Intravitreal ranibizumab improves macular sensitivity in patients with central retinal vein occlusion and macula edema

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Abstract

**Background:** Patients with central retinal vein occlusion (CRVO) and macular edema often are treated by intravitreal ranibizumab injection (IRI). The role of changes in macular sensitivity in the positive effects of IRI on visual functions is unclear. Therefore, we assessed the relationship between macular sensitivity and improvement of visual functions.

**Methods:** We included 15 eyes of 15 patients with treatment-naïve CRVO and followed patients for 6 months after pro re nata IRI. IRI was repeated if the central macular thickness was greater than or equal to 300 µm. Microperimetry-3 was used to measure macular sensitivity within the central 1-mm, 3-mm, and 6-mm fields before and monthly for 6 months after IRI.

**Results:** IRI significantly improved mean macular sensitivity over time within the central 1-mm, 3-mm, and 6-mm fields (all \( P < 0.001 \)). None of the fields showed significant differences in the change of mean macular sensitivity between patients with little improvement in best corrected visual acuity (BCVA; i.e., in patients with a change in logarithm of the minimum angle of resolution [logMAR] BCVA < 0.3) and those with marked improvement in BCVA (change in logMAR BCVA > 0.3). The mean macular sensitivity before IRI showed correlations with the improvement of macular sensitivity in every field.

**Conclusion:** These findings suggest that IRI improves macular sensitivity in patients with CRVO and macular edema independent of any improvement in BCVA and that macular sensitivity before treatment is associated with improvement of macular sensitivity after treatment.

**Keywords:** Central retinal vein occlusion, Macular edema, Ranibizumab, Macular sensitivity

Background

Central retinal vein occlusion (CRVO) and branch retinal vein occlusion (BRVO) are frequently found in people with type 2 diabetes, hypertension, hyperlipidemia, and arteriosclerosis, among other diseases. A common feature of CRVO and BRVO is macular edema, which causes visual impairment. Vascular occlusion research showed that vascular endothelial growth factor (VEGF) is associated with macular edema in CRVO [1], and anti-VEGF injection is used to treat macular edema in patients with CRVO and BRVO. Several randomized clinical trials have demonstrated better visual prognosis with repeated intravitreal anti-VEGF treatment [2–4].

Once macular edema has resolved, visual acuity returns to a relatively good level in most patients. However, visual functions can remain poor in some patients and affect reading speed [5] and contrast sensitivity [6], for

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example. Therefore, besides visual acuity, visual functions should also be assessed after treatment for macular edema, and studies have found that retinal sensitivity, in particular macular sensitivity, is suitable for evaluating visual function [7, 8].

Some research has been performed on macular sensitivity after anti-VEGF therapy for macular edema in CRVO and BRVO [9, 10]. However, it remains unclear whether macular sensitivity plays a role in improvement of visual functions after treatment of macular edema. Therefore, we evaluated macular sensitivity in patients with CRVO before and after intravitreal injection of the anti-VEGF ranibizumab and examined the relationship between macular sensitivity and improvement of visual functions.

**Methods**

**Subjects**

This retrospective study was performed in consecutive patients with treatment-naïve, non-ischemic CRVO and visual impairment due to macular edema who underwent intravitreal ranibizumab injection (IRI) at the Department of Ophthalmology, Hachioji Medical Center, Tokyo Medical University, Tokyo, Japan. Inclusion criteria were as follows: no history of intravitreal injections of anti-VEGF or other drugs; no previous treatment with intraocular corticosteroids, retinal photocoagulation, pars plana vitrectomy, or laser; age over 30 years; less than 3 months between symptom onset and initial examination; logarithm of the minimum angle of resolution (logMAR) best corrected visual acuity (BCVA) between 0.15 and 1.3; central macular thickness (CMT) greater than 300 μm as measured by optical coherence tomography (OCT); and follow-up of at least 6 months. Exclusion criteria were ischemic CRVO, defined as at least 10 disc areas of capillary non-perfusion [11]; macular ischemia; diabetic retinopathy due to type 2 diabetes; coexisting ocular disease (i.e., epiretinal membrane or glaucoma); and systemic disorders other than hypertension or hypercholesterolemia. After patients had provided informed consent, IRI was performed via the pars plana with a 30-gauge needle, 3.5 mm posterior to the limbus.

