Plasma Concentrations of Efavirenz and Nevirapine among HIV-Infected Patients with Immunological Failure Attending a Tertiary Hospital in North-Western Tanzania

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Abstract

Background: Sub-therapeutic and supra-therapeutic plasma concentrations of antiretrovirals are the significant causes of treatment failure and toxicity respectively among HIV-infected patients. We conducted this study to determine the pattern of efavirenz and nevirapine plasma drug concentrations among adult HIV-infected patients with immunological failure attending at a tertiary hospital in North-western Tanzania.

Materials and Methods: A cross-sectional study was conducted among adult HIV-infected patients with immunological failure who have been on either efavirenz or nevirapine based antiretroviral regimen for more than 6 months. Patients were serially enrolled through routine Care and Treatment Clinic (CTC) activities. Plasma drug concentrations for efavirenz and nevirapine were determined by high performance liquid chromatography (HPLC) and Gas Chromatography (GC) respectively. Demographic, clinical and laboratory data such as viral load and CD4 counts were collected. Data analysis was done using STATA 12.

Results: Of the 152 patients with immunological failure enrolled, the sub-therapeutic, therapeutic and supra-therapeutic plasma antiretroviral drug concentrations were found in 43/152 (28.3%), 76/152 (50.0%) and 33/152 (21.7%) respectively. Half of the patients were outside therapeutic window with either sub-therapeutic or supra-therapeutic plasma ARV drug concentrations. There was a significant difference in distribution of ARV adherence (p-value<0.001), NRTI backbone (p-value = 0.039), HIV stage (p-value = 0.026) and viral load (p-value = 0.007) within sub-therapeutic, therapeutic and supra-therapeutic ARV plasma drug concentrations.

Conclusion: There is a wide inter-individual variability of plasma ARV concentrations among HIV patients with immunological failure, with a large proportion of patients being outside therapeutic window. This variability is significantly based on ARV adherence, NRTI backbone, viral load and HIV stage. Routine therapeutic drug monitoring (TDM) could assist identifying these patients early and making timely correction to avoid virological failure, poor immunological outcome and prevent associated drug toxicities. Nonetheless, ARV adherence should be strictly emphasized on HIV patients with immunological failure.

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Introduction

The primary aim of antiretroviral therapy (ART) is to durably suppress the viral replication to undetectable levels to allow a satisfactory immune recovery [1,2]. This is achieved by a long term use of ART at therapeutic concentrations [3]. An exposure of the virus at sub-therapeutic concentrations is likely to cause insufficient suppression of the virus and a probable selection of the resistant strains which will ultimately reduce the efficacy and durability of the ART [4]. Despite the overall success of ART, still many areas are reporting an inadequate virological suppression and drug toxicity complications [5]. The sub-therapeutic ART concentrations stand to be the main cause of poor therapeutic...
outcome [6–12]. On the other hand supra-therapeutic concentrations of ART are frequently associated with toxicities. Previous studies have indicated that between 30–40% of the patients assumed to have drug-related toxicities have abnormally high plasma concentrations of ARTs [4,6,13–17].

Efavirenz and nevirapine are the core and first line non-nucleoside reverse transcriptase inhibitors (NNRTI) commonly used to treat HIV infection [17–19]. Therapeutic drug monitoring (TDM) in patients on ART including these agents has suggested being beneficial in terms of efficacy and toxicity [17,18]. TDM is an approach by which the course of therapy for a patient is guided by measurements of plasma drug concentrations enabling physicians to optimize ART drug effectiveness and to avoid drug-related toxicity [4,20,21]. A number of studies have documented a better treatment outcome in patients whose treatment was TDM guided than those whose concentrations were not monitored [22,23]. In this regard TDM is useful in assessing adherence, investigating drug-drug interactions between antiretroviral drugs or with co-medications, preventing some ART drug toxicities, adjusting the dosage in particular populations, and increasing ART efficacy of some drugs in naive patients [4,9,12,20,22,24]. TDM of ART agents could also be useful in provision of timely dosage adjustments to avoid drug sub-therapeutic or toxic concentrations [12,22,25]. Despite this fact TDM is not done in our setting and therefore the proportions of adult HIV patients with sub-therapeutic, therapeutic and supra-therapeutic ARV plasma concentrations are not known. Therefore it was the aim of this study to determine the pattern of plasma concentrations of the commonly used ART regimens containing efavirenz and nevirapine, among adult HIV-infected patients with immunological failure.

