Commentary

Sense and nonsense of corticosteroid administration in the treatment of ocular toxoplasmosis

There is no consensus about the use of corticosteroids for the treatment of ocular toxoplasmosis. In clinical practice, corticosteroids are usually given in combination with antiparasitic drugs to reduce the inflammatory reaction during active chorioretinitis and to minimise tissue damage. Specifically, in the case of pericentral location, this presumed effect might help to retain central visual acuity and limit scotomas. However, the use of corticosteroids can lead to progression of the underlying infectious disease, as repeatedly demonstrated by the occurrence of fulminant ocular toxoplasmosis following both systemic and periocular administration (Fig 1). On the other hand, low dose corticosteroid monotherapy has been administered without severe side effects.

To our knowledge, a well defined study of the effect of corticosteroid monotherapy in ocular toxoplasmosis as well as the additional value of corticosteroids as adjuvant therapy has not been performed. Thus, the use of corticosteroids for the treatment of ocular toxoplasmosis is controversial, and the timing and dosages during the course of the disease are not well defined.

During an acute infection with *Toxoplasma gondii*, oocysts ingested by the host give rise to the extracellular forms (tachyzoites), which actively penetrate host cells and replicate; subsequently the disease enters the chronic stage when cysts are formed. The signal for formation of cysts is not known; the onset and the quality of the host immune response may be an important factor.

The mechanism of chronic recurrent ocular toxoplasmosis is unknown; it is presumed that the multiplication of parasites, liberated from the cysts, causes damage to adjacent retinal tissue. The hypothesis of autoimmuneinduced, directed against retinal S-antigen, has not been proved unequivocally. In fact, the interaction between *T gondii* and the competent immune system does not lead to parasite elimination, but to a reduction in parasite load and changes in morphology and surface antigen expression.

**Host response and effect of corticosteroids on T gondii infection**

T cell mediated immunity plays a major role in resistance against toxoplasmosis and probably also controls brain cyst formation; B cell mediated humoral immunity is not protective. CD8+ T cells are the principal mediators of resistance against acute toxoplasmosis by means of their cytotoxic activity and the secretion of cytokines, including interferon γ (IFN-γ), which plays a crucial role in the response to *T gondii.*

The effects of corticosteroids on the immune system are complex, and not completely understood. The anti-inflammatory and immunosuppressive effects of corticosteroids are linked to changes in cellular immunity and attributed mainly to inhibition of communication among leukocytes responsible for production and action of cytokines. In low doses, corticosteroids dramatically inhibit exudation of plasma and accumulation of leukocytes at sites of inflammation and suppress cellular immune response. As a consequence they influence the production of IFN-γ and IL-2 by T cells, the effect of IFN-γ on macrophages, and the interaction of IL-2 with its receptor on activated T cells. The effect on cellular immune response is dose dependent. Corticosteroids have a minor effect on antigen stimulated B cells and plasma cells. Higher doses are required to retard the processing and presentation of antigens by monocytes and macrophages, to inhibit natural killer cell function, and to inhibit the release of tumour necrosis factor by activated macrophages. The compromised leucocyte function counteracts the intracellular destruction of parasites and may lead to unrestrained replication of parasites.

An inadequate cell mediated immunity (specifically in patients with AIDS and during iatrogenic immunosuppression) leads to severe, sometimes fatal toxoplasmic infections and is associated with reactivations of latent disease. Although intact tissue cysts were reported not to evoke an inflammatory response, shedding of parasites from tissue cysts was described. It might be possible that, in patients with an insufficient cellular immunity, these escaped parasites may replicate and cause reactivation. Aging is also associated with a decline in cell mediated immunity (especially CD8+ T cells).

**Effect of corticosteroids on toxoplasmosis: animal studies**

A lethal effect of corticosteroids in acute toxoplasmosis was found in the majority of animal studies (mice and rabbits); however, the therapy also induced less effective resistance to subsequent reactivations. In contrast with untreated animals, in corticosteroid treated mice with acute toxoplasmosis, persistence of free parasites together with cysts in the brain and lung tissue was observed. Furthermore, cyst formation in the brain...
Four patients received systemic and periocular steroids. Chorioretinitis without associated old scars. Systemic symptoms. With corticosteroids exhibited no changes in their ocular or systemic symptoms. In the past, eight eyes of patients treated with high dose systemic corticosteroids without an antiparasitic shield anyway. An absence of retinal scars, suggesting a primary infection, was noted for 9/16 of these patients (61%).

