Increased early activation of CD56\textsuperscript{dim}CD16\textsuperscript{dim/-} natural killer cells in immunological non-responders correlates with CD4\textsuperscript{+} T-cell recovery

Qiu-Yue Zhang\textsuperscript{1,2}, Xin Zhang\textsuperscript{1,2}, Bin Su\textsuperscript{1,2}, Li-Feng Liu\textsuperscript{1,2}, Xiao-Dong Yang\textsuperscript{1,2}, Bin Tang\textsuperscript{1,2}, Huan Xia\textsuperscript{3}, Ping Ma\textsuperscript{3}, Tong Zhang\textsuperscript{1,2}, Hao Wu\textsuperscript{1,2}

\textsuperscript{1}Center for Infectious Diseases, Beijing Youan Hospital, Capital Medical University, Beijing 100069, China; \textsuperscript{2}Beijing Key Laboratory for HIV/AIDS Research, Beijing Youan Hospital, Capital Medical University, Beijing 100069, China; \textsuperscript{3}Department of Infectious Diseases, Nankai University Second People’s Hospital, Tianjin 300192, China.

Abstract

Background: Natural killer (NK) cells play a critical role in suppressing human immunodeficiency virus-1 (HIV-1) infection, but knowledge on whether and how NK cells affect immune reconstitution in HIV-1-infected individuals who receive antiretroviral therapy (ART) is limited.

Methods: We performed a case-control study with 35 healthy individuals and 66 HIV-1-infected patients including 32 immunological non-responders (INRs) with poor CD4\textsuperscript{+} T-cell recovery (<500 cells/μL after 4 years of ART) and 34 immunological responders (IRs) with improved CD4\textsuperscript{+} T-cell recovery (>500 cells/μL after 4 years of ART). NK cell phenotype, receptor repertoire, and early activation in INRs and IRs were investigated by flow cytometry.

Results: A significantly higher proportion of CD56\textsuperscript{dim}CD16\textsuperscript{dim/-} NK cells was observed in INRs than IRs before ART and after 4 years of ART. The number of CD56\textsuperscript{dim}CD16\textsuperscript{dim/-} NK cells was inversely correlated with CD4\textsuperscript{+} T-cell counts in INRs before ART ($r = -0.344$, $P = 0.050$). The more CD69-expressing NK cells there were, the lower the CD4\textsuperscript{+} T-cell counts and ACD4, and these correlations were observed in INRs after ART ($r = -0.416$, $P = 0.019$; $r = -0.509$, $P = 0.003$, respectively). Additionally, CD69-expressing CD56\textsuperscript{dim}CD16\textsuperscript{dim/-} NK cells were more abundant in INRs than those in IRs ($P = 0.018$) after ART, both of which had an inverse association trend towards significance with CD4\textsuperscript{+} T-cell counts. The expression of the activating receptors NKG2C, NKG2D, and Nkp46 on CD56\textsuperscript{dim}CD16\textsuperscript{dim/-} NK cell subsets were higher in IRs than that in INRs after 4 years of ART ($P < 0.01$). Strong inverse correlations were observed between CD69 expression and NKG2C, NKG2A-NKG2C+, NKG2D, and Nkp46 expression on CD56\textsuperscript{dim}CD16\textsuperscript{dim/-} NK cells in INRs after ART ($r = -0.491$, $P = 0.004$; $r = -0.434$, $P = 0.013$; $r = -0.405$, $P = 0.021$; $r = -0.457$, $P = 0.008$, respectively).

Conclusions: INRs had a larger number of CD56\textsuperscript{dim}CD16\textsuperscript{dim/-} NK cells characterized by higher activation levels than did IRs after ART. The increase in the CD56\textsuperscript{dim}CD16\textsuperscript{dim/-} NK cell subset may play an adverse role in immune reconstitution. Further functional studies of CD56\textsuperscript{dim}CD16\textsuperscript{dim/-} NK cells in INRs are urgently needed to inform targeted interventions to optimize immune recovery.

Keywords: HIV-1 infection; Immunological non-responders; Natural killer cells; Immune reconstitution

Introduction

Antiretroviral therapy (ART) can effectively suppress human immunodeficiency virus (HIV) replication, increase HIV-infected patients’ peripheral blood CD4\textsuperscript{+} T-cell counts, and dramatically reduce acquired immunodeficiency syndrome (AIDS)-related morbidity and mortality. However, 15% to 30% of patients fail to recover their CD4\textsuperscript{+} T-cell counts despite the complete suppression of viral replication. These individuals are identified as immunological non-responders (INRs).\textsuperscript{[1-3]} INRs exhibit severe immune dysfunction and have a higher rate of AIDS and non-AIDS events, such as metabolic syndrome, liver disease, nephropathy, cardiovascular disease, non-AIDS-related malignancies, and HIV-1-related neurocognitive disorder than complete immunological responders (IRs).\textsuperscript{[1-2,4,5]} The reason for immunological non-response is incompletely understood. It is currently believed that decreased hematopoiesis in the bone marrow and insufficient thymic and lymph node outputs lead to a reduction in the production of CD4\textsuperscript{+} T cells, and residual viral replication, sustained immune activation, and disturbed in cytokine levels can all cause CD4\textsuperscript{+} T-cell apoptosis and depletion.\textsuperscript{[6-8]} However, the current preliminary research on how dysfunction in the innate immune system affects the immune reconstitution in HIV-infected patients is limited.