**Materials and Methods**

**Study design and setting**

This was a hospital based cross-sectional study which was conducted between April 2011 and March 2012 at Bugando Medical Centre (BMC) at Care and Treatment Center (CTC) in Mwanza, Tanzania. BMC is a 1000-bed capacity, tertiary and teaching hospital for the North Western Tanzania. The hospital serves around 13 million people and CTC activities is one of the core part of outpatient activities, which started in 2004, and currently it serves more than 10,000 patients, of whom about 3,600 are active on ARTs. The permission to conduct this study was obtained from the Catholic University of Health and Allied Sciences (CUHAS)/BMC joint ethics review board. The written consent was obtained from all participants.

**Study population, patients’ enrolment and data collection**

The study population was adult HIV patients on either efavirenz or nevirapine based regimen for more than 6 months with immunological failure attending BMC CTC. All patients were treated with the standard dose of nevirapine 200 mg twice daily and efavirenz 600 mg once daily. Inclusion criteria were adult HIV positive patients diagnosed as per WHO guidelines, age over 18 years with immunological failure as per WHO guideline and still on first line. Pregnant women and patients co-treated with anti-tuberculosis medications and other concomitant medications known to interact with NNRTI [26] were excluded.

Immunological failure was diagnosed if the patient met one of the following criteria: i) persistent CD4 below 100 cells/mm³, ii) a drop of CD4 cell count below baseline pre-treatment level, or iii) a drop of CD4 cell count of 50% from peak on treatment value all in the absence of an ongoing co-infection and after a minimum of 6 months of ART. For criteria ii and iii, the CD4 cell count must also fall below 200 cells/mm³ to qualify as immunologic failure [27]. After giving consent a structured questionnaire was used to collect information regarding demographic data, body mass index (BMI), date of diagnosis of HIV, date of ART initiation, regime and adherence. ART adherence level in the last 30 days was assessed using pill counts [28]. The pill counts were performed by the study pharmacist, who counted the number of remaining pills at each drug refill visit. Pill count-based adherence was assessed using the formula [Adherence $= (\text{Number of pills dispensed} - \text{Number of pills returned} \times 100)/\text{Number of pills prescribed} \times \text{Number of days between pharmacy visits}]$. Good adherence was defined as a value ≥95% pills whereas poor adherence was defined as a value <95%. The patients were instructed to have their medication at night and come the following morning for blood sample collection before taking their next ART dose. Two blood samples were drawn, one for viral load which was done at BMC main laboratory and the other sample was sent to Germany for TDM to determine the plasma concentrations of efavirenz and nevirapine.

**Sample collection, processing and analysis**

Patients were instructed to have medication at night and come the following morning for blood sample collection before taking their next ART dose to determine their antiretroviral plasmatic trough concentrations for nevirapine and the mid concentrations for efavirenz. For each patient, 5 ml of whole blood was collected in plasma EDTA bottles for TDM, approximately 8 to 12 hours after the last dose of ART, just before the next dose was due. The samples were immediately centrifuged at 3000 rpm for 3 minutes to obtain plasma that was transferred into cryovials. The cryovials were stored at −20°C before shipment. The samples were packed and shipped to Germany in cold boxes with cooling packs maintaining a temperature of −30°C. The plasma concentrations of efavirenz (EFV) and nevirapine (NVP) were determined using High Performance Liquid Chromatography (HPLC) [HPLC Beckman Coulter System Gold] and Gas Chromatography (GC) [GC 6890; Agilent Technology] respectively as described previously [29,30]. The well-established HPLC/GC method used in this study to determine plasma concentrations of non-nucleoside reverse transcriptase inhibitors (NNRTI) is highly specific and sensitive. The limit of detection (LOD) of nevirapine was determined at 2 ng/ml, the lower limit of quantification (LLQ) of nevirapine was reached at a concentration of 10 ng/ml. For efavirenz the LOD was 3 ng/ml, and the LLQ was 25 ng/ml. Additional 5 ml of whole blood was collected in a tube supplemented with EDTA (BD Biosciences) for plasma preparation and sent to BMC main laboratory for viral load analysis using COBAS AmpliPrep/COBAS TaqMan (Roche molecular systems, USA) according to manufacturer’s guidelines as described previously [31].

