Bilateral and multiple sub-internal limiting membrane hemorrhages in a familial retinal arteriolar tortuosity patient by Valsalva-like mechanism: an observational case report

Ting Chen†, Hongmei Zheng†, Yunyun Wang, Junyi Hu and Changzheng Chen*

Abstract

Background: Bilateral and multiple Valsalva-related sub-internal limiting membrane (ILM) hemorrhages in a familial retinal arteriolar tortuosity (FRAT) patient is rare, and we treated this patient by both observation and Neodymium yttrium aluminum garnet (Nd: YAG) laser membranotomy methods.

Case presentation: A 13-year-old female student presented with sudden visual loss and central scotoma in both eyes after running 800 m at the school gym. The examination revealed six sub-ILM hemorrhages with the biggest hemorrhage measuring approximately 1.5-disc diameters (DD) in the right eye and two sub-ILM hemorrhages with the biggest one measuring 5.5 DD in the left eye. The patient was diagnosed as having Valsalva retinopathy associated with FRAT. Nd: YAG laser membranotomy was performed at the biggest hemorrhages and the rest hemorrhages were treated with observation in both eyes. The visual acuity recovered to 20/16 in the right eye and 20/20 in the left eye. Epiretinal membrane (ERM) formation was observed in the left eye.

Conclusions: Nd: YAG laser could be considered for treating premacular hemorrhage in FRAT patient especially when a quick vision recovery was needed. This is the first reported case of a FRAT patient suffering from bilateral and multiple Valsalva-related sub-ILM hemorrhages which were treated by both observation and Nd: YAG laser treatment.

Keywords: Familial retinal arteriolar tortuosity, Valsalva, Neodymium yttrium aluminum garnet, Observation

Background

Familial retinal arteriolar tortuosity (FRAT) is a disorder characterized by pronounced tortuosity of the second-order and third-order retinal arterioles in the macular and peripapillary areas while sparing of the first-order arteriole and the whole venous system. The observation of male to male inheritance as well as occurrence in parents and their children in the absence of consanguinity suggest an autosomal dominant inheritance in FRAT [1]. FRAT patients are often asymptomatic but some may experience transient vision loss due to intra- or preretinal hemorrhages following physical exertion or minor trauma [1, 2]. The long-time visual prognosis is usually excellent.

The Valsalva maneuver is performed by forceful exhalation against a closed glottis causing an increase in intraabdominal and intrathoracic pressure during activities such as coughing, vomiting, straining, or weightlifting. When the increased pressure transmitted to the eye,
the spontaneous rupture of superficial perifoveal retinal capillaries may happen. It can be bilateral but the unilat-
eral characteristic is more common [3]. The hemor-
rhages in most cases resolve spontaneously over several
weeks.

Here, we presented a case of a FRAT patient suffering
from Valsalva-related retinal hemorrhages was treated
with Neodymium yttrium aluminium garnet (Nd: YAG)
laser membranotomy and observation methods. This
case was unusual as it underwent multiple sub-internal
limiting membrane (ILM) hemorrhages in both eyes in
FRAT patient by a Valsalva-like mechanism, as well as
both observation and Nd: YAG laser treatment were
chosen in this FRAT patient.

