

Oral presentation

Open Access

HIV-1-Specific T Cell Function During Acute HIV-1 Infection

Marcus Altfeld*‡

Address: Research Center and Infectious Disease Division, Massachusetts General Hospital and Division of AIDS, Harvard Medical School, Boston, Massachusetts 02129, USA

Email: Marcus Altfeld* - maltfeld@partners.org

* Corresponding author ‡Presenting author

from 2005 International Meeting of The Institute of Human Virology
Baltimore, USA, 29 August – 2 September 2005

Published: 8 December 2005

Retrovirology 2005, **2**(Suppl 1):S104 doi:10.1186/1742-4690-2-S1-S104

HIV-1-specific CD8+ T cells in primary infection are associated with the dramatic decline of peak viremia to the viral set point, while their antiviral activity in chronic infection is less apparent. Here, we comparatively analyzed functional properties of HIV-1-specific CD8+ T cells in primary and chronic infection, and demonstrate that the functional avidity and TCR affinity of HIV-1-specific CD8+ T cells was consistently higher in primary infection than in chronic infection. The change of TCR affinities between primary and chronic infection was linked to an almost complete switch in the clonotypic composition of epitope-specific CD8+ T cells, resulting from the preferential loss of high-avidity CD8+ T cell clones. These data suggest that the initial recruitment of high-avidity HIV-1-specific CD8+ T cell may contribute to the control of HIV-1 viremia during primary infection, while their selective elimination during the subsequent disease process contributes to the loss of immune control during chronic infection.