously reported in MDDCs (25, 26) and as shown in
Fig. 4. However, LCs showed slower kinetics of first-phase trans-
fer and also a delayed onset of second-phase transfer, which was
inhibited by AZT. Thus, the processes of viral transfer in
MDDCs and eLCs differ only in their kinetics. As previously
observed in MDDCs, the CXCR4 using strain NL43 showed only
first-phase but not second-phase transfer in eLCs. This is expected
because CXCR4 expression is very low on immature MDDCs and
LCs, and these cells do not support productive replication of ×4
strains (47).

These results clarify the apparent contradictions in reports of first
and second phase transfer and the controversial role of langerin in
HIV infection, for which there is no consensus. First, comparing
these results to previous studies of first-phase vesicular uptake and
transfer (26, 27), they are consistent with those using isolated
CD34+ stem cells (35) but not with others (36). CD34+ stem cell-
derived LCs express MR and langerin, and consequently, the role
of langerin could not be distinguished from MR (35). Our study
used authentic eLCs, which do not express MR and can only bind
the virus via langerin (19). In addition, our study used cell-free virus
and authentic isolated immature eLCs like others (36), but
there were several clear differences, particularly in their low levels of
HIV inoculum, the different clones of anti-langerin Ab used and
LC dissociation by trypsin, which unlike collagenase used in this
study, cleaves off the HIV binding domain of membrane-bound
CD4 (44). With respect to viral inocula, we used a range of HIV
MOIs from low (0.1) to high (10), similar to those used previously
to demonstrate two-phase transfer in MDDCs and blood myeloid
DCs (26, 27). Higher titers of HIV are required for de novo in-
fec tion of DCs compared with macrophages and T cells because
the majority of the HIV inoculum is destroyed by vesicular up-
take, probably in an undefined endocytic pathway (30). For an
optimal single cycle of HIV infection, we have found that >90%
of DCs need to take up the virus, and >5% of cells within the cell
sheet are then infected (26). This need for high inocula appears to
be a consistent property of all immature DCs when infected with
HIV, partly because of constitutive SAMHD1 expression (48) and
inducible restriction factors (49). If the MOI is too low, observa-
ions for over a week are necessary to detect infectious virus.
Exposure of DCs to high titer HIV inocula is, however, physio-
logic as high levels of virus (burst size) are released from T cells,
and 100–10,000 infectious particles/cell (50, 51) are potentially
available to contact LCs, especially in an ulcerated anogenital
epithelium. Comparison of eLCs isolated using trypsin rather than
collagenase showed that eLCs isolated by trypsin were able to
engage in first- but not second-phase transfer as shown by GFP
detection when JLTR cells were added at 2 hpi but not 96 hpi of
eLCs (Fig. 1F).

Furthermore, because HIV-gp120 binds to cell surface langerin
(19, 52), we examined the oligomeric status of cell surface langerin
on Mutz-3 LCs. We showed that langerin can be expressed
as a homotrimer, similar to that of the soluble extracellular domain
ECR (53) and that this enhances its ability to bind HIV, with no
evidence of covalent langerin binding to CD4 or CCR5. These
studies guided the choice of soluble langerin homologs to test as
inhibitors of HIV binding to eLCs, including the trimeric ECD and
the key-binding domain, the CRD. Both inhibited HIV binding
and transfer, as did certain mAbs to langerin. The mAb to langerin
CRD used in our transfer inhibition studies was carefully selected
and based on our studies examining inhibition of oligomeric
langerin–HIV gp140 interactions in view of previous reports that
some of these Abs do not block either these interactions or viral
transfer (31, 38). This was also the reason for using three types of
inhibitors.

