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Periocular capillary hemangioma: management practices in recent years

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Purpose: To present a case series on the management options for capillary hemangiomas involving the eyelid and orbit.

Methods: This is a retrospective chart review of clinically diagnosed capillary hemangioma cases involving the periocular region treated at two local eye institutions. The patients’ demographics and clinical presentation – including visual acuity, refractive error, periorbital and orbital examinations, and ultrasound and magnetic resonance imaging findings – were reviewed. The clinical progression, modalities of treatment, and treatment outcomes were studied.

Results: Sixteen cases of capillary hemangiomas involving the eyelid and orbit were studied. The mean age at consultation was 9.6 months (range: 1 month–72 months). The majority were females (75%), with 50% presenting as upper-eyelid hemangiomas and the remaining as lower-eyelid (38%) and glabellar (12%) lesions. Combined superficial and deep involvement was common (64%). Cases whose lesions were located at the upper eyelid or superior orbit led to amblyopia (25%). Fifty-six percent of cases (9/16) were managed conservatively, and 44% (7/16) underwent treatment with either single-agent (n = 4) or combined treatments (n = 3).

Conclusion: Close monitoring of visual development and prompt institution of amblyopia therapy for children with periocular capillary hemangiomas generally preserve vision. Extensive lesions that affect the visual axis require local and systemic treatments, alone or in combination, in order to reduce the size and impact of lesions on the eyeball, to reduce induced refractive error and visual occlusion, and to prevent the development of amblyopia, in order to achieve good visual outcomes.

Keywords: vascular, vascular malformation, bleomycin, propranolol, prednisolone, amblyopia

Introduction

Periocular capillary hemangiomas are common benign vascular tumors seen in children. Histologically, they are characterized by proliferating endothelial cells. They generally present within a few weeks after birth and undergo a rapid proliferating phase, followed by a period of relative quiescence, regressing after a few years into an involution phase.¹⁻⁹ The clinical course can be variable. Demographically, these cases occur commonly among females and have not been reported to vary by race, ethnicity, or geographic location.¹⁻⁶,⁸,¹⁰

Haik et al¹ and others classified these lesions according to their characteristics on clinical presentation (eg, strawberry nevus, bluish discoloration), while Rootman¹¹ presented another classification based on the level of involvement (eg, superficial, subcutaneous, deep orbital, or combined).¹,² Hemangiomas of the eyelid and orbit have also been studied according to their size. Schwartz et al² found that half of the

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cases with hemangiomas exceeding a widest diameter of one centimeter would require treatment. Regardless of the method of classification, hemangiomas can present at any single location or at a combination of locations within the periorcular region.\textsuperscript{1,3,11}

Acceptable indications for intervention may include rapidly enlarging lesions, obstruction of the visual axis, significant induced astigmatism, and cosmetic concerns. Here, a short series based on local experiences is presented, along with some of the current treatment modalities. The modalities available are intralesional and systemic steroid, bleomycin, interferon-alpha, vincristine, cyclophosphamide, topical timolol maleate, oral propranolol, and surgical excision.\textsuperscript{3,5,7–20}

Materials and methods

This is a retrospective chart review of 16 clinically-diagnosed capillary hemangioma cases that presented as red or bluish elevated vascular lesions in the periorcular region seen at our institutions. This study has obtained approval from the institutional review board.

The parameters reviewed include age at presentation; size and location of the lesion; best corrected visual acuity using preferential looking tests, Kay pictures, Snellen, or HOTV charts (the top row of letters tested) (depending on the patient’s age); and the results of cycloplegic refraction, orbital, and eyelid examinations. Imaging studies such as magnetic resonance imaging (MRI), magnetic resonance angiography, and ultrasound were studied when available. The management of each patient was reviewed, including indications, modes of treatment, and patients’ responses and outcomes. The clinical progression, related complications, and final visual outcome at end of follow-up were studied. The effectiveness of treatment was ascertained when either partial or complete reduction in the size of the hemangioma was observed, based on clinical photographs and measurements.

Results

Sixteen cases of capillary hemangiomas involving the eyelids and orbits were reviewed. The mean age at consultation was 9 months (range: 1 month–72 months) with a median of 4 months. Seventy-five percent (n = 12) were females, and 25% (n = 4) were males. Eighty-one percent (13) were born full term, while 19% (3) were delivered prematurely (<36 weeks age of gestation) (Table 1).

