

SIMPLIFICATION TO SINGLE-DRUG REGIMEN WITH A RITONAVIR-BOOSTED PROTEASE INHIBITOR FOR HIV PATIENTS

Sánchez Gundín J., Flor García A., Recuero Galve L., Martí Gil C., Martínez Valdivieso L., Barreda Hernández D.
Pharmacy Department. Virgen de la Luz Hospital, Cuenca (Spain).



Background

Single-drug regimens (SDR) with a ritonavir-boosted protease inhibitors (PI) could potentially be a regimen simplification to avoid nucleoside reverse transcriptase inhibitor (NRTI) toxicities in patients carrying human immunodeficiency virus (HIV) who fulfill several requirements:

- ✓ virologic suppression,
- ✓ high level of medication adherence,
- ✓ no previous PI virologic failure,
- ✓ high CD4 count level (>100 cell/mcL).

Purpose

To evaluate effectiveness and security of SDR with ritonavir-boosted lopinavir (Lp/r) and ritonavir-boosted darunavir (Dr/r) in HIV-positive patients pre-treated with three-drug regimens (TDR) including NRTI.

Material and methods

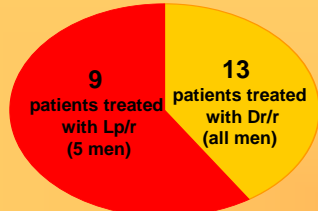
Retrospective observational study of HIV-positive patients with **treatment switches from TDR to SDR** in a second-level hospital.

Data were collected from the Farmatools-Dominion®-Programme and medical records.

Variables included:

- ✓ sex,
- ✓ age,
- ✓ duration of previous TDR,
- ✓ plasma viral load (PVL) pre- and post-treatment switching,
- ✓ PI virologic failure,
- ✓ CD4 cell count before switching,
- ✓ months of SDR to date (June '11-September '14).

Results



Patient characteristics

- Mean age at the time of the study: 48 ± 6 years.
- 4 patients were HIV/Hepatitis C virus co-infected.

2 patients with Lp/r

2 patients with Dr/r

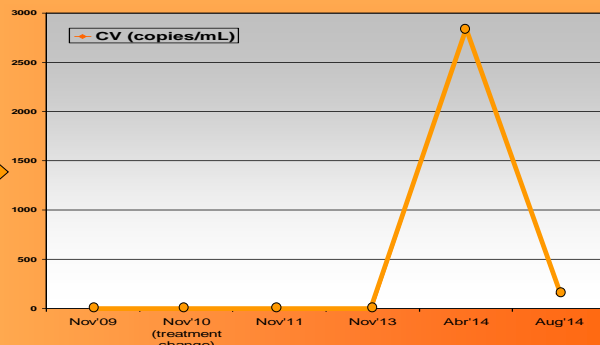
All subjects:

- had been treated with TDR during minimum 12 months prior to treatment change,
- basal PVL was undetected for at least 6 months before switching,
- basal PVL remained undetectable during the entire study,
- no presented previous PI virologic failure,
- the medium CD4 counts at treatment switch were normal (825±583 cell/mcL),
- were treated with SDR for a median period of 22 months.

1 exception

Adherence and tolerance were considered successful before and after switching.

Patient with confirmed viral rebound which led to treatment re-intensification with two NRTI included in the previous TDR.



Conclusions

- ✓ SDR with a ritonavir-boosted PI might be an alternative as effective as traditional combinations.
- ✓ It involves a clear benefit for HIV-positive patients because it provides a treatment simplification with minor toxicity and minor interaction number.