Expression levels of P-glycoprotein in peripheral blood CD8+ T lymphocytes from HIV-1-infected patients on antiretroviral therapy

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Abstract. In this study, we aimed to measure P-glycoprotein (P-gp) expression in CD8+ T lymphocytes of HIV-1-infected patients, to investigate how P-gp levels are affected by antiretroviral therapy (ART) in HIV-1 infection, and to assess the value of using P-gp expression to predict virologic response to ART. Peripheral blood mononuclear cells (PBMCs) were obtained from a cohort of HIV-1-infected patients in China: 140 patients on ART, and 49 ART-naïve patients. We also enrolled 24 healthy blood donors as the controls. The expression levels of P-gp in CD8+ T cells of HIV-1-infected patients were evaluated by quantitative reverse transcription PCR, ELISA and flow cytometry. A high inter-individual variability was observed in the CD8+ T cells of both HIV-1-infected patients and healthy donors; however, the expression levels of P-gp were significantly higher in the HIV-1-infected group compared to the ART-naïve group. The relative proportion of P-gp+CD8+ T cells inversely correlated with the blood CD4+ T cell count in the HIV-1-infected patients on ART (r=-0.3343, P=0.0375). Groups of both good and poor responders showed significantly elevated levels of P-gp+CD8+ T cells. The percentage of P-gp+CD8+ T cells appeared to provide a sensitive estimate of the virologic response to ART compared to the CD4+ T cell count. Our results suggest that P-gp expression varies among HIV-1-infected patients, but is significantly higher in HIV-1-infected patients on ART. The overexpression of P-gp is involved in ART initiation during HIV-1 infection, and P-gp+CD8+ T cells may be an additional criterion for the evaluation of the antiretroviral response to ART.

Introduction

P-glycoprotein (P-gp), the product of the multidrug resistance protein 1 (MDR1) (or ABCB1) gene in humans, is one of the most important and interesting members of the numerous, highly conserved, widespread membrane transporters of the ATP-binding cassette superfamily (1,2). P-gp has been found to be constitutively expressed in epithelial cells of tissues with excretory function, such as the intestine, liver, kidneys, lungs, the blood-brain barrier and placenta, as well as in blood cells such as lymphocytes, where P-gp expression may limit tissue penetration of its substrates (3). The role of P-gp expression and function in cells of the immune system in the treatment of HIV-1 infection has become a focus of research in recent years. P-gp-expressing cells, such as CD4+ lymphocytes, the primary target of HIV-1 infection, show limited intracellular accumulation of antiretroviral drugs (4,5). Therefore, the affinity of antiretroviral drugs to P-gp and the expression level of P-gp and other transporters in lymphocyte subpopulations may be contributing factors in the response to antiretroviral therapy (ART).

Although preliminary results have shown that P-gp is expressed in peripheral blood CD4+ cells, there has been controversy whether antiretroviral drugs inhibit P-gp, while studies on P-gp expression in peripheral blood CD8+ cells of HIV-1-infected patients receiving antiretroviral regimens are limited. It is possible that the in vivo interaction of ART drugs with P-gp may reduce the intracellular drug concentrations, resulting in the insufficient suppression of HIV-1 replication, providing additional diagnostic evidence for the virologic response to

Key words: P-glycoprotein expression, HIV-1, CD8+ T cells, drug resistance
ART during HIV-1 infection (6,7). Therefore, this study was carried out to measure the expression of P-gp in peripheral CD8+ T lymphocytes in HIV-1-infected patients, to investigate how the levels of P-gp are affected by ART in HIV-1 infection, and to assess the value of using P-gp expression to predict the virologic response to ART in a cohort of HIV-1-infected patients in China.

Materials and methods

Study subjects. Ninety-two HIV-1-infected patients on ART [58 patients treated with zidovudine (AZT) + lamivudine (3TC) + efavirenz (EFV) and 34 patients treated with AZT + 3TC + nevirapine (NVP)] and 49 HIV-1-infected ART-naïve patients who were consecutively treated at Lanzhou General Hospital (Lanzhou, China) and Tangdu General Hospital (Xi’an, China) participated in this study to compare the effects of ART on the levels of P-gp in CD8+ and CD4+ T cells. The patients were monitored with blood tests at each clinical visit, scheduled at baseline and every 3 months after the initiation of therapy.

We selected another 48 HIV-1-positive patients for which virologic and immunological data were available from baseline to 24 months after therapy, in order to assess the value of P-gp in predicting virologic response to ART. This group (26 patients treated with AZT + 3TC + EFV and 22 patients treated with AZT + 3TC + NVP), hereafter termed as the extreme group, was divided into 2 subgroups: good virologic responders were defined as patients who maintained a viral load below the detection limit of the assay (<40 copies/ml) at 6 months of therapy and at 12 months of follow-up. Poor virologic responders were defined as patients who had detectable viral loads (>40 copies/ml) at 6 and 12 months. For the patients in each group, we analyzed changes in CD4+ T cell count and viral load every 3 months during the follow-up and until 24 months.