We included 15 eyes of 15 patients (9 men, 6 women) who had macular edema due to CRVO and had been followed up for 6 months after their initial treatment. The mean (SD) age of the sample was 58.5 (11.9) years. In all patients, CRVO had been diagnosed by fundus and spectral-domain OCT (SD-OCT) examinations.

![Fig. 1](image.png)

**Fig. 1** Measurement of retinal sensitivity by microperimetry. The figure shows a typical macular sensitivity map obtained with the MP-3 system. The mean macular sensitivity was calculated within the central 1-mm, 3-mm, and 6-mm fields. The MP-3 system tested the foveal region (central 1-mm field) at 5 points and the macular regions (central 3-mm and 6-mm fields) at 17 points and 29 points, respectively.
After diagnosis, all patients were evaluated monthly and treated with IRI (Lucentis; 0.5 mg in 0.05 ml; Genentech, Inc., South San Francisco, CA) pro re nata. IRI was repeated if the central macular thickness, including any serous retinal detachment [SRD], was greater than or equal to 300 µm. The number of additional IRI injections was recorded as a measure of recurrence.

Functional mapping by microperimetry

We assessed functional mapping by microperimetry at baseline and once a month until 6 months after IRI. Microperimetry combines digital fundus imaging with automated perimetry and was performed with the MP-3 microperimeter (Nidek, Gamagori, Japan). In all patients, the pupil diameter was larger than 4 mm, as is required for measurements with the MP-3. The MP-3 measurement was carried out with the 4–2 full threshold staircase strategy and the standard Goldmann III stimulus size on a background luminance of 31.4 asb. The maximum luminance of the MP-3 is 10,000 asb, which results in a stimulus dynamic range of 0 to 34 dB. The size of the fixation target was adjusted according to the patient’s visual acuity. One of the advantages of the MP-3 perimeter is that it automatically compensates for any refractive error in the patient’s eye [12]. Retinal sensitivity maps were obtained with the MP-3 program, which examines the central 20 degrees of the macula and uses a different number of stimulus locations depending on the area being investigated. Macular sensitivity within the central 1 mm was defined as the mean retinal sensitivity of 5 stimulus locations; macular sensitivity within the central 3 mm, as the mean retinal sensitivity of 17 stimulus locations; and macular sensitivity within the central 6 mm, as the mean retinal sensitivity of 29 stimulus locations (Fig. 1). The central 1-mm field corresponded to the foveal region, and the central 3-mm and 6-mm fields, to the macular region.

Clinical parameters

All patients underwent a comprehensive ocular examination before the start of the study (baseline) and at every follow-up visit, i.e. at 1, 2, 3, 4, 5, and 6 months after IRI. BCVA was measured as the decimal visual acuity by using the Landolt Chart and converted to the

Table 1  Clinical characteristics and macular sensitivity before and after intravitreal ranibizumab injection in patients with central retinal vein occlusion and macular edema

|                              | Before treatment | 6 months after treatment | P value |
|------------------------------|-----------------|--------------------------|---------|
| Age (years)                  | 58.5 ± 11.9a    | 52.5 ± 12.3a             |         |
| Gender (female/male)         | 6/9             | 6/9                      |         |
| Duration of macular edema (days) | 34.5 ± 23.4a   | 25.5 ± 12.3a             |         |
| Hypertension                 | 11              | 10                       |         |
| Systolic Blood pressure (mmHg) | 138 ± 11.3a    | 138 ± 11.3a              |         |
| Diastolic Blood pressure (mmHg) | 87.1 ± 10.3a   | 87.1 ± 10.3a             |         |
| Hyperlipidemia               | 7               | 7                        |         |
| Number of injections         | 2.7 ± 1.4a      | 2.7 ± 1.4a               |         |
| Injection type (1–2 injections/3 or more injections) | 7/8            | 6/8                      |         |
| BCVA (logMAR)                | 0.52 ± 0.39a    | 0.52 ± 0.39a             | 0.018   |
| Central macular thickness (µm) | 721 ± 207a      | 721 ± 207a               |         |
| Macular sensitivity within 1 mm (dB) | 12.8 ± 5.68a   | 12.8 ± 5.68a             | 0.007   |
| Macular sensitivity within 3 mm (dB) | 16.2 ± 5.39a   | 16.2 ± 5.39a             | 0.002   |
| Macular sensitivity within 6 mm (dB) | 17.5 ± 5.16a   | 17.5 ± 5.16a             | <0.001  |
| Presence of SRD              | 11              | 4                        | 0.011   |