Thus, the role of langerin in mediating first-phase transfer is
clearly shown by inhibition of such transfer with both an Ab and
soluble langerin homologs. Two studies of LCs in genital epi-
thelium in evivo explants also claimed to demonstrate first-phase
transfer: Ballweber et al. (8) inoculated vaginal explants with cell-
free virus and demonstrated that emigrating LCs showed no in-
tegrated HIV at 48 hpi but were still able to transfer virus to
T cells. They interpreted this appropriately as first-phase transfer.
As we show in this study, the kinetics of HIV transfer from LCs

<table>
<thead>
<tr>
<th>Seen in Band</th>
<th>Calculated Mass</th>
<th>Δ</th>
<th>Sequence</th>
<th>Protein Name</th>
<th>Accession Number</th>
<th>Position</th>
</tr>
</thead>
<tbody>
<tr>
<td>Monomer 943.48</td>
<td>0.016</td>
<td>K.RFYVPESEP...</td>
<td>Langerin</td>
<td>Q9UJ71</td>
<td>321–328</td>
<td></td>
</tr>
<tr>
<td>Monomer 1012.49</td>
<td>0.003</td>
<td>R.FM*GTISDV...</td>
<td>Langerin</td>
<td>Q9UJ71</td>
<td>67–75</td>
<td></td>
</tr>
<tr>
<td>Monomer 1126.65</td>
<td>-0.053</td>
<td>R.ANAQIQILTR...</td>
<td>Langerin</td>
<td>Q9UJ71</td>
<td>133–142</td>
<td></td>
</tr>
<tr>
<td>Monomer 1391.78</td>
<td>-0.089</td>
<td>K.TAGGLTPIWGLT...</td>
<td>Langerin</td>
<td>Q9UJ71</td>
<td>245–257</td>
<td></td>
</tr>
<tr>
<td>Monomer 1513.79</td>
<td>-0.004</td>
<td>R.QNDILVPVISQGSRK...</td>
<td>Langerin</td>
<td>Q9UJ71</td>
<td>188–200</td>
<td></td>
</tr>
<tr>
<td>Monomer 1546.69</td>
<td>-0.095</td>
<td>K.TW3SAKQFCEVR...</td>
<td>Langerin</td>
<td>Q9UJ71</td>
<td>215–226</td>
<td></td>
</tr>
<tr>
<td>Monomer 1669.89</td>
<td>-0.097</td>
<td>K.RQNDILQVISQGSRK...</td>
<td>Langerin</td>
<td>Q9UJ71</td>
<td>187–200</td>
<td></td>
</tr>
<tr>
<td>Monomer 2097.99</td>
<td>0.007</td>
<td>R.NSHTLSVTSEQPQFLYK.T</td>
<td>Langerin</td>
<td>Q9UJ71</td>
<td>227–244</td>
<td></td>
</tr>
<tr>
<td>Oligomer 943.4427</td>
<td>-0.034</td>
<td>K.RFYVPESEP...</td>
<td>Langerin</td>
<td>Q9UJ71</td>
<td>321–328</td>
<td></td>
</tr>
<tr>
<td>Oligomer 1012.4899</td>
<td>0.003</td>
<td>R.FM*GTISDV...</td>
<td>Langerin</td>
<td>Q9UJ71</td>
<td>67–75</td>
<td></td>
</tr>
<tr>
<td>Oligomer 1126.5927</td>
<td>-0.053</td>
<td>R.ANAQIQILTR...</td>
<td>Langerin</td>
<td>Q9UJ71</td>
<td>133–142</td>
<td></td>
</tr>
<tr>
<td>Oligomer 1513.6927</td>
<td>-0.004</td>
<td>R.QNDILVPVISQGSRK...</td>
<td>Langerin</td>
<td>Q9UJ71</td>
<td>188–200</td>
<td></td>
</tr>
<tr>
<td>Oligomer 1842.8927</td>
<td>-0.044</td>
<td>R.SWEESTLNIQPFLK...</td>
<td>Langerin</td>
<td>Q9UJ71</td>
<td>143–158</td>
<td></td>
</tr>
<tr>
<td>Oligomer 2097.9927</td>
<td>0.007</td>
<td>R.NSHTLSVTSEQPQFLYK.T</td>
<td>Langerin</td>
<td>Q9UJ71</td>
<td>227–244</td>
<td></td>
</tr>
</tbody>
</table>

Table I. Tryptic peptides identified by mass spectrometry in bands containing monomers and oligomers of langerin.

