Detection of cytokine mRNA production in infiltrating cells in proliferative vitreoretinopathy using reverse transcription polymerase chain reaction

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Abstract

Aims—To determine whether the infiltrating cells in the vitreous and subretinal fluid of patients with proliferative vitreoretinopathy (PVR) express messenger RNA for various cytokines found in this condition.

Methods—The presence of mRNA coding for HPRT, IL-6, IL-1β, IL-8, and TNFα was investigated in 20 vitreous and subretinal fluid (SRF) samples from patients with PVR by reverse transcriptase polymerase chain reaction (RT-PCR). 16 samples from patients with retinal detachment and macular holes were used as controls.

Results—HPRT was detected in all samples of PVR and in 11 (69%) control cases. Patients with PVR demonstrated mRNA for the cytokines tested more often than controls. The difference was statistically significant.

Conclusion—The presence of mRNA encoding for IL-6, IL-1β, IL-8, and TNFα is significantly detected by RT-PCR in vitreous and SRF samples of patients with PVR, indicating local production of these cytokines by vitreous and SRF cells.

Proliferative vitreoretinopathy (PVR) is a process of cellular proliferation and contraction that complicates rhegmatogenous retinal detachment. PVR is the most common cause of failure when the primary break has been appropriately treated.¹ ² The exact pathogenic mechanisms involved in the formation of PVR are not completely understood. However, five distinct stages appear to be important in its development including breakdown of the blood-retinal barrier (BRB), chemotaxis and cellular migration, cellular proliferation, membrane formation, and contraction.³ ⁴ ⁵

It is believed that different cytokines, produced as a consequence of immune or inflammatory reactions, are involved in the pathogenesis of PVR.⁶ ⁷ Cytokines such as interleukin 1 (IL-1), interleukin 6 (IL-6), tumour necrosis factor α (TNFα), interferon γ (IFNγ), interferon 8 (IFN8), and monocyte chemotactic protein 1 (MCP-1) have been detected in the vitreous samples as well as epiretinal membranes obtained from PVR patients.⁸ ⁹ ¹⁰ ¹¹ ¹² ¹³ ¹⁴ ¹⁵

It is not known whether these cytokines are produced locally by the cells infiltrating the vitreous or diffuse into the vitreous cavity following a breakdown of the BRB. We therefore tested the infiltrating cells in vitreous obtained from patients with PVR, for mRNA encoding for HPRT (housekeeping gene), IL-1β, IL-6, TNFα, and IL-8.

Materials and methods

VITREOUS AND SRF SPECIMENS

Twelve vitreous and eight subretinal fluid (SRF) samples were obtained by vitrectomy from 18 patients with PVR (both vitreous and SRF were obtained from two patients). Ten vitreous samples obtained from patients with macular hole and six vitreous samples from patients with retinal detachment (RD), not complicated by PVR, were examined as a control group. The presence of mRNA coding for HPRT, IL-6, IL-1β, TNFα, and IL-8 was investigated by reverse transcriptase polymerase chain reaction (RT-PCR).

The severity of PVR was graded according to the criteria of the Retina Society Terminology Committee.¹² Samples were obtained through conventional three port closed vitrectomy technique by manual suction before opening the infusion line. SRF samples were obtained during the drainage procedure with a Charles flute needle connected to a 1 ml syringe with a side port. Samples with vitreous haemorrhage were excluded from the study. The study complied with the declaration of Helsinki.

mRNA EXTRACTION AND cDNA SYNTHESIS

Cellular pellets were obtained from the samples by centrifugation and mRNA extracted using Qiagen RN easy (UK) method following the manufacturer’s procedure. Eluted RNA was made up to 50 µl with DEPC water and cDNA was prepared using Oligo-(dT) priming in ready to go cDNA synthesis tubes (Pharma-
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Vitreous samples. Control (PBMC), M = mimic (2nd positive control), N = negative control, 1–6 represent samples from patients with PVR. Std = 100 bp DNA, ND = not determined, P = positive control (PBMC). M = mimic (2nd positive control), N = negative control, 1–6 represent vitreous samples.

The χ² test (statistics program, spss for Windows) was used to test for significance between presence of cytokines in the PVR samples versus the control group.

Results

HPRT mRNA was detected in all samples from PVR (100%) and in 11 of 16 control samples (69%). Of the 20 samples of PVR, IL-6 mRNA was detected in 16 samples (80%), (10/12 vitreous (83%) and 6/8 SRF (75%)). IL-1β mRNA was detected in 12 samples (60%) (6/12 vitreous (50%) and 6/8 SRF (75%)). TNFα mRNA was detected in 10 samples (50%) (6/12 vitreous (50%) and 4/8 SRF (50%)). IL-8 mRNA was detected in 15 samples (75%) (9/12 vitreous (75%) and 6/8 SRF (75%)). In all, seven samples were positive for all cytokines tested. Six of the seven patients who tested positive for all cytokines had PVR graded between C2 and D1.