BCVA Best-corrected visual acuity, log MAR Logarithm of the minimum angle of resolution, *Mean ± standard deviation (SD)
Fig. 2 (See legend on previous page.)
logMAR. Fluorescein angiography (Heidelberg Engineering, Heidelberg, Germany) was performed at the baseline visit to determine the area of capillary non-perfusion. At the follow-up examinations, visual acuity was measured with the logMAR chart (5 m; NEITZ LVC-10, Tokyo, Japan) and CMT was measured with Spectralis OCT (Heidelberg Engineering, Heidelberg, Germany). The Spectralis OCT mapping images were generated with the currently available Spectralis software. CMT was defined as the thickness of the central 1-mm field on the OCT mapping image and as the distance between the inner limiting membrane and the retinal pigment epithelium (including any SRD) and was automatically measured by the computer software. We used the images obtained with OCT to quantify CMT and evaluated the presence of SRD.

Changes in clinical parameters
To assess improvement of vision, we calculated the change in BCVA over 6 months by subtracting the value at the 6-month follow-up from the value at baseline (i.e., before IRI). To assess macular sensitivity, we calculated the percentage changes in macular sensitivity (%ΔMS), as follows: %ΔMS = (MS<sub>pre</sub>−MS<sub>6 months</sub>) / MS<sub>pre</sub> × 100 = (1 − MS<sub>6 months</sub> / MS<sub>pre</sub>) × 100, where MS<sub>pre</sub> and MS<sub>6 months</sub> represent the sensitivity values at baseline and 6 months after IRI, respectively.

Statistical analysis
All analyses were performed with SAS System 9.4 software (SAS Institute Inc., Cary, North Carolina, USA). Results are presented as the mean ± SD or as the frequency. We compared unpaired continuous variables with an unpaired Student’s t test and paired continuous variables with a paired Student’s t test. One-way or two-way repeated measures analysis was performed with the use of mixed model for evaluating the macular sensitivity. Two-tailed P values of less than 0.05 were considered to indicate statistical significance.

Results
Clinical characteristics
Table 1 shows the clinical characteristics, visual functions, and OCT parameters before and 6 months after IRI. The mean number of injections in 6 months was 2.7 ± 1.4; 7 patients had 1 or 2 injections, and 8 had 3 or more injections. Eleven patients had SRD before treatment, but only 4 of these patients had it 6 months after treatment. BCVA and CMT improved significantly after IRI (P = 0.018 and P = 0.001, respectively). The presence of SRD before IRI was not significantly correlated with the number of injections or the type of injection regimen (1–2 injections or 3 or more injections), but the presence of SRD 6 months after IRI was significantly correlated with both the number of injections and the type of injection regimen (P = 0.026 and P = 0.029, respectively).

Figure 2 shows the monthly change in macular sensitivity from baseline to 6 months after IRI. Over time, mean macular sensitivity significantly improved within the central 1-mm, 3-mm, and 6-mm fields (all P < 0.001; Figs. 2A-C). We found no significant differences in the changes of mean macular sensitivity within the central 1-mm, 3-mm, and 6-mm fields between patients with little improvement in BCVA (change in logMAR BCVA < 0.3) and those with marked improvement in BCVA (change in logMAR BCVA > 0.3) (P = 0.551; and 6-mm field, P = 0.746; Figs. 3A-C).