As the control group for assessing the normal levels of measured cells, we included 24 healthy blood donors who were recruited at the same hospital. The local ethics committee approved the study, and all participants, after having been fully informed of the study procedures, provided written informed consent prior to enrollment.

Measurement of blood CD4+ T cell counts. The T lymphocyte subsets of all patients were measured by a 4-colour FACSCalibur cytometer (BD Biosciences, San Jose, CA, USA). The absolute CD4+ T cell number was calculated from the product of white cell counts. All measurements were made using the same flow cytometer and sample preparation over the duration of the study.

Virologic assessment. Batch viral load measurements were performed at the end of the study period on plasma samples which were stored at -80°C. The COBAS® TaqMan 48 Analyzer (Roche Diagnostics, Basel, Switzerland) was used for the detection and quantification of HIV RNA in the plasma, following the manufacturer’s instructions. RNA was extracted from 500 µl of plasma, following a generic nucleic acid isolation protocol, based on the adsorption of RNA molecules on a glass fiber and elution with an aqueous solution. We calculated the HIV-1 RNA titer in the tested samples based on the HIV-1 signal, the signal of the HIV-1 quantification standard and the lot-specific calibration constants provided with the COBAS TaqMan HIV-1 test. Reported values for the upper limit of quantification (ULQ) and the lower limit of quantification (LLQ) were 10,000,000 (7xlog10) and 40 (1.6xlog10) copies/ml, respectively.

Isolation of peripheral blood mononuclear cells (PBMCs). PBMCs were isolated from venous blood samples and separated with centrifugation on a Ficoll-Hypaque (Sigma-Aldrich, St. Louis, MO, USA) density gradient. Following isolation, the PBMCs were immediately cryopreserved in RPMI-1640 medium supplemented with 10% FBS (both from Gibco-Invitrogen, Carlsbad, CA, USA) and 10% dimethyl sulfoxide (Merck, Amsterdam, The Netherlands) and stored in a liquid nitrogen tank. The frozen PBMCs were thawed with a step-by-step, gradual dilution method, as previously described (8). Cell viability was >90%, as assessed by a trypan blue exclusion assay.

Quantification of MDR1 mRNA expression by quantitative reverse transcription PCR (qRT-PCR). Total RNA from 3x10^6 PBMCs was extracted and purified using an RNeasy Mini Kit (Qiagen, Hilden, Germany) according to the instructions provided by the manufacturer. For first-strand cDNA synthesis, 0.5 µg of total RNA and 1 µl of oligo(dT)18 primer were incubated with 2 µg of a 10-nM dNTP mix, 20 U RNase inhibitor, 4 µg of 5X reaction buffer and 200 U M-MuLV reverse transcriptase (RT) (Revert Aid First Strand cDNA Synthesis kit; Fermentas, Hanover, MD, USA), in a total volume of 20 µl, for 1 h at 42°C. Negative controls were obtained by replacing the enzyme with water.

Quantitative PCR was performed on a LightCycler® instrument (Roche Diagnostics, Meylan, France) using the SYBR Premix Ex Taq II detection kit (Takara Bio Inc., Otsu, Japan). PCR was performed in a 20 µl of solution, consisting of 2.0 µl of the cDNA sample, 10.0 µl of 2X SYBR Premix Ex Taq II buffer (containing Taq DNA polymerase, reaction buffer and deoxynucleotide triphosphate mixture), 6.4 µl of nuclease-free water and 0.8 µl of each primer (10 µM). The primers for the amplification of MDR1 were described previously (9), and their sequences are shown in Table I. Cycling parameters were as follows: 30 sec at 95°C, followed by 40 cycles of 5 sec at 95°C, 20 sec at 60°C and 0 sec at 95°C, followed by 15 sec at 65°C and 0 sec at 95°C. Relative mRNA quantities were calculated using the 2-ΔΔCt method (10), using the housekeeping gene coding for the TATAA-box binding protein (TBP) for normalization, and are represented as fold changes relative to the samples with the lowest mRNA expression. To compare the mRNA expression level of MDR1 between the subjects or subject groups, the ratio of the MDR1 transcript level to the TBP transcript level was calculated. The mean value (duplicate measurements) was shown in each case. The PCR products were subjected to gel electrophoresis to ensure the size of the specific amplicons of interest and to confirm the absence of non-specific PCR products. To further confirm their sequences, the resultant PCR products were further sequenced. The specificity of each reaction was also assessed by carrying out melting curve analysis, to ensure that only one product was present. All samples were amplified in triplicate.

