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### **► To cite this version:**

Christine Danel, Raoul Moh, Gilles Peytavin, Amani Anzian, Albert K. Minga, et al.. Lack of indinavir-associated nephrological complications in HIV-infected adults (predominantly women) with high indinavir plasma concentration in Abidjan, Côte d'Ivoire.. *AIDS Research and Human Retroviruses*, 2007, 23 (1), pp.62-6. 10.1089/aid.2006.0038 . inserm-00173523

**HAL Id: inserm-00173523**

**<https://inserm.hal.science/inserm-00173523>**

Submitted on 20 Sep 2007

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# **Lack of indinavir-associated nephrological complications in HIV-infected adults (predominantly women) with high indinavir plasma concentration in Abidjan, Côte d'Ivoire**

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## **Abstract**

### **Objective**

To report the tolerance of indinavir combined with ritonavir (IDV/r 800/100mg) twice daily (bid) in sub-Saharan African HIV-infected adults.

### **Design**

Prospective cohort study.

### **Methods**

HAART-naïves patients started zidovudine plus lamivudine plus IDV/r 800/100mg bid. Follow-up included: standardised documentation of morbidity; CD4+ cell count, creatininemia, plasma HIV-1 RNA, and IDV minimal plasma concentration ( $C_{min}$ ) measurements at month-1 (M1), M3 and M6.

### **Results**

70 HIV-1 infected adults (68 women, median CD4 235/mm<sup>3</sup>) started HAART. At M6, 63% had undetectable viral load, and the median gain in CD4 since baseline was +128/mm<sup>3</sup>. During the first six months, 21 patients experimented 23 treatment modifications (reduction to IDV/r 400/100mg bid: n=11; switch for efavirenz: n=11; zidovudine replaced by stavudine: n=1), including 22 for digestive intolerance and one for severe anaemia. At M1, M3 and M6, 67, 59 and 48 patients were still receiving IDV/r 800/100 mg bid, of whom 70%, 72% and 60% had IDV  $C_{min}$  above 5 ng/mL, respectively. In these patients, at M1, M3 and M6, the mean ( $\pm$  SD) IDV  $C_{min}$ , were 3431  $\pm$  3835ng/ml, 2288  $\pm$  2116ng/ml and 1543  $\pm$  2398ng/ml, respectively. There was no renal insufficiency of any grade, and no symptom of urinary stones.

### **Conclusion**

The IDV/r 800/100mg bid containing regimen led to high IDV  $C_{min}$ , and high rate of digestive intolerance. There was a surprising lack of nephrological side-effects during the 6 months of follow-up, supporting the hypothesis that nephrological tolerance of IDV might be higher in sub-Saharan African individuals than in American or European ones.

**MESH Keywords** Adult ; Cohort Studies ; Cote d'Ivoire ; Female ; Follow-Up Studies ; HIV Infections ; blood ; drug therapy ; virology ; HIV Protease Inhibitors ; adverse effects ; blood ; therapeutic use ; HIV-1 ; isolation & purification ; HIV-2 ; isolation & purification ; Humans ; Indinavir ; adverse effects ; blood ; therapeutic use ; Male ; RNA, Viral ; blood ; Retrospective Studies ; Time Factors ; Treatment Outcome ; Viral Load

## **Introduction**

The benefits of pharmacokinetically enhanced protease inhibitor (PI) regimens using ritonavir (RTV) as a booster have long been demonstrated [1, 2]. RTV boosting reduces the number of drug intake – thus also reducing the treatment cost –, and diminishes the constraints of food and water requirements, especially for Indinavir (IDV) [3]. RTV-boosted PI (PI/r) regimens are currently rarely employed in sub-Saharan Africa, because RTV soft capsules need to be stored below 25°C. Although World Health Organisation (WHO) experts recognize that PI/r regimens represent valuable alternatives to non nucleoside reverse transcriptase inhibitors (NNRTI)-based regimens [4, 5], no experience of their use has been reported in sub-Saharan Africa so far.

We report here the 6 month-efficacy and tolerance of a highly active antiretroviral treatment (HAART) based on an indinavir/ritonavir (IDV/r) 800/100 mg twice daily (bid) regimen in a cohort of HIV-1 infected patients in Abidjan, the economic capital city of Côte d'Ivoire, West Africa.