*Data indicate whether the peptide was identified in a gel slice containing monomeric langerin or oligomeric langerin.
*Data were calculated monoisotopic masses of uncharged peptides.
*Δ indicates data were calculated as difference between the experimentally derived mass of the precursor and the calculated mass of the peptide.
*M* indicates oxidized methionine. Amino acids flanking the peptide on both sides are shown separated from the peptide sequence by a full stop.
*Given are the accession numbers for the SwissProt database.
*Data indicate the position of the peptide in the unprocessed precursor obtained from the SwissProt database.
are slower compared with MDDCs, and no HIV was detected in supernatants until 72–96 hpi. Therefore, perhaps the presence of integrated HIV DNA should be assessed at later time points before concluding that vaginal LCs do not support de novo infection. The transfer that they observe on days 7–16 could still be “second phase.” Ganor et al. (4) infected inner foreskin explant cultures with either cell-free or cell-associated virus and showed rapid uptake of HIV by LCs and formation of LC–T cell conjugates within 1 h. However, transfer of HIV within these conjugates was assumed rather than demonstrated. Neither studies were able to assess the role of langerin. The exact mechanism for the longer duration of first-phase transfer in LCs needs to be investigated. Although first-phase viral transfer has been clearly shown to occur from VCCs in MDDCs, rapid degradation over 6 h also occurs (Fig. 4) (26), suggesting an interaction between the VCC and the endocytic pathway, which has not yet been defined. This may differ somewhat between the two cell types, resulting in slower degradation and longer preservation of infectious viruses in LCs.

For second-phase or de novo infection and transfer, our results are consistent with previous reports from the Blauvelt and Kawamura laboratories (34), except for the role of langerin. Using CCR5 inhibitors (PSC-RANTES) or eLCs from CCR5Δ32

FIGURE 6. Inhibiting HIV transfer from LCs to T cells using soluble langerin homologs and neutralizing mAb to langerin. (A) A representative native PAGE gel showing the oligomeric states of purified soluble langerin CRD and ECD used in blocking experiments. (B) Cytotoxicity of soluble langerin ECD and CRD. Mutz-3 LCs were seeded in 96-well plates at 50,000 cells/100 μl media, and both forms of soluble langerin were added individually at the final concentrations of 0.1, 0.2, 0.5, 1 and 2 μM. After 72 h, Mutz-3 LCs were counted using trypan blue to assess viability (n = 3). (C) eLCs were 1) HIV_{null} treated (HIV), 2) treated with HIV that was preincubated with either ECD (HIV+ECD) or CRD (HIV+CRD) for 1 h at 37˚C, 3) treated with mAb to langerin CRD (clone 817G7) for 30 min at 4˚C before HIV_{null} infection (HIV+anti-lang), or 4) treated with the reverse transcriptase inhibitor AZT (HIV+AZT) for 30 min before HIV_{null} infection. Cells were infected at an MOI of 0.5 for 2 h at 37˚C. ILTR cells were added after eLCs were thoroughly washed three times in PBS either at 2 hpi (first-phase transfer) or at 96 hpi (second-phase transfer). Data shown represent the mean GFP⁺ ILTR ± SE from three separate experiments derived from different donors eLCs using flow cytometry after 96 h postcoculture (***p = 0.02, ****p = 0.005). (D) Mutz-3 LCs were similarly treated with inhibitors as described in (C), and CD4⁺ T cells were added after infected Mutz-3 LCs were washed three times in PBS. Intracellular p24 staining was assessed after 96 h postcoculture (n = 3, ***p = 0.02 for both ECD and anti-langerin). (E) ILTR cells were inoculated with either HIV or HIV-ECD at an MOI of 0.2 or 1 for 2 h at 37˚C, followed by three washes in PBS. Data shown represent the mean GFP⁺ ILTR ± SE (n = 3, *p = 0.04).
heterozygotes, they reported that CD4 and CCR5 are of key importance in productive HIV infection of LCs and in subsequent transfer to T cells. However, their conclusion that there was no role for langerin in concentrating or transferring virus to these receptors was based on a single application of mannan (14, 18, 34) and one report of anti-langerin Ab (14) (clone 122DS/DCGM4) as inhibitors. These studies used eLCs infected in situ in epidermal explants, which then emigrated 3 d. The apparent disparity with our results may be explained in Fig. 6. Blocking second-phase transfer required sustained presence of the langerin Ab in the culture, not a single dose prior and at the time of viral application. Presumably this is due to HIV persistence beyond the activity of the Ab both in vitro culture and within epidermal explants. Furthermore, other clones of anti-langerin CRD Ab are claimed to be more potent inhibitors (36). The results with the ECD and CRD inhibitors indicate that the role of langerin is similar to that of DC-SIGN in enhancing both phases of viral transfer. As a control, ECD langerin did not impair direct HIV infection of T cells, indicating the inhibition was specific and not just steric hindrance of HIV binding to CD4/CCR5.