Detection of P-gp by ELISA. The CD8+ T cells were purified by a negative selection method, using magnetic-activated cell sorting (MACS; Miltenyi Biotec GmbH, Bergisch Gladbach,
Germany) according to the manufacturer’s instructions. The relative levels of P-gp in the CD8+ T cells were quantified using ELISA kits (Cusabio Biotech, Wuhan, China) according to the manufacturer’s instructions. Each assay was carried out in triplicate wells.

Quantification of P-gp expression by flow cytometry. To quantify P-gp protein expression, flow cytometry analysis was performed on the stored PBMC samples using fluorochrome-conjugated antibodies specific for the surface markers, CD3, CD4, CD8 and P-gp. Approximately 1x10⁶ PBMCs, diluted in 2 ml phosphate-buffered saline (PBS) containing 1% FBS were transferred to 5-ml sterile tubes. The cells were harvested following centrifugation at 1,500 rpm for 10 min at 4°C. The following antibodies were used for staining: anti-CD3-PerCP, anti-CD4-FITC, anti-CD8-APC (all from BD Biosciences) and mouse anti-human ABCB1 PE-conjugated monoclonal antibody (eBioscience, San Diego, CA, USA). The cells were incubated and stained at 4°C in the dark for 30 min, and were then analyzed with a 4-colour FACSCalibur cytometer. Image acquisition was performed using CellQuest software (BD Biosciences), and data analyses were performed using FlowJo software version 8.6 (Tree Star Inc., Ashland, OR, USA). Isotype control antibodies were used to separate the infected from the control cells in the FITC, PerCP, PE and APC fluorescence channels.

Rhodamine 123 (Rh123) efflux assay. P-gp activity was determined by estimating the efflux of Rh123 (Sigma-Aldrich), a fluorescent dye that is a P-gp substrate, as previously described (11). Briefly, 200 µl of purified cell suspension was incubated with Rh123 (final concentration, 200 ng/ml) for 20 min under 5% CO₂ at 37°C. After washing, the cells were incubated in Rh123-free medium in the presence or absence of cyclosporine A (CsA; Sigma-Aldrich), a P-gp-specific competitive inhibitor (final concentration, 1,000 ng/ml), for 30 min under 5% CO₂ at 37°C. Finally, the cells were washed in CsA-containing RPMI-1640 medium and stained with phycoerythrin (PE)-conjugated, anti-CD3 and anti-CD8 antibodies (both from BD Biosciences). After staining, the cells were washed twice, resuspended in PBS and kept at 4°C in the dark for 20 min.

The data were analyzed with a FACSCalibur flow cytometer using CellQuest software. The efflux of Rh123 from the PBMCs was assessed by analyzing the changes in cellular fluorescence in the presence or absence of CsA. The Rh123 fluorescence emits at a wavelength of 488 nm and was detected on fluorescence channel (FL)1 of the flow cytometer. PE fluorescence was detected on FL2. The forward- and side-scatter were used to gate the cell population of lymphocytes. For each sample, 10,000 events were collected. The data were analyzed using the Kolmogorov-Smirnov (KS) statistic, which measures the difference between 2 distribution functions and generates a D-value, ranging from 0 to 1.0. A higher D-value indicates a higher difference between the distribution functions, and thus, a strong expression of P-gp.

Statistical analysis. Data were summarized by computing the median and interquartile range (IQR). Undetectable viral loads were assigned a value of 20 RNA copies/ml (average of maximum and minimum undetectable values). Differences between independent groups were tested for statistical significance using the Mann-Whitney U test or the Wilcoxon signed-rank test for paired data (time-paired data in each group).

For each virologic and immunologic parameter, sensitivities, specificities, positive and negative predictive values were assessed as to their ability to define the response to treatment in the entire cohort (e.g., the sensitivity of high levels of activated cells in identifying patients with a detectable viral load during the follow-up months). The performance of these evaluations was studied by receiver operating characteristic (ROC) curves. ROC curves depict the true positive rate (sensitivity) vs. the false positive rate [1 minus the specificity (1-specificity) or true negative rate]. The area under the curve (AUC) was used to measure discrimination, i.e., the ability of the test to correctly classify good and poor responders. An AUC of 1 represents a perfect test [sensitivity = 1 (100%), specificity = 1 (100%)]. Cut-off values with the highest discrimination capacity between good and poor responders were established. Confidence intervals (CI) of 95% were used to depict uncertainty in the estimates of sensitivity and specificity. All statistical analyses were performed using SPSS 16.0 software for Windows (SPSS Inc., Chicago, IL, USA). The Spearman correlation coefficient was calculated to assess the association between the proportion of P-gp-expressing CD8+ T cells and other virologic or immunological parameters. Coefficients of variation (%) were expressed as (SD/mean) x100. A value of P<0.05 was considered to indicate a statistically significant difference.