## Patients and Methods

In December 2002, a randomised trial (Trivacan ANRS 1269) of structured treatment interruptions (STI) was launched in Abidjan [6]. The trial protocol was approved by the ethics committee of the Ministry of Health of Côte d'Ivoire and the institutional review board of the French Agency of Research on AIDS (ANRS, Paris, France). Inclusion criteria in the trial have been previously described [7]. Between inclusion and randomisation in the STI arms, patients received at least 6 months of continuous HAART. We describe here the 6-month outcomes of treatment-naïve patients who started an IDV/r-based regimen at inclusion in the pre-randomisation phase of the trial.

At inclusion, and every month thereafter, patients received 35 days of treatment, i.e. 70 tablets of Duovir<sup>®</sup> (ZDV 300 mg and 3TC 150, Cipla LTD, Mumbai, India), 140 tablets of Crixivan<sup>®</sup> (IDV 400 mg, Merk Sharp and Dohme, Haarlem, The Netherlands) and 70 soft capsules of Norvir<sup>®</sup> (RTV 100 mg, Abbott, Zwolle, The Netherlands). At inclusion, patients were recommended to ingest a minimum of two liters of water per day. They were asked to return to their study centre at day seven, at month-1 (M1) and every month thereafter. During these monthly visits, questionnaires were administered to record adherence to treatment during the previous four days. Patients had free access to the study clinics whenever they had a medical problem. Symptoms were managed following standardized algorithms of treatment [8].

At inclusion, M3 and M6, blood cell count (MaxM<sup>®</sup> coulter, Beckman, Paris, France), CD4 count (True Count<sup>®</sup> technique on FACScan<sup>®</sup>, Becton Dickinson, Aalst-Erembodegem, Belgium), plasma HIV-1 RNA (real-time PCR on Taq Man technology Abi Prism 7000, Applied Biosystems, limit of detection at 300 copies/mL) [9], serum liver enzymes and creatinemia were measured. Additional samples were collected at M1, M3 and M6 for plasma and serum depository (storage in a -80°C deep-freezer). Patients were asked not to take their morning drugs before month-1, month-3 and month-6 blood sample collections. Blood collections were thus performed early in the morning, in patients whose last drug intake was on the previous evening.

In August 2005, all deep-frozen stored samples were sent to France for IDV and ZDV plasma concentration measurements by means of high performance liquid chromatography coupled with UV-detection (respective LOQ 5 and 10 ng/mL with previously described methods) [10,11]. Patients with a ZDV plasma concentration above 10 ng/mL were suspected to have taken their treatment within the preceding eight hours. Consequently, their IDV plasma concentrations were considered as non-minimal concentrations (non-C<sub>min</sub>). For all other patients, the IDV concentration was classified as minimal concentration (C<sub>min</sub>). The value of 10 ng/mL eight hours after the last drug intake was chosen according to the pharmacokinetic characteristics of ZDV in HIV-infected patients with normal renal function [12].

## Results

From December 2002 to August 2003, 70 HIV-1 infected or HIV-1+2 infected patients started ZDV-3TC-IDV/r. Baseline and follow-up characteristics of patients are shown in table 1.

During the first six months of follow-up, no patient died or was lost to follow-up. From M1 to M6, all percentages of actual/scheduled monthly visits were above 92%. At M1, M2, M3, M4, M5 and M6, respectively 16%, 17%, 19%, 9%, 10% and 7% of the patients declared that they had missed at least one antiretroviral drug intake during the previous four days. The median gain in CD4 since baseline was +124/mm<sup>3</sup> (interquartile range [IQR] +49;+218) at M6. At M3 and M6, 45 (64%) and 44 (63%) patients had plasma HIV RNA below 300 copies/mL, respectively.

During the first six months, 21 patients (31%) experimented 23 HAART regimen modifications, including 21 patients with 22 regimen modifications for digestive intolerance (reduction to IDV/r 400/100mg bid: n=10; switch for efavirenz: n=10; reduction to 400/100mg bid followed by switch for efavirenz: n=1) and one patient with a modification for severe anaemia (zidovudine replaced by stavudine). All digestive intolerance events were severe vomiting, including four with admission to the day care hospital for intravenous treatment, two with associated sub-icterus, and one with grade 3 elevation of transaminases. There was no other grade 3 or grade 4 hepatitis.

No renal insufficiency of any grade and no symptom evocative of uro-nephrological lithiasis were documented. The median creatinine clearance was estimated at 114 ml/min (IQR 103;132) at baseline and 100 ml/min (80;119) at M6 (median difference between baseline and M6: -12 ml/min, IQR -32;+ 0.0).

