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Recurrence of Macular Hole Retinal Detachment after Intravitreal Ranibizumab Injection for the Treatment of Choroidal Neovascularization from the Remaining Macular Hole Edge

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Key Words

Macular hole · Retinal detachment · Intravitreal ranibizumab injection · Anti-VEGF · Lucentis · Choroidal neovascularization

Abstract

Purpose: To report a case who had recurrence of macular hole retinal detachment (MHRD) after intravitreal ranibizumab injection (IVR) for the treatment of choroidal neovascularization (CNV) that arose from the damaged retinal pigment epithelium of the remaining macular hole (MH) edge, which had been successfully treated by pars plana vitrectomy (PPV) 15 years previously.

Case Report: A 67-year-old man with previous PPV for MHRD secondary to high myopia in the right eye had been under observation for 15 years after surgery. The retina had been successfully attached, but the MH remained open. He had CNV which arose from the remaining MH edge. IVR was performed for the treatment of CNV. One month after the injection, CNV was contracted but recurrence of MHRD occurred. PPV with an additional internal limiting membrane peeling, removal of the CNV membrane and 20% SF6 gas tamponade was performed. One year after the last surgery, his right retina was attached and the MH was closed successfully.

Conclusion: We propose that patients who undergo IVR should be carefully maintained and followed up for possible complications including the recurrence of MHRD.

Introduction

Intravitreal anti-vascular endothelial growth factor (VEGF) agent therapy for age-related macular degeneration (AMD), proliferative diabetic retinopathy (PDR) and neovascular glaucoma (NVG) is widely used. On the other hand, retinal pigment epithelium (RPE) tears [1–3] and worsening of tractional retinal detachment (RD) [4–6] have been reported as a result of the contractile forces of regressing neovascular membranes after intravitreal anti-VEGF agent injection. However, the mechanism of construction of the fibrovascular membranes remains to be elucidated.

In this report, we present a case who had recurrence of macular hole retinal detachment (MHRD) after intravitreal ranibizumab injection (IVR) for the treatment of choroidal neovascularization (CNV) that arose from the damaged RPE of the remaining macular hole (MH) edge.

Case Report

A 67-year-old man with previous pars plana vitrectomy (PPV) for MHRD secondary to high myopia in the right eye (OD) had been under observation for 15 years after the surgery. Slit lamp examination revealed pseudophakia in OD and mature cataract in the left eye (OS). Funduscopy revealed attached retina and remaining MH in OD. The fundus condition in OS was invisible due to the mature cataract. Best corrected Landolt ring chart visual acuity (BCVA) was from 0.1 to 0.2 in OD and 30 cm finger motion in OS. The patient presented with sudden vision loss in OD. On presentation, his BCVA was 0.1. Funduscopy revealed a disciform RPE atrophy formation at the bottom of the remaining MH and CNV development around the edge of MH from the superior to temporal side (fig. 1a, d, f). His BVCA was not worsening but his subjective symptom was getting worse day by day. IVR was performed after 14 days from the onset for the treatment of CNV. One month after the injection, he presented with visual disturbance in OD. Funduscopy examination showed CNV contraction to the superior edge and recurrence of MHRD from the inferior edge of the remaining MH (fig. 1b, g). Emergency PPV with an additional internal limiting membrane peeling, removal of CNV and 20% SF₆ gas tamponade was performed on the same day. Extraction of CNV was performed under the patient's written informed consent, the approval of the institutional review board in Kobe Kaisei Hospital and the Tenets of the Declaration of Helsinki. One year after the last surgery, his right retina was reattached and MH was closed successfully (fig. 1c, e, h). His BVCA was improved at 0.3.

Comment

RPE tears and worsening of tractional RD after intravitreal anti-VEGF agent therapy have been reported [1–6]. Several reports hypothesized that intravitreal anti-VEGF agent therapy may cause severe contract force of the neovascular membrane followed by RPE tear in AMD [1–3] and tractional RD in PDR [4–6]. It is possible that the contract force of the regressing CNV might be increased after IVR, thereby causing RD from the remaining MH edge in our case.

In our case, we performed emergency PPV with CNV removal to fix the detached retina because the visual acuity in OS was bad and the MHRD in OD was expanding. For the treatment of CNV, it might be one of the options to continue IVR because IVR is thought less invasive compared with CNV removal during PPV. But if we perform additional IVR first, it is very important to make the CNV inactive before the following PPV because IVR itself may cause recurrence of MHRD due to the contract force of the CNV again. We must follow-up this patient for at least another month after the injection

to confirm the effect of IVR. Unfortunately, we had not a moment to lose in this case. We performed PPV with CNV extraction and achieved both retinal reattachment and the quieting effect of the CNV.

In summary, we propose that patients who undergo IVR should be carefully maintained and followed up for possible complications including the recurrence of MHRD.

Fig. 1. Before IVR, fundus photograph suggested CNV development from the remaining MH edge (**a**). FA demonstrated leakage from CNV (**d**). OCT depicted subretinal CNV (**f**). One week after IVR, fundus photograph (**b**) and OCT (**g**) showed contraction of CNV and recurrence of MHRD. One year after PPV, retina was reattached and MH was closed successfully (**c**). OCT showed MH closure and retina reattachment (**h**). CNV removal and chorioretinal atrophy after PPV was confirmed by FA (**e**).

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