Human natural killer (NK) cells provide powerful innate defenses against virus-infected cells and tumor cells. Their function is crucial at the time of infection and can impact the
The expression and functional relevance of several activating and inhibitory receptors are pathologically modified by high levels of viral replication. \[^{13,14}\]

NK cell effector function during the immune response against HIV infection has been evaluated in different cohorts: HIV controls (who maintain lower levels of HIV-1 replication in the absence of ART), slow progressors, and HIV-1-exposed seronegative individuals who remain uninfected despite repeated exposure to the virus. \[^{15,16}\] The distribution and activation of NK cell subsets are also relevant in the context of mother-to-child HIV-1 transmission, as they influence the susceptibility to perinatal infection. \[^{17}\]

The role of NK cells in immune reconstitution has not been elucidated. HIV-1 has developed various strategies of interaction with NK cell receptors and their ligands. The HIV-1 Nef protein reduces the cell-surface expression of NKG2D ligand, MICA, ULBP1, and, especially, ULBP2, thus impairing NKG2D-dependent killing by NK cells. Such aberrations limit NK cell immune surveillance and allow the development of opportunistic infections and cancers. Despite virus countermeasures, HIV-1-infected T cells express higher NKG2D levels than non-infected cells and are highly susceptible to killing by NK cells through NKG2D activation. \[^{18,20}\] The population expressing another activating receptor of NK cells, the natural cytotoxicity receptor Nkp44, the ligand of which is expressed on the surface of uninfected CD4\(^+\) T cells, is expanded, consequently rendering these T cells sensitive to NK cell-mediated killing. \[^{21-23}\] In addition to the complex effects on NK cell receptors, HIV infection also causes INR patients to have an increase in the CD56\(^{bright}\) NK cell sub-population but causes a poor response to cytokine stimulation in terms of Nkp44 upregulation and interferon (IFN)-γ production, and CD56\(^{bright}\) NK cells from INRs show relatively high cytotoxicity against autologous activated CD4\(^+\) T cells. \[^{24}\] Hence, all of these mechanisms could be involved in CD4\(^+\) T-cell depletion in HIV-1 infection and in impaired immunological recovery in ART-treated patients. In addition, HIV causes upregulation of the expression of inhibitory NK receptors, leading to impairments in NK cell-mediated lysis of virally infected cells. \[^{25}\] Notably, reduced expression/function of natural cytotoxicity receptors can also be detected in NK cells from HIV-infected patients. \[^{14}\] ART reverses the effects of HIV infection on NK cells; however, there is no consensus on the degree to which the suppression of HIV replication restores the NK cell subset distribution and function. \[^{26-28}\] Several changes in the NK cell receptor repertoire and functionality are observed during HIV infection; some of these changes have been related to progression, but others are still under study. These complex and possible changes may represent one of the pathogenic events associated with the drastic reduction in CD4\(^+\) T-cell counts.

To better understand the role of NK cells in INRs, in this study, we focused on the changes in the NK cell phenotype and receptor repertoire in a cohort of INRs compared to IRs and health controls (HCs). Our results showed that the CD56\(^{dim}\)CD16\(^{dim}\) subset was expanded in the INR group compared to the IR group both before and after ART. CD56\(^{dim}\)CD16\(^{dim}\) NK cells exhibited a relatively high degree of early activation even though the suppression of viral replication by ART is negatively correlated with CD4\(^+\) T-cell counts. The expansion of the CD56\(^{dim}\)CD16\(^{dim}\) subset may play a diverse role in immune reconstitution.

### Methods

#### Ethical statement

All the participants came from the Beijing Primo Clinical Cohort, which ran from 2011 to 2012. They provided written informed consent for participation in the study. This study and other related experiments were approved by the Beijing Youan Hospital Research Ethics Committee (No. 2017-13), and informed written consent was obtained in accordance with the Declaration of Helsinki. The study was carried out in accordance with approved guidelines and regulations.

