



**HAL**  
open science

## **HIV triggers immunoregulatory dendritic cells and regulatory T cells through the non-canonical NF- $\kappa$ B pathway**

Olivier Manches, Melissa Victoria Fernandez, Joël Plumas, Laurence Chaperot, Nina Bhardwaj

### **► To cite this version:**

Olivier Manches, Melissa Victoria Fernandez, Joël Plumas, Laurence Chaperot, Nina Bhardwaj. HIV triggers immunoregulatory dendritic cells and regulatory T cells through the non-canonical NF- $\kappa$ B pathway. *Retrovirology*, 2012, 9 (Suppl 2), pp.P184. inserm-00731775

**HAL Id: inserm-00731775**

**<https://inserm.hal.science/inserm-00731775>**

Submitted on 13 Sep 2012

**HAL** is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



POSTER PRESENTATION

Open Access

# HIV triggers immunoregulatory dendritic cells and regulatory T cells through the non-canonical NF- $\kappa$ B pathway

O Manches<sup>1\*</sup>, MV Fernandez<sup>1</sup>, J Plumas<sup>2</sup>, L Chaperot<sup>2</sup>, N Bhardwaj<sup>1</sup>

From AIDS Vaccine 2012

Boston, MA, USA. 9-12 September 2012

## Background

HIV stimulates plasmacytoid dendritic cell (pDC) through TLR7 and induce the secretion of high levels of IFN $\alpha$ . pDC stimulated by HIV also upregulate the expression of the enzyme indoleamine 2,3 dioxygenase (IDO). IDO is critical for the induction of regulatory T cells (Treg) by HIV-activated pDC. We investigated the molecular mechanisms of IDO induction and its consequences for Treg function.

## Methods

The cells used were purified primary pDC and the GEN pDC cell line. A combination of siRNA knock-down, immunoprecipitation of TLR signaling pathway molecules, IDO promoter engineering and chromatin immunoprecipitation was used to determine the molecular mechanisms of IDO induction in pDC. To analyze Treg function and interaction with conventional DC (cDC), blocking antibodies to CTLA-4 and CTLA-4-Ig were used.

## Results

We demonstrate that HIV induces activation of the non-canonical NF- $\kappa$ B pathway in pDC, and is essential for IDO induction. TLR7 triggering induces recruitment of TRAF3 to the TLR-MyD88 complex, followed by release of NIK and phosphorylation of IKK $\alpha$ . Activation of the non-canonical NF- $\kappa$ B pathway culminates in p52/RelB nuclear translocation and binding to the IDO promoter.

Furthermore, IDO-expressing pDC trigger the generation of Treg, which dampen cDC activation through CTLA-4. CTLA-4 also induces IDO expression in cDC

in a NIK-dependent fashion, allowing cDC to induce Treg from naïve CD4<sup>+</sup> T cells.

## Conclusion

The non-canonical NF- $\kappa$ B pathway plays a central role in regulating IDO expression in pDC and cDC upon HIV infection, and may be a potential target for regulating Treg activity in chronic or acute HIV infection.

## Author details

<sup>1</sup>NYU Cancer Institute, New York City, NY, USA. <sup>2</sup>Inserm, U823, Immunobiologie et Immunothérapie des Cancers, La Tronche, France.

Published: 13 September 2012

doi:10.1186/1742-4690-9-S2-P184

Cite this article as: Manches et al.: HIV triggers immunoregulatory dendritic cells and regulatory T cells through the non-canonical NF- $\kappa$ B pathway. *Retrovirology* 2012 **9**(Suppl 2):P184.

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



<sup>1</sup>NYU Cancer Institute, New York City, NY, USA  
Full list of author information is available at the end of the article