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RANTES Production from CD4+ Lymphocytes Correlates with Host Genotype and Rates of Human Immunodeficiency Virus Type 1 Disease Progression

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In this study, we sought to determine whether the characteristics of chemokine production or chemokine receptor expression before HIV-1 infection predicted the postinfection disease course and how these were related to the presence or absence of a Δ32CCR5 allele.

Patients and Methods

Study patient selection. On the basis of the availability of pre- and postinfection peripheral blood samples (time of infection was
documented by measurements of plasma HIV-1 RNA), we selected 56 men who were enrolled in the Chicago component of the Multicenter AIDS Cohort Study, a natural history study of men at risk for HIV-1 infection. Study participants were stratified by the level of HIV-1 RNA in plasma 6 months after the extrapolated time of infection and by the rate of declining CD4+ T cell numbers over time. Patients were excluded from the study if the isolated virus strain used CXCR4 for virus entry [11]. We selected an additional 20 men who remained uninfected to serve as control subjects.

**In vitro infectivity assay.** Viable cryopreserved peripheral blood mononuclear cells (PBMC) were thawed, were stimulated with 5 μg/mL of phytohemagglutinin (PHA), and were carried in culture medium (RPMI 1640 supplemented with antibiotics and 10% fetal bovine serum) containing 100 U/mL interleukin-2. Ten days after culture, the cells were enriched for CD4+ or CD8+ lymphocytes by use of the corresponding immunomagnetic beads. The in vitro virus infection assays were done as described elsewhere [2]. All PHA-stimulated PBMC culture supernatants were monitored on day 10 for the presence of p24, to exclude endogenous HIV-1 activity.

**Chemokine and chemokine receptor expression levels.** RANTES production levels were measured from purified CD4+ and CD8+ lymphocytes. Enriched CD4+ and CD8+ lymphocytes were cultured separately at 1.0 × 10⁶ cells/mL for 4 days after separation, and RANTES levels were determined by use of a commercially available immunoassay (R&D Systems). The CCR5 and CXCR4 cell surface expression levels were determined by fluorescence-activated cell sorter (FACS) analysis, using monoclonal antibodies 2G12 and 2D7, respectively, as described elsewhere [5]. CCR5 and CXCR4 mRNA expression levels were determined by use of real-time kinetic reverse transcriptase–polymerase chain reaction (PCR) assays. In brief, internal sequence-specific probes linked to 2 fluorescent dyes (a 5′ reporter and a 3′ quencher) were added to the PCR mix before initiating amplification of the specific target sequences. The efficiency of amplification was determined by coamplification of an exogenously added unique RNA standard of known copy number prepared by in vitro transcription with DNase treatment to ensure the absence of contaminating DNA.

**Statistical analysis.** Fisher’s exact (2 × 2 tables) and χ² (N × N tables) tests were used to determine the statistical significance of differences in the distribution of categorical variables between groups. The nonparametric Wilcoxon rank sum test was used to determine the statistical significance of differences in continuous variables among the different groups. Correlation among variables was evaluated by use of Spearman’s nonparametric test. Significance was established at P < .03.

**Results**

To determine whether an array of chemokinome or chemokine receptor phenotypes correlated with markers of disease progression and/or the Δ32CCR5 genotype, we measured virus infectivity and levels of CCR5 and CXCR4 mRNA and RANTES in CD4+ and CD8+ lymphocyte cultures. For our study material, we isolated CD4+ and CD8+ lymphocytes from PBMC from patients before the extrapolated time of infection and compared our in vitro assay findings to the postinfection markers of disease progression for each corresponding donor. For the Δ32CCR5 comparison, we also incorporated a group of uninfected control subjects. Because of variations in the number of recovered activated lymphocytes, not all parameters could be studied for all of the study patients. All data points available have been included in presentations and data analysis.

**HIV-1 infectivity of CD4+ lymphocytes in vitro.** We sought to determine whether there were differences in either R5 or X4 virus replication in CD4+ lymphocytes isolated before infection. There was no association between replication of R5 or X4 viruses and plasma HIV-1 RNA levels or the slope of CD4+ lymphocyte decline (data not shown). We also found no association between infectivity profiles and the different CCR5 genotypic subgroups.

**CCR5 and CXCR4 expression levels on CD4+ lymphocytes.** Since cell surface expression profiles of CCR5 are associated with R5 virus replication levels in vitro, we measured the cell surface expression levels of CXCR4 and CCR5 on uninfected cells in vitro-[6]. We tested PHA-activated bulk PBMC and cells enriched for CD4+ lymphocytes on days 4, 7, and 10, by standard FACS analysis, using monoclonal antibodies 2G12 and 3A9. CCR5 and CXCR4 mRNA expression levels also were analyzed on PHA-activated PBMC and CD4+-enriched lymphocytes. No associations were found for cell surface expression or mRNA expression patterns with the levels of viral RNA in plasma or rates of CD4 cell decline (data not shown) or between CCR5 expression patterns and the Δ32CCR5 allele.

**RANTES levels in CD4+ and CD8+ lymphocyte cultures.** Because we showed previously that the level of RANTES in the culture supernatant of activated CD4+ lines and clones correlates with levels of MIP-1α and MIP-1β and because RANTES is the most potent inhibitor of HIV-1 replication in vitro [12], we measured the levels of this CC chemokine in the culture supernatants of activated CD4+ and CD8+ lymphocytes isolated from preinfection PBMC. We observed a wide range of values for the RANTES levels in PHA-activated CD4+ lymphocyte culture supernatants (figure 1). When we examined the RANTES levels according to the Δ32CCR5 allele, we found a statistical difference (P = .01) between patients who were heterozygous for this allele (mean, 1305 pg/mL) and CCR5 wild-type patients (mean, 790 pg/mL; figure 1).

