Unsuccessful surgical excision of optic nerve drusen

Magdalena Pfriem · Hans Hoerauf

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Abstract

Background To report the case of failed surgical excision of optic nerve drusen (OND).
Methods Case report. A 53-year-old woman presented with bilateral OND leading to progressive field defects and LE light perception and RE 20/25 vision. A vitrectomy was performed on the legally-blind left eye to test the resectability of OND.
Results The removal of superficial OND failed because the singular-appearing superficial OND presented as one large mass with multiple excrescences on its surface. Retinal vessels passed through it, which made an excision impossible.
Conclusion OND cannot always be removed surgically, due to their variable consistency and hardness.

Keywords Optic nerve drusen · Vitrectomy · Visual field · Surgical technique

Introduction

Optic nerve drusen (OND) are acellular, calcific deposits occurring in 3.4 to 24 % of the population [1]. Changes in axoplasmatic transport and the axonal degeneration of optic nerve fibres are suspected of being involved in OND formation [2, 3]. Most patients are subjectively asymptomatic, but visual acuity, preceded by severe visual field defects, is severely impaired in a small number of patients [1]. There is still no definitively effective treatment of OND [1]. The successful surgical removal of superficial OND was recently reported [4]. In contrast, and to demonstrate the heterogeneity of cases, we present a patient whose surgical excision of OND failed.

Case report

A 53-year-old woman presented with progressive visual field defects in the right eye, and long-standing visual loss in the left eye. Initial examination revealed best-corrected visual acuity of 20/25 RE, light perception LE, a severely constricted visual field in the RE <10° and an unremarkable slit-lamp examination OU. Dilated ophthalmoscopic examination revealed bilateral superficial OND, which appeared as multiflobular, yellowish-white, sharp-edged nodules and anomalous arterial and venous branchings, increased tortuosity, and cilioretinal arteries (Fig. 1).

To test the resectability of OND and to gain experience, we decided to perform a vitrectomy with surgical excision on the blind left eye before considering surgical manipulations on the right eye with the risk of losing the remaining small central visual field, leading to complete blindness. After informing the patient about the procedure’s experimental character and obtaining informed consent, we conducted a standard three-port pars plana vitrectomy. We then performed a small retinotomy at the nasal edge of the optic nerve. Despite numerous efforts using different vitreoretinal surgical instruments such as blades, forceps and fine scissors, removal of the superficial OND failed. During surgery, we became aware that the small, multiple, and singular-appearing superficial ONDs presented as the multilobulated surface of one calcified crust covering the complete optic nerve head, with central retinal vessels...
passing through (Fig. 2). It was impossible to split the 
very hard crust into small pieces and to remove 
considerable parts of it. Moreover, radial neurotomy as 
described in the literature was impossible due to the 
hardness of the OND material [5]. Manipulations were 
stopped and the eye was filled with air.

Discussion

Axonal degeneration, ischaemia within the optic nerve 
head, and OND pressing down on prelaminar nerve fibres 
have been proposed as pathogenetic mechanisms in OND 
disease [1, 3].

A recent publication reported on a good functional 
outcome after vitrectomy and radial optic neurotomy 
(RON) for OND, suggesting that RON leads to mechanical 
decompression with subsequent optic nerve fibre recovery, 
and that the excision of prelaminar deposits is an effective 
strategy in OND disease [5]. However, our case shows that 
surgical removal or decompression is not possible in all 
cases; our patient was considerably older than theirs.

A histological study in 1981 demonstrated that the 
number and size of superficial and buried OND vary 
greatly [3]. Furthermore, their size and density increase 
with age, due to continuous calcium apposition [1]. Rather 
than revealing multiple small deposits, our patient’s OND 
formed a single, large solitary mass with central retinal 
arteries and veins passing through it. Our observation 
concurs with a report by Kapur et al., who reported on 
the incomplete surgical excision and bisection of OND in a 
59-year-old patient “because retinal vessels were observed 
to be emanating from the mass” [4]. In contrast to their 
experience, in our patient the excision or even splitting of 
this large, extremely hard OND crust without damaging 
adjacent structures was not feasible.

Thorough fundus biomicroscopy may help identify candi-
dates for surgery. Other diagnostic methods such as auto-
fluorescence fundus photography, ultrasound B and optical 
coherence tomography are available, but they do not enable us 
to assess the size, formation and hardness of OND.

In conclusion, this case report demonstrates that OND 
removal is not feasible in every case. Their variability in 
consistency and hardness makes them unpredictable, thus 
we cannot know in advance which patient is an appropriate 
candidate for surgery. Patient age seems to be an important 
factor associated with the composition of OND, but also 
early surgery could perhaps yield superior results. In 
accordance with Kapur et al., we propose that other surgical 
or ablating laser techniques must be developed to achieve 
safe removal of extremely hard material, while avoiding 
damage to adjacent, functionally important and sensitive 
structures such as optic nerve fibres and retinal vessels [4].

Conflict of interest  None

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