**Data management and analysis**

Data were managed using Epi Data 3.1 (CDC Atlanta, US) and analysis was done using STATA version 12 (College Station, Texas, US). ARV drug concentrations were recorded as continuous variables. Based on the reference ARVs therapeutic ranges, defined as 1000–4000 ng/ml for EFV and 3400–8000 ng/ml for NVP [12], we defined 3 categories of ARV plasma drug concentrations: sub-therapeutic (below the lower therapeutic range limit), therapeutic (within the therapeutic range), and supra-therapeutic (above the higher therapeutic ranged limit) [12]. Categorical variables were summarized as proportion and their
significance of the difference in distribution within the categories of ARV plasma drug concentrations was assessed using Pearson’s Chi-square test or Fisher’s exact test where appropriate. We used probability plots and Shapiro-Wilk normality test to assess the normality of continuous variables. Parametric continuous data were summarized as mean with standard deviation and the significance of difference in means within categories of ARV plasma drug concentrations was assessed using one way analysis of variance (ANOVA). Non-parametric continuous data were summarized as median with interquartile range and the difference in medians within the categories of ARV plasma drug concentrations was compared using Kruskal-Wallis equality-of-populations rank test. Inter-individual pharmacokinetic variability was evaluated through the coefficient of variation calculated as the quotient of the standard deviation divided by the mean plasma drug concentrations ×100. In determining the median and the inter-individual pharmacokinetic variability, patients with plasma drug concentrations below the lower limit of quantification of the assay (25 ng/ml and 10 ng/ml for efavirenz and nevirapine respectively) were arbitrarily considered as having a level of 24 ng/ml for efavirenz and 9 ng/ml for nevirapine. In all analyses the difference was considered significant if a p-value was less than 0.05.

Results

A total of 152 HIV infected adult patients with immunological failure were enrolled in the study. Of these 79/152 (52.0%) were using nevirapine based regimen whereas 73/152 (48.0%) were on efavirenz based regimen. The ART regimens used were Zidovudine+Lamivudine+Nevirapine 46/152 (30.3%), Zidovudine+Lamivudine+Efavirenz 45/152 (29.6%), Stavudine+Lamivudine+Nevirapine 31/152 (20.4%), Tenofovir+Emtricitabine+Efavirenz 28/152 (18.4%) and Tenofovir+Emtricitabine+Nevirapine 2/152 (1.3%). The duration of use of these regimens ranged from 7 to 72 months. The mean age was 40.8±10.0 years with most patients 107/152 (70.4%) being females (Table 1). Of the 152 patients with immunological failure, 121/152 (79.6%) were in WHO clinical stage 2 or 3. Good adherence was observed in 84.2% (128/152) of patients while a viral load ≥400 copies/µL was observed in 44.7% (68/152). There were 8/152 (5.3%) patients co-infected with either hepatitis B or C virus (HBV or HCV) infection, of these 7 had HBV and one had HBC. The median [interquartile range] plasma concentrations of efavirenz and nevirapine were 2112 [1349–3452] ng/ml and 4915 [2326–7044] ng/ml respectively (Table 1).

Of the 152 patients enrolled, the sub-therapeutic, therapeutic and supra-therapeutic plasma antiretroviral drug concentrations were observed in 43/152 (28.3%), 76/152 (50.0%) and 33/152 (21.7%) respectively. Half of the patients were outside therapeutic window with either sub-therapeutic or supra-therapeutic plasma ARV drug concentrations. Based on the ARV regimens, sub-therapeutic plasma concentrations were more common among patients using nevirapine than those using efavirenz, 33.4% (28/79) versus 20.3% (15/73). Supra-therapeutic plasma antiretroviral drug concentrations were slightly lower among patients using nevirapine than those using efavirenz, 20.3% (16/79) versus 23.3% (17/73). These differences were not statistically significant (Table 2). Of the 43 patients with sub-therapeutic plasma antiretroviral drug concentrations, 17 (39.5%) had concentrations below the detection limit of the HPLC/GC. Of these, 12 were using efavirenz and 5 were using nevirapine.

