Diffusion-weighted MRI in acute posterior ischemic optic neuropathy

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Abstract

Blindness following surgery, especially cardiac surgery, has been reported sporadically, the most common cause being ischemic optic neuropathy. The role of MRI in the diagnosis of this condition is not well established. We present a case of postoperative posterior ischemic optic neuropathy that was diagnosed on diffusion-weighted MRI.

Key words: Postoperative vision loss; ischemic optic neuropathy; diffusion-weighted MR imaging

Introduction

Perioperative vision loss¹ due to ischemic optic neuropathy may present as sudden blindness after surgeries such as cardiac surgery, head and neck surgeries, spine surgeries and major abdominal surgeries.³ Anterior ischemic optic neuropathy is easily diagnosed by fundoscopic findings, but posterior ischemic optic neuropathy (PION) is difficult to diagnose.¹

Postoperative PION may occur without any other ischemic effect in the eye or central nervous system¹ and may be unilateral or bilateral. In cases of bilateral PION, a high index of suspicion is necessary for diagnosis as the optic nerves initially appear normal and no afferent pupillary defect may be evident.³

The utility of imaging in ischemic optic neuropathy was not known until recently.⁴ We report a case of PION that was diagnosed by diffusion-weighted MRI.

Case Report

A 50-year-old man presented to the hospital with sudden loss of vision in his right eye, followed by loss of vision in the left eye on the next day. He had undergone coronary bypass surgery and mitral valve replacement 15 days before the presentation, and the immediate postoperative period had been uneventful.

MRI was performed with a 1.5T Siemens Symphony® MRI scanner (Siemens Erlangen, Germany). A standard quadrature coil (head coil) was used. The section thickness for diffusion-weighted MRI was 5mm, with 0.5mm gap, matrix: 128 × 128, and field of view: 30 cm. MRI of the brain was normal. However, it showed symmetric high signal intensity in the optic nerves on diffusion imaging [Figure 1A], with decreased apparent diffusion coefficient (ADC) [Figure 1B], indicating acute ischemic injury/infarction. The ADC was low (25–35) in both optic nerves. The anterior-most part of the optic nerves, close to fundus, did not show diffusion restriction. Conventional spin-echo sequences were unremarkable [Figure 1C].

The patient was managed conservatively but he did not regain vision even after 2 months.

Discussion

Vision loss due to ischemia can be broadly classified⁵ into anterior ischemic optic neuropathy and posterior ischemic optic neuropathy as the blood supply to the anterior part
and posterior part of the optic nerves is different and as the mechanism of ischemia is also different.

The anterior portion of the optic nerve, which includes the optic disc and the small part of the optic nerve within the scleral canal, is supplied mainly by the short posterior ciliary arteries via the choriocapillaries around the optic disk or through branches that form an anastomotic microvascular ring around the optic nerve. Rarely, there is a double arterial supply, with pial branches of the ophthalmic artery also providing blood supply to this section of the optic nerve. Because of this rich blood, this part is less prone to ischemia.

The posterior portion of the optic nerve has a peripheral vascular supply from pial branches of the ophthalmic artery. The central retinal artery often supplies branches to the central fibers. However, the blood flow in the posterior optic nerve is significantly lower than that in the anterior portion.

Hence, this part of the optic nerve (i.e., the segment between the optic foramen and the the point where the central retinal artery enters the nerve) is relatively more susceptible to ischemia during the postoperative period, as the only supply is through centripetal pial vessels that are easily compressible.

Posterior optic nerve ischemia\(^1\) however is less common than anterior ischemic optic neuropathy. PION has been reported after various surgeries like laparotomy and, cardiac, neck and spinal surgeries.\(^2\) The exact cause is not clear; proposed theories include severe hypotension and anemia and, rarely, other factors like infection, congenital absence of central retinal artery, venous occlusion, etc.

Only five reports of MRI findings of ischemic optic neuropathy\(^3\) could be found in the literature after an extensive search. Our patient is the only the second case to be reported of postoperative/perioperative PION\(^4\) showing restricted diffusion. Other reported cases of ischemic optic neuropathy with restricted diffusion were due to mucormycosis.\(^5\)

## Conclusion

Radiologists should be aware of this entity and attention should be paid to the diffusion-weighted images, where the ischemic or infarcted optic nerve shows restricted diffusion. Clinicians should be encouraged to refer patients for MRI if PION is suspected.

## References


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**Figure 1 (A-C):** Diffusion-weighted (b1000) MRI (A) shows hyperintense signal in the optic nerves (arrows). ADC (apparent diffusion coefficient) image (B) shows restricted diffusion (arrows) in the optic nerves. STIR-coronal image (C) of the orbits posterior to the globes shows normal optic nerves.