



## P/ Impact of LPS on IFN- $\gamma$ secretion by human T-cell lines

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**Background and Aim:** Lipopolysaccharide (LPS), an outer membrane component of Gram-negative bacteria, has an important role in inflammation including acute lung inflammatory responses. Furthermore modulatory effects of LPS on inflammatory cytokines have been shown. Interferon gamma (IFN- $\gamma$ ) as a T helper type 1 (Th1) cytokine is a key regulator of inflammatory responses. Moreover IFN- $\gamma$  plays an essential role in defense against intracellular pathogens. This study was conducted to evaluate the impact of LPS on IFN- $\gamma$  secretion by human leukemic (Jurkat and Molt-4) T cell lines in vitro.

**Methods:** Jurkat and Molt-4 cells were cultured in whole RPMI-1640 media. The cells were distributed at a density of  $2 \times 10^6$  cell/ml. The cells were stimulated with different concentrations of lipopolysaccharide (LPS) (1-4  $\mu$ g/ml) for 48 hours. Then the cell-conditioned medium was used for IFN- $\gamma$  assay. The different groups were compared by the analysis of variance.

**Results:** LPS did not show any significant effect on IFN- $\gamma$  production in human leukemic T cells compared with unstimulated cells.

**Conclusion:** According to the results of this study, LPS did not have any significant effect on IFN- $\gamma$  secretion in human leukemic T cells. So LPS-induced inflammatory responses in lung and other tissues may be due to LPS impacts on other inflammatory mediators. Further studies about LPS effects on IFN- $\gamma$  expression in another immune cells in vitro as well as in vivo are warranted.

**Key words:** LPS, T- cells, IFN- $\gamma$