<table>
<thead>
<tr>
<th>Race</th>
<th>Number of cases</th>
<th>%</th>
</tr>
</thead>
<tbody>
<tr>
<td>Caucasian</td>
<td>1</td>
<td>6</td>
</tr>
<tr>
<td>Chinese</td>
<td>9</td>
<td>56</td>
</tr>
<tr>
<td>Indian</td>
<td>2</td>
<td>13</td>
</tr>
<tr>
<td>Malay</td>
<td>3</td>
<td>19</td>
</tr>
<tr>
<td>Others</td>
<td>1</td>
<td>6</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Number of cases</th>
<th>%</th>
</tr>
</thead>
<tbody>
<tr>
<td>No of patients</td>
<td>16</td>
</tr>
<tr>
<td>Male/female</td>
<td>1:3</td>
</tr>
<tr>
<td>Median age at consultation</td>
<td>4 months (1–72 months)</td>
</tr>
<tr>
<td>Preterm</td>
<td>3</td>
</tr>
<tr>
<td>Term</td>
<td>13</td>
</tr>
</tbody>
</table>

Vision, refraction, and amblyopia

The mean logarithm of the minimum angle of resolution (logMAR) visual acuity was 0.77 at the first visit and 0.46 upon the most recent examination (Table 2). Of those patients with recorded cycloplegic refraction, a majority of the children less than 2 years of age were hyperopic, with mean spherical refractions of +1.83 and +1.00 on the first and last visits, respectively. All cases whose lesions were located in the upper eyelid had astigmatism, with a mean cylinder of -1.714 (range: 0.75–4.25).

Twenty-five percent (4/16) of patients (mean age of 4 months) were found to be amblyopic during the follow-up period. The majority (75%) were amblyopic because of anisometropia, and a quarter (25%) were amblyopic because of occlusion of the visual axis; all of them had lesions located in the upper eyelid. No cases with lower-eyelid lesions had amblyopia. Two cases presented with proptosis, globe displacement, and limitation in eye movement due to lesions extending deep into the orbit.

Fifty-six percent (9/16) were managed conservatively. Of the remaining 44%, treatment was either with a single agent (57%) or a combination (43%) (Table 3).
Management

The mean follow-up was 19 months (range: 1.5–72 months), and 12% (n = 2) were lost to follow-up. Fifty-six percent (nine) were managed conservatively, which involved regular spectacle correction and amblyopia therapy, by taping and/or patching, as well as close follow-up. By two years of age, seventy-eight percent (7/9) showed no table regression (according to clinical photos and clinical examinations) in the size of the hemangioma (Figure 2).

Forty-four percent (7/16) required medical or surgical intervention. Indications were deep orbital extension, occlusion of the visual axis, and cosmetic concerns.

The treatments we employed included: (1) steroid therapy, (2) surgical excision, (3) oral propranolol, and (4) bleomycin (received at other institutions), either singly or in combination (Table 3). Among those that received treatment, 57% (four) were treated with a single agent (oral propranolol, prednisolone, or surgery), and regression was noted in three of them. One developed respiratory distress after propranol therapy, and medication was immediately terminated.

The remaining 43% (three) received a combination of two to three agents (oral prednisolone and intralesional triamcinolone:betamethasone; oral propranolol and intralesional bleomycin, or surgical excision and topical timolol maleate). Regression was observed in two of them. One case that underwent surgical excision for a small superficial hemangioma involving the left medial upper eyelid had a recurrence after 2 weeks. Application of topical timolol maleate 0.5% eyedrops twice daily was advised. However, in this case, there was no change in the size of the lesion after 1 month of eyedrop application. One patient treated with oral prednisolone and intralesional triamcinolone:betamethasone experienced a complication of orbital cellulitis, which was subsequently treated with intravenous antibiotics and drainage of the subperiosteal abscess.

Propranolol

Three patients were treated with oral propranolol. Reduction in the size of the lesion was observed in two out of the three cases.

Table 2 Mean visual acuity, and sphere and cylinder refraction at first and last visits

<table>
<thead>
<tr>
<th></th>
<th>First visit</th>
<th>Latest visit</th>
</tr>
</thead>
<tbody>
<tr>
<td>UCVA (logMAR)</td>
<td>0.77 (range: 0.1 to 1.68)</td>
<td>0.46 (range: 0.1 to 0.67)</td>
</tr>
<tr>
<td>Mean sphere</td>
<td>+1.83 (range: +0.25 to +3.50)</td>
<td>+1.00 (range: −0.50 to +2.50)</td>
</tr>
<tr>
<td>Mean cylinder</td>
<td>−1.63 (range: −0.75 to −4.25)</td>
<td>−1.28 (range: 0.00 to −4.00)</td>
</tr>
</tbody>
</table>

Abbreviations: UCVA, uncorrected visual acuity; logMar, logarithm of the minimum angle of resolution.

