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Poster Session VI

HIV infection - clinical profile

THE PROGNOSIS OF HAART-TREATED HIV INFECTED WOMEN IN RESOURCE-LIMITED SETTINGS FROM 1998-2012 IN BELGRADE, SERBIA

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Objectives: Gender differences, mostly due to differences in access to highly active antiretroviral therapy (HAART) and/or innate biological factors, combined with scarce data from clinical studies concerning the efficacy and side effects of HAART solely in women still remain relevant questions in HAART era. Therefore we conducted a study to determine possible factors influencing prognosis and outcome of HIV-infected women in resource-limited settings.

Methods: A cross-sectional study was performed on HIV infected women, who initiated HAART between 1st January 1998 and 31st December 2012, at the HIV/AIDS Center of the University Hospital for Infectious and Tropical Diseases in Belgrade, Serbia, with regular clinical and laboratory check-ups and obtained consent.

Results: A total of 230 women were followed for 8.2 ± 3.4 years (range 1-12). During the follow-up, 26 patients died. The mean age of the patients at HAART initiation was 37 ± 9.7 years. Clinical AIDS at presentation was observed in 43.9% of the patients, while 80% of them had CD4 cell counts below 200 cell/mm³. Univariate and stepwise multivariate analysis have shown that the progression to death was associated with basal CD4 counts below 100 cells/ μ L (OR 3.0 95% CI 1.7-8.4, $P=0.02$) and HCV co-infection (OR 2.6 95% CI 1.0-6.6, $P=0.03$). However the NNRTI based regimens and good adherence to HAART (OR 0.2, 95% CI 0.09-0.6, $P=0.005$ and OR 0.3, 95% CI 0.1-0.66, $P=0.03$, respectively), all prevented death. Although in patients with sustained viral suppression the CD4 counts varied significantly, these did not affect overall survival ($p=0.21$, log rank)

Conclusion: A significant number of participants were late-presenters with advanced immunodeficiency, in whom hepatitis co-infection and low nadir CD counts affected overall survival. However in case of optimal viral suppression up to a mean 8 years of treatment, regardless of the level of immune recovery, their prognosis may be fairly good even in a resource limited settings.