Response to a Dolutegravir-based Regimen in an HIV-Infected Woman with Multiple Comorbidities and a Highly Resistant Strain

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Abstract HIV-infected patients have a higher burden of comorbidities than the general population and drug-drug interactions limit the choice of antiretroviral compounds, also limiting the possibility of drug sequencing in case of toxicity or failure. In our patient, affected by HIV-related pulmonary hypertension (HIV-PAH), the use of sildenafil and ambrisentan excluded the possibility of using two of the main classes of antiretrovirals, protease inhibitors (PIs) and non-nucleoside reverse transcriptase inhibitors (NNRTIs). Having failed a combination regimen based on raltegravir, while awaiting the results of genotypic testing for drug resistance she was switched to dolutegravir, a brand new integrase strand transfer inhibitor (INSTI) with high genetic barrier, and in one month her HIV-1 viremia dramatically dropped to undetectable levels. The genotypic test revealed resistance to all the drugs she was taking including dolutegravir. The backbone was changed to a dual regimen including rilpivirine and viral suppression was maintained at three months. Unprecedented pharmacokinetic data are provided for the two antiviral drugs in combination with sildenafil and ambrisentan.

Keywords: HIV, PAH, dolutegravir, sildenafil, ambrisentan, resistance

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1. Introduction

Pulmonary arterial hypertension (PAH) is a rare disease, that left untreated has an estimated median survival of 2.8 years [1]. Survival, however, has dramatically improved in the last decade due to advances in treatment options [2]. Among HIV-infected subjects with PAH, survival has improved from 58%, 32%, and 21% at 1, 2, and 3 years, respectively, in the 1990s, to 88%, 72%, and 63% at 1, 3, and 5 years, respectively [3]. Antiretroviral therapy improves survival, but specific PAH treatment limits the choice of antiretrovirals. Moreover, antihypertensive drugs may cause liver damage or circulatory effects that can sum to antiretroviral therapy side effects and toxicity. Also, in several subjects, the presence of HCV coinfection or liver decomposition adds the burden of porto-pulmonary shunts with porto-pulmonary arterial hypertension. Therefore an HIV-infected subject is a particularly delicate person, in whom any error may lead to serious consequences.

