Bilateral branch retinal artery occlusion in a child with nephrotic syndrome

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DESCRIPTION

A 5-year-old girl was referred to our retina clinic with the chief problem of painless diminution of vision in both eyes for 1.5 years, more so in the last 1 month. Medical records showed that she had been on oral steroid and immunomodulator (cyclosporine) for nephrotic syndrome (NS) (biopsy proven minimal change disease) for last 6 months. She was also under treatment for dyslipidaemia. At presentation, her best corrected visual acuity was 20/400 in the right eye and 20/40 in the left eye. The pupils were round, regular and reacting to light in both the eyes without any afferent pupillary defect. Slit lamp examination showed an unremarkable anterior segment with an intraocular pressure of 17 and 14 mm Hg in the right and left eyes, respectively. Colour vision with Ishihara pseudoisochromatic chart was unremarkable in the left eye, while she could not even read the demo plate in the right eye. Fundus examination of both the eyes showed temporal disc pallor, bright internal limiting membrane reflex over posterior pole except nasally over the areas of retinal thinning and retinal pigment epithelial (RPE) alteration between the disc and fovea (figure 1A,B). There were sclerosed arterioles just inferotemporal to optic disc in both eyes better appreciated in autofluorescence imaging (figure 1C,D). Horizontal trans foveal line scan revealed inner retinal thinning with preservation of outer retinal layers and thickened choroid in both the eyes (figure 2A,B). The entire nasal half of the macula was found to be thinned in the macular thickness map in both the eyes (figure 2C–F). Fundus fluorescein angiography could not be done because of preexisting renal pathology. Based on the above, she was diagnosed to be a case of resolved branch retinal artery occlusion in both the eyes. The girl was referred to the treating physician who advised subcutaneous low molecular weight heparin based on systemic coagulation profile.

NS, a common renal disorder in childhood, with a worldwide incidence between 1.52 and 16.9/100 000/year, is characterised by a deranged renal filtering system.¹ The clinical and biochemical features are attributed to the heavy proteinuria > 40 mg/m²/hour, resulting in hypoalbuminemia (<2.5 g/dL), hypercholesterolemia and anasarca. Minimal change disease is the most common cause of NS in paediatric population. Though the exact pathogenesis remains unknown, autoimmunity plays an important role in altering the integrity of the glomerular basement membrane leading to intense proteinuria and intravascular protein depletion.² Next to infection, thromboembolism is often

Figure 1 Colour fundus photographs of bilateral eyes show temporal optic disc pallor, areas of retinal thinning with bright internal limiting membrane reflex nasal to fovea, sclerosed arterioles temporal to disc with localised thinning and retinal pigment epithelial (RPE) alteration in the area between the disc and fovea at the posterior pole (A and B) and sclerosed arterioles could be better appreciated in autofluorescence image (C and D).

considered as the most significant life-threatening complication of NS.³ Hypovolemia, haemoconcentration, dyslipidaemia, urinary loss of anticoagulants like antithrombin III, proteins C and S, increase in circulating fibrinogen, factors V and VIII, and diminished fibrinolytic activity leads to a hypercoagulable stage in 97% of these patients. Though most of the thromboembolic events occur

Figure 2 Swept source optical coherence tomography of bilateral eyes shows the presence of symmetrical bilateral thinning of inner retinal layers nasal to fovea along the papillomacular bundle with preservation of outer retinal layers (A and B) with features being better appreciated in retinal thickness map (C–F). ETDRS, Early Treatment Diabetic Retinopathy Study; ILM, Internal Limiting Membrane; OS, left eye; RPE, retinal pigment epithelium.

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Images in...

within the first 6 months of the disease, it can occur at any time during the disease course.⁴

Ocular manifestations of NS, including posterior subcapsular cataract, increased intraocular pressure, ptosis and keratitis, are mostly related to prolonged corticosteroid usage.⁵ Retinal vascular involvement has rarely been reported and could be secondary to hypertension or arterial or venous thromboembolism. Sinha *et al* has reported a case of bilateral combined central retinal artery and vein occlusion in a 3-year-old child with NS with good visual recovery after systemic treatment.⁶ However, a possible diagnosis of Purtscher-like retinopathy was suspected by Dwivedi *et al* for the above reported case in their letter to editor.⁷ Dwivedi *et al* have reported a case of Purtscher-like retinopathy in a 12-year-old male child with steroid-dependent NS with multiple episodes of relapse.⁸

The visual outcome depends on the type and extent of retinal vasculature involved, timing of presentation and prompt intervention, if any. While a large embolus can block central retinal artery, the smaller ones can block a branch retinal artery, precapillary arteriole or distal capillaries leading to branch retinal artery occlusion, Purtscher flecken or cotton wool spot,

respectively. Central retinal artery occlusion can cause profound visual impairment (and early presentation), whereas a branch retinal artery or smaller order retinal vascular occlusion can at times remain asymptomatic leading to late presentation. In such cases, indirect evidences like selective thinning of inner retinal layers or sclerosed arterioles can help in the diagnosis as in the present case. To the best of our knowledge, bilaterally symmetric branch retinal artery occlusion in a child with NS has never been reported in the literature (Medline search with key words 'branch retinal artery occlusion' and 'nephrotic syndrome'). Paediatricians should be aware of these sequelae. Any child with NS and visual complaints should be promptly referred to an ophthalmologist for a detailed evaluation. Patients diagnosed with fresh or old vascular occlusion should be kept under adequate prophylaxis and follow-up for preventing a similar thromboembolic event happening in the eye or any other organs in the body.

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Patient's perspective

We are happy that our daughter's disease could be diagnosed at an early stage so as to avail of immediate further consult and treatment. We have been apprised about my condition, its natural course and possible complications. The need for regular follow-up and self-awareness of visual loss has been explained to us in a language that we can easily understand.

Learning points

- All patients of nephrotic syndrome (NS) presenting with visual complaints must be evaluated thoroughly.
- Retinal artery occlusion in a child with NS is mostly associated with a systemic hypercoagulable state. Urgent physician consultation is mandated to avoid multiorgan complications secondary to thromboembolism.

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