#### Study participants

Sixty-six HIV-1-infected patients from the Primo cohort of men who have sex with men that were collected at the Beijing Youan Hospital between 2011 and 2012 were enrolled: 32 INRs with poor CD4\(^+\) T-cell recovery (<500 cells/μL after four years of ART) and 34 IRs with immunological restoration (>500 cells/μL after four years of ART). We also included 35 HCs with ages similar to the patients. All participants were divided into 5 groups: HCs, INRs before ART (INR pre-ART), INRs after ART (INR-ART), IRs before ART (IR pre-ART), and IRs after ART (IR-ART). All subjects without ongoing co-infections or other diseases and their characteristics are summarized in Table 1.

#### Cell-surface staining and flow cytometry analysis

Peripheral blood mononuclear cells (PBMCs) were thawed in a water bath at 37°C for one minute, washed and resuspended in Roswell Park Memorial Institute (RPMI) medium containing 10% v/v fetal calf serum. Antibodies
Parameters | Healthy controls | INRs | IRs | INRs vs IRs |
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Number of subjects | 35 | 32 | 34 | NS |
Mean age (years) | 32 (26–53) | 34 (21–57) | 33 (23–55) | NS |
CD4+ T-cell counts (cells/μL) | | | | |
Pre-ART | NA | 207.00 (100–313) | 227.00 (130–313) | 0.021 |
ART | NA | 380.00 (163–497) | 680.00 (505–1240) | <0.0001 |
CD8+ T-cell counts (cells/μL) | | | | |
Pre-ART | NA | 664.50 (255–1716) | 766.50 (331–1928) | NS |
ART | NA | 570.00 (229–1192) | 919.00 (515–1909) | <0.0001 |
CD4/CD8 ratio | | | | |
Pre-ART | NA | 0.25 (0.12–0.76) | 0.28 (0.11–0.67) | NS |
ART | NA | 0.62 (0.34–1.31) | 0.70 (0.45–1.36) | NS |
Viral load (log10, copies/mL) | | | | |
Pre-ART | NA | 4.41 (1.97–5.26) | 4.06 (1.70–5.97) | NS |
ART | NA | ND | ND | |

Data are presented as n or median (range). \(p\) values were determined by Mann-Whitney U two-tailed \(t\)-test. NA: Not applicable; NS: Not significant; ND: Not detected.

Routine blood CD4+ T-cell counts (cells/μL) were measured by four-color flow cytometry using the human CD45, CD3, CD4, and CD8 cell markers (BD Biosciences) in whole peripheral blood samples from each patient processed with a fluorescence activated cell sorter (FACS) lysing solution (BD Biosciences) according to the manufacturer’s instructions. The plasma HIV-1 viral load (copies per milliliter of plasma) was quantified by real-time PCR (Abbott, Des Plaines, IL, USA). The sensitivity of viral RNA detection using this assay is 40 copies/mL of plasma (BD Biosciences).

Statistical analysis
Statistical analyses were performed using GraphPad Prism 6.0 software (GraphPad Software, Inc., La Jolla, CA, USA). The statistical significance between the two groups was calculated using the Mann-Whitney U test, and correlations were determined by the Spearman rank correlation test, with \(r\) being the Spearman correlation coefficient. A \(p\) value < 0.05 was considered statistically significant.

Results
Demographic and clinical characteristics of the subjects enrolled in this study
We enrolled chronically HIV-1-infected patients undergoing ART for 4 years who were divided into two groups based on CD4+ T-cell recovery: a group of 32 INRs with <500 CD4+ T cells/μL and a group of 32 IRs with >500 CD4+ T cells/μL. Thirty-five age-matched HCs were also included in this study. The CD4+ T-cell counts of the IRs were higher than those of the INRs \((P = 0.021\) before ART; \(P < 0.0001\) after ART). The CD8+ T-cell counts of the INRs were similar to those of the IRs before ART, but at the time of analysis after ART, the CD8+ T-cell counts in the IR group were higher than those in the INR group \((P < 0.0001)\). There was no difference in the ratio of CD4/CD8 between the two groups before or after ART. The plasma HIV-1 viral load did not differ significantly between the IR and INR groups before ART. After 4 years of ART, all HIV-infected patients achieved viral suppression, and the plasma HIV-1 viral load could not be detected. No differences in demographics or treatment duration were present between the groups [Table 1].

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INR patients have an increased proportion of CD56dimCD16dim^− NK cells

PBMCs were analyzed by flow cytometry with initial gating on CD3^− peripheral blood lymphocytes (PBLs) and subsequent analysis based on CD56 and CD16 expression levels, identifying NK cell subsets as CD56^brightCD16^dim^−, CD56^dimCD16^dim^−, CD56^dimCD16^bright, and CD56^−CD16^bright (Figure 1A, presented in gates P1, P2, P3, and P4, respectively).

