

## The Effect of Interferon Beta in HLA-G Expression on Monocyte in Diabetes Type1

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### Abstract

**Background and objectives:** Dendritic Cells are the most important of antigen presenting cells with an effective role in immune tolerance. This study, aims to clarify the role of IFN-  $\beta$  in induction on dendritic cells derived by monocyte in diabetes type1 to evaluate the T cells response to beta cell specific antigenic molecule.

**Material and Methods:** In this research, peripheral blood mononuclear cells were isolated by phiCole and then dendritic cells generated from blood monocytes, in Seven days, by adding granulocyte-monocyte colony stimulating factor and interleukin-4 with or without IFN-beta. mRNA was extracted by dendritic cells and cDNA was produced by reverse transcriptase enzyme. Then, Specific polymerase chain reaction for HLA-G was performed. In addition, Tcell proliferation with a mixed Leukocytic reaction evaluated between dendritic cell and T by means of MTT.

**Results:** based on the results, IFN- $\beta$  induces HLA-G molecule on dendritic cells. In addition, T cell proliferation responses in mixed leukocyte culture show significance difference between Case and control  $p < 0.05$ . T cell proliferation was inhibited in their co-culture system affected by IFN- $\beta$

**Conclusion:** In this study, we show that dendritic cells-treated IFN- $\beta$  with expression of HLA-G molecule inhibited T cell proliferation and so, our results suggest that some of the IFN-  $\beta$  regulatory effects with expression of HLA-G can probably prevent from beta cell destruction.

**Key words:** dendritic cells, Interferon Beta, Human Leukocytic Antigen-G.