Results

Patient characteristics. A total of 92 patients were enrolled in the initial analysis, with more than half (79.7%) being male (Table II). The median age of the HIV-1-infected patients on ART was 35 years (IQR, 22-55), the median age of the HIV-1-infected ART-naive patients was 33 years (IQR, 19-46), and there were no significant differences in age among

Table I. Primers used for qRT-PCR.

<table>
<thead>
<tr>
<th>Gene</th>
<th>Primer sequence</th>
<th>Length (bp)</th>
<th>GenBank Accession no.</th>
</tr>
</thead>
<tbody>
<tr>
<td>TBP</td>
<td>F: TGCACAGGAGCCAAAGATGAA R: CACATCAGACTCCCAACCA</td>
<td>132</td>
<td>NM_003194</td>
</tr>
<tr>
<td>MDR1</td>
<td>F: CACCCGACCTACAGATGATG R: GTGCCATTGACTGAAAGAA</td>
<td>81</td>
<td>NM_000927</td>
</tr>
</tbody>
</table>

TBP, TATAA-box binding protein; MDR1, multidrug resistance protein 1; F, Forward, R, reverse.
Table II. Demographic and immunological parameters of HIV-1-infected patients enrolled in this study.

<table>
<thead>
<tr>
<th>Parameters</th>
<th>HIV-1-infected patients on ART</th>
<th>HIV-1-infected ART-naïve patients</th>
<th>Healthy blood donors</th>
</tr>
</thead>
<tbody>
<tr>
<td>No.</td>
<td>92</td>
<td>49</td>
<td>24</td>
</tr>
<tr>
<td>Age (years)</td>
<td>35 (22-55)</td>
<td>33 (19-46)</td>
<td>28 (25-33)</td>
</tr>
<tr>
<td>Gender ratio (male/female)</td>
<td>74/18</td>
<td>38/11</td>
<td>14/10</td>
</tr>
<tr>
<td>CD4+ T cell count, cells/µl</td>
<td>217 (3-615)</td>
<td>103 (6-231)</td>
<td>721 (620-839)</td>
</tr>
<tr>
<td>Viral load, log_{10} copies/ml</td>
<td>2.2 (1.6-4.2)</td>
<td>3.8 (2.6-5.5)</td>
<td>N/A</td>
</tr>
</tbody>
</table>

Values are expressed as the median (interquartile range), apart from the number (no.) and gender ratio. N/A, not applicable. ART, antiretroviral therapy.

Table III. Demographic and immunological parameters of the 48 HIV-1-infected patients, defined as the extreme group.

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Good virologic responders</th>
<th>Poor virologic responders</th>
<th>Healthy blood donors</th>
</tr>
</thead>
<tbody>
<tr>
<td>No.</td>
<td>20</td>
<td>28</td>
<td>24</td>
</tr>
<tr>
<td>Age (years)</td>
<td>40 (33-43)</td>
<td>42 (34-41)</td>
<td>28 (25-33)</td>
</tr>
<tr>
<td>Gender ratio (male/female)</td>
<td>12/8</td>
<td>18/10</td>
<td>14/10</td>
</tr>
<tr>
<td>CD4+ T cells/µl</td>
<td>269.5 (210.3-339.8)</td>
<td>321 (228.3-341.3)</td>
<td>721 (620-839)</td>
</tr>
<tr>
<td>Viral load, log_{10} copies/ml</td>
<td>4.5 (4.2-4.8)</td>
<td>4.6 (4.5-4.9)</td>
<td>N/A</td>
</tr>
</tbody>
</table>

Values are expressed as the median (interquartile range), apart from the number (no.) and gender ratio. N/A, not applicable.

the different groups. The median CD4+ T cell count of the HIV-1-infected patients on ART was 217 cells/µl (IQR, 3-615), and the median HIV RNA level was 2.2 log_{10} copies/ml (IQR, 1.6-4.2). The median CD4+ T cell count of the HIV-1-infected ART-naïve patients was 103 cells/µl (IQR, 6-231), and the median HIV RNA level was 3.8 log_{10} copies/ml (IQR, 2.6-5.5).

Of the 48 patients that had received ART for 2 years, 20 were categorized as good virologic responders and 28 as poor virologic responders. The median ages were 40 years (IQR, 33-43) for the good responders, 42 years (IQR, 34-41) for the poor responders, and 28 years (IQR, 25-33) for the healthy blood donors. Median CD4+ T cell counts at baseline were estimated at 269.5 cells/µl (IQR, 210.3-339.8) for the good responders, 321 cells/µl (IQR, 228.3-341.3) for the poor responders (P>0.05), and 721 cells/µl (IQR, 620-839) for the healthy blood donors (P<0.05). Plasma RNA viral load at baseline was similar for good and poor responders [4.5 log_{10} copies/ml (IQR, 4.2-4.8) and 4.6 log_{10} copies/ml (IQR, 4.5-4.9), respectively, P>0.05] (Table III). The viral load of the good responders was maintained below the detection limit for at least 12 months. By contrast, in the poor responders, median viral loads were 3.0 log_{10} and 3.9 log_{10} copies/ml after 6 and 12 months of therapy, respectively.