Using transcriptomic analysis, we recently reported that Mutz-3 cells are different from authentic eLCs (54). However, these model cells are useful in examining HIV binding to surface langerin and in assessing the role of langerin, especially when it is impossible to obtain sufficient eLCs for biochemical studies. After cytokine manipulation, Mutz-3 LCs expressed high levels of CD4 and langerin but lacked the expression of DC-SIGN, MR, and cell surface CCR5. In addition, intracellular CCR5 RNA was much lower than in eLCs. Thus, as shown in Fig. 3, Mutz-3 LCs could only be productively infected with VSV-G pseudotyped and not with wild-type HIV. This explains the absence of the second phase transfer by Mutz-3 LCs at any HIV concentration (MOIs of 4, 1, 0.5, and 0.05), in contrast to a previous report (55). HIV was used in this study as it was previously shown a higher affinity for CCR5 than any other commonly used HIV strain (43).

The relative importance of first versus second phase transfer from LCs to T cells in vivo or ex vivo requires further study despite the recent suggestions from foreskin and vaginal explant studies that first phase is more important (4, 8), contrasting with previous data from Kawamura et al. using PSC-RANTES (34) and recently Maraviroc (56). Recent reports of transmitted R5 HIV-1 strains suggest they are T- rather than M-tropic (57), which might suggest first phase transfer to be more important. However, second-phase transfer also would result in passage through mucosal T cells, so the latter cells remain as the filter even after transmission from DCs (57).

In the absence of a HIV vaccine, the search for effective microbicides to protect against sexual transmission remains an important approach to prevent infection. Currently, the only microbicide reported to be clinically effective is the reverse transcriptase inhibitor, Tenofovir, which showed a 39% decrease in infection rate (58). It is therefore likely that combination therapy with microbicides will be required to improve efficacy, similar to systemic antiviral therapy and/or chemoprophylaxis. Such a strategy should include inhibitors of HIV entry into mucosal DCs. Prior to this study, HIV transmission by LCs to T cells has been reported to be either independent (2, 8, 59) or dependent on productive LC infection (15, 16, 59, 60), and the controversy surrounding the function of langerin during the early stages of infection has led to uncertainty in designing and testing langerin inhibitors as potential anti-HIV microbicides. Thus, these two temporal phases of HIV uptake and transfer have been shown sequentially in authentic eLCs and that both are at least partially mediated by langerin. Furthermore, we showed that soluble langerin ECD and CRD and also mAb to langerin were potent in inhibiting first- and second-phase viral transfer to T cells. Perhaps a combination of specific inhibitors may be required to block both the interaction between HIV envelope and trimeric langerin on the eLC surface and also the interaction between HIV envelope and CD4/CCR5, thereby optimally inhibiting both the viral transfer from VCC and productive infection. In addition, the equivalent inhibitory efficacy of the smaller CRD monomer to the trimeric ECD is promising for the development of synthetic inhibitors.