Figure 1: The CD56dimCD16dim^− NK cell subset is enlarged in INR patients. (A) The gating strategy used to identify NK cell subsets by flow cytometry and the frequencies of gated populations are shown for a representative HC. Lymphocytes were gated according to forward and side scatter dot plots. Single cells were gated according to forward height and side scatter forward area. Dead cells were excluded by staining with Live/Dead fixable viability stain 510. NK cells were defined within the CD3-gate on the basis of the expression of CD16 and CD56. NK cells were divided into four subsets: CD56^brightCD16^dim^− (P1), CD56^dimCD16^dim^− (P2), CD56^dimCD16^bright (P3) and CD56^−CD16^bright (P4). (B) Frequencies of NK cells derived from peripheral blood lymphocytes. (C) Percentages of the different natural killer (NK) cell subsets relative to the total NK cell population (100%). The frequency of CD56^brightCD16^dim^− cells in the total NK cell population was higher in INRs than that in IRs and HCs. (D) and (E) Correlation between CD56dimCD16dim^− NK cell subset with CD4^+ T-cell counts and viral load in INR pre-ART group. The Mann-Whitney U test was used for statistical analysis. Horizontal lines indicate mean values, and error bars represent the mean ± standard error of mean. Correlations between two variables were analyzed in nonparametric Spearman’s rank correlation tests, and p<0.05 was considered statistically significant. HCs: Health controls; INRs: Immunological non-responders; IR: Immunological responder; NK: Natural Killer; PBL: Peripheral blood lymphocytes; VL: Viral load.
To further delineate the potential role of NK cells in immune reconstitution, we analyzed their proportions according to the immune reconstitution status. In our study, before ART, the frequency of NK cells among PBLs in INRs was lower than that among HCs (Figure 1B, INR pre-ART vs. HC, P = 0.029), but it was restored by ART. No differences were observed in the frequency of NK cells between HCs and IRs.

Significant differences existed in the CD56<sup>dim</sup>CD16<sup>bright</sup> and CD56<sup>dim</sup>CD16<sup>dim</sup> subsets between groups, but there were no differences in the CD56<sup>bright</sup>CD16<sup>dim</sup>- and CD56<sup>dim</sup>CD16<sup>bright</sup> subsets [Figure 1C]. Interestingly, both before ART and after ART, the frequency of CD56<sup>dim</sup>CD16<sup>dim</sup>- NK cells was significantly higher in INRs than that in IRs (INR pre-ART vs. IR pre-ART, P = 0.019; INR-ART vs. IR-ART, P = 0.011). Compared to that in the HC group, the frequency of CD56<sup>dim</sup>CD16<sup>dim</sup>- NK cells in both the IR and INR groups was increased with HIV-1 infection (INR pre-ART vs. HC, P = 0.003; INR-ART vs. HC, P < 0.001; IR-ART vs. HC, P = 0.019). With the prolongation of HIV infection, the frequency of CD56<sup>dim</sup>CD16<sup>dim</sup>- NK cells was increased and not restored by ART (INR pre-ART vs. INR-ART, P = 0.008; IR pre-ART vs. IR-ART, P = 0.02). The CD56<sup>dim</sup>CD16<sup>bright</sup> subset in INRs was lower than that in HCs before ART (INR pre-ART vs. HC, P = 0.006). After 4 years of ART, the CD56<sup>dim</sup>CD16<sup>bright</sup> subset in both INRs and IRs was lower than that in HCs (INR-ART vs. HC, P = 0.001; IR-ART vs. HC, P = 0.006), but there was no difference between the INRs and IRs. These results implied that NK cell subsets were only partially recovered after ART. Of note, in the INR pre-ART group, correlation analysis revealed that the increase in the CD56<sup>dim</sup>CD16<sup>dim</sup>- NK cell subset was inversely correlated with CD<sup>+</sup> T-cell counts (r = -0.344, P = 0.050, Figure 1D) and also inversely correlated with viral load (r = -0.4495, P = 0.010, Figure 1E). We did not find any significant correlations in other groups.

**INRs with an increase in the CD56<sup>dim</sup>CD16<sup>dim</sup>- NK cell subset display higher immune activation**

NK cell activation was confirmed by measuring CD69 upregulation in this study. We found that CD69 expression in the overall NK cell population was higher in HIV-infected patients than that in HCs (INR pre-ART vs. HC, P < 0.0001; INR-ART vs. HC, P = 0.012; IR pre-ART vs. HC, P = 0.001; Figure 2A). However, if HIV-infected patients were stratified into IRs and INRs, the INRs had a significantly higher frequency of CD69 expression than the IRs after ART and HCs (INR-ART vs. IR-ART, P = 0.038; Figure 2A). In the CD56<sup>dim</sup>CD16<sup>dim</sup>- NK cell subset, the frequency of CD69 expression in the INR-ART group was still significantly higher than that in the IR-ART and HC groups (INR-ART vs. IR-ART, P = 0.018; INR-ART vs. HC, P = 0.003; Figure 2E). These results suggest that the CD56<sup>dim</sup>CD16<sup>dim</sup>- NK cell subset is activated.