Correlations of visual functions with BCVA, macular sensitivity, and CMT
Table 2 shows the correlations of BCVA, macular sensitivity, and CMT before and after IRI with the improvement of visual functions at 6 months after IRI. BCVA at baseline was significantly correlated with the improvement of BCVA (P < 0.05); mean macular sensitivity in the central 1-mm at baseline was significantly correlated with the improvement of macular sensitivity in the central 1-mm, 3-mm, and 6-mm fields (P < 0.001, and P < 0.001, respectively); mean macular sensitivity in the central 3-mm at baseline was significantly correlated with the improvement of macular sensitivity in the 1-mm, 3-mm, and 6-mm fields (P < 0.001, and P < 0.001, respectively); mean macular sensitivity in the central 6-mm at baseline was significantly correlated with the improvement of macular sensitivity in the 1-mm, 3-mm, and 6-mm fields (P < 0.001, and P < 0.001, respectively); and CMT at baseline was significantly correlated with the improvement of macular sensitivity in the 3-mm and 6-mm fields (both P < 0.05, respectively).
Fig. 3 (See legend on previous page.)
In addition, CMT at 6 months after IRI showed a significant correlation with the improvement of BVCA (P<0.05). The number of injections (1 or 2 injections or 3 or more injections) and type of injections also were not significantly correlated with any changes in visual functions.

**Discussion**

To evaluate whether macular sensitivity plays a role in changes in visual functions after treatment of macular edema, this study evaluated macular sensitivity in patients with CRVO before and after IRI. Regarding visual functions, we found that not only visual acuity but also macular sensitivity within the central 1-mm, 3-mm, and 6-mm fields improved significantly after IRI. This finding suggests that, after anti-VEGF treatment, not only the fovea but also the larger macular area should be evaluated because treatment improves edema of the entire macula in CRVO. Furthermore, it indicates that—when assessing the visual prognosis of CRVO patients with macular edema—it may be relevant to evaluate not only visual acuity (an indicator of foveal function) but also macular sensitivity (an indicator of retinal sensitivity in the larger macular area). Thus, microperimetry with the MP-3 may demonstrate the effectiveness of anti-VEGF treatment for macular edema associated with CRVO more accurately than measurements of visual acuity alone. This hypothesis is supported by our earlier finding that retinal thickness and retinal volume are related to visual acuity and retinal sensitivity in patients with CRVO and macular edema [8].

Interestingly, we found no significant differences in the changes of mean macular sensitivity within the central 1-mm, 3-mm, and 6-mm fields between patients with little improvement in BCVA and those with marked improvement. This result suggests that improvement in macular sensitivity is independent of improvement in visual acuity. Some studies indicated that macular sensitivity in patients with age-related macular degeneration and retinitis pigmentosa is associated with vision-related quality of life [13, 14]. Our result that macular sensitivity improved even in patients with little improvement in visual acuity may help to explain these earlier findings. Furthermore, we found no significant correlations between the number of injections (as a measure of recurrence) and the improvement of BCVA and macular sensitivity.

We also found that BCVA before IRI was significantly correlated with the improvement of BCVA, that macular sensitivity before IRI was significantly correlated with the improvement of macular sensitivity in all 3 central fields and that CMT before IRI was significantly correlated with the improvement of macular sensitivity in

| Table 2 | Correlations between improvement of visual functions after treatment and ocular parameters in patients with central retinal vein occlusion |
| Parameters | Improvement of BCVA | Improvement of macular sensitivity within 1 mm | Improvement of macular sensitivity within 3 mm | Improvement of macular sensitivity within 6 mm |
|------------|----------------------|-----------------------------------------------|-----------------------------------------------|-----------------------------------------------|
|            | P value (correlation coefficient) | P value (correlation coefficient) | P value (correlation coefficient) | P value (correlation coefficient) |
| **Before IRI** | | | | |
| Duration of macular edema, days | 0.689 (−0.11) | 0.502 (−0.19) | 0.355 (−0.26) | 0.301 (−0.29) |
| BCVA (logMAR) | 0.002 (0.72) | 0.942 (0.02) | 0.136 (0.40) | 0.100 (0.44) |
| Macular sensitivity within 1 mm, dB | 0.344 (−0.26) | 0.005 (−0.68) | <0.001 (−0.80) | <0.001 (−0.82) |
| Macular sensitivity within 3 mm, dB | 0.159 (−0.38) | 0.011 (−0.64) | <0.001 (−0.84) | <0.001 (−0.87) |
| Macular sensitivity within 6 mm, dB | 0.221 (−0.34) | 0.022 (−0.59) | <0.001 (−0.81) | <0.001 (−0.85) |
| Central macular thickness, μm | 0.093 (0.45) | 0.059 (0.50) | 0.005 (0.68) | 0.007 (0.66) |
| Presence of SRD | 0.434 | 0.700 | 0.999 | 0.952 |
| **6 months after IRI** | | | | |
| Central macular thickness, μm | 0.036 (−0.55) | 0.139 (−0.40) | 0.172 (−0.37) | 0.233 (−0.33) |
| Presence of SRD | 0.728 | 0.206 | 0.208 | 0.255 |
| Number of injections | 0.075 (−0.47) | 0.457 (−0.21) | 0.407 (−0.23) | 0.497 (−0.19) |
| Injection type | 0.071 | 0.905 | 0.597 | 0.667 |