2. Case Report

A 47 year old hispanic female was diagnosed and treated for Hodgkin lymphoma in 2000, underwent total hysterectomy for uterine cancer in 2006, and in September 2010 had an episode of pneumonia, after which pulmonary arterial hypertension (PAH) was diagnosed. Looking for a possible aetiology, an HIV antibody test was performed and resulted positive in December, 2010. The patient had never tested for HIV before and had had heterosexual contacts. Her plasma HIV-1 RNA was 34.030 copies/mL and she had 174 CD4+ T lymphocytes/mm3. Her mean pulmonary arterial pressure (PAPs) was 100 mmHg at cardiac ultrasound, right heart catheterization showed severe pulmonary hypertension (84 mmHg) with increased vascular and total – systemic and pulmonary - resistance, with a slightly reduced cardiac output (3.25 L/min). CT scan, x-rays, scintigraphy and blood work did not reveal other causes. A thyroid suspect nodule was secondarily found during the assessments for PAH. Referred to our hospital, combination antiretroviral therapy (cART) with tenofovir/emtricitabine plus raltegravir was started on December 13, 2010, together with anti-hypertensive treatment with sildenafil 20 mg q8h, while the thyroid nodule was aspirated, yielding the diagnosis of papillary thyroid carcinoma, that was surgically removed.
Radiotherapy was performed and subsequently the patient started levothyroxine 100 mg/day. One year later a cardiac ultrasound revealed a hypokinetic right ventricle with further increase of pulmonary pressure (112 mmHg) and initial rectilinearization of the interventricular septum. Ambrisentan 5 mg/day was added with important reduction of hypertension (PAPs 45 mmHg, July, 2012). The patient was classified as NYHA III. The situation stabilized in the following years, but in summer 2014 her adherence to therapy worsened and she did not present for her scheduled blood work in July. In October, 2014, a stress test revealed serious functional aerobic deficit with an important deficiency of the pulmonary perfusion, and the 6 minutes’ walking test had worsened (400 meters). The blood work performed the 19th of November, 2014 showed immunovirologic failure, with HIV RNA rising to 4661 copies/mL and CD4+ T cells falling to 292/mmC. A genotypic test was requested and in the meanwhile, having very little choice available, only raltegravir was changed to dolutegravir 50 mg q12h, waiting for the results to make a rational treatment change. At the visit of January, 7, 2015, the genotypic test revealed the presence of K65R and M184V resistance-associated mutations (RAMs) in rev causing high-level resistance to all nucleoside/nucleotide analogues except zidovudine while the sequencing of the integrase region yielded RAMs L74M, G140S and Q148H, conferring high-level resistance to all INSTIs; both the TRUGENE Guidelines™ Rules 17.0 and the Stanford HIVdb algorithms were concordant in data interpretation (Figure 1). Despite this apparent complete resistance to the modified regimen – as well as to the previous one, the HIV-1 RNA plasma level had decreased in four weeks to completely undetectable, therefore it was decided to maintain dolutegravir, switching from tenofovir/emtricitabine to rilpivirine, the only NNRTI free from drug-drug interactions with her comediations. At week 12, the patient maintains optimal viral suppression, her immunity is recovering (+24 CD4+ T cells/mmc, +2.5%, Figure 2) and the subject tolerates the new regimen very well. Plasma minimum concentrations (Cmin) of dolutegravir (12h) was quite high, while that of rilpivirine (24h) was perfectly in the range. No ECG alterations were recorded, as dolutegravir does not impact on QT prolongation, however some concern may be raised (Figure 3). No cancer recurrence has developed up to date. The 6 minutes’ walking test has improved again (from 400 to 435 meters).

3. Discussion

The pathogenesis of HIV-related PAH is complex and not entirely understood. Many factors, as viral proteins, HIV-related inflammation, coinfections with other microorganisms, and genetic susceptibility were considered to contribute to HIV-related PAH. Recently, a direct role of HIV infection to endothelial damage through the release of different inflammatory mediators such as endothelin-1 has been suggested [4]. Even the effect of antiretroviral therapy on pulmonary artery pressure has been an area of controversy. A recent study suggested the possible role of antiretroviral therapy in improving pulmonary artery pressure in HIV/AIDS patients if started at early stages (WHO classes I and II). Any significant effect was observed at more advanced stages [5]. In our
case report, the patient showed a worsening in pulmonary perfusion and in the 6 minutes’ walking test at the moment of viral failure and an improvement after viral suppression with the new INSTI, suggesting a possible direct virologic role in PAH evolution. However, since viral failure was a consequence of reduced compliance to HIV medications, a poor compliance to specific drugs may provide a simpler explanation.

Figure 2. Evolution of HIV-1 RNA, CD4 cell counts, resistance mutations and pulmonary arterial pressure

Vasodilator and anti-endothelin receptor A drugs are essential in the treatment of PAH, but can limit the choice of antiretroviral drugs, due to possible drug-drug interactions. Sildenafil is a phosphodiesterase 5 inhibitor (PDE5 inhibitor) commonly used in PAH at high doses. At these doses is contraindicated with antiretroviral regimens based on protease inhibitors (PI) or non nucleoside inverse transcriptase inhibitors (NNRTI). Coadministration with PIs is expected indeed to increase sildenafil blood concentrations, possibly resulting in adverse reactions [6]. On the other hand, coadministration with NNRTIs (except rilpivirine) may lead to a decrease in sildenafil blood concentrations, with possible reduction in clinical response. This is true especially for etravirine, that leads to a decrease in sildenafil AUC by 57% and in Cmax by 45% [7]. No data are available for efavirenz and nevirapine, but similar pharmacokinetic interactions are expected. Ambrisentan and bosentan are endothelin receptor A (ETa) antagonists used for PAH, metabolized by the hepatic cytochrome 3A (CYP3A). Thereby, the concomitant use of antiretrovirals inhibiting CYP3A, as for example PIs, or inducing CYP3A, as for example NNRTIs, may lead to an increase or reduction in ETa antagonists blood concentration [8]. Moreover, antihypertensive drugs may cause liver toxicity or circulatory effects that can sum to antiretroviral therapy side effects [9].