**Early activation of NK cells was negatively related to CD4<sup>+</sup> T-cell counts**

The immune activation of NK cells is responsible for the poor immune reconstitution of INRs. To further determine the role of NK cell activation in HIV infection with different degrees of immune restoration after long-term ART and viral suppression, we analyzed the correlation between the expression of CD69 on NK cells and CD4<sup>+</sup> T-cell counts. In the current study, there was a significantly negative association between the percentage of CD69<sup>+</sup> NK cells and peripheral CD4<sup>+</sup> T-cell counts in the INR-ART group (r = -0.416, P = 0.019; Figure 2B). The number of CD4<sup>+</sup> T cells was increased after 4 years of ART (ΔCD4) among INRs and negatively correlated with the frequency of CD69<sup>+</sup> NK cells (r = -0.509, P = 0.003; Figure 2D). However, a trend toward a negative relationship between the percentage of CD69<sup>+</sup> NK cells and CD4<sup>+</sup> T-cell counts, ΔCD4, was observed in the IR-ART group, but this relationship was not statistically significant (r = -0.054, P = 0.762; r = -0.098, P = 0.578; Figures 2C and 2E).

The correlation between the percentages of CD69<sup>+</sup> expressing CD56<sup>dim</sup>CD16<sup>dim</sup>- NK cells and CD4<sup>+</sup> T-cell counts in HIV-infected patients tended to negative; however, this correlation was not statistically significant [Figures 2G and 2H]. Notably, ΔCD4 in the INR-ART group was significantly inversely correlated with the percentages of CD69<sup>+</sup> expressing CD56<sup>dim</sup>CD16<sup>dim</sup>- NK cells (r = -0.363, P = 0.034), but this relationship was not seen in the INR group [Figures 2I and 2J]. Thus, a strong CD56<sup>dim</sup>CD16<sup>dim</sup>- NK cell activation capacity may have an adverse effect on INR CD4<sup>+</sup> T-cell recovery after ART.

The frequency of CD56<sup>dim</sup>CD16<sup>dim</sup>- NK cells in INRs was positively correlated with the early activation of NK cells

To further determine the correlation of the relatively high frequency of CD56<sup>dim</sup>CD16<sup>dim</sup>- NK cells to NK cell activation, we analyzed the correlation between the frequencies of CD69<sup>+</sup> NK cells and CD56<sup>dim</sup>CD16<sup>dim</sup>- NK cells. In the current study, the frequency of CD56<sup>dim</sup>CD16<sup>dim</sup>- NK cells was positively correlated with the early activation of NK cells in both INRs and IRs before and after ART (r = 0.360, P = 0.042, INR pre-ART; r = 0.381, P = 0.034, INR-ART; r = 0.602, P = 0.0002, IR pre-ART; r = 0.401, P = 0.018, IR-ART; Figures 3A–3D). These findings suggest that a higher level of CD69 expression on NK cells is correlated with a higher frequency of CD56<sup>dim</sup>CD16<sup>dim</sup>- NK cells in HIV-infected patients. This also further confirms the negative role of CD56<sup>dim</sup>CD16<sup>dim</sup>- NK cells in immune reconstitution.

**Expression of NK activating receptors (NKG2C, NKG2D, and Nkp46) on the CD56<sup>dim</sup>CD16<sup>dim</sup>- NK subset was higher in IRs than that in INRs after ART**

NK cell functionality is coordinated by a dynamic balance between inhibitory and activating signals upon encountering a target cell. Previous data have shown that abnormal expression of NK activating and inhibitory receptors is associated with impaired NK cell function. We focused on the receptor spectrum in CD56<sup>dim</sup>CD16<sup>dim</sup>- NK cells, including the inhibitory receptor NKG2A and activating receptors NKG2C, NKG2D, Nkp30, Nkp44, and Nkp46 (Figure 4).

First, we analyzed receptor expression on total NK cells. Before ART, the frequency of NKG2D expression on NK cells was lower in INRs than IRs (INR pre-ART vs. IR pre-ART, P = 0.003; INR-ART vs. IR-ART, P = 0.034). However, after ART, the frequency of NKG2D expression on NK cells was lower in INRs than IRs (INR-ART vs. IR-ART, P = 0.042; Figures 3A and 3B). These findings suggest that a lower level of NKG2D expression on NK cells is correlated with a lower frequency of CD56<sup>dim</sup>CD16<sup>dim</sup>- NK cells in HIV-infected patients. This also further confirms the negative role of CD56<sup>dim</sup>CD16<sup>dim</sup>- NK cells in immune reconstitution.