Sub-therapeutic drug concentrations were significantly more common (as supra-therapeutic was less common) among patients with poor ARV adherence, NRTI backbone comprising Stavudine+Lamivudine (d4T+3TC), advanced HIV stage and those with high viral loads than their counterparts. Generally, there was a significant difference in distribution of ARV adherence rate (p-value < 0.001), type of NRTI backbone (p-value = 0.039), viral load (p-value = 0.007) and WHO HIV stage (p-value = 0.026) within the categories of ARV plasma drug concentrations (sub-therapeutic, therapeutic and supra-therapeutic). Table 2 summarizes the significance of the difference in distribution of various patients’ characteristics within the categories of ARV plasma drug concentrations. The inter-individual variability was higher among patients using efavirenz based therapy than those using nevirapine based therapy (120.9% versus 88.7%). Generally, there was a wide inter-individual variability of plasma ARV concentrations among HIV patients with immunological failure using efavirenz and nevirapine in routine clinical practice as summarized in table 3.

Discussion

This study has demonstrated a presence of a wide inter-individual variability of plasma ARV concentrations among HIV patients with immunological failure in routine clinical practice, with a large proportion of patients being outside therapeutic window. This emphasizes that clinicians are often confronted with treatment failure or side-effects, and are in need of methods to evaluate drug exposure among these patients. The finding of higher inter-individual variability among patients using efavirenz based therapy than those using nevirapine based therapy was also observed in a study done in Italy [32]. However, our inter-individual variability was higher than that observed in Italy for both antiretroviral drugs (120.9% and 88.7% versus 85.1% and 50.1% respectively) [32].

In this study sub-therapeutic ARV plasma concentrations were detected in 28.3% of patients. Our findings are similar to that obtained in the study done in Netherlands among patients at a risk of treatment failure in a routine clinical care, in which 27.4% of the plasma concentrations were classified as having sub-therapeutic ARV plasma concentrations [21]. However, our findings are slightly higher than that from previous studies done in Uganda and Italy, in which the overall sub-therapeutic ARV concentrations were found in 14.3% and 16.9% respectively [33,34]. This difference in prevalence might be due to the fact that in our study all participants had immunological failure, which was not the case in the Ugandan and Italian studies. Furthermore, our prevalence of sub-therapeutic ARV are lower than that from a study done British Columbia, in which the overall sub-therapeutic ARV concentrations were reported in 41.8% of patients with immunological failure [35]. This high prevalence could be attributed by the fact that all participants in the British Columbian study had a CD4 cell count less than 50 cells/µL. On the other hand, supra-therapeutic ARV plasma concentrations were detected in 21.7% of patients. This prevalence is comparable to that reported from Uganda where 23.9% of the patients had supra-therapeutic ARV plasma concentrations [33], nevertheless all these observations embrace comparable consequence [17,20] in clinical practice of HIV medicine.

The observations from this study are of paramount clinical relevance especially in resource-limited setting like ours. For the first time in Tanzania, we have demonstrated a presence of a wide inter-individual variability of plasma ARV concentrations and a significant association between good adherence and therapeutic ARV plasma concentrations among HIV-infected patients with immunological failure. We found that the proportion of patients with sub-therapeutic ARV plasma concentrations significantly increased with poor ART adherence, NRTI backbone comprising
Stavudine+Lamivudine (d4T+3TC), increasing viral loads and advancing HIV stage. This finding is similar to that from previous studies done in Uganda and Italy [33,34].

Therapeutic drug concentrations are a key to successful ART [7,36], as any low drug concentrations observed in patients on ART has extrapolative of a failure to achieve an immediate virological success and a longer term immunological failure [31,37]. We found that the proportion of patients with sub-therapeutic ARV plasma concentrations was significantly high in patients with high viral loads ($\geq 400$ copies/µl) than those with low

