In vitro effects of polyvalent immunoglobulin for intravenous use

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High dose intravenous immunoglobulin (IVIg) therapy is used in the treatment of a wide range of auto-immune or presumed immune-mediated neurological disorders. A beneficial effect of IVIg has been reported in Guillain-Barré syndrome,1 chronic inflammatory demyelinating polyneuropathy,2,3 myasthenia gravis,4 multiple sclerosis,5 multifocal motor neuropathy,6-12 intractable childhood epilepsy,13 and dermatomyositis.14-15

The mechanisms of response to IVIg treatment in these disorders is not fully understood. Firstly, adequate models to study the effects of IVIg are lacking. Secondly, treatment with IVIg has probably multiple effects on induction and amplification, as well as the effector phase of the immune response.

Effects of IVIg on the effector phase of the immune response

Most studies on IVIg have dealt with the effector phase of the immune response. It has been demonstrated that IVIg has the following effects in vivo: inhibition of autoantibody production,16 neutralisation of circulating pathogenic antibodies,17-22 and decrease of antibody dependent cellular cytotoxicity (ADCC) by Fc receptor blockade.22 In vitro it was shown that IVIg inhibits pokeweed mitogen stimulated antibody production,23-26 contains anti-idiotypic antibodies against multiple auto-antibodies,27 contains antibody dependent cellular cytotoxicity (ADCC),28-30 suppresses ADCC,30,32 decreases natural killer cell function,33 and interferes with complement-activation.34,35 These mechanisms are dependent on the presence of the intact IgG molecule, which has a half-life of approximately three weeks in vivo.36 However, the observed long term effect of IVIg treatment, in some diseases for up to three months, is not fully explained by the modes of action already mentioned. IVIg may therefore modulate auto-immunity at other levels of the immune response.

Effects of IVIg on the induction and amplification phase of the immune response

ANTIPROLIFERATIVE EFFECTS

The effects of IVIg on the induction and amplification of the immune response in which the CD4+ helper T-cell interacts with antigen presenting cells (in a way that stimulatory signals are activated) has not yet been explored. Interactions of IVIg with T-cells have been scarcely investigated.37 An enhancement of CD8 positive suppressor T-cell function in vivo has been reported.38-40 Furthermore, IVIg appears to interfere with the cytokine network. In vitro administration of IVIg results in induction of the interleukin-1 receptor antagonist and downregulation of the production of IL-1 and tumour necrosis factor-a. Moreover, antibodies against IL-1a have been demonstrated in IVIg.41-43

We recently showed inhibitory effects of IVIg on antigen-specific and non-specific stimulated lymphocytes.44 Subsequent experiments showed that IVIg added to cultures of various autonomously growing cell lines (including the haematopoietic cells HL-60, Daudi, Molt-4 and K562, B-cell hybridomas, EBV transformed B-cells and neuroblastoma cell lines of various human and non-human origin) also ceased with proliferation. We demonstrated that IVIg interfered with DNA-synthesis and protein metabolism.3-H-thymidine uptake of activated cells was measured, after incubation with IVIg for various intervals. IVIg interferes with 3-H-thymidine uptake, in a dose-dependent fashion, regardless of the stimulus. To confirm these observations, cells cultured with and without IVIg, were pulse-labelled with bromodeoxyuridine (BrdU) and subsequently analysed in an immunofluorescence-assay. The BrdU labelling studies showed that cells were arrested in the G1 - G2 - and no longer progressed into the S-phase. Similar results were obtained with normal human myoblasts and fibroblasts in culture. Parallel experiments with similar concentrations of human albumin did not show these effects. There was no apparent cytotoxic effect, as removal of IVIg within an interval of several days and reculturing in fresh medium, resulted in normal proliferation of K562, HL-60, EBV transformed B-cells, human myoblast and fibroblast. Further experiments are needed to exclude that a proportion of cells are induced to apoptosis.

GROWTH FACTOR SIGNALLING

A possible explanation for this broad proliferative arrest could be the interference of IVIg with essential growth factors. II-1a, II-2, II-3
Putative effects of IVIg on induction, amplification and effectors phase of the immune response.

**Induction and activation phase**
Effect on signal transduction
- Inhibitory effect on antigen-specific stimulated lymphocytes
- Inhibitory effect on antigen non-specific stimulated lymphocytes (PHA, PWM)
- Enhancement of CD8 positive suppressor T-cell function
- CD4 helper T-cell activation

**Amplification phase**
- Effect on co-stimulatory cells
- Effect on accessory cells
- Interference with the cytokine network

**Proliferation and maturation phase**
- Interference with growth factors (receptors and production)
- Interference with DNA synthesis
- Interference with protein metabolism
- Interference with antibody production

**Effectors phase**
- Binding to FcγRII, III receptors
- Neutralisation of circulating pathogenic antibodies
- Decrease of ADCC by Fcγ receptor blockade
- Decrease of natural killer cell function
- Interference with complement activation

**Conclusion**
Our results show that, in addition to the F(ab')-mediated anti-idiotypic neutralisation of autoantibodies, the Fc-mediated immunoregulation of B-cell function and the depression of monocyte effector ADCC-function.

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