Exposure to HIV-1 Altered CCR7-Mediated Migration of Monocytes: Regulation by PGE$_2$

Sandra C Côté, Stamatoula Pasvanis and Nancy Dumais*

Département de Biologie, Faculté des Sciences, Université de Sherbrooke, Sherbrooke (QC), Canada

*Corresponding author: Nancy Dumais, Faculté des Sciences, Université de Sherbrooke, Sherbrooke (QC), J1K 2R1, Canada, Tel: +1-819-821-8000, Fax: +1-819-821-8049, E-mail: nancy.dumais@usherbrooke.ca

Abstract

Monocytes play critical roles in human immunodeficiency virus type-1 (HIV-1) pathogenesis. During HIV-1 infection, proinflammatory molecules such as prostaglandin E$_2$ (PGE$_2$) are observed at elevated levels in infected individuals. Mono-Mac-1 cells as well as freshly isolated monocytes were exposed to pseudotyped, R5 or dual tropic HIV-1 particles. CCR7 expression was determined by FACS analysis while chemotaxis assays were performed to show CCR7 functionality. We demonstrated that PGE$_2$ enhanced CCR7 surface expression on monocytes. However, monocytes exposition to R5 or dual tropic HIV-1 particles did not highly modulate surface expression of CCR7. On the other hand, monocytes exposition to HIV-1 impaired the CCR7 dependent migratory capacity to CCL19. The addition of PGE$_2$ to HIV-1-exposed monocytes restored the CCR7 dependent migration to levels similar to PGE$_2$ treated unexposed monocytes. Monocytes acquired a functional responsiveness to CCL19 after exposure to PGE$_2$ only when gp120 is expressed on viral particles. Finally, HIV-1-exposed monocytes that migrated in a Transwell system efficiently transmitted the infection to sensible cells. Collectively, we show that PGE$_2$ is essential for the CCR7-dependent migration of monocytes exposed to HIV-1.

Keywords

Monocytes, HIV, CCR7-dependent migration, PGE$_2$

Background

Transmission and pathogenesis of the Human Immunodeficiency Virus type-1 (HIV-1) are interconnected during all stages of the disease with cells of the monocyte/macrophage lineage [1-3]. Monocytes are constantly exposed or are targets of HIV-1 and their migratory behaviors are implicated in the dissemination of the infection throughout the host [4-8]. Monocytes arising from quiescent and infected cells of the myeloid precursor lineage in bone marrow have been demonstrated as a source of residual HIV DNA [1,7,9-12]. Despite treatment with antiretroviral therapies, monocytes can harbor latent HIV-1 proviral DNA during all stages of the disease forming a long-lived reservoir for the virus [4] and appropriate stimulation and/or differentiation can reactivate productive replication [6,13,14]. Infected circulating monocytes can migrate to peripheral tissues such as the brain [13,15], lung [16], lymphatic system [17], bone marrow [9], and kidney [18] but the implicated mechanism is poorly understood.

The chemokine receptor CCR7 plays an important role in the migration behavior of immune cells. CCL19 and CCL21 are the natural ligands of CCR7, which is not only expressed on DCs [19] but also on T and B cells [20-22] as well as on monocytes [23]. Mice deficient in CCL19, CCL21 or CCR7 demonstrate defective DC trafficking and altered immune response [24,25]. In human monocytes and DCs, prostaglandin E$_2$ (PGE$_2$) a pleiotropic immunomodulatory molecule, exerts multiple effects on both CCR7 expression level and functionality [23,26,27].

Since it has been shown that monocytes are actively implicated in HIV-1 pathogenesis and serum level of PGE$_2$ is a hallmark of HIV progression [28-30], we asked whether PGE$_2$ modulates CCR7 expression and functionality on HIV exposed monocytes. We investigated the cell surface expression of CCR7 in monocytes exposed to R5 or dual tropic strains of HIV-1. In parallel, chemotaxis assays were performed to establish the influence of HIV-1 exposition on CCR7 functionality. We also sought to determine whether the presence of PGE$_2$ could modify the migratory capacity of monocytes exposed to HIV-1. Our results provide evidence that PGE$_2$ restores functional CCR7 expression on HIV-exposed human monocytes allowing their migration toward the CCR7 ligand CCL19. Expression of CCR7 on HIV-exposed monocytes in response to PGE$_2$ may have significant implications for HIV-1 dissemination to secondary lymphoid organs and/or peripheral tissues.