Table 2 shows the number of patients who: (1) were still receiving IDV/r 800/100 mg bid, (2) had been reduced to IDV/r 400/100mg bid, and (3) had been switched for a non IDV-containing regimen at M1, M3 and M6. Figure 1 describes IDV plasma C<sub>min</sub> and non-C<sub>min</sub> in patients still receiving IDV/r 800/100 mg bid.

## Discussion

Within our cohort study of sub-Saharan African adults starting a IDV/r 800/100 mg bid containing regimen, the percentage of patients with an undetectable viral load at month-6 and the mean gain in CD4 after 6 months of follow-up were in the lower bracket of those previously described in industrialized countries [13 –15 ]. Meanwhile: (i) the percentage of patients with severe digestive intolerance leading to IDV/r dosage reduction or to IDV/r replacement by another drug reached 30%; (ii) the mean IDV concentration in patients still on the 800/100 mg bid dosage decreased within time, suggesting that a significant proportion of patients who were not asked by the medical team to reduce the IDV/r dosage to 400/100 mg bid probably decided by themselves to interrupt the treatment; (iii) IDV  $C_{min}$  were in the upper bracket of those previously reported in patients receiving the 800/100 mg bid regimen [3 ,14 ,16 ,17 ]; (iv) In spite of these high concentrations, there was no episode of uro-nephrological side-effects.

To our knowledge, this is the first time that an experience with IDV/r is reported in sub-Saharan Africa. In industrialized countries, urinary stones have been reported to occur in 18% to 33% of patients on IDV/r 800/100 mg bid [13 ,14 ]. IDV nephrological toxicity has been associated with a high urinary pH, a low diuresis, a high concentration of the parent drug or its metabolites in urine, an increasing area under the curve, an IDV  $C_{min}$  above 500 ng/ml, a low lean body mass, warm weather environments, and the concomitant use of aciclovir [16 ,18 ,19 –22 ]. In tropical areas with high temperatures and high levels of humidity, the IDV concentration in urine would have been expected to be higher, thus resulting in higher uro-nephrological toxicity [22 , 23 ]. Furthermore, the urinary pH of HIV-infected adults living in West Africa has been reported to be higher than that of European adults, and this should also result in a higher frequency of IDV related nephrolithiasis in Africa [24 ]. Finally, most of our patients were women, contrary to most previous studies on IDV/r in industrialized countries. Previous studies demonstrated a higher IDV apparent bioavailability in females than in males or suggested a sex-related metabolism of IDV, but this is not likely to explain the lack of nephrological complications in our study [25 ,26 ].

One limitation of our study is the 6-month follow-up, compared with higher durations of exposure to IDV in previous studies in industrialized countries [3 ,14 ,16 ,17 ]. However, the risk of IDV-associated renal complications has been reported to be higher in patients with IDV treatment < 74 weeks than in those with longer exposure [21 ], and indinavir crystalluria is frequently observed soon after initiation of therapy [27 ]. Furthermore, our patients were followed up under cohort conditions, with no patients lost to follow up and with free access to a study clinic at any time. Under these circumstances, the frequency of urinary stones is not likely to have been greatly underestimated. Previous studies showed an increasing risk of urinary stones in Caucasian US patients [28 ] and a low incidence of IDV nephrological side effects in a population predominantly American with African ancestry [15 ]. Following these reports, our data support the hypothesis of a lower frequency of IDV-associated urinary stones in sub-Saharan African adults.

In our study, we were unable to assess the efficacy and tolerance of the IDV/r 400/100 mg dosage, because of the reduced number of patients who switched from the IDV/r 800/100 mg bid to the 400/100 mg bid dosage and of the consecutive reduced followed-up time on the 400/100 mg bid dosage. In industrialized countries, IDV/r is now given at the 400/100 mg bid dosage [29 ,30 ], provided that plasma concentration measurement-driven dosage adaptations are possible. In most sub-Saharan African areas, therapeutic drug monitoring is not routinely available. Given the low digestive tolerance – leading to low adherence -, and despite of the good nephrological tolerance of the 800/100 mg bid dosage in our study, 400/100 mg bid should probably be the recommended first-line dosage of IDV/r in sub-Saharan Africa. However, this remains to be carefully evaluated in terms of efficacy and tolerance.

## Acknowledgements:

This study was supported by the French Agence Nationale de Recherches sur le SIDA et les Hépatites Virales (ANRS), and the Ivoirian Ministry of Public Health within the collaborative Programme PAC-CI.