**BCVA** Best-corrected visual acuity, **IRI** Intravitreal ranibizumab injection, **log MAR** Logarithm of the minimum angle of resolution, **SRD** Serous retinal detachment
the 3-mm and 6-mm fields. These findings suggest that treatment improved visual functions in patients with poor baseline values for BCVA, CMT, and/or macular sensitivity. On the other hand, CMT at 6 months after IRI was significantly correlated with improvement of BCVA but not with the improvement of macular sensitivity, suggesting that improvement of CMT after IRI does not affect improvement of macular sensitivity but does affect improvement of BCVA. A recent study found that resolution of subretinal fluid 6 months after anti-VEGF treatment was associated with improved retinal sensitivity [15], indicating that once macular edema improves, macular sensitivity may be less likely to worsen.

This study was limited by the small sample size, which meant that the repeated measures analyses were underpowered. Consequently, the clinical relevance of the study findings should be interpreted with caution, and large, prospective studies are required on macular sensitivity after anti-VEGF therapy in patients with macular edema due to CRVO.

**Conclusions**

This study demonstrated that, over 6 months, IRI significantly improves mean macular sensitivity within the central 1-mm, 3-mm, and 6-mm fields in patients with CRVO and macular edema and that changes in mean macular sensitivity show no significant differences between patients with little improvement in BCVA and those with marked improvement. Furthermore, mean macular sensitivity before IRI shows significant correlations with improvement of macular sensitivity in the 3 fields mentioned above. These findings suggest that IRI improves macular sensitivity independent of any improvement in BCVA and that macular sensitivity before treatment is associated with improvement of macular sensitivity. Because improvements in macular sensitivity may increase quality of life even in patients who show no improvement in BCVA, these findings are of great relevance in clinical practice and warrant further evaluation in prospective studies in larger samples.

**Abbreviations**

BCVA: Best-corrected visual acuity; BRVO: Branch retinal vein occlusion; CMT: Central macular thickness; CRVO: Central retinal vein occlusion; IRI: Intravitreal ranibizumab injection; LogMAR: Logarithm of the minimum angle of resolution; OCT: Optical coherence tomography; SD-OCT: Spectral-domain OCT; SRD: Serous retinal detachment; VEGF: Vascular endothelial growth factor.

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**Authors’ contributions**

H.N., and M.S. were involved in the design and conduct of the study. Collection and management of the data were done by T.O., A.N., M.A., A.T., and H.N., while analysis and interpretation of the data were performed by K.Y., and M.S.

**Preparation of the first draft of the manuscript was done by T.O., and H.N., and review and approval of the manuscript was performed by H.G., and M.S. All authors have read and approved the final manuscript.**

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**Availability of data and materials**

The data that support the findings of this study are not publicly available because they contain information that could compromise the privacy of research participants and because ethics committee approval for release of the data was not obtained.

**Declarations**

**Ethical approval and consent to participation**

This study was conducted at the Department of Ophthalmology of Tokyo Medical University Hachioji Medical Center and approval was obtained from Institutional Review Board of Tokyo Medical University Hachioji Medical Center (IRB No. H-132). The procedures of the study conformed to the tenets of the Declaration of Helsinki and all patients gave written informed consent before enrollment.