Case presentation
A 13-year-old female student was admitted to our de-
partment because of sudden visual loss and central scot-
oma in both eyes after running 800 m at the school gym
for 1 week. She had no history of ocular, systemic dis-
ease or trauma. The presenting best corrected visual
acuity (BCVA) was 20/100 in the right eye and 20/160
in the left eye. Her intraocular pressures were 20 mmHg
in the right eye and 18 mmHg in the left eye. Her
pupillary response, ocular motility, external and anterior
segment examinations were unremarkable. Dilated
fundus examination revealed six retinal hemorrhages
with the biggest hemorrhage measuring approximately
1.5-disc diameters (DD) in the right eye and two retinal
hemorrhages with the biggest one measuring 5.5 DD in
the left eye. The hemorrhages showed a double-ring
sign, with the outer ring representing the subhyaloid
bleed and the inner ring representing the sub-ILM
hemorrhage. Several areas of intraretinal hemorrhages
located in and around the posterior pole were also ob-
 served. Moreover, fundus examination of both eyes
showed the tortuosity of second and higher order arteri-
oles which were consistent with FRAT (Fig. 1). Her
mother has similar arteriolar tortuosity in the fundus
(Fig. 2). Spectral-domain optical coherence tomography
(SD-OCT; Spectralis HRA + OCT; Heidelberg Engineer-
ing, Heidelberg, Germany) showed a dome-shaped lesion
with a hyper-reflective band and a hypo-reflective area
beneath consistent with blood at the macula in both
eyes. From the OCT, the hemorrhages were confined
below the ILM (Fig. 1). Findings from fundus fluorescein
angiography (FFA) in both eyes showed the blocked
fluorescence from the retinal hemorrhages, and the sec-
ond- and third-order retinal arteriolar tortuosity with
corkscrew shape in the posterior pole, without vascular
filling defects or hyperfluorescent lesions. No lesions
were detected in the venous vasculature and first-order

Fig. 1 Fundus photographs, FFA and SD-OCT results of a student in both eyes. Fundus photograph showed multiple retinal hemorrhages, as well
as second- and third-order retinal arteriolar tortuosity in the macula and peripapillary area in the right eye (a) and the left eye (d). FFA of the
right (b) and the left (e) eye revealed retinal arteriolar tortuosity with the blocked fluorescence from the retinal hemorrhages. SD-OCT
demonstrated a dome-shaped lesion with a hyper-reflective band and a hypo-reflective area beneath consistent with blood in the right (c) and
the left eye (f)
arteries (Fig. 1). The blood pressure, biochemical test, blood cell count and coagulation were normal. Based on the clinical presentation and her history, the ultimately diagnosis of this patient was Valsalva retinopathy associated with FRAT.

As a student, she and her family request a quick vision recovery. Comparing the potential risks, benefits and costs between surgery and laser, Nd:YAG laser was chosen. Nd:YAG laser (Ultra Q Reflex, Ellex, Australia) with a Goldmann three-mirror contact lens was performed at the lower part of the maximum premacular hemorrhages in both eyes at the same time. The energy was 5–6 mJ with single pulse. Consequently, the hemorrhages were rapidly trapped into the vitreous cavity. Five days after the membranotomy, the premacular hemorrhages had drained mostly with mild vitreous hemorrhages, and her visual acuity was improved to 20/16 in the right eye and 20/20 in the left eye. At 7 weeks, the visual acuity of both eyes were stable. All retinal hemorrhages were resolved completely with little grayish vitreous hemorrhages in both eyes and the ILM was wrinkled in the left eye (Fig. 3). OCT showed subtle loss of foveal contour in both eyes and in the left eye, a laser perforation in the ILM and hypo-reflective premacular cavity was observed (Fig. 3).

**Discussion and conclusions**
FRAT is a rare disease (OMIM %180,000), with the exact etiology is unknown yet. Diagnosis of FRAT is based on the clinical feature of the second and third order tortuosity of arterioles in the peripapillary and macular areas. Moreover, when there is a family history or a positive fundus result of one family member, the diagnosis is more definite. FRAT is usually asymptomatic and the visual symptoms occur with the retinal hemorrhage. However, the hemorrhage is not associated with the severity of the arteriolar tortuosity in FRAT [1]. The hemorrhages are often correlated with mild trauma, physical exertion, or other activities that could result in elevated central venous pressure (Valsalva effect) [4].

