CHARACTERISTICS OF INDIVIDUALS WITH HETEROSEXUALLY- ACQUIRED COMPARED WITH HOMOSEXUALLY- ACQUIRED HIV AND IMPLICATIONS FOR CLINICAL PRACTICE.

Authors:

<u>Herbert SE¹</u>, Puhr R², Lewis DA^{3,4}, Varma R^{2,5}, Couldwell DL³, Petoumenos K², Law M², Templeton DJ^{1, 2,4}; on behalf of the Australian HIV Observational Database (AHOD)

¹RPA Sexual Health, Sydney Local Health District, Sydney, ²The Kirby Institute, UNSW Sydney, ³Western Sydney Sexual Health Centre, ⁴Sydney Medical School, University of Sydney, ⁵Sydney Sexual Health Centre, South Eastern Sydney Local Health District, Sydney, NSW, 2000.

Background:

Little is known about the clinical characteristics of Australian individuals who acquire HIV heterosexually.

Methods:

We compared patients enrolled in AHOD since 1999 who reported only heterosexual (Het-HIV) or only homosexual (Hom-HIV) exposure as likely mode of HIV acquisition. Multivariate models were adjusted *a priori* for age, sex, country of birth, hepatitis B (HBV) and C (HCV) serology, CD4 count and viral load at treatment initiation, and year of combination antiretroviral therapy (cART) initiation.

Results:

806 Het-HIV and 3125 Hom-HIV patients were included and contributed 5,287 and 25,731 person-years of follow-up, respectively. Median age at diagnosis was 33.6 vs 33.3 years for Het-HIV vs. Hom-HIV. Compared with Hom-HIV, Het-HIV were more often born outside Australia (51.4% vs 30.5%, p<0.001), had lower median CD4 counts at both diagnosis (307 vs 465 cells/μL, p<0.001) and cART initiation (277 vs 320, p<0.001), and were less likely to have had past exposure to HCV (6.6% vs 9.1%, p=0.045) but not HBV (p=0.808). Previous AIDS diagnoses were no different between groups (p=0.323). Among those with Het-HIV, time to viral suppression was no different (aHR 1.08, 95% CI 0.95-1.24), although the risk of virological failure was lower in Het-HIV (aHR 0.77; 95% CI 0.61-0.98). Among Het-HIV, the risk of loss-to-follow-up (LTFU) was higher (p=0.009) and the risk of all-cause mortality was lower (aHR 0.62; 95%CI 0.39-1.00). Overall, Het-HIV achieved similar immunological recovery despite cART initiation at lower CD4 counts, although CD4 reconstitution was lower for Het-HIV among the subgroup who started cART after 2006.

Conclusions:

Compared with Hom-HIV, Het-HIV had lower CD4 counts at diagnosis and cART initiation, lower risk of virological failure and lower risk of all-cause mortality. However, a higher rate of LTFU among Het-HIV suggests greater efforts may be required for this group to maintain engagement in care.

Disclosure:

The Australian HIV Observational Database is funded as part of the Asia Pacific HIV Observational Database, a program of amfAR, The Foundation for AIDS Research; and is supported in part by grant no. U01AI069907 from the U.S. National Institutes

of Health's National Institute of Allergy and Infectious Diseases, the Eunice Kennedy Shriver National Institute of Child Health and Human Development, the National Cancer Institute, the National Institute of Mental Health, and the National Institute on Drug Abuse, and by unconditional grants from Merck Sharp & DoHome; Gilead Sciences; Janssen-Cilag; ViiV Healthcare. The Kirby Institute is funded by the Australian Government Department of Health, and is affiliated with the Faculty of Medicine, UNSW Australia. The content is solely the responsibility of the authors and the views expressed in this publication do not necessarily represent the position of the Australian Government or the official views of the U.S. National Institutes of Health or other funders.