Expression of MDR1 transcripts. The relative mRNA levels for the MDR1 gene in the HIV-1-infected patients on ART, the HIV-1-infected ART-naïve patients and the healthy donors are shown in Fig. 1A. The MDR1 gene was detected in all the groups analyzed. The mRNA levels of MDR1 were low in both the HIV-1-infected ART-naïve patients and the healthy blood donors, and were significantly lower than those observed in the HIV-1-infected patients on ART. As regards the demographic and immunological characteristics of the patients, MDR1 expression did not correlate with age, gender, or the initial white blood cell count.

Expression of P-gp in CD8+ T cells of HIV-1-infected patients. P-gp expression in the CD8+ T cells was detected by ELISA in the 92 HIV-1-infected patients on ART, the 49 HIV-1-infected ART-naïve patients and the 24 healthy blood donors (Fig. 1B). The levels of P-gp in the HIV-1-infected patients on ART were significantly higher than those observed in the HIV-1-infected ART-naïve patients and the healthy blood donors.

The expression of P-gp was further quantified by flow cytometry in the same groups. As shown in Fig. 2, a high inter-individual variability was observed in the samples from the 3 groups, with coefficients of variation (CV) approaching 40%. Notably, the highest degree of variability was observed for the HIV-1-infected patients on ART (45.5%).

Determination of the MDR1 gene levels. To determine the levels of MDR1 gene expression, the RNA was isolated from PBMCs, and reverse transcription-polymerase chain reaction (RT-PCR) was performed to quantitate the mRNA levels of MDR1. The expression levels of MDR1 were significantly higher in the HIV-1-infected patients on ART (22.40±8.101) compared with the HIV-1-infected ART-naïve patients (19.57±7.124) and the healthy blood donors. As regards the demographic and immunological characteristics of the patients, MDR1 expression did not correlate with age, gender, or the initial white blood cell count.

Comparison of P-gp activity in the CD8+ T cells between the HIV-1-infected patients and healthy donors. In order to
evaluate the ability of P-gp to actively pump drugs from the cytosol and plasma membrane into the extracellular space, we analyzed the amounts of extruded Rh123 via calculations of the D-value in CD8+ cells. We then compared the P-gp activity in peripheral CD8+ cells between healthy donors and HIV-1-infected patients. P-gp activity tended to be higher in the HIV-1-infected patients on ART, and statistically significant differences were noted in P-gp activity (Fig. 3) between the HIV-1-infected patients on ART and the HIV-1-infected ART-naïve patients, as well as between the HIV-1-infected patients on ART and the healthy donors (P<0.001 and P<0.001, respectively).

Correlation of P-gp+CD8+ T cell with blood CD4+ T cell count. To investigate whether the increase in P-gp expression in peripheral blood CD8+ T lymphocytes correlates with the blood CD4+ T cell count, we measured the CD4+ T cells in the recruited patients. Due to the limited availability of the material, it was not possible to evaluate the CD4+ T cell numbers for all the HIV-1-infected patients; the number of CD4+ T cells was measured in 39 HIV-1-infected patients on ART. We found a significant inverse correlation between P-gp+CD8+ T cell and CD4+ T cell counts in the HIV-1-infected patients on ART (Fig. 4). The correlation between these 2 cell populations was also investigated in the HIV-1-infected ART-naïve group and the healthy donors group, but no significant correlation was found (P>0.05, data not shown).

Correlation between expression of P-gp and virologic parameters. Spearman’s correlation analysis was used to determine whether the expression levels of P-gp are associated with the level of plasma HIV RNA. The results revealed no evidence of the prognostic value of P-gp expression in HIV-1 infection, since no correlation was observed between P-gp expression
and the level of plasma HIV-RNA in HIV-1-infected patients on ART (P>0.05, data not shown).

Changes in numbers of P-gp*CD8+ T cells among good and poor virologic responders. In the good-responder group, 6 the median levels of CD4+ T cells increased steadily from 269.5 cells/µl at baseline to 497 cells/µl at 6 months (P<0.0001), to 512.5 cells/µl at 12 months (P<0.0001), to 460 cells/µl at 18 months (P=0.001), and to 486.5 cells/µl at 24 months (P<0.0001). (B) In the poor responders, CD4+ T cell levels after 6, 12 and 18 months of therapy [370 (239.5-462.3); 361.5 (265.8-526.3); 398 (259.3-585) cells/µl] were similar to values observed at baseline [321 (228.3-341.3) cells/l] (P=0.16; P=0.168; P=0.051), then increased to 462.5 cells/µl at 24 months (P=0.012). Boxplots represent the median, interquartile range (boxes) and the 5-95% data range (whisker caps). Healthy blood donors (HD) were used as controls. ART, antiretroviral therapy.