Methods

Ethics statement

The study was approved by the local ethics committee (Comité d’Éthique de la Recherche sur l’Humain, Centre Hospitalier Universitaire de Sherbrooke, Université de Sherbrooke (04-116-R5)) and written consent was obtained from all participants.

Cells

Mono-Mac-1 cells, an acute peripheral monoblastic leukemia derived cell line (German Collection of Microorganisms and Cell Cultures, Braunschweig, Germany) was cultured in RPMI 1640 (Wisent, St-Bruno, QC, Canada) supplemented with 10% heat-inactivated fetal bovine serum (FBS) (Wisent), non essential amino acids (NEAA) (Wisent), 1 mM sodium pyruvate (Wisent), 100 U/L penicillin, and 100 μg/ml streptomycin (Wisent). Human embryonic kidney 293T cells (American Type Culture Collection (ATCC)


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Manassas, VA) and the TZM-bl HIV-1 indicator cell line (NIH AIDS Research and Reference Reagent Program) were cultured in Dulbecco’s modified Eagle’s medium (DMEM) (Wisent) supplemented with 10% heat-inactivated FBS, NEAA, 1 mM sodium pyruvate, 100 IU penicillin, and 100 μg/ml streptomycin. TZM-bl is an indicator cell line derived from HeLa cells, which express the HIV-1 receptors CD4, CXCR4, and CCR5 and carry a stably integrated luciferase reporter gene under the control of HIV-1 long terminal repeat (LTR) [31]. Human monocytes were isolated from healthy donors’ blood using Lymphocyte Separation Media as recommended by manufacturer (Wisent) and left to adhere on culture plates for 1 hour at 37°C in RPMI 1640 supplemented as described above. Suspension cells were removed by two washes with phosphate buffered saline (PBS) and adherent cells were detached using Cell Dissociation Media (Wisent), washed with PBS, and used immediately.

**Production of viral stocks**

Pseudotyped HIV-1 particles were obtained by co-transfection of pNL4-3-LucE’R+ with pcDNA1/Amp-based expression vectors coding HIV-1 ADA (R5), JR-FL (R5) envelope proteins or pHCMV-G vesicular stomatitis virus envelope glycoprotein G (VSV-G) under the control of the human cytomegalovirus promoter. The infectious molecular clones pcDNA1/89.6 and pNL4-3-balenv were also used. All those plasmids were kindly provided by Dr. M. J. Tremblay (Centre de Recherche en Infectiologie, CHUL, Québec, Canada). The virus-containing supernatants were collected 48 hours after transfections with FuGENE® 6 (Roche Applied Science, Laval, QC), filtered through a 0.45-μm cellulose acetate, and normalized for virion content using a p24 ELISA as described previously [32].

**Exposition of monocytes to HIV-1**

Freshly isolated blood monocytes and Mono-Mac-1 were exposed to HIV-1 strains using 100 ng p24 per million cells for 1 hour at 37°C, washed with PBS, and seeded at 1 × 10⁶ cells per ml in RPMI 1640 supplemented. Cells were then left untreated or treated with 1 μM PGE₂ (Sigma Aldrich, Oakville, ON, Canada).

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**Figure 1: PGE₂ enhances CCR7 surface expression on HIV-1 exposed monocytes**

Mono-Mac-1 cells were exposed to the indicated pseudotyped (A) or replication competent (B) HIV-1 clones and left untreated or treated with 1 μM PGE₂ for 24 hours. FACS analysis was performed on cells labeled by a CCR7-specific antibody (black line) or corresponding isotype control (gray line). Percentages of positive cells are indicated. Results represent 1 out of 5 independent experiments with similar results. (C) Blood isolated monocytes were exposed to the indicated HIV-1 replication competent clones and left untreated or treated with 1 μM PGE₂ for 24 hours. Cells were double-stained for CD14 and CCR7 using specific antibodies and analyzed by flow cytometry. Quadrants were positioned based on corresponding isotypes. Results are from 1 of 4 different donors with similar responses.
Chemotaxis assays

