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Case Report

Management of Coats' disease with Razumab™ (World's first biosimilar ranibizumab) and laser photocoagulation: A case report

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ABSTRACT

We report a case of a 19-year-old Indian male with stage 2 Coats' disease who was managed with monthly intravitreal biosimilar ranibizumab injections (n=3) along with laser photocoagulation in the right eye. Post-treatment, significant reductions were seen in the exudation, angioma-like lesions, central macular thickness and edema in the retina. At the last follow-up at 9 months after intravitreal biosimilar ranibizumab initiation, distant vision improved from best corrected visual acuity (BCVA) of 5/60 at baseline to 6/18, and near vision from 'not able to read' to N/12 in the right eye.

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1. Introduction

Coats' disease, a rare idiopathic unilateral ophthalmic condition, is associated with development of retinal vasculature (telangiectasia) causing exudative retinopathy (intra- and sub-retinal exudation), which may result in retinal detachment leading to vision loss.¹ The therapeutic armamentarium of Coats' disease include laser photocoagulation, cryotherapy, surgery and triamcinolone.² Anti-vascular endothelial growth factor (VEGF) agents, including ranibizumab, have also demonstrated beneficial effects.³ Razumab™, the world's first biosimilar ranibizumab, has demonstrated efficacy and safety in several macular disorders.^{4,5} Limited data exists on Coats' disease from Indian subcontinent. We report a case of an Indian patient with Coats' disease managed using biosimilar ranibizumab.

2. Case Report

A 19-year-old man presented with diminution of vision in both eyes since 6 months. There was no past history of any

significant ocular disease.

On examination, the best corrected visual acuity (BCVA) in the right eye was 5/60 with counting of fingers close to the face (CFCF) in the left eye. Examination of the fundus demonstrated a large area of sub-retinal exudation and multiple angioma-like lesions in the center of exudation in the entire fundus in both eyes (Figure 1). The fundus fluorescein angiography (FFA) demonstrated leaking aneurysm-like vascular anomalies (Figure 2) and optical coherence tomography (OCT) showed collection of intraretinal and subretinal fluid in both eyes (Figure 3). The central macular thickness (CMT) was 691μ and 317μ in right and left eye, respectively. The diagnosis was made as stage 2 Coats' disease.

Monthly intravitreal biosimilar ranibizumab was administered for 3 months in the right eye followed by 3 sittings of laser photocoagulation. In the left eye, 3 sittings of laser photocoagulation were given whereas biosimilar ranibizumab was not administered on patient's unwillingness due to the poor visual prognosis.

After the first intravitreal biosimilar ranibizumab injection, his BCVA improved to 6/24, and edema was reduced to some extent; no significant improvement seen

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after the second injection. After 3 months, significant improvement was demonstrated in vision with BCVA of 6/18, and a significant reduction in the exudation, edema in retina and angioma-like lesions (Figure 4) was reported in the right eye. Also, a reduction in the CMT from 691 μ to 339 μ was observed and the OCT demonstrated uneventful recovery in the right eye (Figure 5). In the left eye, the vision remained same whereas CMT increased from 317 μ to 672 μ .

The patient's last follow-up done in October 2018 demonstrated improved distant vision of 6/18 from the baseline (January 2018) vision of 5/60 and improved baseline near vision from not able to read to N/12 vision in the right eye. There were no improvements in the distant or near visions in the left eye.

Fig. 1: Fundoscopy images showing sub-retinal exudation and multiple angioma-like lesions together with exudates and telangiectasis as well as dilation of retinal vessels (A) right eye (B) left eye

Fig. 2: FFA showing several leaking aneurysms (A) right eye (B) left eye

3. Discussion

Anti-VEGF agents have modernized the treatment of several retinal diseases in adults including diabetic retinopathy and are also used in children.^{6,7} Bevacizumab, an anti-VEGF, has been studied extensively in managing Coat's disease but its use is associated with a risk of vitreoretinal fibrosis and tractional retinal detachments, hence, should be used with caution.⁸ Ranibizumab, an anti-VEGF, has been approved for the treatment of several macular disorders including wet age-related macular degeneration (AMD), retinal vein occlusion (RVO), diabetic macular edema (DME), and myopic choroidal neovascularization (CNV).^{9,10} Ranibizumab has also been evaluated for

Fig. 3: OCT imaging showing intraretinal and subretinal fluid accumulation (A) right eye (B) left eye

Fig. 4: Fundoscopic imaging at 3 months post-treatment with intravitreal biosimilar ranibizumab showing reduction in exudation in right eye

Fig. 5: OCT imaging at 3 months post-treatment demonstrating (A) recovery in the right eye (B) worsening in the left eye due to presence of fluid

the treatment of Coats' disease in children in previous reports.^{3,11,12} We report the successful management of Coats' disease with adjuvant intravitreal biosimilar ranibizumab injection along with laser photocoagulation. The use of anti-VEGF agents may enhance the activity of laser therapy by decreasing the vascular permeability of the capillary endothelial cells.¹³

The treatment goal in the Coats' disease is focused at managing the telangiectatic vessels to control and resolve intra and sub-retinal exudates and eventually vision salvation.² The patient, in the present case-scenario, demonstrated vision recovery in the right eye where both intravitreal biosimilar ranibizumab and laser therapy were employed compared to no improvement in the left eye, which was managed using laser therapy only. Furthermore, biosimilar ranibizumab treatment showed reduction in intraretinal and subretinal exudation, multiple angioma-like lesions, telangiectasis as well as in the dilation of retinal vessels in our patient. There were no injection-related ocular or systemic adverse events during treatment and follow-up. Thus, the adjuvant administration of intravitreal biosimilar ranibizumab was appropriate in the present case-scenario.

Intravitreal ranibizumab combined with other ablative therapies was effective in visual acuity improvement from logMAR BCVA of 1.4 ± 0.7 to 1.1 ± 0.5 ($P < 0.001$) in patients ($n=17$) with Coats' disease. Also regression of telangiectasia and resolution of the exudate were observed. There were no severe ocular or systemic side-effects reported.¹² Intravitreal ranibizumab plus conventional amblyopia treatment resulted in transient restoration of the hemato-retinal barrier and suppression of neovascularization amongst patients with Coats' disease. This was propounded on the basis of resolution of iris neovascularization within 2 weeks of treatment with anatomical success.⁸ Giannakopoulos and colleagues reported the long-term results of Coats' disease after treatment with ranibizumab and demonstrated a visual acuity of 20/20 with no exudates or edema at 20 months of follow-up; the patient was stable at 6 years of follow-up.¹³

Biosimilar ranibizumab with laser photocoagulation significantly reduced the exudation, angioma-like lesions, CMT and edema in retina, and improved visual acuity in a patient with Coats' disease. The current case provides insights for biosimilar ranibizumab based options for Coats' disease, which needs to be further studied.

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6. Conflict of Interest

Dr. Shashikant Sharma is an employee of Intas Pharmaceuticals Ltd, Ahmedabad, India.

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