| Table 1. Distribution of patients’ characteristics among 152 participants. |
|---------------------------------------------------------------|
| **Patient Characteristic** | **Number (%)/Mean±SD/Median [IQR]** |
| **Antiretroviral based regimen** |  |
| Efavirenz | 73 (48.0) |
| Nevirapine | 79 (52.0) |
| **Mean age in years** | 40 ± 10.0 |
| **Gender** |  |
| Female | 107 (70.4) |
| Males | 45 (29.6) |
| **Median BMI in Kg/M²** | 22.2 [20.5–24.8] |
| **Median antiretroviral concentrations (ng/ml)** |  |
| Efavirenz | 2112 [1349–3452] |
| Nevirapine | 4915 [2326–7044] |
| **NRTI backbone** |  |
| AZT+3TC | 91 (59.9) |
| D4T+3TC | 31 (20.4) |
| TDF+FTC | 30 (19.7) |
| **Median Duration on ART in months** | 40 [26–48] |
| **ARV Adherence level** |  |
| Good | 128 (84.2) |
| Poor | 24 (15.8) |
| **Median Enrolment CD4 counts (cell/µl)** | 200 [133–288] |
| **Viral Load (copies/µl)** |  |
| $\geq 400$ | 68 (44.7) |
| <400 | 84 (55.3) |
| **WHO HIV stage** |  |
| Stage 1 | 11 (7.2) |
| Stage 2 | 63 (41.4) |
| Stage 3 | 58 (38.2) |
| Stage 4 | 20 (13.2) |
| **Hepatitis B/C virus co-infection** |  |
| No | 8 (5.3) |
| Yes | 144 (94.7) |
| **Plasma ARV Drug level** |  |
| Sub-therapeutic | 43 (28.3) |
| Therapeutic | 76 (50.0) |
| Supra-therapeutic | 33 (21.7) |

*SD = Standard deviation; IQR = Interquartile range; CD4 = Cluster of differentiation; BMI = Body mass index; ARV = Antiretroviral; AZT = Azidothymidine (Zidovudine); 3TC = Lamivudine; TDF = Tenofovir; FTC = Etmicitabine; D4T = Stavudine.
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viral loads (<400 copies/μl) [39.7% versus 19.0%]. The presence of high rates of sub-therapeutic ARV concentrations among adult patients implies that these patients are standing a high risk of inadequate viral suppression and a subsequent potential of developing and accumulating resistant viral strains [4,38], if these drug concentrations are not corrected timely [39]. On the other hand the patients with supra-therapeutic plasma NNRTI, are at a high risk of developing drug toxicity [16-18] which has also been

Table 2. Comparison of distribution of patients’ characteristics within plasma antiretroviral drug concentrations (sub-therapeutic, therapeutic and supra-therapeutic) among 152 participants.

| PATIENT CHARACTERISTIC | PLASMA DRUG CONCENTRATIONS | p-value |
|------------------------|-----------------------------|---------|
|                        | SUB-THERAPEUTIC (n = 43)    | THERAPEUTIC (n = 76) | SUPRA-THERAPEUTIC (n = 33) |
| Antiretroviral based regimen |                             |                     |                           |
| Efavirenz              | 15 (20.5)                   | 41 (56.2)           | 17 (23.3)                 | 0.122 |
| Nevirapine             | 28 (35.4)                   | 35 (44.3)           | 16 (20.3)                 |       |
| Mean Age (years)       | 38.3 ± 10.4                 | 40.8 ± 9.7          | 44.0 ± 9.8                | 0.053 |
| Gender                 |                             |                     |                           |
| Female                 | 34 (31.8)                   | 50 (46.7)           | 23 (21.5)                 | 0.311 |
| Male                   | 9 (20.0)                    | 26 (57.8)           | 10 (22.2)                 |       |
| Median BMI (Kg/M²)     | 22.9 [21.1–27.2]            | 22.1 [20.6–24.6]    | 21.6 [19.0–24.2]          | 0.299 |
| NRTI backbone          |                             |                     |                           |
| AZT+3TC                | 22 (24.2)                   | 48 (52.7)           | 21 (23.1)                 | 0.039 |
| D4T+3TC                | 15 (48.4)                   | 9 (29.0)            | 7 (22.6)                  |       |
| TDF+FTC                | 6 (20.0)                    | 19 (63.3)           | 5 (16.7)                  |       |
| Median ART duration (months) | 36 [24–48] | 39 [27–48] | 45 [33–48] | 0.535 |
| ART Adherence          |                             |                     |                           |
| Good                   | 26 (20.3)                   | 70 (54.7)           | 32 (25.0)                 | <0.001 |
| Poor                   | 17 (70.8)                   | 6 (25.0)            | 1 (4.2)                   |       |
| Viral Load (copies/μl) |                             |                     |                           |
| <400                   | 16 (19.0)                   | 44 (52.4)           | 24 (28.6)                 | 0.007 |
| ≥400                   | 27 (39.7)                   | 32 (47.1)           | 9 (13.2)                  |       |
| WHO HIV Stage          |                             |                     |                           |
| Stage 1 or 2           | 18 (24.3)                   | 45 (60.8)           | 11 (14.9)                 | 0.026 |
| Stage 3 or 4           | 25 (32.1)                   | 31 (39.7)           | 22 (28.2)                 |       |
| Hepatitis B/C virus co-infection | 39 (27.1) | 73 (50.7) | 32 (22.2) | 0.399 |
| No                     | 4 (50.0)                    | 3 (37.5)            | 1 (12.5)                  |       |
| Yes                    |                             |                     |                           |