Chemotaxis of HIV-1 exposed blood isolated monocytes and Mono-Mac-1 was measured by migration through a polycarbonate filter with 5- or 8-µm pore size respectively, in 24-wells transwell chambers (Millipore, Nepean, ON, Canada). The lower chamber contained 500 µl of a 300 ng/ml dilution of CCL19 in medium without PBS (R&D Systems, Minneapolis, MN, USA), or medium alone as a spontaneous migration control, and the upper chamber contained 5 × 10⁵ cells in 100 µl of medium. Blood isolated monocytes containing chambers were incubated for 2 hours at 37°C and Mono-Mac-1 containing chambers were incubated for 4 hours at 37°C. A 100 µl aliquot of the cells that migrated to the bottom chamber were used for the infectivity assay or mixed with 200 µl of a 2% paraformaldehyde solution and counted using BD FACSCalibur flow cytometer (BD Biosciences) by acquiring events for a fixed period of 60 seconds using CellQuest software (BD Biosciences). The percentage of migrated cells was calculated as follows: the number of migrated cells in response to media only was subtracted to the number of migrated cells to CCL19 and this number was reported on the total input of cells.

Infectivity assays

Reporter TZM-bl cells were seeded at 10⁴ cells/well in 96-well plates 12 hours before the assay. A 100-µl aliquot for the bottom well of the migration assay (see above) was added to the TZM-bl cells in triplicate. Cells were co-cultured for 2 hours and then the media was replaced with fresh complete media. The cells were incubated for an additional 48 hours at 37°C and analyzed for luciferase activity [33].

Flow cytometry

Monocytes were collected and washed twice with ice-cold PBS. The Fc receptors were blocked using 1 µg purified IgG (Santa Cruz Biotechnology, Santa Cruz, CA, USA) for 10 minutes on ice. Mono-Mac-1 cells were stained with a phycoerythrin (PE)-conjugated anti-CCR7 antibody (R&D Systems). Blood isolated monocytes were double-stained with PE-conjugated anti-CCR7 and a Peridinin chlorophyll protein complex (PerCP)-conjugated anti-CD14 antibody (BD Biosciences, Mississauga, ON, Canada) or were incubated with corresponding isotypes as negative controls. Cells were then washed twice with ice-cold PBS supplemented with 3% bovine serum albumin (BSA) and fixed for 30 minutes on ice in 2% paraformaldehyde. Fluorescence was read using a BD FACSCalibur flow cytometer (BD Biosciences) and results were analyzed using CellQuest software (BD Biosciences).

Statistical analysis

Each experiment was performed at least three times in triplicate. Statistically significant differences between experimental groups were evaluated using a paired t-test. Computations were carried out using GraphPad PRISM version 5.0b statistical software.