Although toxoplasmic chorioretinitis is more common during the second and third decades of life, 16/26 patients (61%) were older than 50 years. Severe toxoplasmic retinitis resembling herpetic retinitis was recently described in seven elderly patients. The authors hypothesised that the susceptibility of older patients to severe toxoplasmic chorioretinitis may be due to the previously mentioned age related decrease in immunological resistance, which might also have played a role in our patients. It is noteworthy that the initial diagnosis was acute retinal necrosis in four of the 10 present cases (40%). The advanced age of the patients together with severe and atypical retinitis, may explain why toxoplasmic retinitis was not suspected and initially misdiagnosed. In the past, eight eyes of patients treated with corticosteroids were examined histologically with consistent results—large areas of necrotic tissue with multiple cysts, located between the necrotic retina and remnants of normal tissue. In the necrotic regions were numerous free parasites without an inflammatory reaction.

Evaluation of the efficacy of corticosteroids as an adjunct to antiparasitic regimens has never been performed. Studies addressing the efficacy of multiple therapies were not designed for evaluation of the effects of corticosteroids. However, no adverse effects of additional corticosteroids were noted. No differences in onset of inactivity were seen between patients treated with pyrimethamine and sulphadiazine with or without added corticosteroids, even at high doses in a study by Fajardo et al. In a meta-analysis of three studies, involving 134 patients altogether (antiparasitic therapy without corticosteroids in 16 cases and with corticosteroids in 118 cases), no striking differences were found in the duration of activity. Although occasionally a poor ocular outcome was reported, this might be explained by the bias of added corticosteroids in a selected group of patients with extremely severe inflammation. Similar visual outcomes were found for patients treated with antiparasitic drugs with and without corticosteroids in a retrospective non-randomised study of 75 patients.

In conclusion, our results confirm the case report based assumption that in ocular toxoplasmosis corticosteroids, whether given systemically or periocularly, should be administered with a shield of antiparasitic drugs, especially if the patients are elderly or exhibit evidence of a primary infection. Owing to its frequent atypical clinical features, it is important to include the possibility of ocular toxoplasmosis in the differential diagnosis of uveitis and to consider the use of corticosteroids, even in the absence of a specific diagnosis.

<table>
<thead>
<tr>
<th>Evidence for primary aquired infection</th>
<th>Corticosteroid administration</th>
<th>Retinal scars</th>
<th>Legal blindness of affected eye</th>
</tr>
</thead>
<tbody>
<tr>
<td>No of patients</td>
<td>Age at onset</td>
<td>Laboratory No (%)</td>
<td>Clinical* No (%)</td>
</tr>
<tr>
<td>Literature survey†</td>
<td>16</td>
<td>10 (63)</td>
<td>9 (56)</td>
</tr>
<tr>
<td>Present study</td>
<td>10</td>
<td>6 (60)</td>
<td>7 (70)</td>
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<tr>
<td>Total</td>
<td>26</td>
<td>16 (62)</td>
<td>16 (62)</td>
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</tbody>
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*Chorioretinitis without associated old scars.
†See references 2-8; three patients received additional medication (two azathoprine, one sulphasoxazole respectively); five patients received systemic and periocular steroids.
‡Four patients received systemic and periocular steroids.
NS= not specified.

Effect of corticosteroids on toxoplasmosis: human studies

Various case reports provide information on the effects of corticosteroids in human ocular toxoplasmosis. In the past, the number of recurrences were not different for 10 patients with presumed toxoplasmic chorioretinitis treated with low dose systemic corticosteroids compared with 10 patients treated with antiparasitic drugs together with corticosteroids. In contrast, destructive and disseminated ocular toxoplasmosis was repeatedly demonstrated in case reports of presumably immunocompetent patients (n=16) after administration of corticosteroids without a shield of antiparasitic drugs (Table 1). We examined 10 additional patients with fulminant ocular toxoplasmosis following monotherapy with corticosteroids (Table 1).

Many of corticosteroid treated patients were initially not recognised as having toxoplasmosis (recent series 4/10; literature 8/16; total 12/26); in 4/26 patients (2/10 in the present series) corticosteroid therapy was given for an unrelated systemic disease. The remaining patients (10/26) were known to have toxoplasmic retinitis but received corticosteroids without an antiparasitic shield anyway.

Legal blindness (visual acuity less than 0.1) in at least one eye developed in 19 of 26 reported patients (73%). Recurrent disease occurred subsequently in four of our 10 patients (40%); the number of recurrences among the previously reported 16 patients is unknown. The dosages of administered corticosteroids varied between 20 mg and 150 mg per day (or were not specified). The eventual relation of ocular disease to the dosage of corticosteroids used cannot be assessed and therefore it is not possible to define something as a "safe dose".
toxoplasmosis in the differential diagnosis for elderly patients. Further research is indicated to evaluate the beneficial effect of corticosteroids in combination therapy.

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