# Spontaneous retrobulbar haemorrhage in idiopathic thrombocytopenic purpura

Filipa Caiado de Sousa, <sup>1</sup> Joana Pinto Medeiros, <sup>1</sup> Raquel Marques, <sup>1</sup> Carlos Marques-Neves<sup>1,2</sup>

<sup>1</sup>Department of Ophthalmology, Hospital de Santa Maria, Lisbon, Portugal

<sup>2</sup>Department of CECV, Universidade de Lisboa Associacao para a Investigacao e Desenvolvimento da Faculdade de Medicina, Lisboa, Portugal

## **Correspondence to** Professor Carlos Marques-

Neves, bombordo.seven@gmail.com

Accepted 5 December 2017

### **DESCRIPTION**

An 81-year-old male patient presented to the emergency department with complaints of intense and active subconjunctival haemorrhage, eyelid ecchymosis and proptosis of the right eye (optic disc, OD) with 1 hour of evolution (figure 1). The patient indicated no related pain, headache or recent head trauma. He referred to have reduced vision of the same eye 20 years prior to this event, which he could not explain. His medical history presented an idiopathic thrombocytopenic purpura (ITP), which had been treated with azathioprine for the past 4 years. Visual acuity (VA) of OD was 20/200 (same as previously), with no ocular motility restrictions or afferent pupillary defect. The intraocular pressure (IOP) OD was 22 mm Hg. Fundus examination revealed an OD of normal appearance, with no signs of posterior compression. A non-contrast CT of the brain and orbit revealed thickening and densification of the periorbital soft tissues, relative proptosis of OD and intraconal retro-orbital densification compatible with a right retrobulbar haematoma (figure 2). Further analytical study revealed a thrombocytopenia of  $7 \times 10^9$ /L. We performed local compression and irrigation with cold saline solution, and successfully stopped the haemorrhage.

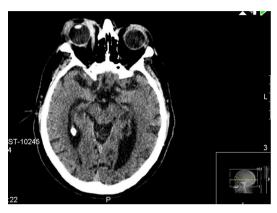
The haematology and internal medicine departments were contacted to provide assistance and carefully evaluate the risk of potentially fatal extraocular bleeding sites.

No platelet transfusion was recommended due the high risk of peripheral destruction. The previous chemotherapy treatment was adjusted, and the patient was started on oral corticosteroids.

Although neuroimaging suggested retrobulbar haemorrhage, the ocular exam showed no signs of compression, the patient reported no recent



**Figure 1** Subconjunctival haemorrhage, eyelid ecchymosis and proptosis of the right eye.



**Figure 2** CT of the brain and orbit showing thickening and densification of periorbital soft tissues, relative proptosis of the right eyeball and intraconal retroorbital densification compatible with right retrobulbar haematoma.

VA loss, and given the haemorrhagic risk of lateral canthotomy and cantholysis, these procedures were not performed.

The patient was re-evaluated daily by ophthal-mologists and haematologists, with a complete ophthalmological examination including evaluation of best corrected VA, IOP measurements, detailed fundoscopic examination and anterior segment inspection to insure normal reabsorption of the haemorrhage, while also closely monitoring the ITP.

Two days later, there was an increase in the patient's platelet count  $(11\times10^9/L)$ , a significant improvement of the proptosis and subconjunctival haemorrhage, with an IOP of 18 mm Hg and no signs of optic compression.

Managing this condition posed a challenge due to the patient's marked thrombocytopenia platelet count, for which there is currently no specific reversal agent. Consequently, our care focused on stabilising the systemic disease and monitoring for orbital compartment syndrome.

Spontaneous retrobulbar haemorrhage (SRH) is a very rare event. The causes can be either local or systemic, such as orbital vascular anomalies, underlying coagulopathy, haemophilia, chronic pharmacological anticoagulation, pregnancy, toxic conditions and uncontrolled hypertension. There are few recorded cases.

One case report described an SRH event in a patient taking rivaroxaban, with an IOP of 32 mm Hg. A canthotomy was performed resulting in bleeding from the canthotomy site and increased ocular bleeding. The patient was given intravenous



**To cite:** Sousa FC, Pinto Medeiros J, Marques R, et al. BMJ Case Rep Published Online First: [please include Day Month Year]. doi:10.1136/bcr-2017-223028

# Images in...

prothrombin complex to reverse the anticoagulatory effect of the drug, and no further surgical treatment was recommended, with no deterioration of VA.<sup>3</sup>

In conclusion, we consider that the treatment of patients with underlying coagulopathy should be multidisciplinary in

## **Learning points**

- ► Signs and symptoms of a spontaneous retrobulbar haemorrhage (SRH) are identical to those of traumatic or postoperative aetiology. SRH may occur in patients with an underlying systemic coagulopathy.
- ► The medical evaluation consists of checking for: afferent pupillary defect, increased intraocular pressure, pulsations of the central retinal artery, choroidal folds and loss of vision.
- ▶ If optic neuropathy is present, immediate orbital pressure relieve with a lateral canthotomy should be performed. However, if surgery involves a high risk and there is no evidence of ocular ischaemia or compressive optic neuropathy, the patient can be monitored with serial examinations.

nature. In this case, under exclusive medical treatment, the patient remained stable and demonstrated no VA impairment.

**Contributors** FCS: substantial contribution to conception and design, acquisition of data, analysis and interpretation of data; drafting the article; final approval of the version to be published. JPM: substantial contribution to conception and design; revising the article for important intellectual content; final approval of the version to be published. RM and CMN: substantial contribution to conception and design; drafting and revising the article for important intellectual content; final approval of the version to be published.

Competing interests None declared.

Patient consent Obtained.

**Provenance and peer review** Not commissioned; externally peer reviewed.

© BMJ Publishing Group Ltd (unless otherwise stated in the text of the article) 2017. All rights reserved. No commercial use is permitted unless otherwise expressly granted.

#### REFERENCES

- 1 Rajabi MT, Hassanpoor N, Parsa R, *et al.* Spontaneous retrobulbar hemorrhage in a patient with breast cancer: a case report. *J Curr Ophthalmol* 2016;28:48–51.
- 2 Kwon JH, Song YJ, Choi SS, et al. Spontaneous intraorbital hemorrhage: a case report. J Korean Neurosurg Soc 2008;44:156–8.
- David B, Michael M, et al. Case report-a 79 year old man experienced continued bleeding following canthotomy/cantholysis for a spontaneous retrobulbar hemorrhage. Emergency Medicine 2016;48:309–12.

Copyright 2017 BMJ Publishing Group. All rights reserved. For permission to reuse any of this content visit http://group.bmj.com/group/rights-licensing/permissions.

BMJ Case Report Fellows may re-use this article for personal use and teaching without any further permission.

Become a Fellow of BMJ Case Reports today and you can:

- ► Submit as many cases as you like
- ▶ Enjoy fast sympathetic peer review and rapid publication of accepted articles
- ► Access all the published articles
- ► Re-use any of the published material for personal use and teaching without further permission

For information on Institutional Fellowships contact consortiasales@bmjgroup.com

Visit casereports.bmj.com for more articles like this and to become a Fellow