Predictive value of immunological markers for the response to treatment. We examined whether the absolute CD4+ T cell count or the relative proportion of P-gp*CD8+ cells can be used for discriminating the response to treatment between the good- and poor-responder groups. For each parameter, we plotted the true positive rate (sensitivity) and the false positive
The AUC was used to assess and compare the discriminatory power of the 2 markers in the good- and poor-responder groups (Table IV). The AUC of an ideal test equals 1, whereas the AUC of a poor test equals 0.5. In the present study, the P-gp\(^{+}\)CD8\(^{+}\) cell percentage was the best predictor of virologic response with an associated AUC value equal to 0.885, 0.846, 0.814 and 0.852 at 6, 12, 18 and 24 months, respectively. Absolute CD4 counts did not show any discriminatory value for the response to treatment, with associated AUC values close to 0.5 and P>0.5 for all months, apart from the 6th month (Table IV and Fig. 8). The use of P-gp mean fluorescence intensity (MFI) as a marker did not improve the discrimination between the two groups.

In Figure 8, the receiver operating characteristic (ROC) curve of blood CD4\(^{+}\) T cell counts at 6 months in the extreme group (n=48) is shown. The correlation between sensitivity (sens) and 1-specificity (spec) of each immunological parameter to predict good or poor response to treatment was examined. ROC curves were constructed by plotting the true positive rate (sens) against the false positive rate (1-spec) for all possible cut-off values, displayed along the curve. The smallest cut-off value in the curve is the minimum observed value minus 1 and the largest cut-off value is the maximum observed value plus 1. All the other cut-off values are the average of 2 consecutive observed values. The area under the curve (AUC) is 0.782. The optimal operating point (op) i.e., the cut-off yielding the highest test accuracy, is associated with a relatively high false negative rate (sens, 60%; spec, 92.9%). This point lies on a 45° line closest to the northwest corner (sens, 100%; spec, 0%) of the ROC curve. For more efficient screening of poor response to treatment, a maximal sensitivity point may be preferred by using 31.515% as the optimal operating point (sens, 100%; spec, 57.1%).

In Figure 9, the receiver operating characteristic (ROC) curve of P-gp\(^{+}\)CD8\(^{+}\) T cell percentages at 6 months in the extreme group (n=48) is shown. The area under the curve (AUC) is 0.885. The optimal operating point (op) i.e., the cut-off yielding the highest test accuracy, is associated with a relatively high false negative rate (sens, 64.3%; spec, 92.9%). This point lies on a 45° line closest to the northwest corner (sens, 100%; spec, 0%) of the ROC curve. For more efficient screening of poor response to treatment, a maximal sensitivity point may be preferred by using 31.515% as the optimal operating point (sens, 100%; spec, 57.1%).
sensitivity and specificity of the discriminatory assay compared to the P-gp+CD8+ T cell percentage (data not shown).

A range of cut-off values between the optimal operating point (i.e., the cut-off value yielding the best trade-off between sensitivity and specificity) and the maximal sensitivity point (highest possible sensitivity value) was identified from the ROC curve of P-gp+CD8+ percentages at 6 months of ART. The results are presented in Table V.

### Table V. Sensitivity and specificity of P-gp+CD8+ cell percentages associated with different cut-off values in predicting good and poor virologic response to antiretroviral therapy (ART) in the HIV-1-infected group (extreme group; n=48).

<table>
<thead>
<tr>
<th>Treatment time point</th>
<th>Criterion for choice of cut-off</th>
<th>Cut-off (%)</th>
<th>Sensitivity</th>
<th>Specificity</th>
</tr>
</thead>
<tbody>
<tr>
<td>Month 6</td>
<td>Optimal operating point</td>
<td>50.550</td>
<td>64.3</td>
<td>92.9</td>
</tr>
<tr>
<td></td>
<td>Maximal sensitivity point</td>
<td>31.515</td>
<td>100</td>
<td>57.1</td>
</tr>
<tr>
<td>Month 12</td>
<td>Optimal operating point</td>
<td>56.500</td>
<td>100</td>
<td>85.7</td>
</tr>
<tr>
<td></td>
<td>Maximal sensitivity point</td>
<td>56.500</td>
<td>100</td>
<td>85.7</td>
</tr>
<tr>
<td>Month 18</td>
<td>Optimal operating point</td>
<td>63.415</td>
<td>64.3</td>
<td>85.7</td>
</tr>
<tr>
<td></td>
<td>Maximal sensitivity point</td>
<td>50.500</td>
<td>100</td>
<td>42.9</td>
</tr>
<tr>
<td>Month 24</td>
<td>Optimal operating point</td>
<td>63.270</td>
<td>78.6</td>
<td>78.6</td>
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<tr>
<td></td>
<td>Maximal sensitivity point</td>
<td>50.615</td>
<td>100</td>
<td>42.9</td>
</tr>
</tbody>
</table>

AUC, area under the curve; ROC, receiver operating characteristic; CI, confidence interval. *Statistically significant differences.

