Central retinal artery occlusion and optic neuropathy secondary to platelet rich plasma injection: a case report

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Dear Editor,

I am Dr. Norashikin Maslan, from the Department of Ophthalmology of University Kebangsaan Malaysia Medical Centre in Kuala Lumpur, Malaysia. I am writing to present a case report on central retina artery occlusion (CRAO) and optic neuropathy secondary to platelet rich plasma (PRP) injection.

In recent times, PRP injection is increasingly being used in medical specialties especially in aesthetic industry. PRP injection delays ageing with the presence of cytokines and growth factors in it\(^1\)\(^-\)\(^2\). It is administered either topically, superficially or through deep dermal injections\(^3\). The injection promotes the production of type 1 procollagen (skin fibroblast) and hyaluronic acid which in turn improves the skin texture, firmness and color homogeneity\(^4\). There are, however, some minor adverse effects of PRP such as discomfort, small prick bleeding and swelling that have been reported\(^4\). Although many have noted its efficacy\(^4\)\(^-\)\(^5\), we would like to highlight a visually threatening complication caused by pure PRP injection in this report.

CASE PRESENTATION

A 48-year-old woman with no known medical illness arrived with a sudden onset loss of vision in left eye (OS) immediately after receiving a PRP injection on the glabellae region by an aesthetic practitioner. The monocular loss of vision was accompanied by an ocular discomfort. It is important to also note that her previous PRP injection, however, was uneventful and patient denies skin filler injections.

Patient was referred to a nearby ophthalmology clinic within 3h after the initial complaint. Oral medication was given to lower her intraocular pressure (IOP). Her vision did not improve regardless, so she decided to seek treatment at our center.

She arrived in 9h after the loss of vision. Her vision was 20/20 in right eye (OD) and 20/60 OS. There was a grade 3 relative afferent pupillary defect (RAPD) OS. Anterior segment examination OS was unremarkable with an IOP of 16 mm Hg. A fundus examination revealed generalized edematous retina with dull foveal reflex but no cherry red spot present (Figure 1A). Humphrey visual field (HVF) 24-2 test showed generalized constricted visual field OS with normal finding OD (Figure 4A).

Four days after the PRP injection, the patient developed left periorbital hematoma associated with worsening of vision, occasional headache, and numbness over the glabella site. Her visual acuity dropped to 10/200 in OS. There was subconjunctival hemorrhage with anterior chamber cells of +4. The fundus was hazy but no vitritis was seen. MRI orbit showed perineural enhancement of the optic nerve (Figure 5). The magnetic resonance venography (MRV) was normal and there was no retro-orbital hemorrhage. She was therefore treated for optic neuropathy and covered with intravenous methylprednisolone 250 mg q.i.d. for 3d followed by an oral prednisolone (1 mg/kg·d) for 11d.

Fortunately, her vision improved slowly. Three months after the event, visual acuity has improved to 20/20 OS, but RAPD remained positive (grade 2) and red color saturation was 8/10 OS. Her periorbital hematoma resolved. Anterior segment...
examination was unremarkable, but optic disc OS became pale with an increase cupping to 0.9 (Figure 1B). Repeated OCTA showed improvement in perfusion and vessel density of the SCP (Figure 6). Her latest HVF also showed significant improvement (Figure 4B).

**DISCUSSION**

Treatment with PRP is generally safe and effective. While it appears simple to administer, devastating intravascular complications can still occur. Periocular vascular events are particularly important to be aware of as immediate intervention may restore the ocular circulation and prevent permanent blindness. Inadvertent intravascular injection of materials, either fillers, PRP or local anesthetic medications can cause retrograde flow to the original artery resulting in the obstruction of the retinal vascular lumen\(^1\). The inner retina layer is supplied by central retinal artery and occlusion will result in ischemia of the inner retina layer, leading to edema and subsequently atrophy. In CRAO, some vision may be preserved since the choroidal circulation also supplies part of the inner retinal tissue. Experiments on animal models have demonstrated irreparable retinal damage occurs after 105 min of vascular occlusion\(^1\). Visual loss following filler injections has been reported although its occurrence is very rare. Facial injections can be performed at various locations such as at the glabella, nasolabial fold, or periorbital region. These regions are rich in
blood supply. Injections at the glabella is commonly associated with compression or obstruction of blood vessel because the supraorbital/supratrochlear artery which supplies this area does not have a strong collateral circulation[9].

Blood supply to the face is from branches of the external and internal carotid arteries. Collaterals between the anterior and posterior ethmoidal arteries medial to the orbit, between the supraorbital and infraorbital artery anteriorly and the and posterior ethmoidal arteries medial to the orbit, between the periorbital and orbital region. Angular artery supplies the zygomaticotemporal branches laterally provides a rich supply to the periorbital and orbital region. Angular artery supplies the dorsal part of the nose and inferomedial fossa of the orbit[9]. Connections between the terminal branches of angular artery with supraorbital/supratrochlear artery plays an important role in establishing anastomosis[8].

In our case, the patient received PRP injections at the glabella area which is an area known to have arterial anastomosis. Hu et al[10] highlighted three essential concomitant factors that eventually lead to spread into the ophthalmic circulation which are retrograde movement of material, forceful injection with high pressure and adequate amount of injectable material. Following that, the embolus can travel along the central retinal artery causing occlusion of the retinal vessels.

Currently, limited literature suggested a correlation between CRAO and optic neuropathy and to our knowledge, this is the first case of unilateral CRAO with optic neuropathy after a PRP injection. In this patient, optic neuropathy was diagnosed based on perineural optic nerve enhancement seen on MRI. Optic neuropathy may be caused by either ischemia or inflammation, leading to disturbance of the blood-optic nerve barrier. We postulate that the retrograde flow of the PRP material may have affected both the central retinal artery and the posterior ciliary artery that mainly supplies the intra-orbital and intraocular portion of the optic nerve, resulting in impairment of small nutrient vessels feeding the nerve by ischemic or inflammatory process and consequently causing optic neuropathy.

Our case study shows that OCTA is a useful tool to establish reduced perfusion and vessel density, conforming to the study by Yang et al[10] that vessel density of the SCP and a 300 μm area around the foveal avascular zone is significantly reduced in retinal artery occlusion. OCTA is also a non-invasive alternative diagnostic tool to monitor progression of patients with vascular occlusions[11].

To conclude, our case highlights a potentially blinding complication of PRP injection around the ocular region. Even though it is a rare complication, ophthalmologist need to be aware of this due to its devastating complication. Prompt recognition and appropriate intervention may help prevent irreversible visual loss.

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