*NRTI = Nucleoside reverse transcriptase inhibitor, ARV = Antiretroviral; BMI = Body Mass Index; AZT = Azidothymidine (Zidovudine); 3TC = Lamivudine; TDF = Tenofovir; FTC = Etricitabine; D4T = Stavudine.

Table 3. Inter-individual variability for Efavirenz and Nevirapine among 152 participants.

| Antiretroviral drug | Number of patients | Mean plasmatic drug concentrations ± SD in ng/ml | Inter-individual Coefficient of variation (%) |
|---------------------|--------------------|-----------------------------------------------|---------------------------------------------|
| Efavirenz           | 73                 | 3539.2 ± 4831.5                               | 120.9                                       |
| Nevirapine          | 79                 | 5448.7 ± 4831.5                               | 88.7                                        |

*SD = Standard Deviation

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reported as a common cause of non-compliance and discontinuation of their medications [6,14]. Moreover it is well documented that drug toxicity happens commonly among patients with supra-therapeutic than among those with normal drug (therapeutic) concentrations [4,6,13,14,16,17]. In this study 13.2% of patients with supra-therapeutic plasma drug concentrations also had high viral loads. This minor proportion of patients with supra-therapeutic and yet had high viral loads might be harboring HIV drug resistant strains. So both sub-therapeutic and supra-therapeutic ARV concentrations are clinically very important in the current era of HIV medicine. However this is a great challenge in Tanzania and other resource-limited settings where TDM is not done. Therefore, it is difficult to diagnose patients with sub-therapeutic and supra-therapeutic ARV status in order to make appropriate corrections to improve virological outcome of our patients. Since this study has demonstrated that a good adherence among patients with immunological failure is significantly associated with therapeutic ARV plasma level, strict emphasis on ARV adherence on this study population could be very helpful.

Conclusion

There is a wide inter-individual variability of plasma ARV concentrations among HIV patients with immunological failure in routine clinical practice, with a large proportion of patients being outside therapeutic window. This variability is associated with ARV adherence, NRTI backbone, viral load and HIV stage. Routine therapeutic drug monitoring (TDM) could assist identifying these patients early and making timely correction to avoid immediate virological failure, long term poor immunological outcome and prevent associated drug toxicities. Good adherence is associated with therapeutic ARV plasma concentrations; therefore ARV adherence should be strictly emphasized on HIV patients with immunological failure.

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Author Contributions

Conceived and designed the experiments: CK BRK SEK GWK HK. Performed the experiments: DWG CK BRK. Analyzed the data: BRK RK JK GWK HK. Interpreted the data: CK BRK SEK GWK HK. Manuscript critically: DWG CK BRK SEK GWK HK. Conceived and designed the experiments: DWG CK BRK SEK GWK HK. Performed the experiments: DWG CK BRK. Analyzed the data: BRK RK. Interpreted the data: DWG CK BRK. Wrote the paper: DWG BRK SEK JK. Edited and reviewed the manuscript critically: DWG CK BRK RK.

