

Switching to dual therapy in elderly and multiexperienced patients: profile of a reference service in Brazil



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BACKGROUND

With the current availability of drugs with greater potency, tolerability and genetic barrier, interest in antiretroviral-sparing strategies to reduce toxicity, regimen complexity and costs has resurfaced (1). Current studies have already shown that this appears to be a safe option (2)(3)(4), but little has been studied to date in long-lived and multi-experienced populations.

MATERIAL AND METHODS

To evaluate the profile of patients who switched to dual regimens (DTG + 3TC or DTG + DRV/r or DRV/r + 3TC) at the ADEE 3002 outpatient clinic. Retrospective analysis with data collected from April 2021 to December 2023 from PLWHA followed at the ADEE3002/HCFMUSP outpatient clinic, São Paulo. The patients evaluated were switched to the dual regimen having been undetectable for at least 6 months and with no reported resistance. Data were retrieved from medical records.

RESULTS

The ADEE3002 outpatient clinic currently has 430 active patients, of which 34 are eligible for our analysis. The main characteristics of this population analyzed are: Men 29/34 (85.29%), mean age 55.6 years, mean time of HIV infection 18.5 years, mean CD4 nadir 327.44, previous diagnosis of advanced HIV in 10/34 (29.41%) and previous opportunistic infection in 7/34 (20.6%).

The average time of exposure to ARV was around 16 years, the average number of previous regimens was 4.12, exposure to integrase inhibitors 20/34 (58.8%), exposure to protease inhibitors 21/34 (61.76%). Only 8/34 (23.5%) of patients did not have any comorbidity. Among the main comorbidities were dyslipidemia 19/34 (55.9%), renal dysfunction 16/34 (47%), systemic arterial hypertension 14/34 (41.2%), type II diabetes 7/34 (20.6%), psychiatric comorbidities 6/34 (17.6%), lipodystrophy 6/34 (17.6%), osteopenia or osteoporosis 4/34 (11.8%), neurological sequelae 4/34 (11.8%).

RESULTS

After 12 months of exchange, 32/34 (94.11%) remained undetectable. No virological failure or need to change the regimen was detected in the patients analyzed. CD4 T lymphocyte values remained without significant changes.

CONCLUSIONS

Even in long-lived and multi-experienced populations, dual therapy regimens with DTG + 3TC or DRV/r + 3TC or DTG+DRV/r appear to be safe options in the management of comorbidities and adverse effects in PLWHA undergoing viral suppression without prior resistance.

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Abbreviations: Dolutegravir (DTG); Lamivudine (3TC); Darunavir / Ritonavir (DRV/r); People living with HIV AIDS (PLWHA)