Results

Surface expression of CCR7 on HIV-1 exposed human monocytes is modulated by PGE₂

We initially characterized the surface expression of CCR7 on HIV-1 exposed monocytes without or with PGE₂ treatment (Figure 1). Our results demonstrated that exposition to pseudotyped viruses did not considerably alter cell surface expression of CCR7 compared to mock-treated cells (Figure 1A). However the addition of PGE₂ to HIV-1-exposed Mono-Mac-1 was associated with an increase of CCR7 expression compared to PGE₂-treated uninfected cells. The number of CCR7+ cells increased of 20.5% in uninfected cells and 9.7% and 15.4% in cells exposed to NL4-3-Luc+E/R+Adaenv or NL4-3-Luc+E/R+Adaenv+Adaenv, respectively. Next, Mono-Mac-1 cells were exposed to replication competent viruses (NL4.3balaenv and 89.6) and FACS analysis revealed that CCR7 expression was slightly affected by the viral exposition (Figure 1B), with uninfected cells being 54.5% CCR7+ and 61.7% and 59.0% for NL4.3balaenv and 89.6, respectively. The addition of PGE₂ to replication competent HIV-1-exposed Mono-Mac-1 cells also augmented CCR7 cell surface expression as observed with cells exposed to pseudotyped viruses. PGE₂, treatment increased CCR7 expression by 25.0% in uninfected cells and 17.8% and 21.1% in cells exposed to NL4.3balaenv and 89.6, respectively. Since, Mono-Mac-1 cells are an immortalized cell line whose growth properties have been altered by transformation, we repeated this series of experiments with freshly isolated blood monocytes from healthy donors. As above, we determined the extent to which HIV-1 exposure and PGE₂ treatment affects CCR7 cell surface expression in freshly isolated blood monocytes (Figure 1C). In mock-treated cells, 9.7% of CD14+ cells expressed CCR7. Exposition to NL4.3balaenv and 89.6 reduced CCR7 expression to 4.78% and 5.22%, respectively. Stimulation of uninfected blood monocytes with PGE₂ increased CCR7 cell surface expression for uninfected cells (14.41%), as well as blood monocytes exposed to HIV-1 (NL4.3balaenv; 12.65% and 89.6: 10.09%). This degree of increased CCR7 surface expression is functionally relevant since other reports have shown that similar percentages of CCR7 expression enhance cell migration in response to CCL19 in the presence of PGE₂[23,26,27]. Collectively, these data demonstrate that addition of PGE₂, invariably increased cell surface expression of CCR7 in both Mono-Mac-1 cells and freshly isolated blood monocytes. In contrast, HIV-1 exposure did not significantly modulate the expression of CCR7 on monocytes. Indeed in Mono-Mac-1, exposition to pseudotyped HIV-1 particles slightly decreased cell surface expression of CCR7 while CCR7 levels were unchanged in Mono-Mac-1 cells. In freshly isolated blood monocytes exposition to R5 or dual tropic HIV-1 particles did not significantly change CCR7 expression.

PGE₂ enhanced CCR7-dependent migration to CCL19 of infected monocytes

We next assessed CCR7 function by measuring the migratory response of cells to the chemokine CCL19. The chemotaxis assays showed that PGE₂, substantially increased Mono-Mac-1 cell migration in response to the CCR7 ligand CCL19 (Figure 2), as previously observed [23]. Incubation of Mono-Mac-1 cells with pseudotyped HIV-1 particles (NL4-3-Luc+E/R+Adaenv or NL4-3-Luc+E/R+Adaenv+Adaenv) resulted in specific migration of 6.9 ± 3.8% for NL4-3-Luc+E/R+Adaenv and 5.9 ± 3.8% for NL4-3-Luc+E/R+Adaenv+Adaenv, which were similar to the migration behaviour of uninfected cells, 6.2 ± 3.5% (Figure 2A). In contrast, exposition to fully infectious HIV-1, NL4.3balaenv or 89.6, induced a significant augmentation in the percentage of migrating cells compared to the unexposed Mono-Mac-1 cell control (Figure 2B). Interestingly, PGE₂, treatment of cells exposed to either pseudotyped or fully infectious viruses increased the percent of migrating cells to levels similar to unexposed Mono-Mac-1 cells (Figure 2A and Figure 2B). We demonstrated that the observed migration was CCR7 specific by showing a loss of migration upon blocking CCR7 (data not shown).

Binding of chemokines to their receptors causes a characteristic increases in cytosolic calcium, which is one of the earliest biochemical events that occur in response to chemokines [34]. Thus, we hypothesized that treatment of cells with the potent intracellular Ca²⁺ chelator, BAPTA/AM [35], would block the migration-promoting actions of PGE₂. In cells treated with PGE₂, we observed 19.7 ± 6.9% cell migration but pretreatment with 25 µM BAPTA/AM reduced migration to 2.9 ± 1.6% (Figure 3) without affecting cell viability (data not shown). These observations establish a primary role for intracellular calcium signaling in the PGE₂-induced migration of Mono-Mac-1 cells in response to CCL19.