References

1. Palella FJ Jr, Delaney KM, Moorman AC, Loveless MO, Fuher J, et al. (1998) Declining morbidity and mortality among patients with advanced human immunodeficiency virus infection: HIV Outpatient Study Investigators. N Engl J Med 339(13):885–60.
2. Raboud JM, Montaner JS, Conway B, Rae S, Reiss P, et al. (1998) Suppression of viral plasma load below 20 copies/ml is required to achieve a long-term response to therapy. AIDS 12(13):1619–24.
3. Gandhi M, Anelli N, Bacchetti P, Gange SJ, Anastos K, et al. (2004) Protease inhibitor levels in hair strongly predict virologic response to treatment. AIDS 23(4):471–8.
4. Bossi P, Peytavin G, Ait-Mohand H, Delaugerre C, Kooza N, et al. (2004) GENOPHAR: a randomized study of plasma drug measurements in association with genotypic resistance testing and expert advice to optimize therapy in patients failing antiretroviral therapy. HIV Med 5(5):352–9.
5. Kredo T, Van der Walt JS, Siegfried N, Cohen K (2009) Therapeutic drug monitoring of antiretrovirals for people with HIV. Cochrane Database Syst Rev (3):CD007262.
6. Marzolini C, Telenti A, Decosterd LA, Greub G, Biollaz J, et al. (2001) Antiretroviral adherence should be strictly emphasized on HIV patients with immunological failure in routine clinical practice. J Acquir Immune Defic Syndr 24(2):109–32.
7. Raboud JM, Montaner JS, Conway B, Rae S, Reiss P, et al. (1998) Suppression of viral plasma load below 20 copies/ml is required to achieve a long-term response to therapy. AIDS 12(13):1619–24.
8. Kredo T, Van der Walt JS, Siegfried N, Cohen K (2009) Therapeutic drug monitoring of antiretrovirals for people with HIV. Cochrane Database Syst Rev (3):CD007262.
9. Clevenbergh P, Moudy S, Seller P, Badia E, Gervoni J, et al. (2004) Improving ARV adherence, NRTI backbone, viral load and HIV stage. J Acquir Immune Defic Syndr 37(4):367–73.
10. Langmann P, Schirmer D, Váth T, Desch S, Zilly M, et al. (2002) Rapid determination of nevirapine and efavirenz in plasma of patients failing antiretroviral treatment. Antivir Ther 10(4):489–98.
11. Kisser JJ, Anderson PL, Gerber JG (2005) Therapeutic drug monitoring: a clinician’s point of view.Curr HIV Res 2(4):309–21.
12. Murphy RL, Sammoudoss JP, Lamson M, Hall DB, Myers M, et al. (1999) Antiviral effect and pharmacokinetic interaction between nevirapine and indinavir in persons infected with human immunodeficiency virus type 1. J Infect Dis 179(3):1116–23.
13. Kontorinis N, Dieterich DT (2003) Toxicity of non-nucleoside analogue reverse transcriptase inhibitors. Semin Liver Dis 23(2):109–79.
14. Rendon A, Nunez M, Jimenez-Nacher I, Gonzalez de Requena D, Gonzalez- Sánchez J, et al. (2003) Urological complaints in relation to indinavir plasma concentrations in HIV-infected patients. AIDS 17(4):473–8.
15. Clevenbergh P, Moudy S, Seller P, Badia E, Gervoni J, et al. (2004) Improving HIV infection management using antiretroviral plasma drug levels monitoring: a clinician’s point of view. Curr HIV Res 2(4):309–21.
16. Murphy RL, Sammoudoss JP, Lamson M, Hall DB, Myers M, et al. (1999) Antiviral effect and pharmacokinetic interaction between nevirapine and indinavir in persons infected with human immunodeficiency virus type 1. J Infect Dis 179(3):1116–23.
17. Kontorinis N, Dieterich DT (2003) Toxicity of non-nucleoside analogue reverse transcriptase inhibitors. Semin Liver Dis 23(2):109–79.
18. Dahari K, Ensou MH (2007) Efavirenz and nevirapine in HIV-1 infection: is there a role for clinical pharmacokinetic monitoring? Clin Pharmacokinet 46(2):109–22.
19. van Leth F, Lange JM (2006) The use of the non-nucleoside reverse transcriptase inhibitors nevirapine and efavirenz in the treatment of patients with a chronic HIV-1 infection. Ned Tijdshr Geneeskd. 150(31):1719–22.
20. Molto J, Clotet B (2004) Therapeutic drug monitoring of antiretroviral agents. J HIV Ther 9(4):75–8.