Acknowledgments

We thank Dr. Richard Barnett for providing us with the skin specimens, Dr. Eve Diefenbach for assistance with mass spectrometry, Dr. Saskia J.A.M. Santegoets (Department of Pathology, VU University Medical Center, Amsterdam, the Netherlands) for providing us with the Mutz-3 cell line, and Dr. Xin Wang for her assistance with cell sorting that was performed at the Westmead Research hub Flow Cytometry Centre, which is supported by the Westmead Millennium Institute, the Children’s Medical Research Institute, the Kid’s Research Institute, the National Health and Medical Research Council, and the Cancer Institute New South Wales.

Disclosures

The authors have no financial conflicts of interest.

References

2564

INHIBITION OF HIV-1 TRANSFER FROM LCs


Figure S1. Purity of isolated eLCs which co-express CD1a and langerin. A) After collagenase treatment of epidermal sheets, the isolated cells were stained for CD1a, CD45, CD3 and CD14. eLCs were sorted by excluding the CD3⁺CD14⁺ populations and collecting the CD1a⁺CD45⁺ population on a FACSVantage fluorescence-activated cell sorter to a purity of >98%. B) Human skin epidermis was stained with CD1a (BD, Pharmingen) and langerin (R&D Systems; Minneapolis, MN, USA). eLCs co-express CD1a (green) and langerin (red). Nuclei were stained with DAPI (blue; Invitrogen). Fluorescently labeled cells were visualized with an inverted Olympus IX-70 microscope (DeltaVision Image Restoration Microscope; Applied Precision/Olympus) using a numerical aperture oil immersion lens (1.4 or 1.43) and a photo-metrics CoolSnap QE camera. The representative z-series were deconvoluted, the pictures overlaid, colocalization and significance were performed using SoftWoRx software (version 3.4.5, Universal Imaging Corporation). Bars indicate 15 µm.
Figure S2. Viability, proliferation and viral replication of JLTRs. A) Viability was assessed by flow cytometry using 50 µg/mL DAPI (Life Technologies, Australia) staining at 1, 3, 5 and 6 days post JLTR passage. Filled histograms represent JLTRs treated with ethanol (dead cells, positive control for DAPI stain) and open histograms show JLTR viability (90%±7) in the culture. B) Proliferation was assessed by flow cytometry using 5 µM cell trace carboxyfluorescein succinimidy Ester CFSE (Life Technologies, Australia) staining on day 2, 3, 4, 5 and 6 post passage. Open histogram represents unstained JLTRs (negative control) and filled histograms represent JLTRs stained with CFSE and how the CFSE intensity decrease as cells were proliferating over 6 days. C) HIV_{Bal} infection (MOI=0.1) of JLTRs was assessed by flow cytometry by measuring GFP fluorescence intensity by flow cytometry on days 2, 3, 5 and 6 post infection.
Fig S3: Soluble ECD langerin homolog does not inhibit DC antigen presentation measured as T cell alloproliferation. T cell alloproliferation was determined by carboxyfluorescein succinimidyl ester (CFSE) dilution using mature and immature Mutz-3 LCs. Mutz-3 LCs were matured for 48 h in maturation mix consisting of 50 pg/mL IL-1β (R&D Systems; Minneapolis, MN, USA), 5 U/mL IL-6 (R&D Systems), 50 pg/mL TNF-α (R&D Systems) and 5 ng/mL PGEβ (Sigma; Milwaukee, WI, USA) in 0.1% bovine serum albumin (BSA, Sigma) in PBS. Briefly, allogeneic PBMCs were stained with 5 µM final concentration CSFE (Molecular Probes; Eugene, OR) for 10 minutes at 37°C, rescued with equal volume of 100% FBS and washed with RPMI 1640 supplemented with 10% fetal bovine serum. These PBMCs were added to Mutz-3 LCs (mature and immature) treated with different concentrations of ECD (and CRD; data not shown) at a ratio of 1 Mutz-3 LCs:10 PBMCs and incubated at 37°C for 5 days. PBMCs with or without PHA were added to Mutz-3 LCs as positive or negative controls, respectively. Cells were subsequently stained for CD4. CFSE and CD4 expression were assessed by flow cytometry.