Discussion

Drug interactions between anti-HIV drugs, as well as viral resistance to ART drugs, represent a considerable limitation to the safety and effectiveness of ART. As regards HIV-1 infection, several studies have reported that a number of ATP-dependent transport proteins, such as P-gp, breast cancer resistance protein (BCRP), multidrug resistance-associated protein (MRP)1, MRP4 and MRP5, are able to transport anti-retroviral drugs (12-15). Although evidence for the central role of other transporters in the pharmacokinetics of the interaction with antiretroviral drugs is accumulating, the role of P-gp is far from being elucidated, particularly in large clinical cohorts of patients with HIV-1 infection.

HIV-1 infections are treated with HIV-protease inhibitors (PIs), nucleosides (NRTIs), non-nucleosides (NNRTIs) and nucleotide reverse transcriptase (NtRTIs) inhibitors. The combined administration of different drugs in ART improves the disease outcome, while increasing the likelihood of drug interactions. P-gp represents an important site for drug interactions and the induction of its activity can reduce the effectiveness of drugs that are P-gp substrates. PIs are well established inducers of P-gp (16-19), while data on P-gp modulation by NRTIs and NNRTIs are limited and often conflicting, mainly due to the difficulties in comparing induction results from different assays.

The induction of P-gp expression by AZT in tumor cell lines was found by Gollapudi and Gupta (20). Their study demonstrated that HIV-1-infected T cells, as well as monocytic cell lines, have increased levels of P-gp, while AZT accumulates in significantly lower quantities in these cells compared to uninfected cells (20). Likewise, a decrease in AZT accumulation in P-gp-overexpressing CEM VBL100 cells with a corresponding

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**Table IV.** AUC, P-values and 95% CI corresponding to each parameter analyzed on the ROC curve for its discriminating value between good and poor responders within the HIV-1-infected group (extreme group; n=48).

<table>
<thead>
<tr>
<th>Treatment time points</th>
<th>Parameters</th>
<th>AUC</th>
<th>P-value</th>
<th>95% CI</th>
</tr>
</thead>
<tbody>
<tr>
<td>Month 6</td>
<td>CD4+ T cell count</td>
<td>0.782</td>
<td>0.021</td>
<td>0.592-0.972</td>
</tr>
<tr>
<td></td>
<td>P-gp+CD8+ T cells (%)</td>
<td>0.885*</td>
<td>0.001*</td>
<td>0.766-1.004</td>
</tr>
<tr>
<td>Month 12</td>
<td>CD4+ T cell count</td>
<td>0.736</td>
<td>0.053</td>
<td>0.527-0.945</td>
</tr>
<tr>
<td></td>
<td>P-gp+CD8+ T cells (%)</td>
<td>0.946*</td>
<td>&lt;0.001*</td>
<td>0.854-1.039</td>
</tr>
<tr>
<td>Month 18</td>
<td>CD4+ T cell count</td>
<td>0.607</td>
<td>0.380</td>
<td>0.375-0.840</td>
</tr>
<tr>
<td></td>
<td>P-gp+CD8+ T cells (%)</td>
<td>0.814*</td>
<td>0.005*</td>
<td>0.652-0.975</td>
</tr>
<tr>
<td>Month 24</td>
<td>CD4+ T cell count</td>
<td>0.600</td>
<td>0.412</td>
<td>0.359-0.841</td>
</tr>
<tr>
<td></td>
<td>P-gp+CD8+ T cells (%)</td>
<td>0.852*</td>
<td>0.002*</td>
<td>0.713-0.991</td>
</tr>
</tbody>
</table>

AUC, area under the curve; ROC, receiver operating characteristic; CI, confidence interval. *Statistically significant differences.
decline in the antiviral efficacy of the drug was observed in another study (21).

Weiss et al (22) demonstrated the induction, with different potencies, of P-gp activity by a number of NRTIs and NNRTIs, such as EFV and NVP. In the Caco-2 cell line, the NNRTIs NVP, EFV and delavirdine, which are not P-gp substrates, were shown to induce the expression and activity of P-gp, with NVP being the most potent inducer compared with the other 2 NNRTIs (23). Another study also reported the P-gp-inducing properties of delavirdine, EFV and NVP (24); however, results from studies aiming to assess whether these NNRTIs are also substrates and/or inhibitors of P-gp are conflicting (25,26).