We also compared the in vitro RANTES secretion results with the postinfection plasma viral RNA measurements and rates of CD4+ lymphocyte decline. When comparing the RANTES secretion profiles for preinfection CD4+ lymphocytes, we found a negative statistical correlation for viral RNA measurements in plasma (r = −0.4; P = .01) and a positive correlation for rates of CD4 lymphocyte decline (r = 0.5; P = .002). No statistically significant association was noted between plasma HIV-1 RNA levels or CD4 cell slopes and RANTES secretion levels from CD8+ lymphocytes (data not shown). When we separated the patients into those with high versus low levels of RANTES (as determined on the basis of the median value, 764 pg/mL), we found statistically significant dif-
Figure 1. RANTES production levels from CD4+–enriched lymphocytes isolated from noninfected CC chemokine receptor 5 (CCR5) wild-type (CCR5-wt/wt) and Δ32CCR5 heterozygous patients.

Figures between the 2 groups (figure 2). We found differences for both mean plasma HIV-1 RNA levels (mean, 56,643 and 3755 copies/mL for low and high secretors, respectively; \( P = .007 \)) and mean rates of CD4+ lymphocyte decline (mean, −109 and −4 cells/mL per year for low and high secretors, respectively; \( P = .005 \)). Of interest, when we removed the Δ32/CCR5 patients from the analysis, we still obtained statistical significance when comparing RANTES secretion levels from the CD4+ lymphocytes with both plasma viral RNA measurements and rates of CD4 cell decline, which suggests that the presence of this allele alone cannot account for the result (\( P < .01 \)).

Discussion

We showed previously that CC chemokine levels and CCR5 expression levels correlated with HIV-1 R5 virus infectivity in vitro and, therefore, wanted to determine whether these factors correlated with in vivo markers of disease progression [5, 6]. To do so, we analyzed several chemokine and chemokine receptor profiles in ex vivo–activated CD4+ and CD8+ cell populations generated from PBMC obtained 9–12 months before HIV-1 infection and compared the results with in vivo markers of disease progression after infection. From the parameters that we measured, we identified that patients with the higher CD4+ lymphocyte culture levels of RANTES before infection had lower plasma HIV-1 RNA levels and slower rates of CD4+ lymphocyte decline after infection. Higher levels of RANTES in the supernatants of cultured CD4+ lymphocytes also were observed in patients who were heterozygous for the Δ32CCR5 allele, a finding in support of our previous report [5].

We used pre- rather than postinfection samples, to eliminate the possibility that HIV-1 replication could skew the RANTES production profiles by altering the lymphocyte subsets being propagated in culture. Although it is unlikely, we cannot rule out the possibility that the exposure of these patients to HIV-1 antigen encountered during the period before the presumed time of infection in some way influenced our results. However, we observed no difference in the spectrum of RANTES secretion patterns between the patients who became infected and nonexposed control subjects (data not shown). In addition, mitogen activation of the cells with PHA is likely to have obscured any influence of HIV-1–specific lymphocyte responses.

No correlations were observed between virus infectivity profiles, cell surface CCR5/CXCR4 or CCR5/CXCR4 mRNA expression levels, and plasma HIV-1 RNA levels or CD4+ T lymphocyte decline. This may be a reflection on the duration and method of the stimulation employed in these experiments.

The results presented here support a number of studies describing an association between CC chemokine production levels from activated PBMC and clinical course [7, 8]. A previous study, strongly supported by our own findings, demonstrated that CD4+ lymphocyte clones, but not CD8+ lymphocyte

Figure 2. Plasma human immunodeficiency virus–1 RNA levels (copies/mL) 9–12 months after infection (A) and decline of CD4+ cells (cells/mL/year; B) dependent on the median RANTES production level (764 pg/mL) from CD4+–enriched lymphocytes.
clones, from long-term nonprogressing patients secreted higher levels of CC chemokines than did clones derived from rapid progressors [8]. Taken together, these studies support the concept that chemokine production from CD4+ lymphocytes can protect the lymphocytes from infection and presumably from destruction. As an added benefit, the protection of CD4+ cells by CC chemokines may allow for the maintenance of a vigorous anti–HIV-1–specific CD4+ T lymphocyte response. This may lead to better control of viral replication through the induction and maintenance of effective neutralizing antibody and cytotoxic T lymphocyte responses [13].

A recent study has reported high levels of RANTES, MIP-1α, and MIP-1β being secreted from HIV-1 antigen-specific CD8+ lymphocytes, which inversely correlates with the corresponding infected individuals’ HIV-1 load measurements [14]. Of interest, as with our own findings described here, this study showed higher cellular production levels of the CC chemokines in patients who were heterozygous for the Δ32CCR5 allele. These results suggest that a higher production of CC chemokines may be beneficial in maintaining lower levels of virus replication through the induction of a more effective anti–HIV-1–mediated immune response. Whether patients with the higher preinfection levels of CD4+ cell CC chemokine production correspond to patients who subsequently develop higher postinfection HIV-1 antigen–specific CD8+ production levels remains to be determined. Our study adds considerable weight to the phenomenon that chemokine production levels are genetically predetermined and can correlate with the control of HIV-1 replication in vivo, although via which mechanism is still unknown.

References