Among antiretrovirals, the INSTI class presents a favorable metabolic and pharmacokinetic profile, except for elvitegravir that needs to be boosted by the CYP3A4 inhibitor cobicistat. Raltegravir and dolutegravir (DTG) may represent thereby a good choice in patients with comorbidities, to limit the potential risk of drug-drug interactions and the amount of toxicity.

In clinical trials on naïve patients, 50% of subjects failing raltegravir developed raltegravir RAMs, and strains harboring more than two RAMs were rare [10]. The presence of single INSTI RAMs (Q148H/R/K, N155H, Y143R/H/C, E92Q e T66I) does not reduce the in vitro susceptibility to dolutegravir [11,12], however, when combined with at least two secondary mutations, a Q148 RAM leads to a 5-10 fold (or more) susceptibility reduction to DTG. The Viking study is a phase III randomized clinical trial, in which ARV experienced
patients harboring raltegravir and/or elvitegravir resistance were randomized to DTG 50 mg twice daily or placebo plus the failing regimen for 7 days (functional monotherapy phase) [13]. At day 8, all patients switched to open-label DTG 50 mg BID plus optimized background therapy (≥ 1 fully active drug). The study resulted in a day 8 plasma HIV-1 RNA mean change from baseline of −1.43 log10 c/mL and in a 69 % virologic suppression rate (VL <50 c/mL) at week 24. Baseline INI resistance (Q148 + ≥2 associated mutations) and viral load were highly significant predictors for week 24 response. The odds of achieving HIV-1 RNA <50 c/mL were 63% lower for every 2-fold increase in DTG fold change in 50% inhibitory concentration (FC) and about 80% lower for every 10-fold increase in baseline HIV-1 RNA. The odds of achieving virologic undetectability resulted moreover 96% lower in subjects with virus harboring Q148 + ≥2 mutations compared with those with no evidence of Q148 mutations.

Our patient’s genotypic test showed mutations conferring resistance to almost all nucleoside analogues (65R, 184V) and mutations in the integrase region (148H, 65R, 184V) and mutations in the IN region (108N, 109P) that led to a reduction in the sensitivity to DTG. Of note, only 8 patients in the Viking study had total resistance to nucleoside analogues (backbone genotypic susceptibility score = 0) and virologic success at 24 weeks was achieved in 4 of the 8 subjects. As seen in the first days of the Viking study, in our case report DTG has led after 1 month to optimal suppression of viral replication (with a 2.11 log10 copies/mL drop in viremia), despite the almost total suppression of viral replication (with a 2,11 log10 inhibitory concentration (IC) and about 80% lower for every 10-fold increase in baseline HIV-1 RNA. The odds of achieving virologic undetectability resulted moreover 96% lower in subjects with virus harboring Q148 + ≥2 mutations compared with those with no evidence of Q148 mutations.

Once resistance to nucleoside analogues was acknowledged, our patient was switched to a dual treatment with dolutegravir and rilpivirine, allowing the maintenance of virologic suppression. This appears to be a simple regimen, very promising for the DTG high genetic barrier and for the pharmacokinetic profile of both drugs, that may deeply improve the patient’s compliance and quality of life in the future [14]. Dolutegravir Cmin was more than twice the expected level, and this suggests a significant change in blood concentration.

4. Conclusion

Patients affected with HIV-PAH have a limited treatment choice as compared with the general HIV-infected population and a higher mortality rate. Dolutegravir, due to its high genetic barrier and low potential for drug-drug interactions, is a very interesting option in this setting. Rilpivirine, too, was not affected by the use of specific PAH drugs. Possible future long-acting formulations of these drugs are therefore adequate to this subgroup of patients.

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