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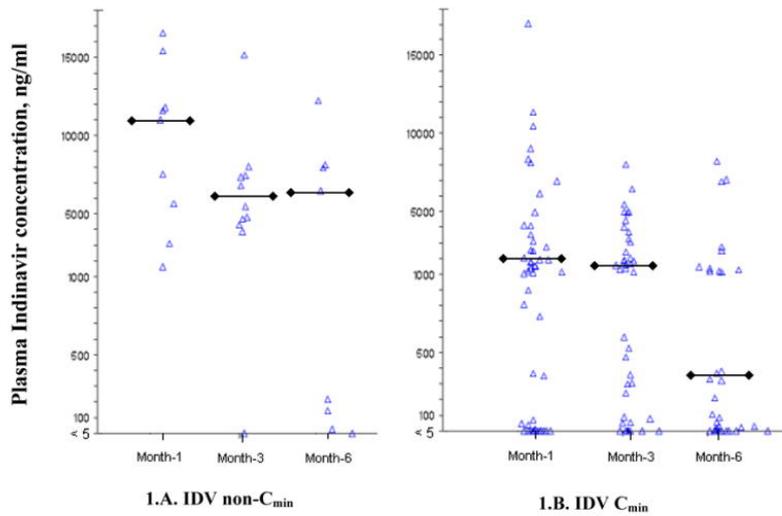
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**Figure 1**

Plasma IDV concentration at M1, M3 and M6 in patients treated with IDV/r (800/100 mg bid) 1A: non-IDV  $C_{min}$  (patients with ZDV plasma concentration equal or above 10 ng/mL) 1B: IDV  $C_{min}$  (patients with ZDV plasma concentration below 10 ng/mL) Each plot represents one IDV concentration measurement

: median value among patients with IDV concentration  $\geq 10$  ng/ml C: concentration; IQR: interquartile range



	IDV Non- $C_{min}$			IDV $C_{min}$		
	Month-1	Month-M3	Month-M6	Month-1	Month-M3	Month-M6
IDV $C < 5$ ng/ml, n (%)	0 (0%)	1 (9%)	1 (13%)	16 (30%)	13 (28%)	16 (40)
IDV $C \geq 5$ ng/ml, n (%)	9 (100%)	10 (91%)	7 (87%)	37 (70%)	33 (72%)	24 (60)
Median, ng/ml	11024	5490	6349	1939	1663	376
IQR, ng/ml	5681-11795	4327-7470	85-8068	1051-4137	471-3730	72-1422

**Table 1**

Baseline and follow-up characteristics

**Baseline**

Male/Female, n (%)	2/68	(3/97%)
Age (years), median (IQR)	31	(29 – 36)
Dual VIH 1+2 infected, n (%)	3	(4%)
WHO clinical stage, n (%)		
1	15	(21%)
2	32	(46%)
3	18	(26%)
4	5	(7%)
CD4 count (/mm <sup>3</sup> ), median (IQR)	235	(194 – 313)
Plasma HIV RNA (log <sub>10</sub> /ml), median (IQR)	4.9	(4.1–5.4)
Haemoglobin level (g/l), median (IQR)	11.3	(10.4 – 12)
Creatinine clearance (ml/min), median (IQR)	113.7	(102–132)
Positive serum HBs antigen, n (%)	7	(10%)

**Follow-up**

Cumulative (person-months)	427	
Per patients (months), median (IQR)	6.1	(6.1-6.1)
Plasma HIV RNA at M		
log <sub>10</sub> /ml, median (IQR)	0.0	(0.0–3.9)
< 300 copies/mL, n (%)	44	(63%)

n: number of patients; %: percentage; IQR: interquartile range; g/l: gramme per litre; /mm<sup>3</sup> : per cubic millimeter**Table 2**

number of patients with IDV/r 800/100 and IDV/r 400/100 bid-containing regimens at month-1, month-3 and month-6

HAART regimen	Number of patients at		
	Month-1	Month-3	Month-6
IDV/r 800/100 bid-containing regimen	67	59	49
ZDV plasma concentration unavailable	5	2	1
ZDV plasma concentration ≥ 10 ng/ml *	9	11	8
ZDV plasma concentration < 10 ng/ml **	53	46	40
IDV/r 400/100 bid-containing regimen	0	3	10
Non IDV-containing regimen	3	8	11

\* Patients with detectable ZDV plasma concentrations were suspected to have taken their antiretroviral drugs within the preceding eight hours. For these patients, the IDV plasma concentration was classified as non-C<sub>min</sub> (see figure 1A )\* Patients with undetectable ZDV plasma concentrations were confirmed not to have taken their antiretroviral drugs within the preceding eight hours. For these patients, the IDV plasma concentration was classified as C<sub>min</sub> (see figure 1B ).