Only a small proportion of samples from the control group tested positive for cytokine mRNA. Four out of 16 (25%) were positive for IL-6, one (6.25%) for IL-1β, one (6.25%) for TNFα, and three (18.75%) for IL-8 (Table 1). This difference in the presence of cytokine mRNA between samples from the PVR group and the control group, for each cytokine tested, was highly significant (p = 0.00097, 0.00085, 0.00080, and 0.00463, for IL-6, IL-1β, and the control group, for each cytokine tested, respectively).

Discussion

The results obtained from RT-PCR analysis of vitreous and SRF samples from patients undergoing vitrectomy for PVR provide evidence for the expression of mRNA for IL-6, IL-1β, TNFα, and IL-8.

Previous studies have been based on the detection of protein secretion and quantitation using bioassay and enzyme labelled immunosorbent assay (ELISA) techniques. Bioassays which measure functional properties, depending on the responsiveness of indicator cells lines, are technically restrictive. These techniques require in vitro stimulation, which could result in distortion of the true in vivo response. Techniques based on ELISA are highly specific but do not distinguish between biologically active or inactive substances. Furthermore, these assays determine the extracellular presence of a secreted interleukin, reflecting the net outcome of produced, absorbed, and degraded cytokine.
Our approach to the role of cytokines in PVR pathogenesis was to directly detect cytokine mRNA from cells within the vitreous and SRF. This approach provides information on the possibility of local production of the cytokines in question, within the vitreous cavity, by the infiltrating cells. In addition, analysis of mRNA can detect the presence of a cytokine which may be absent in culture supernatants due to uptake and utilisation by cells bearing receptors. Thus, mRNA profiles can provide a fairly accurate account of potential cytokine production. On the downside, however, the presence of cytokine mRNA does not guarantee that it will be translated. Potential regulation at transcriptional, post-transcriptional, translational, and post-translational levels may occur.

To our knowledge the only work to detect the presence of cells expressing mRNA coding for IL-6, IL-1β, TNFα, and IL-8 in epiretinal membranes from patients with PVR was carried out by Limb et al. They used an in situ hybridisation technique which, although sensitive, is technically very demanding to quantify. Owing to the different distribution of cytokines between the vitreous and epiretinal membranes in patients with PVR, the above study data did not provide information on cytokine production in the vitreous. This information was obtained by our approach. Both studies, however, have a similar drawback in that the demonstration of mRNA does not necessarily imply the presence of the protein product. RT-PCR was used because of its sensitivity, specificity, and its particular value where a limited amount of tissue is available and small amounts of cells express the gene of interest.

The majority of cells present in the vitreous samples of patients with PVR are capable of producing cytokines in vitro. RPE cells, upon activation, release IL-6, IL-8, and granulocyte macrophage colony stimulating factor. Gliolymphocytes produce many different cytokines including IL-1, IL-6, TNFα, and IFNγ. The actual role of cytokines in the pathogenesis of PVR is not clear, but their involvement in regulating immune and inflammatory responses is well described. In the retinal microenvironment, they could contribute substantially to the occurrence of PVR.

This study, like that of Kauffmann et al. and Elner et al., demonstrated a consistent presence of IL-6 and IL-8 in samples obtained from PVR patients. The definitive role of IL-8 in the pathogenesis of PVR is not clear, but owing to its chemotactic activity for leucocytes, it was postulated that TNFα and IL-1β, which were not directly chemotactic for leucocytes, exert their chemotactic effect through induction and release of secondary cytokines including IL-8 and MCP-1. Both IL-1 and IL-6 mediate inflammatory cell functions such as cell growth, migration, and differentiation. They overlap in their functions, and IL-1 is a potent inducer of IL-6 expression in many cell types.

Although some patients with macular holes had epiretinal membrane (ERM), the macular hole group did not show the same cytokine profiles as did the PVR patients. This may be due to difference in the composition of ERM in macular holes. ERM is primarily a glial reaction with few inflammatory cells and lymphocytes.

In conclusion, our data indicate that mRNA encoding for IL-6, IL-1β, TNFα, and IL-8 is significantly detected by RT-PCR in vitreous and SRF samples of patients with PVR, indicating probable local production of these cytokines by vitreous and SRF cells and their possible role in pathogenesis of the disease. Further work is needed to correlate the level of mRNA to respective cytokines in the vitreous and to severity of the disease and prognostic outcome of surgery.

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