Tenoforv Plasma Concentrations According to Companion Drugs: a Cross-Sectional Study of HIV-Positive Patients with Normal Renal Function


Unit of Infectious Diseases, University of Torino, Department of Medical Sciences, Torino, Italy; Department of Infectious Diseases, Divisione A, Ospedale Amedeo di Savoia, ASLTO2, Torino, Italy

As the risk of tenofovir-associated renal toxicity has been found to be proportional to the drug plasma concentration, our aim was to measure the determinants of tenofovir plasma exposure in HIV-positive patients with normal renal function. A cross-sectional analysis was conducted in HIV-positive patients chronically receiving tenofovir-containing highly active antiretroviral therapies (HAARTs). Patients on tenofovir-containing antiretroviral regimens, presenting 22 to 26 h after drug intake, having estimated glomerular filtration rates above 60 ml/min, reporting high adherence to antiretroviral medications (above 95% of the doses), and signing a written informed consent were included. Plasma tenofovir concentrations were measured through a validated high-performance liquid chromatography–mass spectrometry (HPLC/LC-MS) method. The tenofovir trough concentrations in 195 patients (median, 50 ng/ml, and interquartile range, 35 to 77 ng/ml) were significantly associated with the estimated glomerular filtration rate, body mass index, and third-drug class (protease-containing versus protease-sparing regimens) (with the highest exposure in unboosted-atazanavir recipients). The results of multivariate analysis showed that the third-drug class and the weight/creatinine ratio were independent predictors of tenofovir trough concentrations. This cross-sectional study shows that tenofovir trough concentrations are predicted by the weight/creatinine ratio and by the coadministered antiretrovirals, with protease inhibitors (whether boosted or unboosted) being associated with the highest plasma exposure. These data, previously available in healthy subjects or for some drugs only, could be useful for designing strategies to manage tenofovir-associated toxicity, since this toxicity has been reported to be dose dependent.

A. Calcagno, Andrea. calcagno@unito.it.

MATERIALS AND METHODS

Patients on TDF-containing antiretroviral regimens were consecutively enrolled at the Department of Infectious Diseases of the University of Torino and at the Amedeo di Savoia Hospital, Torino, Italy, between September 2010 and January 2011. The protocol was approved by the local ethics committee. Patients were included if they had taken TDF 22 to 26 h before, reported high adherence to antiretroviral medications (above 95% of the doses), presented no concomitant renal disease, and signed a written informed consent. Patients diagnosed with chronic kidney disease (defined by estimated creatinine clearance below 60 ml/min), on hemodialysis, or affected by diabetes mellitus were excluded from this study. Tenofovir trough concentrations were measured through a validated high-performance liquid chromatography–mass spectrometry (HPLC/LC-MS) method with a limit of detection of 2 ng/ml. eCLCR was calculated using the Cockcroft-Gault formula. The results are expressed as median values and interquartile ranges; nonparametric tests (Spearman, Mann-Whitney, and Kruskal-Wallis) were used for all the analyses (comparisons and correlations), while a multivariate linear regression analysis.
was used to evaluate the effects of several covariates (with a \( P \) value of <0.20 at bivariate analysis) on tenofovir plasma exposure.

### RESULTS

One hundred ninety-five adult HIV-positive patients (68.2% male) were enrolled in this study; they were mainly of Caucasian ethnicity (166 [85.1%]). The median values and interquartile ranges for age, body mass index (BMI), plasma creatinine, and eCLCR were 45 years (39 to 50), 23.1 kg/m\(^2\) (21.1 to 25.8), 0.96 mg/dl (0.84 to 1.08), and 84.4 ml/min (70.2 to 102.9). Patients were cotreated with unboosted atazanavir (ATV) (34 patients [17.4%]), with a boosted protease inhibitor (118 patients [60.5%]); 54 patients on ATV/ritonavir [ATV/r], 30 on lopinavir/ritonavir [LPV/r], and 23 on darunavir/ritonavir [DRV/r], with non-nucleoside transcriptase inhibitors (NNRTIs) (32 patients [16.4%];16 on nevirapine [NVP] and 13 on efavirenz [EFV]), and with raltegravir (11 patients [5.6%]). The demographic and treatment characteristics of the study participants are summarized in Table 1, stratified by third-drug class.