**Consent to publication**

Written informed consent for publication of clinical images was obtained from the individual patients. The details of consent to publication were included in the consent form for participation in this study.

**Competing interests**

The authors declare that they have no competing interests.

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**References**

1. Noma H, Funatsu H, Mimura T, Harino S, Hori S. Vitreous levels of interleukin-6 and vascular endothelial growth factor in macular edema with central retinal vein occlusion. Ophthalmology. 2009;116:87–93.

2. Heier JS, Campochiaro PA, Yau L, Li Z, Saroj N, Rubio RG, Lai P. Ranibizumab for macular edema due to retinal vein occlusions: long-term follow-up in the HORIZON trial. Ophthalmology. 2012;119:802–9.

3. Heier JS, Clark WL, Boyer DS, Brown DM, Vitti R, Berliner AJ, Kazmi H, Ma Y, Steemer B, Zeitz O, et al. Intravitreal aflibercept injection for macular edema due to central retinal vein occlusion: two-year results from the COPERNICUS study. Ophthalmology. 2014;121:1414–1420. e1411.

4. Larsen M, Waldstein SM, Priglinger S, Hykin P, Barnes E, Gekkieva M, Das Gupta A, Wenzel A, Monés J. Sustained Benefits from Ranibizumab for Central Retinal Vein Occlusion with Macular Edema. 24-Month Results of the CRYSTAL Study. Ophthalmol Retina. 2018;2:134–42.

5. Suher LJ, Bressler NM, Varma R, Lee P, Dolan CM, Ward J, Colman S, Rubio RG. Reading speed improvements in retinal vein occlusion after ranibizumab treatment. JAMA Ophthalmol. 2013;131:851–6.

6. Murakami T, Okamoto F, Sugiuira Y, Monkawa S, Okamoto Y, Hiraoka T, Oshika T. Contrast sensitivity and quality of life following intravitreal ranibizumab injection for central retinal vein occlusion. Br J Ophthalmol. 2021;27:2021–319863.

7. Noma H, Mimura T. Macular sensitivity and morphology after intravitreal injection of triamcinolone acetonide for macular edema secondary to central retinal vein occlusion. Clin Ophthalmol. 2012;6:1901–6.

8. Noma H, Mimura T, Shimada K. Retinal function and morphology in central retinal vein occlusion with macular edema. Curr Eye Res. 2013;38:143–9.
9. Yamaike N, Tsujikawa A, Sakamoto A, Ota M, Kotera Y, Miyamoto K, Kita M, Yoshimura N. Retinal sensitivity after intravitreal injection of bevacizumab for the treatment of macular edema secondary to retinal vein occlusion. Retina. 2009;29:757–67.

10. Papadia M, Misteli M, Jeannin B, Herbelot CP. The influence of anti-VEGF therapy on present day management of macular edema due to BRVO and CRVO: a longitudinal analysis on visual function, injection time interval and complications. Int Ophthalmol. 2014;34:1193–201.

11. Baseline and early natural history report: The Central Vein Occlusion Study. Arch Ophthalmol. 1993;111:1087–95.

12. Balasubramanian S, Uji A, Lei J, Velaga S, Nittal M, Sadda S. Interdevice comparison of retinal sensitivity assessments in a healthy population: the CenterVue MAIA and the Nidek MP-3 microperimeters. Br J Ophthalmol. 2018;102:109–13.

13. Sugawara T, Sato E, Baba T, Hagiwara A, Tawada A, Yamamoto S. Relationship between vision-related quality of life and microperimetry-determined macular sensitivity in patients with retinitis pigmentosa. Jpn J Ophthalmol. 2011;55:643–6.

14. Forshaw TRJ, Parpounas AK, Sørensen TL. Correlation of macular sensitivity measures and visual acuity to vision-related quality of life in patients with age-related macular degeneration. BMC Ophthalmol. 2021;21:149.

15. Khayat M, Perais J, Wright DM, Williams M, Lois N. Anatomic-Functional Correlates in Lesions of Retinal Vein Occlusion. Invest Ophthalmol Vis Sci. 2021;62:10.

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