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pre-ART, \( P = 0.005 \); Figure 4A), and NKp30 expression on NK cells was higher in INRs (INR pre-ART vs. IR pre-ART, \( P = 0.003 \); Figure 4B). However, these differences disappeared after ART. Compared to that in HCs, the expression of NKp46 in IRs and INRs was lower, and this gap actually increased after ART (INR-ART vs. HCs, \( P = 0.007 \); IR pre-ART vs. HC, \( P = 0.002 \); IR-ART vs. HC, \( P < 0.0001 \); Figure 4B). The expression levels of NKp44 on the NK cells of HIV-1-infected patients were very low [Figure 4B] if not undetectable without in vitro
stimulation. There was no difference in NKG2A, NKG2C, and NKp44 expression on NK cells among the groups.

As shown in Figures 4C and 4D, we further investigated and analyzed the expression of NK cell receptors in the CD56<sup>dim</sup>CD16<sup>dim</sup>−/− subset. A comparative evaluation of the phenotype of NK cells among INRs and IRs was performed. Before ART, the frequency of NKG2D-expressing CD56<sup>dim</sup>CD16<sup>dim</sup>− NK cells was lower in the INRs (INR pre-ART vs. IR pre-ART, P = 0.013; Figure 4C), and Nkp30 expression was higher in the INRs (INR pre-ART vs. IR pre-ART, P = 0.012; Figure 4D). After ART, the difference in Nkp30 expression disappeared. However, the levels of NKG2C, NKG2D, and NKp46, the activating receptor of NK cells expressed in the CD56<sup>dim</sup>CD16<sup>dim</sup>− subset, were all higher in the IR-ART group (INR-ART vs. IR-ART, P = 0.016, P = 0.002, and P = 0.025, respectively; Figures 4C and 4D). In addition, these activating receptor levels were all higher in HIV-infected patients after ART, both INRs and IRs, than those in HCs (INR-ART vs. HC, P < 0.0001 and P = 0.001; IR-ART vs. HC, P < 0.0001, P < 0.0001, and P < 0.0001, respectively; Figures 4C and 4D).

The expression of the inhibitory receptor NKG2A was lower in HIV-infected patients than that in HCs before ART (INR pre-ART vs. HC, P = 0.012; Figure 4C), and as the infection was sustained, the expression of NKG2A also increased despite after four years ART (IR-ART vs. HC, P < 0.0001; Figure 4C). There was no difference between INRs and IRs.

The activating receptor NKG2C and the inhibitory receptor NKG2A recognize the same ligand, HLA-E, a non-classical HLA molecule characterized by limited polymorphism. It has been reported that the proportion of NKG2A/NKG2C<sup>+</sup> NK cells is a potential biomarker for predicting HIV disease progression. In our results, NKG2A/NKG2C<sup>+</sup> NK cells was increased in the IR-ART group compared to the HC group (INR-ART vs. HC, P = 0.049; Figure 4E). The frequency of NKG2A/NKG2C<sup>+</sup> CD56<sup>dim</sup>CD16<sup>dim</sup>− NK cells was higher in IR-ART than that in INR-ART (INR-ART vs. IR-ART, P = 0.03; Figure 4F). Compared to HCs,
INR-ART, and IR-ART showed increased levels of NKG2A-NKG2C+ CD56dimCD16dim NK cells; however, were also increased in INR-ART and IR-ART, although there were no differences between HIV-infected patients and HCs at baseline (INR-ART vs. HC, P < 0.0001; IR-ART vs. HC, P < 0.0001; Figure 4F).

In summary, the expression of NK activating receptors (NKG2C, NKG2A-NKG2C+, NKG2D, and Nkp46) on the CD56dimCD16dim NK subset was higher in IRs than that in INRs after 4 years of ART. Although we did not perform further analysis of NK cell inhibitory receptors due to logistical limitations, in the current study, the higher...
activating receptor expression may be reflective of the higher cytotoxicity against HIV infection and better immune reconstitution among IRs.

