Disclaimer

DESCRIPTION: Haemorrhage within the orbit posterior to the orbital septum. Retrobulbar haemorrhages can cause an orbital compartment syndrome thereby compressing the central retinal artery and other vessels with resultant optic nerve and retinal ischaemia and permanent loss of vision. Immediate assessment and management is necessary to reduce the risk of vision loss.

HOW TO ASSESS:

Red Flags:
- Painful, tense exophthalmos with a visual deficit and/or a relative afferent pupillary defect (RAPD) is an indication for an immediate lateral canthotomy and cantholysis
- Do not let detailed history/investigations interfere with immediate management if diagnosis clear
- Irreversible visual loss can occur within 1.5 to 2 hours time from onset of haemorrhage
- Consider possible association of traumatic optic neuropathy in the setting of orbital trauma
- If contacted by another hospital’s emergency department for advice for treatment, discuss the indications for immediate canthotomy/cantholysis

On History:
- Causes of retrobulbar haemorrhage include orbital/lid surgery, retrobulbar/peribulbar injections, facial trauma, anticoagulants, blood dyscrasias

Note: Retrobulbar haemorrhage may have a delayed presentation

On Examination:
Examine for associated injuries, e.g. ruptured globe or penetrating eye injury.

The following signs are present if the retrobulbar haemorrhage is causing visual compromise:
- Decreased visual acuity
- RAPD
- Painful proptosis with limitation of eye movements
- Tense orbit/resistance to retropulsion
- Marked subconjunctival haemorrhage
- Intraocular pressure (IOP) > 40mmHg
- Macula: cherry red spot if associated central retinal artery occlusion

When performing a dilated fundus examination, it is recommended that you dilate the pupil only on the affected side, so an afferent pupillary defect can still be assessed (reverse RAPD).
On Investigation:
- Investigations should not delay the immediate management of a clinically diagnosed retrobulbar haemorrhage with visual compromise.
- Platelet count, coagulation profile if a coagulopathy is suspected or if on an anticoagulant.
- CT orbit (contrast not required). If there is visual loss, CT orbit should not be arranged until after acute management is instituted.

Acute Management:
- If no signs of visual compromise, manage conservatively with close observation. Monitor visual acuity, pupils and IOP.
- Treatment of a retrobulbar haemorrhage with visual compromise requires an emergency lateral canthotomy and cantholysis. Medical management can be commenced concurrently.
- Lateral canthotomy and cantholysis:
  - Betadine® prep and drape
  - Local anaesthetic (2% lignocaine with 1:100,000 adrenaline) into lateral canthus, needle orientated away from globe.
  - Clamp the lateral canthus with artery forceps for ~1min. Pass one tip of the forceps posterior and one anterior to the lateral canthus all the way to the bony orbit. This helps reduce bleeding and displaces oedema for your incision.
  - **Lateral canthotomy**: take blunt tipped scissors and cut the lateral canthus skin all the way to the bony orbit, with the blades straddling the lateral canthus.
  - **Lateral cantholysis** (inferior crus): use forceps to pull the outer edge of the lower lid out to expose the lateral canthal tendon. Then identify the inferior crus of the lateral canthal tendon inferolaterally (can usually strum the fibrous band with closed scissors as it will be taut). Using blunt tipped scissors place one blade on either side of the inferior crus of the lateral canthal tendon. When it is cut the lower lid should become more lax. Confirm by attempting to strum the canthal tendon again. A cantholysis is usually adequate to decompress the orbit If not, consider whether you have properly cut the inferior crus or if you need to cut the superior crus.
  - Apply gentle pressure to the wound as required for haemostasis.
  - Apply chloramphenicol ointment.
  - Lateral canthotomy and cantholysis may be left to spontaneously granulate and heal or may be electively repaired.
Medical management

- Medical management may be considered in conjunction with immediate surgical management. It should not delay surgical treatment if there is visual loss.

- **Acetazolamide**
  - Adult, oral/IV, initial dose of 250 to 500mg. Maintenance, 250mg every 4 hours as required, to a maximum of 1g daily (see [Acetazolamide Prescribing and Administration Guideline](#)).
  - Used to reduce intraocular pressure

- **Mannitol**
  - Adult 1-2g/kg (5-10mL/kg of 20% solution) intravenously over 30 to 60 mins (see [Mannitol Infusion Procedure](#)). Administer with caution to patients with renal/heart disease or severe dehydration.
  - Used to reduce the volume of vitreous

- **Methylprednisolone**
  - Adult 1g IV stat. Consider dose reduction if comorbidities (see [Methylprednisolone Procedure](#)).
  - Used to provide neuroprotection against ischaemia, reduce spasm and oedema of the microcirculation and reduce intra-orbital pressure

- **Other measures:**
  - Admit patient
  - Position in a head up position
  - Monitor visual acuity, IOP and RAPD every 15 minutes for 2 hours, then 30 minutes for the next 4 hours, then hourly.
  - Contact Oculoplastics fellow. Failure of improvement with lateral canthotomy and cantholysis may require formal orbital decompression or surgery to evacuate the haematoma.

Authors:

Jonathan Goh and CPG Working Party

Review date:

20/10/2020
## Evidence Table

<table>
<thead>
<tr>
<th>Author/s</th>
<th>Title</th>
<th>Source</th>
<th>Level of Evidence (I – VII)</th>
<th>Comments</th>
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<tr>
<td>Popat H, Doyle PT, Davies SJ</td>
<td>Blindness following retrobulbar haemorrhage-it can be prevented</td>
<td>The British journal of oral &amp; maxillofacial surgery 2007;45:163-4.</td>
<td>VI</td>
<td></td>
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### The Hierarchy of Evidence

The Hierarchy of evidence is based on summaries from the National Health and Medical Research Council (2009), the Oxford Centre for Evidence-based Medicine Levels of Evidence (2011) and Melynk and Fineout-Overholt (2011).

- **I** Evidence obtained from a systematic review of all relevant randomised control trials.
- **II** Evidence obtained from at least one well designed randomised control trial.
- **III** Evidence obtained from well-designed controlled trials without randomisation.
- **IV** Evidence obtained from well designed cohort studies, case control studies, interrupted time series with a control group, historically controlled studies, interrupted time series without a control group or with case series.
- **V** Evidence obtained from systematic reviews of descriptive and qualitative studies.
- **VI** Evidence obtained from single descriptive and qualitative studies.
- **VII** Expert opinion from clinician, authorities and/or reports of expert committees or based on physiology.
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