Samples were collected at steady state and after a median un-interrupted time of TDF intake of 22.2 months (interquartile range, 8.9 to 56.2). The median tenofovir trough plasma concentration was 50 ng/ml (35 to 77), with a coefficient of variation of 64.6%. Tenofovir trough concentrations were significantly associated with eCLCR (rho, 0.24; \( P = 0.012 \)), BMI (rho, 0.16; \( P = 0.025 \)), and weight/creatinine ratio (rho, 0.24; \( P = 0.001 \)); a borderline association with age emerged (rho, 0.13; \( P = 0.07 \)). The tenofovir concentrations according to the third-drug class were higher in unboosted atazanavir recipients (median value and interquartile range, 65 ng/ml [41.7 to 84.2]) than in patients under treatment with boosted PIs (51.5 ng/ml [34 to 77]), NNRTIs (43 ng/ml [28.2 to 50]), and raltegravir (37 ng/ml [28 to 79]) (Kruskal-Wallis test, \( P = 0.02 \)) (Fig. 1). Dunn’s multiple comparison test identifies as statistically significant the pairwise comparison (with post hoc correction) between atazanavir and NNRTI recipients. Receiving a protease inhibitor (whether with or without ritonavir) was associated with higher tenofovir plasma trough concentrations (54.5 ng/ml [37 to 80] versus 43 ng/ml [28 to 50]) in the NNRTI and raltegravir group; \( P = 0.007 \). No significant differences were found in BMI, plasma creatinine, and eCLCR among these groups; nevertheless, patients on boosted PIs and raltegravir had shorter durations of tenofovir exposure (Table 1). No statistically significant intraclass differences in tenofovir exposure were found among patients on boosted PIs (ATV/r 54 ng/ml [35 to 76], LPV/r 55 ng/ml [40 to 78], and DRV/r 43 ng/ml [29 to 77]; \( P = 0.27 \)) or NNRTIs (NVP 41 ng/ml [29 to 53] and EFV 46 ng/ml [28 to 50]; \( P = 0.67 \)). A multivariate linear regression analysis was performed using the backward exclusion procedure and including age, BMI, eCLCR, weight/creatinine ratio, and third-drug class; the third-drug class (\( P = 0.004 \)), beta, 0.22; 95% inhibitory concentration (IC\(_{95}\), 6.8 at 35.3) and the weight/creatinine ratio (\( P = 0.04 \), beta, –2.02; IC\(_{95}\) = 0.73 to 0.009) were independently associated with tenofovir trough concentrations.

### DISCUSSION

In this cross-sectional study in HIV-positive patients, we report new clinical data on tenofovir plasma PK exposure according to different concomitant drugs. Information on this was mainly derived from data for healthy volunteers, with only a few clinical studies performed on specific tenofovir-antiretroviral combinations in HIV-infected patients (such as lopinavir/ritonavir) (11). Our results further support the hypothesis of a tenofovir dose/

---

**TABLE 1** Demographic and clinical characteristics

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Unboosted atazanavir</th>
<th>Boosted PIs</th>
<th>NNRTIs</th>
<th>Raltegravir</th>
<th>( P ) value*</th>
</tr>
</thead>
<tbody>
<tr>
<td>No. of patients (( n = 195 ))</td>
<td>34</td>
<td>118</td>
<td>32</td>
<td>11</td>
<td>0.56</td>
</tr>
<tr>
<td>Gender (male)</td>
<td>20 (58.8%)</td>
<td>84 (71.2%)</td>
<td>23 (71.9%)</td>
<td>6 (54.5%)</td>
<td>0.56</td>
</tr>
<tr>
<td>Ethnicity (Caucasian)</td>
<td>27 (79.4%)</td>
<td>106 (89.8%)</td>
<td>24 (75%)</td>
<td>9 (81.8%)</td>
<td>0.38</td>
</tr>
<tr>
<td>Age (yr)</td>
<td>44.5 (37.8–50)</td>
<td>44 (39–49.5)</td>
<td>47 (39–54)</td>
<td>43 (41–47)</td>
<td>0.74</td>
</tr>
<tr>
<td>BMI (kg/m(^2))</td>
<td>24.4 (22.7–27.5)</td>
<td>23 (20.5–25.3)</td>
<td>22.7 (21.1–26.5)</td>
<td>23.7 (20.5–29.6)</td>
<td>0.18</td>
</tr>
<tr>
<td>Creatinine (mg/dl)</td>
<td>0.9 (0.84–1.09)</td>
<td>0.96 (0.84–1.07)</td>
<td>1.02 (0.86–1.07)</td>
<td>1.03 (0.94–1.15)</td>
<td>0.24</td>
</tr>
<tr>
<td>eCLCR (ml/min)</td>
<td>89.5 (75.6–104)</td>
<td>83.1 (69.8–102.7)</td>
<td>81 (69–109.2)</td>
<td>83.5 (71–93.7)</td>
<td>0.97</td>
</tr>
<tr>
<td>Duration of tenofovir treatment (mo)</td>
<td>29.1 (18.3–56.5)</td>
<td>17.5 (7–46)</td>
<td>33.9 (7.4–68)</td>
<td>11.2 (1.8–51.4)</td>
<td>0.09</td>
</tr>
</tbody>
</table>

*Variables were compared with Chi-square and Kruskall-Wallis tests.
concentration-dependent toxicity, since TDF PK exposure was found to vary significantly according to the third antiretroviral being concurrently taken, with a magnitude that reproduces the relative risk of TDF-associated nephrotoxicity according to concurrently administered drugs seen in cohort studies. In looking at TDF concentrations, it must be considered that in the three studies associating higher tenofovir exposures to kidney tubular dysfunction, the mid-dose value (approximately 12 h after drug intake) (4–7) was used, while our plasma samples were taken at 24 h.

