Corticosteroid treatment of periorbital haemangioma of infancy: a review of the evidence

T M Ranchod, I J Frieden, D R Fredrick

Aim: To systematically review the literature for corticosteroid treatment of periorbital haemangioma of infancy (HOI) and determine the relative efficacy and safety of oral, topical and intralesional corticosteroids.

Methods: PubMed and the Cochrane Library were queried using keywords, and further articles were obtained by reviewing bibliographies. Inclusion and exclusion criteria were applied to create a subset of literature for analysis.

Results: Systematic review revealed 81 original reports of periorbital HOI cases treated with steroids. Most studies and case series failed to document refractive error or visual acuity before and after treatment. Of cases meeting inclusion criteria, five patients received topical steroids and 25 patients received intralesional steroids. Patients receiving intralesional injections tended to demonstrate reduced astigmatism at follow up after treatment (21 of 28). The lack of studies with relevant objective ophthalmological end points prevented statistical meta-analysis.

Conclusion: Intralesional injections may reduce refractive error, while the efficacy of topical steroids is unclear. Studies measuring objective ophthalmic data before and after treatment are sparse, and more studies are needed to determine the relative efficacy of different steroids. There are insufficient data to estimate the incidence of steroid side effects in patients treated with steroids for periorbital HOI or complications of intralesional injections in particular.

Abbreviations: HOI, haemangioma of infancy; SE, spherical equivalent.
Collins published a study in 1991 in which nine of 15 eyelid haemangioma patients responded to intralesional steroids as measured by the lesion's dimensions.35 The presence or absence of amblyopia was recorded post-treatment, but no ophthalmological end points were recorded before treatment. Willshaw and Dead published a study in 1987 with 13 periorbital capillary haemangioma patients treated with intralesional steroids, but without specifying intraorbital versus extraorbital involvement.36 Dead and Willshaw published a study in 1986 of 24 patients with periorbital haemangiomas, of whom nine were treated with intralesional steroids.37 Refractive error for a subset of patients was measured before treatment, but no objective end points were measured after treatment. A 1992 study by Assaf et al included 12 patients of whom 11 were treated with intralesional steroids and one with oral steroids.38 Both refractive error and visual acuity were measured before treatment, but no end points were measured after treatment. Stigmar et al published a study of 51 patients in 1978, but the number of patients treated with steroids was not specified.39

Several large case series of capillary haemangioma patients contained subsets of patients with facial lesions but either did not specify location (periorbital, intraorbital) or did not break down objective data by location.40-44

Of the cases that met inclusion criteria, five patients in one series were treated with topical steroids and 23 patients from three reports were treated with injected steroids. No patients treated with oral steroids met inclusion criteria for the study. Patients who received intralesional injections tended to demonstrate reduced astigmatism at follow up after treatment (19 of 23). Two out of five patients in the topical steroid series showed reduced astigmatism following treatment. Only one of five topically treated patients had reduced spherical equivalent refractive error after treatment, compared to 16 of 23 patients treated with intralesional steroids.

### Side effects and complications of steroid delivery

Systemic side effects of corticosteroids for treatment of periorbital HOI have been documented for both intralesional and oral steroids. Side effects and complications of steroid delivery were not evaluated by inclusion or exclusion criteria since the vast majority of examples in the literature come from case reports or small series. No existing studies document frequency of side effects or complications of the various steroids.

### DISCUSSION

The most striking result of our study is the very small number of patients whose treatment results could be evaluated objectively. Despite the use of corticosteroids in the treatment of haemangiomas for more than three decades and the high risk of morbidity leading to permanent visual loss, studies with measurable outcomes in this setting are lacking. In those case series which were evaluated, we found insufficient evidence to demonstrate benefits of one corticosteroid over another in patients with periorbital haemangioma of infancy. Similarly, we found insufficient evidence for increased side effects of one steroid over another. We found weak evidence.
that intralesional steroid injections may result in reduced refractive error.

The five cases involving topical steroids generally demonstrated worse spherical equivalent refractive error after treatment, with no clear trend in the astigmatic component alone. The 23 cases involving intralesional injection demonstrated a trend towards reduced astigmatism and reduced spherical equivalent refractive error with treatment. Although one might argue that steroids speeded up involution, the improvement in refractive error cannot be attributed to steroid treatment rather than the natural course of development after spontaneous haemangioma involution. The Motwani series included four patients who were managed conservatively (without steroids, not included in table 3). The authors concluded that visual outcomes were similar in treated and untreated patients, but selection of patients was non-randomised.

Despite reports of successful steroid treatment of periorbital HOI dating back to the 1970s, the evidence for relative efficacy and safety of oral, topical, and intralesional steroids remains scant. Several of the largest studies produced in our literature search were not included in this review because they did not measure objective ophthalmological data before and after treatment. The largest single study meeting criteria, by Morrell and Willshaw, contained only 13 patients with measured objective end points both before and after treatment."