This case is unique as our FRAT patients suffering from bilateral and multiple sub-ILM hemorrhages likely because of a Valsalva effect related to the running activity. It was unclear how FART predisposes to Valsalva retinopathy. Some researchers indicated that the tortuosity of the retinal arterioles may be the result of disruption of the extracellular matrix of the arterial wall [5, 6]. Meanwhile, FRAT is thought to be an autosomal dominant disorder associated with a missense mutation in the Collagen types IV alpha-1 (COL4A1) gene in some cases [7]. COL4A1 is the most abundant and ubiquitous basement membrane protein and is present in the basal lamina of various ocular structures and vascular basement membranes [8, 9]. COL4A1 mutation can affect the collagen secretion and accumulate misfolded proteins [7]. So we hypothesized that when a Valsalva-like mechanism induced an increased venous pressure, the hemorrhage originating from the fragility of the tortuous arterioles in FRAT patients happened.

The visual prognosis is often excellent either in FRAT or in Valsalva retinopathy. Following the absorption of the hemorrhages, the visual acuity usually returns to normal [10]. The primary treatment of FRAT and Valsalva retinopathy are both conservation. However, in some observed cases, irreversible visual impairment may occur because of macular pigmentary changes, epiretinal membrane (ERM) formation, or the toxicity of hemoglobin and iron on retina [11–13]. Other treatment options such as Nd: YAG laser membranotomy [14], vitrectomy [15], intravitreal injection of ranibizumab [16], pneumatic displacement of the hemorrhage by gas and tissue plasminogen activator [17] and argon green laser [18] were recommended in Valsalva retinopathy, while not reported in FRAT patient until now. The other particularity of our case was that this FRAT patient was under observation for the little sub-ILM hemorrhages as well as Nd: YAG for the maximum premacular sub-ILM hemorrhages. Both therapies were
effective for our patient. To our best knowledge, this was the first case report of sub-ILM hemorrhages in FRAT patient treated by Nd: YAG laser membranotomy. Although it has been advocated that if the premacular hemorrhage is < 3 DD, Nd-YAG laser should not be performed [19]. However, other study also suggested that if quicker resolution of symptoms was demanded, Nd: YAG laser is an option for premacular hemorrhage [14]. Chang et al. [20] also advised that if Nd: YAG laser is considered, it should be performed as early as possible. As it is hard to achieve the complete drainage of the clotted blood, despite successful laser perforation. As a student, our patient and her family demanded a quick resolution, so we also gave a Nd: YAG laser treatment for the right eye with the premacular hemorrhage was only 1.5 DD.

Some research showed that the ERM formation and ILM wrinkling may occur as a late complication of Nd: YAG laser therapy [21, 22]. In this case, we also observed the ERM formation in left eye. Although we reported that ERM formation occurred in one eye, it did not indicate that the complication rate of Nd: YAG laser membranotomy for treating premacular hemorrhage in FRAT patient was 50% when the studied sample was only one patient. The presence of persistent hemorrhage beneath the ILM may be the stimulus for ERM formation rather than the Nd: YAG laser treatment itself [22]. ILM’s molecular constituents include members of the laminin, nidogen, collagen IV, and proteoglycan families [23]. The mutation in the COL4A1 gene, which encodes type IV collagen in basement membranes, has been reported in FRAT [7]. Thus, it was possible that the FRAT patient may have a defect ILM due to the genetic abnormality, and it may be related to the unsealed and un-reattached ILM membrane observed in this patient. In addition, the ERM formation may originate from the persistent unsealed and un-reattached ILM membrane [22]. This opinion was verified by our patient by the evidence that the ERM formation only occurred in the left eye.

Fig. 3 Fundus photographs and SD-OCT images 7 weeks after Nd: YAG laser membranotomy. Fundus photos showed that all retinal hemorrhages were resolved in the right (a) and left (c) eye, with the internal limiting membrane (ILM) wrinkling in the left eye (c, arrow). SD-OCT revealed subtle loss of foveal contour in the right (b) and left (d) eye, as well as a laser perforation in the ILM and a sub-ILM hypo-reflective area in the left eye (d, arrow)
eye with unsealed ILM membrane while did not observe in the right eye. And there was investigation showed that any elevations of the ILM may seal and reattach among 2–6 months [24]. According to this view, our patient may have a structure alteration in macular in the left eye observed by OCT. Meanwhile, at the follow-up of 7 weeks, our FRAT patient did not complain of metamorphopsia. We will observe this patient for a long time.