21. de Maat MM, Huitema AD, Mulder JW, Meenhorst PL, van Gorp EC, Mairuhu AT, et al. (2003) Subtherapeutic antiretroviral plasma concentrations in routine clinical outpatient HIV care. Ther Drug Monit. 25(3):367–73.
22. Khow SH, Lloyd J, Dalton M, Bonington A, Hart E, et al. (2006) Pharmacologic optimization of protease inhibitors and non-nucleoside reverse transcriptase inhibitors (PONI)-a randomized controlled trial of therapeutic drug monitoring and adherence support. J Acquir Immune Defic Syndr 41(4):461–7.
23. Leth VF, Kappelhof BS, Johnson D, Losso MH, Boron-Kaczmarska A, et al. (2006) Pharmacokinetic parameters of nevirapine and efavirenz in relation to antiretroviral efficacy. AIDS Res Hum Retroviruses 22(3):232–9.
24. Back D, Ganti G, Fletcher C, Garaffo R, Haulrich R, et al. (2000) Therapeutic drug monitoring in HIV infection: current status and future directions. AIDS Suppl 1:S3–7.
25. Best BM, Gaiocechea M, Witt MD, Liar D, Saar ES, et al. (2007) A randomized controlled trial of therapeutic drug monitoring in treatment-naïve and -experienced HIV-1-infected patients. J Acquir Immune Defic Syndr 46(4):433–42.
26. Bollito M, Acosta E, Burger D, Fletcher CV, Flexner C, et al. (2005) Therapeutic drug monitoring and drug-drug interactions involving antiretroviral drugs. Antivir Ther 10: 469–477.
27. Reynolds SJ, Nakigosi G, Newell K, Nyanabo A, Galwongos R, et al. (2009) Failure of immunologic criteria to appropriately identify antiretroviral treatment failure in Uganda. AIDS. 23(8):697–700.
28. Grossberg R, Gross R (2007). Use of pharmacy refill data as a measure of antiretroviral adherence. Current AIDS Reports 4:187–191.
29. Langmann P, Schirmer D, Vath T, Zilly M, Klinker H (2001) High-performance liquid chromatographic method for the determination of HIV-1 non-nucleoside reverse transcriptase inhibitor efavirenz in plasma of patients during highly active antiretroviral therapy. J Chromatogr B Biomed Sci 785(2):143–6.
30. Langmann P, Schirmer D, Vath T, Zilly M, Klinker H (2001) High-performance liquid chromatographic method for the determination of HIV-1 non-nucleoside reverse transcriptase inhibitor efavirenz in plasma of patients during highly active antiretroviral therapy. J Chromatogr B Biomed Sci 785(2):143–6.
31. Langmann P, Schirmer D, Vath T, Zilly M, Klinker H (2001) High-performance liquid chromatographic method for the determination of HIV-1 non-nucleoside reverse transcriptase inhibitor efavirenz in plasma of patients during highly active antiretroviral therapy. J Chromatogr B Biomed Sci 785(2):143–6.
33. Ahoua L, Guenther G, Pinoges L, Anguzu P, Chaix ML, et al. (2009) Risk factors for virological failure and subtherapeutic antiretroviral drug concentrations in HIV-positive adults treated in rural northwestern Uganda. BMC Infect Dis 9:81.
34. Fabbiani M, Bracciale L, Ragazzoni E, Santangelo R, Cattani P, et al. (2011) Relationship between antiretroviral plasma concentration and emergence of HIV-1 resistance mutations at treatment failure. Infection 39(6):563–9.
35. Alexander CS, Asselin JJ, Ting LS, Montaner JS, Hogg RS, et al. (2003) Antiretroviral concentrations in untimed plasma samples predict therapy outcome in a population with advanced disease. J Infect Dis 188(4):541–8.
36. Young B (2005) Review: mixing new cocktails: drug interactions in antiretroviral regimens. AIDS Patient Care STDS 19(5):286–97.
37. Boulle A, Van Cutsem G, Cohen K, Hilderbrand K, Mathee S, et al. (2008) Outcomes of nevirapine- and efavirenz-based antiretroviral therapy when coadministered with rifampicin-based antitubercular therapy. JAMA 300(5):590–9.
38. Robertson SM, Penzak SR, Pau A (2007) Drug interactions in the management of HIV infection: an update. Expert Opin Pharmacother 8(17):2947–63.
39. Kasang C, Kalluyia S, Majinge C, Stich A, Bodem J, et al. (2011) HIV Drug Resistance (HIVDR) in Antiretroviral Therapy-Naïve Patients in Tanzania Not Eligible for WHO Threshold HIVDR Survey Is Dramatically High. PLoS ONE 6(8): e23091. doi:10.1371/journal.pone.0023091.