Fusiform Intracanalicular Ophthalmic Artery Aneurysm; Case Report and Review of Literature

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A 35-year-old man’s vision had progressively deteriorated over a 3-month period. His left visual acuity was 5/20. Enhanced orbital computed tomographic (CT) scans revealed a fusiform dilatation of the ophthalmic artery in the left optic canal. Cerebral Angiography revealed a fusiform aneurysm on the left ophthalmic artery in the optic canal, measuring 6.2×4.6 mm in size. Four days after admission, visual acuity dropped to hand-motion. Endovascular treatment was chosen and a microcatheter was guided into the proximal segment of the ophthalmic artery. Using 4 detachable coils, parent artery occlusion was done. Three months after the intervention, the visual acuity in his left eye improved to 20/20. Dramatic recovery of visual acuity is exceptional with an ophthalmic artery trunk aneurysm. When an occlusion of the proximal ophthalmic artery is the only treatment option in such a situation, the endovascular occlusion of the proximal ophthalmic artery is quite feasible in the sense that it does not require any optic nerve manipulation.

KEY WORDS: Detachable coil · Fusiform aneurysm · Intracanalicular portion · Ophthalmic artery trunk aneurysm.

INTRODUCTION

"Ophthalmic artery aneurysm" usually refers to a carotid-ophthalmic junction aneurysm. Due to its rarity, only a small number of true ophthalmic artery aneurysms have been reported. Intracanalicular ophthalmic artery aneurysms are even rarer, with only one previously reported case.

Because the chance of rupture is not high, the benign nature of an aneurysm with minor symptoms in this location has been previously noted. However, since the ophthalmic artery trunk runs in intimate proximity to optic nerve, these aneurysms often create vision-related symptoms, which respond poorly to surgical treatments. We report a case of a fusiform intracanalicular ophthalmic artery aneurysm presented with progressive visual loss, which recovered dramatically after coiling.

CASE REPORT

A 35-year-old man with a 3-month history of progressive visual loss was transferred to the neurosurgical department after an initial impression of retrobulbar optic neuropathy at a local clinic. He reported 2 minor traffic accidents, but past medical records and radiologic study results were not available. Visual acuity was 20/20 in the right eye and 5/200 in his left. A relative afferent pupillary defect of the left eye was noted. Ocular pressure was normal. Biomicroscopic and ophthalmoscopic examination revealed no remarkable abnormal findings. Fluorescein angiography did not show any leakage or delayed filling. Time to P1 of flash visual evoked potential was delayed to 159 ms in the left eye compared to 132 ms in the right eye. His visual field was not checkable due to his failing vision. An orbit magnetic resonance image (MRI) revealed a signal void in the left optic canal (Fig. 1A) MR angiography showed an aneurysm in the proximal portion of the left internal carotid artery (ICA) ophthalmic segment (Fig. 1B). An ophthalmic artery aneurysm was suspected. An orbital CT with enhancement revealed a fusiform dilatation of the ophthalmic artery in the left optic canal (Fig. 1C). A widened left optic canal with chondromal bony erosion was observed (Fig. 1C, D).

Digital subtraction cerebral angiography with left carotid injection showed a fusiform ophthalmic artery aneurysm in the intracanalicular segment, measuring 6.2×4.6 mm in size (Fig. 2A). External carotid artery (ECA) collateral from the infraorbital artery was well maintained. Four days after admission, his left visual acuity dropped to hand-motion status. The fusiform nature and the intracanalicular location of the ophthalmic artery aneurysm made...
a usual neck clamping procedure difficult. The only remaining option was the parent artery occlusion either by surgery or endovascular intervention. The effect of obliterating the parent artery by coil seemed equally effective as surgical occlusion. Since it does not require direct manipulation of optic nerve, endovascular treatment was chosen. A 6F sheath was inserted in the right femoral artery and a 6F guiding catheter was placed high to the left cervical ICA. Using a microcatheter (Excelsior 10, Boston Scientific), the proximal ophthalmic artery was selected (Fig. 2B).

With 4 platinum coils (GDC, Boston scientific), the proximal segment of the ophthalmic artery harboring a fusiform aneurysm was obliterated (Fig. 2C). Postintervention angiography revealed complete occlusion of the left ophthalmic artery (Fig. 2D), and choroidal crescent supplied by the external carotid collateral (Fig. 2E).

