



POSTER PRESENTATION

Open Access

Regulatory B cells are induced in untreated HIV-1 infection and suppress HIV-1 specific T cell responses

J Liu^{1*}, W Zhan¹, C Kim¹, E Lee¹, J Cao¹, B Ziegler¹, A Gregor¹, F Yue¹, S Huibner¹, S Macparland¹, K Clayton¹, J Schwartz¹, H Song¹, E Bento², C Kovacs², R Kaul¹, M Ostrowski¹

From AIDS Vaccine 2012
Boston, MA, USA. 9-12 September 2012

Background

Regulatory B cells (Breg), the B cells producing interleukin 10 (IL-10), have been identified in mice and humans. Mouse Breg can suppress innate and T cell responses and are implicated in pathogenesis of some autoimmune diseases and immune evasion of some pathogens. However, the role of Breg in humans is less clear.

Methods

PBMC and gut biopsy samples were obtained from healthy donors and HIV infected individuals. Flow cytometry and Luminex were used to quantify cytokine production. Flow cytometry were used to analyze Breg's phenotype.

Results

Breg were elevated in both peripheral blood and gut tissue of untreated HIV-1 infected individuals and the elevation correlated with viral load in early HIV-1 infection. Breg from HIV-1 infected individuals were CD19⁺TIM-1⁺. Anti-retroviral therapy could reduce elevated Breg frequency. Treatment of B cells from healthy donors with microbial translocation products could differentiate them toward a Breg phenotype. Ex vivo Bregs from HIV-1 infected individuals suppressed cytokine production /degranulation of HIV-1 specific T cells that was in part IL-10 dependent.

Conclusion

Our findings show that Bregs are induced early in HIV-1 infection, which may play a role in inhibiting effective HIV-1-specific T cell responses.

Author details

¹University of Toronto, Toronto, Canada. ²Maple Leaf Clinic, Toronto, Canada.

Published: 13 September 2012

doi:10.1186/1742-4690-9-S2-P102

Cite this article as: Liu et al.: Regulatory B cells are induced in untreated HIV-1 infection and suppress HIV-1 specific T cell responses. *Retrovirology* 2012 **9**(Suppl 2):P102.

Submit your next manuscript to BioMed Central and take full advantage of:

- Convenient online submission
- Thorough peer review
- No space constraints or color figure charges
- Immediate publication on acceptance
- Inclusion in PubMed, CAS, Scopus and Google Scholar
- Research which is freely available for redistribution

Submit your manuscript at
www.biomedcentral.com/submit



¹University of Toronto, Toronto, Canada
Full list of author information is available at the end of the article