While other factors were found to influence tenofovir plasma trough concentrations (age, BMI, and eCLCR, including the previously suggested combined parameter weight/plasma creatinine) (12, 13), the choice of the third-drug class seems to be the only readily modifiable factor. As was found in EUROSIDA, D:A:D; and two additional retrospective surveys, (3, 14–16), the use of boosted PIs together with TDF is associated with the highest risk of renal toxicity, with reduced risk in the case of NNRTI or raltegravir coadministration. Recent data report that tenofovir with either boosted protease inhibitor leads to a greater initial decline in eCLCR than tenofovir with efavirenz and that this decline may be worse with ATV/r than with LPV/r (17). The mechanism by which boosted PIs increase TDF exposure is thought to be dependent on the inhibition of multiple resistance protein (MRP) transporters at the apical side of renal tubular cells, with consequent TDF intratubular accumulation and possible chronic harm to mitochondria and with intestinal P glycoprotein inhibition (and related increased absorption) as a possible complementary mechanism. It is worth noting that the highest exposure was found in unboosted ATV recipients: MRP-2 inhibition both by boosted PIs and atazanavir could contribute to explaining this effect. It is unclear, however, why unboosted atazanavir intake was associated with higher tenofovir concentrations than boosted atazanavir, and we are not able to find a causative reason for this observed effect. In any case, since in the absence of ritonavir, ATV exposure is also lower, such a difference might be attributable to an intrinsically greater inhibition of TDF extrusion from tubular cells by ATV (than by ritonavir), which would thus interact with MRPs without competing with ritonavir for drug transporter binding. The only available data suggest that the addition of unboosted ATV to TDF in healthy volunteers was associated with increased tenofovir plasma concentrations (the geometric mean ratios for area under the concentration-time curve [AUC], maximum concentration of drug in serum [C_max], and C_min, respectively, were 1.24, 1.14, and 1.22) (18).

Some limitations of this study should be highlighted: the cross-sectional design implies that patients with severe tubular and renal toxicities have not been included due to selection bias and inclusion criteria. However, such a selection does limit the risk of confounding the cause with the effect, since no patients with altered creatinine clearance values were studied. Fewer patients were included in the efavirenz and the raltegravir groups due to their habit of taking antiretrovirals at nighttime. Finally, the measurement of adherence to treatment by self-report is not precise, and it is possible that some patients did not take the drug as they had reported.

These data warrant further investigation in order to evaluate the clinical impact in terms of risk of tubular toxicity. Apart from the implicit opportunity to switch tenofovir and/or the companion drug (to an antiretroviral associated with lower TDF PR exposure), the identification of patients with increased tenofovir exposure should drive attention toward the possibility of reducing the TDF dose or the frequency of administration. This dose reduction (one pill every other day) has already been suggested in patients with significant renal impairment and could be justified by the long intracellular half-life of tenofovir diprophosphate. As an alternative (or complementary) approach, the use of probenecid might also be considered, since in a small case series, the drug was found to actually decrease the development of tenofovir-associated tubular damage in patients with hepatitis B virus infection. This effect is likely to be attributable to a significant reduction of TDF uptake by proximal tubular renal cells due to probenecid interference with apical organic anion transporters (OATs), although the data in this report cannot support such a strategy (19).

Considering the prior demonstration of a dose-concentration relationship between tenofovir and renal toxicity, these data confirm real-life clinical grounds that TDF pharmacokinetics also depends on the antiretrovirals being administered concurrently. As a consequence, based on this knowledge, different strategies to reduce the impact of TDF on renal function can be suggested and warrant proper clinical investigation.

ACKNOWLEDGMENTS
A.C. received travel grants and speaker honoraria from Abbott, Janssen-Cilag, Bristol-Meyer-Squibb, ViViV, and Merck Sharp & Dome. S.B. and G.D.P. received travel grants and speaker’s honoraria from Gilead, Abbott, Janssen-Cilag, Bristol-Meyer-Squibb, ViViV, and Merck Sharp & Dome. All other authors declare no potential conflict of interest. This study was funded internally.

REFERENCES