To further investigate the effects of PGE₂ on CCR7-dependant migration of HIV-1-exposed monocytes, we repeated this series of experiments with freshly isolated human blood monocytes. These cells were exposed to different strains of pseudotyped viruses or replication competent HIV-1 clones and then HIV-1-exposed blood-isolated monocytes were left untreated or treated with PGE₂ for 24 hours. Cells were then assessed for CCL19-specific migration as described above (Figure 4). While HIV-1 exposure reduced the number of migrating monocytes, PGE₂ treatment significantly increased specific
migration toward CCL19. Migration analysis of monocytes that were exposed to replication competent viruses showed significantly decreased numbers of migrating cells (Figure 4A compared with Figure 4B). However, the impaired migration observed in HIV-1-exposed cells was reversed by PGE2 treatment, resulting in migration levels corresponding to PGE2-treated uninfected monocytes. Importantly, our results indicate that HIV-1 exposure perturbed CCR7 expression in freshly isolated blood monocytes leading to a generalized defect of CCR7-dependent chemotaxis. This defect was reversed by the addition of PGE2. These data support the above results obtained by flow cytometry analysis of CCR7 surface expression observed in HIV-1-exposed blood monocytes treated with PGE2.

Next, we investigated whether the HIV-1 envelope glycoprotein directly regulated chemotaxis by exposing freshly isolated monocytes to the VSV-G pseudotyped virus NL4-3-LucE'R'/VSVenv. As shown in Figure 5, exposition to virus particles where the gp120 was replaced by the VSV glycoprotein did not significantly reduce the percentage of migrated cells in the absence of PGE2 (uninfected cells: 12.8 ± 3.2% and NL4-3-LucE'R'/VSVenv infected cells: 10.9 ± 6.7%). Our results suggest that the interaction of gp120 with its receptor(s) on monocytes could be implicated in CCR7-driven migration.

HIV-1-exposed monocytes can transfer infection to susceptible cells

Finally, we evaluated whether HIV-1-exposed monocytes that
migrated in response to CCL19 were able to efficiently spread the virus to TZM-bl indicator cells (Figure 6). Treatment of NL4.3balenv-exposed monocytes with PGE2 significantly increased the relative infection index (RLU/number of migrated cells in the lower chamber of the Transwell) compared to untreated HIV-1-exposed monocytes (Figure 6). Together the data demonstrated that PGE2 considerably regulates the capacity of HIV-1-exposed monocytes to migrate toward CCL19 and transmit the virus to uninfected host cells.

Discussion

Monocytes are recognized as multifunctional contributors to immune function but many questions remain to be answered, such as how monocytes acquire trafficking and migration behaviors. Although it is controversial that monocytes are susceptible to HIV-1, several studies have reported that circulating monocytes are indeed infected in vivo and can serve as important reservoirs of HIV-1, especially in patients with late-stage disease and opportunistic infections [1,6-12,36]. Monocytes are regarded as early targets for HIV infection and as an important viral reservoir. Indeed, infected monocytes are important short-lived viral pool that upon entering tissues differentiated into macrophages (reviewed in [37]). On the other hand, HIV infection causes an early increase and a late decrease of CCL19 and CCL21 that are associated with disease progression [38-40], HIV replication in T cells [41,42], as well as
the release of inflammatory mediators [39]. However, the impact of HIV-1 exposure on CCR7 expression in monocytes was still not characterized. Here, we have shown for the first time that exposition of freshly isolated human monocytes to R5 and dual-tropic HIV-1 altered CCR7-dependent migration in response to CCL19.

Firstly, this study demonstrated that HIV-1-exposure of human monocytes impaired CCR7-dependent migration in response to CCL19 and this required gp120 binding to human monocytes. This disruption of CCR7-dependent monocyte chemotactic function in HIV-1 infection may have major consequences on monocyte circulation given that CCR7 play an essential role in immune cells entry to lymph nodes. Due to defective CCR7 functions, monocytes exposed to HIV are likely to enter and circulate within peripheral tissues less efficiently, leading to decreased recognition pattern. The virus, to favour its dissemination, may exploit this mechanism. Furthermore and in concordance with our study, HIV infection was associated with reduced T-cell migration in response to CCL19 [43]. It was also shown that, in spite of preserved CCR7 expression, CCR7-dependent chemotactic responses were significantly decreased within most T-cell subsets from viremic patients, including naïve, central memory, and effector memory CD4+ T cells and naïve, central memory, and effector CD8+ T cells. Further, this defect in T-cell chemotaxis was only partially corrected by effective HAART treatment [43]. Our data are also supported by an investigation identifying a novel defect in CCR7 expression of plasmacytoid dendritic cells (pDCs) in pediatric patients with progressive HIV disease. The failure of CCR7 up-regulation was associated with failure of immune reconstitution and with ongoing plasma virus replication [44]. They proposed that failure to up-regulate CCR7 could be an impediment to the homing of pDCs to lymphoid tissue resulting in impaired maturation of myeloid DCs and impaired T-cell activation. In addition, it has been recently shown that respiratory syncytial virus infection can downregulate CCR7 expression in newborns leading to delayed innate and adaptive immune activation [45]. Taken together, these data suggest that HIV has an important impact on cell circulation but the mechanism of HIV inhibition of CCR7 functionality remains to be fully defined.