**Correlations between CD69 and NKG2C, NKG2A-NKG2C+, NKG2D, and Nkp46 expression on CD56dimCD16dim/- NK cells**

Next, we analyzed the correlations between activation and functional markers on NK cells in HCs and HIV-infected subjects. Interestingly, we found significant negative correlations between NKG2C, NKG2A-NKG2C+, NKG2D, and Nkp46 expression and CD69 expression on CD56dimCD16dim/- NK cells in the INR-ART group (r = -0.491, P = 0.004; r = -0.434, P = 0.013; r = -0.405, P = 0.021; r = -0.457, P = 0.008, respectively; Figures 5A-5D). In the IR-ART group, NKG2C, NKG2A NKG2C+, NKG2D, and Nkp46 expression and CD69 expression on CD56dimCD16dim/- NK cells but not Nkp46 expression was significantly negatively correlated with CD69-expressing CD56dimCD16dim/- NK cells (r = -0.416, P = 0.017; r = -0.424, P = 0.015; r = -0.435, P = 0.010, respectively; Figures 5E-5H). This evidence supports that increased activating receptor (Nkp46, Nkp30, and NKG2D) expression on the CD56dimCD16dim/- subset is accompanied by decreased CD69 expression. Altogether, this pattern indicates a relatively favorable immune reconstitution environment in IR patients. These results also suggest that NK cells may be activated to express these activation and functional markers by different mechanisms in HIV-infected patients.

**Discussion**

This study was performed to investigate the role of NK cells in the immune reconstitution of HIV-1-infected individuals. In the expansion of CD56dimCD16bright NK cells, the frequency of CD56dimCD16bright NK cells was higher than that in IRs both before ART and after four years of ART. The increase in CD56dimCD16bright NK cells was inversely related to CD4+ T-cell counts in the INR pre-ART group. In addition, INRs displayed higher levels of early activation than IRs after ART, and the early activation of NK cells was negatively correlated with CD4+ T-cell counts in HIV-infected individuals. Furthermore, CD56dimCD16dim NK cells were positively correlated with the early activation of NK cells. Accordingly, CD56dimCD16dim NK cells may be involved in the poor immune reconstitution of INRs.

It has been demonstrated that in the acute phase of HIV-1 infection, NK cells expand in the peripheral blood to inhibit HIV replication and can mediate in vivo immune pressure in infected individuals, resulting in the viral escape.[16] NK cell features are affected by high and chronic viremia, including the frequencies, phenotypes, and functions.[17] The number of circulating NK cells appears to be restored as soon as HIV-1 infection enters its chronic stage, but the distribution of the different subsets undergoes pathological redistribution.[18] Several studies have shown that HIV-1 pathologically changes NK cell homeostasis and hampers NK cell antiviral effector functions. These pathogenic events are associated with a pathological redistribution of NK cell subsets that includes the expansion of anergic CD56 CD16bright NK cells with an aberrant repertoire of activating and inhibitory receptors.[19] Recently, the existence of a population of CD56dimCD16dim NK cells was detected but found to be globally reduced in a longitudinal cohort of HIV-1-infected individuals. On CD16 versus CD56 flow cytometry dot plots of fresh human PBLs, the five usual NK cell subpopulations appeared: CD56bright/CD16+, CD56bright/CD16dim, CD56dim/CD16+, CD56dim/CD16dim, and CD56dimCD16bright. From a functional point of view, in HCs and HIV-1-infected patients, the most proficient degranulating subset was CD56dimCD16+ cells and sorted CD56dimCD16dim cells degranulated more than CD56dimCD16bright cells but less than CD56dimCD16+ NK cells. The CD56dimCD16+ population was similarly the subset that most effectively produced IFN-γ, followed by the CD56dimCD16+ subset.[40] Therefore, it can be speculated that the CD56dimCD16dim subset has strong cytotoxicity and the ability to produce IFN-γ. Due to the clinical interest in NK cells, it is therefore highly relevant to precisely identify NK cell populations with specific or pronounced functions. In particular, multiparametric flow cytometry allows in-depth investigation of human immune cells, and clear, univocal identification on the basis of phenotypic traits determined using flow cytometry can help us understand the role of NK cells in disease.[41]

Our data are consistent with previous reports indicating that CD56dimCD16bright NK cells are the largest population of NK cells in PBLs and that there is a decrease in the cytotoxic CD56dimCD16bright NK cell level despite four years of ART.[18,42,43] CD56brightCD16dim/- NK cells are described as the progenitors of CD56dimCD16bright NK cells. In an African adult HIV treatment cohort, researchers found high proinflammatory CD56brightCD16dim- NK cell levels among sub-optimal IRs despite four years of suppressive ART.[44] However, we did not find a difference in the CD56brightCD16dim/- subset between INRs and IRs.