The complications of untreated periorbital HOI have been well documented, including astigmatism, myopia, amblyopia, and strabismus. Several articles have reported on the beneficial effect of corticosteroids in preventing or reducing occlusion of the visual axis. However, a substantial proportion of these studies and case series has failed to document refractive error or visual acuity before and after treatment. Several studies developed customised grading systems in order to stratify patient responses to steroids. In most cases, the grading systems were either subjective or based on data such as change in lesional size, which may or may not cause an objective improvement in visual outcome. Corticosteroids may facilitate haemangioma involution and thereby decrease volume or surface area of the lesion. However, changes in volume or surface area are only relevant to the extent that the haemangioma impinges on the visual axis at some point and involution results in clearing of the axis. Oral steroid

*Table 3 Steroid dosage and refractive error*

<table>
<thead>
<tr>
<th>Author, year</th>
<th>Subjects (total)</th>
<th>Modality: dose*</th>
<th>Refraction before treatment</th>
<th>SE before treatment</th>
<th>Refraction after treatment</th>
<th>SE after treatment</th>
<th>Age (months)</th>
<th>Following (months)</th>
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<tbody>
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<td>Elsas 1994</td>
<td>5 (5)</td>
<td>Top: clobetasol 0.05%</td>
<td>+2.00 to +3.00 x 1.40</td>
<td>+3.50</td>
<td>+5.00 to +1.00 x 2.00</td>
<td>+5.50</td>
<td>14 weeks</td>
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<td></td>
<td></td>
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<td>-1.00</td>
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<td>-1.00</td>
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</tr>
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<td></td>
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<td>+1.00</td>
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<td>-0.50</td>
<td>11 weeks</td>
<td>38</td>
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<td></td>
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<td>+1.00</td>
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<td>-0.75</td>
<td>11 weeks</td>
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<td>Kushner 1982</td>
<td>5 (10)</td>
<td>IL: 12 mg/80 mg, 9 mg/6 mg</td>
<td>+1.75 to +1.00 x 1.80</td>
<td>+2.25</td>
<td>+1.50 sp</td>
<td>+1.75</td>
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<tr>
<td></td>
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<td>IL: 40 mg/6 mg x 2</td>
<td>+1.00</td>
<td>+1.00</td>
<td>+1.50 sp</td>
<td>+1.50</td>
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<td></td>
<td></td>
<td>IL: 40 mg/6 mg x 2</td>
<td>-3.00 sp</td>
<td>+1.00</td>
<td>+0.50 sp</td>
<td>+0.50</td>
<td>0.87</td>
<td>10</td>
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<td>+0.50 sp</td>
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<td>0.87</td>
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<td>Morris 1991</td>
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<td>IL: 20–40 mg/4 mg, up to 4 x</td>
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<td>+1.50 sp</td>
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<td>+0.50</td>
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</table>

*Doses for combination steroid injections are listed as triamcinolone (mg) followed by betamethasone (mg) unless otherwise specified.
SE, spherical equivalent; Top = topical, IL = intralesional. Values in bold are for the unaffected eye, when available."
treatment has been shown to produce a response in cutaneous haemangiomas as measured by surface area, but the goals of ophthalmological treatment differ, at least in part, from the goals of dermatological treatment of HOI.1

Because of the limited numbers of cases meeting our inclusion criteria and the variation in dosage and administration of corticosteroids given, the results are not conducive to a statistical meta-analysis. Since there were no studies comparing different modalities of steroid therapy and no studies of individual modalities large enough to estimate efficacy, no conclusions can be drawn regarding which modality is most efficacious. The studies reviewed indicate that intralascular injections appear efficacious insofar as they may reduce astigmatism and spherical equivalents of refractive error; the efficacy of topical steroids is less clear. However, greater numbers of patients in each group are needed to project an estimate of overall efficacy. A larger study might also clarify how much of observed reduction in refractive error is the result of correction with ageing or the natural course of haemangioma involution as opposed to steroid treatment effects.

Similarly, there are insufficient data to estimate the incidence of corticosteroid side effects for patients treated with steroids for periorbital HOI or complications of intralaminar injection in particular. Numerous case reports have established the need for caution in giving prolonged courses of steroids and the need for preferential looking and standardised protocols of follow-up to avoid vascular occlusion when injecting intralaminar steroids, but the incidence of such complications remains unknown.

A useful determination of relative efficacy and side effects of different steroid modalities for treatment of periorbital HOI could be achieved through a prospective, longitudinal multicentre trial. Objective measurements such as visual acuity, astigmatism, or strabismus must be measured both before and after treatment, with follow up more than 6 months after completion of treatment and concurrent measurements in the unaffected eye. We would recommend checking cycloplegic refractions monthly until stabilisation of haemangioma growth, with subsequent bimonthly and then quarterly checks for 3 years. Visual acuity can be measured by preferential looking, and standardised protocols of occlusion therapy can be used for treatment of amblyopia.

Steroid preparations and dosages must also be standardised.

In an era of evidence based medicine, corticosteroid treatment of periorbital HOI is still based primarily on recommendations made decades ago as well as the personal preferences of the treating paediatric ophthalmologist. The need for objective data to guide future practices is evident, and establishing the relative efficacy and safety of oral, topical and intralaminar steroids through a randomised prospective study would greatly benefit those children with periorbital HOI requiring ophthalmological intervention.

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No financial support was provided for this work and the authors claim no competing interests.

Competing interests: none declared

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Br J Ophthalmol 2005 89: 1134-1138
doi: 10.1136/bjo.2005.070508

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