

Oral presentation

Open Access

Human herpesvirus 8 (HHV-8) “in vitro” infection of human placental histocultures

Mariantonietta Di Stefano*¹, Maria Luisa Calabro², Iole Maria Di Gangi², Santina Cantatore³, Massimo Barbierato⁴, Luigi Chieco-Bianchi⁴, Pantaleo Greco⁵, Loreto Gesualdo¹, Elisabeth Menu⁶ and José Ramon Fiore⁷

Address: ¹Laboratory of Molecular Medicine, University of Foggia, Foggia, Italy, ²Istituto Oncologico Veneto, IRCCS, Immunology and Diagnostic Molecular Oncology, Padova, Italy, ³Laboratory of Histology, School of Medicine, University of Foggia, Italy, ⁴Department of Oncology and Surgical Sciences, Oncology Section, University of Padova, Padova, Italy, ⁵Department of Surgical Sciences, University of Foggia, Foggia, Italy, ⁶Unité de Régulation des Infections Rétrovirales, Institut Pasteur, Paris, France and ⁷Department of Clinical and Occupational Health, University of Foggia, Foggia, Italy

* Corresponding author

from Fourth Dominique Dormont International Conference. Host-Pathogen Interactions in Chronic Infections Paris, France. 13-15 December 2007

Published: 9 April 2008

Retrovirology 2008, **5**(Suppl 1):O4 doi:10.1186/1742-4690-5-S1-O4

This abstract is available from: <http://www.retrovirology.com/content/5/S1/O4>

© 2008 Di Stefano et al.; licensee BioMed Central Ltd.

Background

Most human Herpesvirus infect placental cells and may be harmful in pregnancy, leading to obstetrical and/or neonatal complications. Although a correlation between human herpesvirus 8 (HHV-8) infection and abortion or low birth weight in children has been reported [1,2] presently no information has been published regarding HHV-8 tropism for placenta.

Materials and methods

In this study, a placenta histoculture system was used to evaluate the susceptibility of placental cells to “in vitro” HHV-8 infection. Quantitative detection of HHV-8 was performed by real-time PCR, and virus expression was evaluated by immunohistochemistry for latent and lytic HHV-8 antigens.

Results

Increasing amounts of HHV-8 DNA were detected in placental tissues and culture supernatants and immunohistochemistry analyses demonstrated that both cyto- and syncytiotrophoblasts, as well as placental endothelial cells, expressed latent (see Figure 1) and lytic antigens. In addition, relevant apoptotic phenomena were observed in infected histocultures.

Conclusions

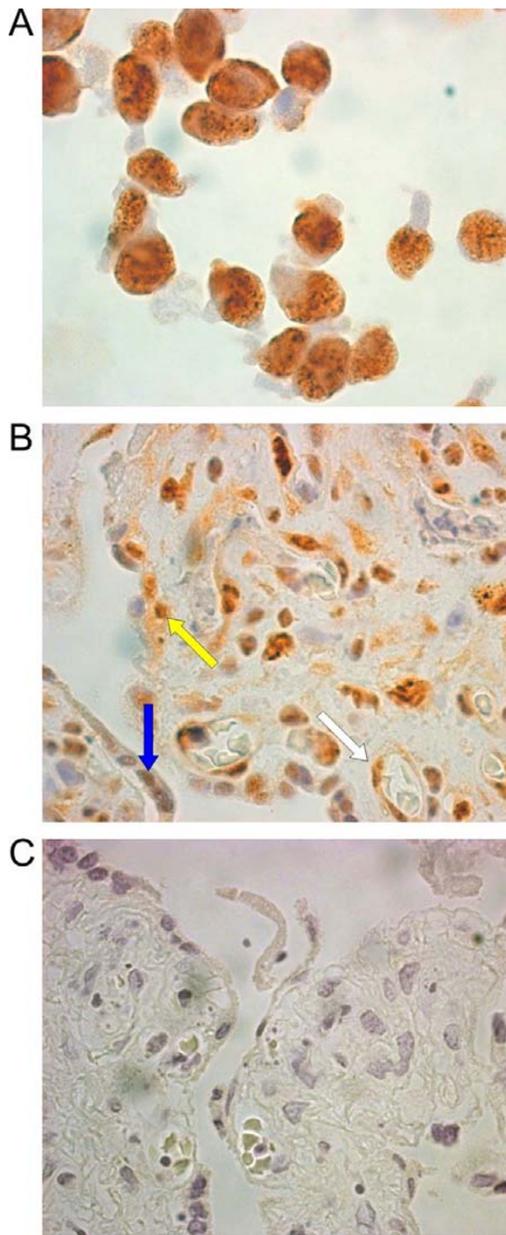
We here demonstrated for the first time that HHV-8, like other human herpesviruses, may productively infect placental cells in vitro, thus providing evidence that this phenomenon might influence vertical transmission and pregnancy outcome in HHV-8-infected women.

Acknowledgements

This work was supported by grants from Istituto Superiore di Sanità (grant n. 50G.3, MLC; grant n. 50G.29, JRF) and from Associazione Italiana per la Ricerca sul Cancro (MLC). MB was the recipient of a fellowship from Associazione Italiana per la Lotta contro le Leucemie, Linfomi e Mieloma (AIL).

References

1. Gaye-Diallo A, Toure AT, Gessain A, Gueye-Ndiaye A, Ndour AN, et al.: **Preliminary study of human Herpesvirus type 8 infection in pregnant women in Dakar (Senegal).** *Bull Soc Pathol Exot* 2001, **94**:231-234.
2. Sarmati L, Ticconi C, Santangelo R, Montano M, Rezza G, et al.: **Does the risk of abortion increases in women with high human herpesvirus-8 antibody titers?** *J Infect Dis* 2003, **188**:173-174.

**Figure 1**

Immunohistochemical detection of the HHV-8 LANA protein in placental histocultures. Specific reactivity was visualized with immunoperoxidase staining using anti-LANA-1 monoclonal antibodies with a DAB developer (brown colour) and haematoxylin counterstaining. (A) HHV-8-infected CRO-AP/3 cells showed a strongly positive nuclear immunostaining. (B) HHV-8-infected placental histocultures showed positive immunostaining in cytotrophoblasts (yellow arrow), syncytiotrophoblasts (blue arrow) and endothelial cells (white arrow). (C) Mock-infected placental histocultures. Original magnifications, X100.