Table 3 Modes of treatment used for periocular capillary hemangiomas in this series

<table>
<thead>
<tr>
<th>Management</th>
<th>n</th>
</tr>
</thead>
<tbody>
<tr>
<td>Single treatment</td>
<td>4</td>
</tr>
<tr>
<td>Combined treatment</td>
<td>3</td>
</tr>
<tr>
<td>Observation</td>
<td>9</td>
</tr>
<tr>
<td>Topical timolol maleate 0.5%</td>
<td>1</td>
</tr>
<tr>
<td>Intraleisonal triamcinolone/betamethasone</td>
<td>1</td>
</tr>
<tr>
<td>Intraleisonal bleomycin</td>
<td>1</td>
</tr>
<tr>
<td>Oral prednisolone</td>
<td>2</td>
</tr>
<tr>
<td>Oral propranolol</td>
<td>3</td>
</tr>
<tr>
<td>Surgical excision</td>
<td>2</td>
</tr>
</tbody>
</table>
One of the cases treated with oral propranolol was a 6-month-old female with amblyopia from mechanical ptosis, with two previous intralesional bleomycin injections, 2 months apart, from another institution. The hemangioma showed remarkable reduction at the time of consultation at our institution (Figure 3). The second case was a preterm female with subcutaneous hemangioma involving the left superomedial orbit with maximal MRI axial dimensions of $1.7 \times 1.9$ cm, which decreased to a clinical measure of $0.8 \times 0.3$ cm after 7 weeks of treatment.

**Discussion**

Capillary hemangiomas represent an abnormal growth of blood vessels characterized by endothelial proliferation. They expand progressively in weeks to months, then involute over the years.\textsuperscript{1,9,11,22} Formerly dense endotheliomatous lesions during the growth phase become replaced by fewer endothelial channels, increased collagen deposition, and fat, as well as inflammatory cells, as lesions regress.\textsuperscript{11}

The clinical diagnosis of capillary hemangiomas is often straightforward. Superficial hemangiomas may affect the lid or conjunctiva and appear as strawberry lesions, scarlet red in color, whereas deeper lesions lie posterior to the orbital septum and present as a subcutaneous bluish hue. Combined presentations have both superficial and deep components. Entities known as complex hemangiomas are found to be multicentric. These hamartomas often present as isolated lesions or, in rare instances, with systemic associations, such as Kassabach-Merritt syndrome (thrombocytopenia) and neurocutaneous PHACE syndrome (a disorder in which a child is born with posterior fossa malformation, cervico-facial hemangiomas, arterial anomalies, cardiac defects, and various eye anomalies).\textsuperscript{11,21,22} None of the patients in our series had associated systemic syndromes.

Imaging studies help support the diagnosis. B-scan and Doppler ultrasonography showed the lesions exhibited high echogenicity, with either smooth lobular or irregular contours, and fast blood flow. Computed tomography showed the lesions as well-defined homogenous or inhomogenous, moderate-to-intense enhancing masses. MRI also showed them to be well defined, with some irregular margins, and with low signal intensity on T1-weighted images versus high signal intensity on T2-weighted images with respect to fat and extraocular muscles. With gadolinium, the capillary hemangiomas demonstrated heterogenous or homogenous enhancement. These lesions could be found in any or in a combination of compartments within the orbit, being intraconal and extraconal, pre- and postseptal.\textsuperscript{11,22}

Indications for the treatment of capillary hemangiomas in the periorbital region are ptosis with visual axis occlusion, deprivational amblyopia, anisometropia, and cosmesis.\textsuperscript{7} Similar to the observations by Schwartz et al,\textsuperscript{2} four of our patients with upper-eyelid lesions, with basal dimensions measuring $>1$ cm, developed amblyopia secondary to anisometropia and mechanical ptosis that occluded the visual axis.

Refractive errors are potentially reversible, especially when spectacle correction and prompt amblyopia therapy are given. However, our study could not correlate this thoroughly, due to missing data and a small sample size.

Observation remains an option in the management of hemangiomas. More than half of our patients in this study were monitored closely for any change in refractive error, and were treated for amblyopia when it was present.