We conducted the Nd:YAG laser membranectomy at the lower part of the maximum premacular hemorrhages. But in our patient, the location of ILM defect observed in the fundus after 7 weeks after the membranotomy was far from the inferior border of the initial sub-ILM hemorrhage, which may indicate that the laser performed at an inappropriate location. The explanation for the location of ILM defect far from the inferior border was due to the tangential traction of the ILM [25].

In conclusion, we reported a rare case of a FRAT patient suffering from bilateral and multiple Valsalva-related sub-ILM hemorrhages which were treated by both observation and Nd: YAG laser treatment. Nd: YAG laser could be considered for treating premacular hemorrhage in FRAT patient especially when a quick vision recovery was needed. We recommend that this study avoid physically intensive performances or Valsalva. The long-time efficacy and safety of Nd: YAG laser in FRAT patient needs further study.

Abbreviations
ILM: Internal limiting membrane; FRAT: Familial retinal arteriolar tortuosity; Nd: YAG: Neodymium yttrium aluminum garnet; DD: Disc diameters; ERM: Epiretinal membrane; BCVA: Best corrected visual acuity; FFA: Fundus fluorescein angiography; COL4A1: Collagen types IV alpha-1

Acknowledgements
The authors would like to thank the patient and her family for agreeing to collect and report the clinical data.

Authors’ contributions
TC, HMZ and CZC collected the clinical information of the patient, analyzed and interpreted the clinical data. TC wrote the drafting of this manuscript. HMZ performed the Nd: YAG therapy. YYZ and HY performed the fundus examinations and OCT evaluation. CZC reviewed and edited the manuscript. All authors read and approved the final manuscript.

Funding
None.

Availability of data and materials
All data and materials supporting our findings are contained within this manuscript.

Ethics approval and consent to participate
The ethics approval was obtained from the Renmin Hospital of Wuhan University and all research conducted according to the tenets of the Declaration of Helsinki. Written informed consent to participate was obtained from the patient’s mother.

Consent for publication
Written informed consent was obtained from the patient’s mother for publication of this case report and all accompanying images.

Competing interests
The authors declare that they have no competing interests.