CD56dimCD16bright cells sequentially progress from an immature population via characteristic loss of inhibitory receptors, such as NKG2A, and gain of CD16, KIR, and CD57 receptors.[45] CD56dimCD16bright NK cells also lose expression of CD16 through metalloprotease-mediated shedding and become CD56dimCD16+, and the latter population demonstrates a higher degranulation ability.[46] Under these conditions, CD56dimCD16dim- NK cells are likely a mixture of relatively immature cells and previously CD56dimCD16bright cells that are on their way to losing CD16 expression due to activation-induced shedding. Recently, a study from France confirmed that the NK cell maturation status of primary infected patients should be considered a relevant marker of an immune process contributing to the early outcome of ART that could help in the management of HIV-infected patients. In that study, HIV-infected individuals were grouped into those with predominantly immature/early differentiated NK cells and those with predominantly mature NK cells. In conclusion, a better early response to ART is observed in patients whose NK cell profile is skewed to maturation at inclusion.[47] According to Mathieu’s investigation, CD56dimCD16+ NK cells may be an intermediate stage between CD56dimCD16dim- and CD56dimCD16bright NK cells.[40] In this study, the higher proportion of immature/early differentiated CD56dimCD16dim- NK cells in INRs might contribute to a poor response to ART and be
disadvantageous for CD4+ T-cell recovery. Phenotypic immaturity does not necessarily correlate with the absence of functionality, and it has been demonstrated that sorted CD56dimCD16dim/- cells more effectively degranulate and produce IFN-γ than CD56dimCD16bright cells. Previous studies have confirmed that CD56brightCD16dim/- NK cells from INRs show relatively high cytotoxicity against autologous activated CD4+ T cells. **Figure 5:** Correlations between CD69 expression and NKG2C, NKGA/NKG2C*, NKG2D, and NKp46 expression on CD56dimCD16dim/- NK cells. (A) NKG2C in INR-ART, (B) NKG2A/NKG2C* in INR-ART, (C) NKG2D in INR-ART, (D) NKp46 in INR-ART, (E) NKG2C in IR-ART, (F) NKGA/NKG2C* in IR-ART, (G) NKG2D in IR-ART, and (H) NKp46 in IR-ART. Correlations between two variables were analyzed with non-parametric Spearman rank correlation tests, and $P < 0.05$ was considered statistically significant. ART: Antiretroviral therapy; INR: Immunological non-responder; IR: Immunological responder; NK: Natural killer.
speculate that the CD56dimCD16dim NK sub-group plays a detrimental role in the recovery of CD4+ T cells.

The pathogenic and clinical relevance of this persistent NK cell activation during the course of immune recovery is still unclear. NK cell activation has been correlated with microbial translocation, a factor associated with poor CD4+ T-cell restoration after long-term virus-suppressive ART. Increased NK cell activation in INRs is inversely correlated with CD4+ T-cell recovery after ART. In this study, we analyzed early NK cell activation by monitoring the expression of CD69. Increased NK cell activation was found in HIV-infected individuals. The percentage of CD69-expressing NK cells among INR-ART was inversely correlated with CD4+ T-cell counts. These findings reveal a novel mechanism by which the alterations in NK cell populations may play an adverse role in CD4+ T-cell recovery in INRs. Further analysis of the function of the CD56dimCD16dim subset in INRs is urgently needed to inform targeted interventions to optimize immune recovery.

Due to different functional and migratory behaviors, the phenotypes of the various NK cell sub-populations in terms of repertoires of activating and inhibitory receptors are not the same. In a study on poor CD4+ T-cell reconstitution evaluating the cell-surface expression of CD56brightCD16dim NK cell-activating receptors (NKP30, NKP44, NKP46, and NKG2D), only slightly higher NKP46 expression was observed in INRs than IRs. However, we did not find a difference in the expression of NKP46 or NKG2A, NKG2C, NKG2D, NKP30, and NKP44 on total NK cells between INRs and IRs after ART. With regard to the CD56dimCD16dim subset, we were surprised to find that NK cell functional marker (NKG2C, NKG2A/NKG2A*, NKG2D, and NKP46) expression in the ART group was higher than that in the INR-ART group. The increased expression of activating receptors may be reflective of the comprehensive functions against HIV infection and better immune reconstitution in INRs. In addition, we found a negative correlation between NKG2C, NKG2A/NKG2A*, NKG2D, and NKP46 expression and CD69 expression on CD56dimCD16dim NK cells in the INR-ART and IR-ART groups. Consistently, the sustained activation of the CD56dimCD16dim subset in INRs might play an inhibitory role in immune reconstitution in this patient population.

Our results imply that after four years of suppressive ART, HIV-associated NK cell dysfunction is only partially rescued, with a high CD56dimCD16dim subset characterized by relatively high activation in INRs both before and after ART. The increase in the CD56dimCD16dim NK cell level was inversely related to CD4+ T-cell counts in the INR pre-ART group. Increased CD56dimCD16dim NK cell activation in INRs was also inversely correlated with CD4+ T cells. These findings reveal a novel mechanism by which the alterations in NK cell populations may play an adverse role in CD4+ T-cell recovery in INRs. Further analysis of the function of the CD56dimCD16dim subset in INRs is urgently needed to inform targeted interventions to optimize immune recovery.

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Conflicts of interest

None.

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