In following day, visual acuity of the left eye improved to finger-count status and the pupillary reflex was prompt. Four days later, his visual acuity increased to 20/25. Fluorescein angiography performed after 1 week demonstrated no delayed filling or ischemic change. Time to P1 of pattern reversal visual evoked potential was still delayed to 130 ms in the left eye compared to 100 ms in the right eye. Three months after the intervention, a control cerebral angiography was performed and showed a well-obliterated proximal ophthalmic artery (Fig. 3). After 2 years, MR angiography showed no evidence of recanalization or regrowth of the aneurysm. His vision was 20/20 in both eyes.

**DISCUSSION**

The ophthalmic artery is composed of three segments; intracranial, intracanalicular and orbital segments. Unlike usual ophthalmic artery aneurysms, which arise from the junction of the ICA trunk and ophthalmic artery, an ophthalmic artery trunk aneurysm is extremely rare. There have been a small number of reported cases of intrabulbar and intracranial segment aneurysms. However, only one case of intracanalicular segment aneurysms has been reported so far.

Previously reported aneurysms in the intracranial and
intraorbital segments have been often accompanied by arteriovenous fistula, arteriovenous malformation, moyamoya disease, evident trauma or with aneurysms in different locations, In the current case, luminal irregularities (Fig. 2A), the ectatic and stenotic segments on the proximal and distal portion of aneurysm (Fig. 2A) suggest common features of the dissecting aneurysm. Although the patient reported two previous traffic accidents, there was no evidence of skull fracture or deformed orbital structure caused by previous traumas. This aneurysm must have enlarged very slowly, slowly enough to cause the total erosion of the ethmoidal wall and a widening of the optic canal (Fig. 1C, D).

The symptoms of a ruptured ophthalmic artery trunk aneurysm depend on its localization. When they rupture, intracranial segment aneurysms cause subarachnoid hemorrhage (SAH), whereas intraorbital segment aneurysms may result in intraorbital bleeding. However, the incidence of bleeding is known to be low, and the benign nature of this aneurysm has previously been commented on. Most commonly, they present with eye-related symptoms, such as visual disturbance, visual field defect and sometimes exophthalmos. Visual disturbance is more likely to develop with an intracanalicular segment aneurysm. In the optic canal, an aneurysm doesn't have enough space to expand, and the intracanalicular optic nerve is known to have a poor blood supply from the superior hypophysial artery, not from the ophthalmic artery itself. Visual disturbance associated with ophthalmic trunk aneurysms responds poorly to treatment. Once it starts, it is difficult to reverse. Of the treated ophthalmic artery trunk aneurysms cases where the initial symptom was visual disturbance, postoperative visual function was better than the preoperative state in only one case, where preoperative visual disturbance had been mild. Two reported cases showed similar postoperative visual acuity results to the preoperative state. With these clinical characteristics, ophthalmic trunk aneurysm associated with the progressive visual disturbance should be treated urgently.

In the current case, the fusiform nature and intracanalicular location precluded the usual neck clipping procedure. The only remaining option was the parent artery occlusion either by surgery or endovascular intervention. There are numerous anastomoses between the external carotid and ophthalmic artery, and this collateral blood flow can prevent ocular ischemia after occlusion of the internal carotid and ophthalmic artery in 90% of cases. However, surgical occlusion of the ophthalmic artery has not been so fruitful in previous reports. The location immediately near the optic nerve makes it difficult to ligate the pathology and unroofing the optic canal may cause additional damage to the nerve. In a case with an intraorbital segment aneurysm reported by Ernemann et al., postoperative angiography showed a choroidal crescent supplied by the external carotid collateral, but the patient's visual function did not improve. Ogawa et al. trapped and resected the intraorbital segment aneurysm. Four months after the surgery, visual acuity was still at finger-count status, even though a branch of the right external carotid artery fed the choroidal crescent. Piché et al. failed to accomplish occlusion of the proximal ophthalmic artery because of the adherence to the inferior aspect of the optic nerve. Later, they occluded the ophthalmic artery with endovascular detachable coils, which yielded a preoperative baseline visual acuity status.

Upon reviewing the aforementioned previous reports, the recovery of visual acuity from hand-motion status is very exceptional. It is not easy to conclude which factors contributed to the reversal of the visual symptoms in the current case. It is believed that endovascular occlusion played a role, because this obviated the optic nerve manipulation.

**CONCLUSION**

We reported a case of a fusiform intracanalicular ophthalmic artery aneurysm presented with decreased visual acuity which improved completely after parent artery occlusion by coiling.

**References**