Received: 8 January 2020 Accepted: 30 March 2020
Published online: 15 April 2020

References
1. Sutter FK, Helbig H. Familial retinal arteriolar tortuosity: a review. Surv Ophthalmol. 2003;48(3):245–55.
2. Bartlett WJ, Price J. Familial retinal arteriolar tortuosity with retinal hemorrhage. Am J Ophthalmol. 1983;95(4):556–8.
3. Duane TD. Valsalva hemorrhagic retinopathy. Trans Am Ophthalmol Soc. 1972;70:298–313.
4. Jones WL. Valsalva maneuver induced vitreous hemorrhage. J Am Optom Assoc. 1995;66(3):301–4.
5. Chyatte D, Reilly J, Tilson MD. Morphometric analysis of reticulin and elastin fibers in the cerebral arteries of patients with intracranial aneurysms. Neurosurgery. 1990;26(3):939–43.
6. Skirgaudas M, Awad IA, Kim J, Rothbart D, Criscuolo G. Expression of angiogenesis factors and selected vascular wall matrix proteins in intracranial saccular aneurysms. Neurosurgery. 1996;39(3):537–47.
7. Zenteno JC, Crespi J, Bueno-Velez B, Sull IA, Basaganyas F, Vela-Segarra J, Diaz-Cascajoa J, Marlieges MT. Next generation sequencing uncovers a missense mutation in COL4A1 as the cause of familial retinal arteriolar tortuosity. Graefes Arch Clin Exp Ophthalmol. 2014;252(11):1789–94.
8. Hann CR, Springett MJ, Wang X, Johnson DH. Ultrastructural localization of collagen IV, fibronectin, and laminin in the trabecular meshwork of normal and glaucomatous eyes. Ophthalmic Res. 2001;33(6):314–24.
9. Ishizaki M, Westerhaus-Larson A, Kino J, Hayashi T, Kao WW. Distribution of collagen IV in human ocular tissues. Invest Ophthalmol Vis Sci. 1993;34(9):2680–9.
10. Ding PC, Chen MT. Retinal arteriolar tortuosity with recurrent retinal hemorrhages—case report. Kochi J Med Sci. 2000;16(7):380–2.
11. De Maeyer K, Van Ginderdeuren R, Postelmans L, Stalmans P, Van Calster J. Missense mutation in COL4A1 as the cause of familial retinal arteriolar tortuosity. Br J Ophthalmol. 2007;91(7):869–72.
12. Xie ZG, Yu SQ, Chen X, Zhu J, Chen F. Macular hole secondary to Valsalva retinopathy after doing push-up exercise. BMC Ophthalmol. 2014;14:98.
13. Li N, Zhu Z, Yi G, Li S, Han X. Valsalva retinopathy in twin-pregnancy: a case report and literature review. Ann J Case Rep. 2018;19:155–9.
14. Narag A, Dhingra N. Valsalva retinopathy. Postgrad Med J. 2017;93(1097):174.
15. Kumar V. Optical coherence tomography changes following vitrectomy for long standing premacular hemorrhage in Valsalva retinopathy. Int J Ophthalmol. 2017;10(11):1779–82.
16. Noorlala B, Zunaina E, Raja-Azmni MN. Successful resolution of Preteretinal Haemorrhage with Intravitreal Ranibizumab. Case Rep Ophthalmol Med. 2016;2016:4164198.
17. Chou YK, Huang YM, Lin PK. Sub-internal limiting membrane hemorrhage treated with intravitreal tissue plasminogen activator followed by octafiuropropane gas injection. Taiwan J Ophthalmol. 2015;5(4):198–201.
18. Tirhissi H, Caglar C, Yilmazbas P, Anayol MA, Sekeroglu MA. Argon green laser for Valsalva retinopathy treatment and long-term follow-up of the internal limiting membrane changes in optical coherence tomography. Korean J Ophthalmol. 2015;29(6):437–8.
19. Bypareddy R, Chawla R, Arad SV, Takkar B. Iatrogenic parafocal macular hole following Nd:YAG posterior hyaloiodotomy for premacular haemorrhage. BMJ Case Rep. 2016;2016:bc6201617234.
20. Chang PY, Wang JK, Yang CH. Spectral-domain optical coherence tomography findings of subintimal limiting membrane hemorrhage in the macula before and after Nd:YAG laser treatment. Taiwan J Ophthalmol. 2015;5(1):33–5.
21. Kwok AK, Lai TY, Chan NR. Epiretinal membrane formation with internal limiting membrane wrinkling after Nd:YAG laser membranotomy for macular hemorrhage. Am J Ophthalmol. 2003;136(4):763–6.
22. Zou M, Gao S, Zhang J, Zhang M. Persistent unsealed internal limiting membrane after Nd:YAG laser treatment for vitreoretinal disease. BMC Ophthalmol. 2013;13:15.
23. Henrich PB, Monnier CA, Halfer W, Haitoglou C, Strauss RW, Lim YR, Loparic M. Nanoscale topographic and biomechanical studies of the human internal limiting membrane. Invest Ophthalmol Vis Sci. 2012;53(6):2561–70.
24. Sabella P, Bottoni F, Staurenghi G. Spectral-domain OCT evaluation of Nd:YAG laser treatment for Valsalva retinopathy. Graefes Arch Clin Exp Ophthalmol. 2010;248(4):599–601.

25. Suren E, Akidan M, Erol MK, Toslak D. Spontaneous macular hole closure after Valsalva retinopathy and Nd:YAG laser treatment. Case Rep Ophthalmol Med. 2018;2018:1762847.

Publisher’s Note
Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.