Retrovirology



Poster presentation

Open Access

Long-term intrapatient evolution during HIV-2 infection Adam MacNeil*1, Jean-Louis Sankalé1, Seema Thakore Meloni1, Abdoulaye Dieng Sarr1, Souleymane Mboup2 and Phyllis Kanki1

Address: ¹Department of Immunology and Infectious Diseases, Harvard School of Public Health, Boston, Massachusetts, USA and ²Cheikh Anta Diop Universite, Dakar, Senegal

* Corresponding author

from 2006 International Meeting of The Institute of Human Virology Baltimore, USA. 17–21 November, 2006

Published: 21 December 2006

Retrovirology 2006, 3(Suppl 1):P38 doi:10.1186/1742-4690-3-S1-P38

© 2006 MacNeil et al; licensee BioMed Central Ltd.

HIV-2 disease progression and transmission is attenuated compared to HIV-1, yet prospective studies examining HIV-2 intrapatient evolution have been limited. We examined viral sequence evolution in the C2V3C3 region of the viral env gene in 8 HIV-2 infected individuals from Dakar, Senegal, over the course of approximately 10 years. To compare results to HIV-1 infection, we reanalyzed data from our previous study that examined intrapatient evolution in HIV-1 infected individuals from the same population. HIV-2 sequences from early and late timepoints were phylogenetically intermixed for all subjects, and no distinct trends were observed in terms of increases or decreases in fragment size or number of N-linked glycosylation sites. In homologous env C2V3 sequence, rates of viral divergence and diversification were slower in individuals infected with HIV-2 than individuals with HIV-1. This data indicates that viral evolution occurs slowly in HIV-2 infection, which is consistent with the slow disease progression observed in HIV-2 infection, and supports the notion that viral evolution may be a relevant correlate for disease progression.