Secondly, because serum levels PGE2 is a hallmark of the progression of the progression of HIV infection and contributes HIV pathogenesis, we also investigated its effect on the migratory capacity of HIV-exposed monocytes. We assessed both the cell surface expression of CCR7 and CCR7 response to its ligand CCL19. As previously observed, we confirmed that PGE2 induced functional CCR7 expression on blood monocytes and Mono-Mac-1 cells, enabling them to respond to and migrate toward CCL19. Interestingly, PGE2-treatment of HIV-exposed blood-isolated monocytes restored their potential to migrate in response to CCL19 to a level comparable to PGE2-treated unexposed monocytes. The cellular and molecular mechanisms responsible for this are being pursued in our laboratory. Since CD16+ monocytes are sensitive to HIV-1 infection and they serve as a major virus reservoir [8,46], restoration of CCR7 function due to PGE2 may influence the pattern of cell recirculation, allowing cells to be directed to lymph nodes. Peyer’s patches as well as other sites [9,13,15-18]. Moreover in our study, we also demonstrate that monocytes exposed to HIV-1 in the presence of PGE2 transfer HIV-1 particles to uninfected susceptible cells in response to CCL19. These findings may have important implications in HIV-1 dissemination to anatomical reservoirs. Indeed, interactions between prostaglandins, leukotrienes and HIV-1 may modulate HIV infection and contribute to the viral dissemination to anatomical reservoirs such as the central nervous system (reviewed in [47]).

The demonstration that CCR7 expression and functionality is restored by PGE2 in monocytes is not a universal phenomenon because chemokines receptors for other chemotactants such as CCR5, CCR2 or CXC4 are not affected by the same manner. The expression of CCR5, the receptor for RANTES, which may affect not only leukocyte migration, but also infectivity by HIV-1, has been shown to be down-regulated by PGE2 in monocytes [48,49] whereas CCR2 and CXCR4 expression were not affected [49]. While Thivierge et al. [48] showed that exposure to PGE2 reduced chemotaxis of monocytes in response to MIP-1β, Panzer and Uguccioni [49] clearly demonstrated that PGE2 enhances the migration in response to MCP-1, RANTES and SDF-1. Altogether, these data indicate that in inflammatory conditions, the up-regulated production of PGE2 by monocytes or tissue cells might favor the migratory capacity towards locally expressed inflammatory chemokines. By extension, monocytes in close contact with HIV-1 or infected by HIV-1 might disseminate the virus throughout the body.

Conclusion

In summary, our findings suggest a complex interplay between HIV and PGE2 for the expression of functional CCR7 on monocytes. In one hand, HIV exposure impairs CCR7-specific migration in monocytes. In the other hand, PGE2, found in serum of HIV-1 infected individuals in concentrations that increase over the time, restores the migratory capacity of HIV-1-exposed monocytes to CCL19. Our data reinforce the concept that PGE2 is a key mediator in CCR7 functionality. Since CCL19 and CCL21 production is induced in early phases of HIV-1 infection and serum level of PGE2 is amplified, circulating HIV-1-loaded monocytes may accumulate in peripheral lymph nodes as well as in other sanctuary sites such as brain where they can spread the infection to other susceptible cells.

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Authors’ contributions

SCC performed the experiments, prepared the figures, analyzed the data and drafted the manuscript. SP carried out the transfections and viruses production. ND conceived the study, participated in its design and coordination and wrote the manuscript. All authors read and approved the final manuscript.

Conflict of interest

The authors declare they have no competing interests.


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Page 7 of 7