Some of our patients received both local and systemic treatments. Oral prednisolone was commonly prescribed as a systemic treatment for extensive lesions, and we likewise
observed regression in the size of the hemangioma (1/2) (Figure 4).

Intralesional steroid injection was the most common form of treatment in the past for capillary hemangioma.5 Weiss observed a reduction in astigmatism of up to 63% within 1 to 14 months after injection.7 In this series, triamcinolone with betamethasone was used as an additional treatment in a case unresponsive to oral prednisolone.

Generally, small and superficial lesions may be effectively managed with topical or intralesional treatments, while larger and deeper hemangiomas require systemic treatments or a combination thereof.5,13 We previously treated patients having relatively superficial and localized lesions with intralesional steroid injections, while for extensive lesions with deep orbital involvement, this was managed through systemic treatments. When one form of treatment was insufficient for reducing the size of the hemangioma, we used a different modality.

Bleomycin is a sclerosing agent and has been widely used as an anticancer drug. We have no local experience with bleomycin in treating hemangiomas. It is believed to act on infantile hemangiomas by reducing vascular endothelial cell proliferation. Sarihan et al were first to report on its application as an intralesional injection for cutaneous hemangiomas.23,24 They noted extensive fibrosis and spontaneous resolution of hemangiomas. One of our patients received intralesional injections of bleomycin at a different institution, as well as oral propranolol.

In recent years, propranolol has produced encouraging results for capillary hemangioma and has become our preferred treatment. We observed dramatic positive responses (2/3) in our patients, except for one who unfortunately developed a systemic complication and had to terminate the treatment.

Systemic complications can be potentially life threatening; one of our patients developed respiratory distress during propranolol treatment. This clearly shows that careful systemic evaluation and patient selection are crucial prior to treatment and that the patient’s family should be informed of the benefits and risks of the therapy.

Although hospital admission with close observation by pediatricians for possible side effects such as hemodynamic changes (eg, bradycardia, hypoglycemia, and respiratory distress)6,18,20 is recommended, outpatient management has been reported by Haider and coworkers to show favorable responses in more than half of patients, with minor side effects.15 Hence, the multidisciplinary collaboration of cardiology, pediatrics, dermatology, and ophthalmology – with pretreatment workups, such as baseline electrocardiography – should remain standard for either inpatient or outpatient intervention.15,16,18

Our study showed there was a favorable response to oral propranolol, with dramatic regression (2/3), except for one who developed respiratory distress. Currently, propranolol is our preferred treatment, at a recommended dose of 1–2 mg/kg/day, given on an outpatient basis when the patient is in good health, or when admitted if otherwise.

To date, several studies have been conducted attesting the efficacy of this nonselective beta-blocker, which has long been used as an antihypertensive agent. It has been speculated that the apparent vasoconstriction may be due to the inhibition of vascular endothelial growth factor and the triggering of endothelial cell apoptosis. In vitro pharmacologic research has shown that propranolol beta adrenergic blockade inhibits several steps in the angiogenic cascade, such as growth factor-induced migration, tubulogenesis, and matrix metalloproteinase secretion of endothelial cells.25

Surgical excision, either complete or partial resection, remains an option for treatment. It is often indicated for sight-threatening hemangiomas that do not respond to medical treatments, or for small well-defined lesions that can be removed completely with minimal morbidity.5,12,13

**Conclusion**

In general, most children with periocular capillary hemangiomas do not require treatment, given their natural capacity for spontaneous involution. The preconditions for intervention are rapidly enlarging lesions, visual axis obscuration, significant induced astigmatism, and cosmetic reasons. For extensive lesions involving the visual axis, local and systemic treatments effectively reduce the size of the lesion, whether they are used alone or in a combination. We recommend systemic treatments such as propranolol for large and deep subcutaneous lesions, especially when these

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**Figure 4** Sagittal MRI of a hemangioma involving the left orbit and pterygopalatine fossa before (left) and 1 year after treatment (right) with oral prednisolone.

**Abbreviation:** MRI, magnetic resonance imaging.
cause visual occlusion, significant refractive error, and globe displacement. Pretreatment, systemic evaluation should be done. Observation is appropriate for small and non-visually threatening hemangiomas. Visual outcomes are generally good when appropriate treatment is given early and existing refractive errors and amblyopia are corrected, which remain of utmost importance. Despite the effectiveness of each treatment, complications exist, and though not frequent, prudence dictates awareness and timely management.

Acknowledgment

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Disclosure

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References