Moreover, the few studies available used different methods, resulting in restricted comparability and ambiguous conclusions. In our study, the recruited patients who were receiving the AZT + 3TC + EFV/NVP treatment showed higher levels of P-gp in their CD8+ T cells compared with ART-naïve patients, which suggests that NRTIs/NNRTIs may have P-gp-inducing properties and may contribute to drug resistance in ART in HIV-1 infection.

The data from the present study indicate that there is a high inter-individual variability in the protein expression of P-gp in both HIV-1-infected patients (on ART or ART-naïve) and healthy blood donors. The degree of variability in CD8+ T lymphocytes was higher in the patients on ART compared with the ART-naïve patients and healthy donors, which suggests that the therapy may contribute to the variability of the expression of this transporter. However, the coefficient of variation of the measured P-gp expression in the CD8+ cells of the HIV-1-infected patients on ART was 29%, which is less than coefficients 61, 40 and 37% reported in previous studies (27-29); these discrepancies are possibly due to different methods used to quantify the expression or to the different study subjects. Although variations in P-gp levels in the CD4+ population in our study were also reduced compared to previous studies (data not shown), the tendency for CD4+ cells to express lower levels of P-gp compared to CD8+ cells is in agreement with previous studies (30,31). Overall, the variability in expression of P-gp in the lymphocyte subpopulations reported in our study is similar or reduced compared to other studies. ART appears to affect the expression of P-gp, since its expression was significantly higher in HIV-1-infected patients on ART compared with HIV-1-infected ART-naïve patients and healthy donors. This result is in agreement with previous in vitro studies demonstrating that the expression of ABC transporters in PBMCs is increased in HIV-1 infection due to ART (32-34).

There was a significant inverse correlation between P-gp/CD8+ T cell counts and CD4+ cell counts in HIV-1-infected patients on ART in our study. A previous study (35) demonstrated that the basal expression of P-gp in different cell populations ranks as follows: CD56+ >CD8+ >CD4+. However, the ratio of CD4+/CD8+ cells is reversed as CD4+ T cells are continuously destroyed by the invading virus in HIV-1 infections, and therefore, the expression of P-gp may vary at different stages of the infection.

In our study, P-gp expression did not correlate with viral load, although the transporter expression in HIV-1-infected patients on ART was higher compared with HIV-1-infected ART-naïve patients and healthy donors, which supports the hypothesis that several factors may be involved in HIV infection. The role of antiretroviral drugs in modulating the expression of ABC transporters has not yet been elucidated. The possibility that both therapy and HIV-1 infection may modulate the transporter expression and translation cannot be ruled out. Moreover, increasing evidence suggests that P-gp directly inhibits the replication of enveloped viruses, and that P-gp expression may inhibit HIV-mediated membrane fusion, as well as subsequent steps in the HIV-1 life cycle (36-38). Overall, we cannot ascertain that the lower viral loads observed in the good virologic responders are related to ART-enhanced P-gp expression.

We also analyzed the changes in the levels of CD4+ T cells, the plasma viral load and the percentages of P-gp/CD8+ T cells throughout 24 months of ART. The suppression of HIV-1 replication following ART induced significant changes in P-gp expression in the majority of patients. Our study provided evidence for a progressive increase in the percentage of P-gp/CD8+ T cells, with steadily increasing CD4+ T cell counts and a sharp decline in the plasma viral load, indicating that P-gp/CD8+ T cells may represent a marker monitoring the response to ART when the viral load decreases to levels lower than the detection limit after antiretroviral intervention.

We used ROC curves to assess the sensitivity and specificity of the percentage of P-gp/CD8+ and CD4+ T cells to predict virologic response. We found that the relative proportion of P-gp/CD8+ T cells is a more efficient predictive marker compared to CD4+ T cell counts. This result confirmed previous findings that the percentage of P-gp/CD8+ T cells constitutes a good marker of virologic response in HIV-1-infected patients on ART.

There are certain limitations to the current study, in particular, the small sample size and the fact that data on the effects of viral suppression on CD8+ T cell counts in patients who did not respond to ART was lacking. Future studies including a group of patients who do not respond to ART are required to provide crucial information on the effects of ART on P-gp expression in HIV-1 infection. Furthermore, additional time-series data on the expression of the transporter in patients and healthy volunteers are required to define the exact effect of drug-to-drug interactions on the expression of P-gp.

In conclusion, the results from the present study suggest that P-gp expression varies among HIV-1-infected patients, but is significantly increased upon ART in HIV-1-infected patients. The overexpression of P-gp is affected during the initiation of ART in HIV-1 infection and the P-gp/CD8+ T cell count may provide an additional criterion for estimating the response to ART. The relative proportion of P-gp/CD8+ T cells appeared to provide a sensitive estimate of the virologic response to ART in comparison to the blood CD4+ T cell count. P-gp expression and its correlation to the virologic and immunological parameters of HIV-infected patients warrants further investigation using a larger study group to determine the effects of P-gp expression on clinical outcome.

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References


