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***Retrospective Study***

**Three-year follow-up of coats disease treated with conbercept and 532-nm laser photocoagulation**

Jiang L *et al*. Conbercept and laser for coats disease

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**Abstract**

BACKGROUND

Coats disease is an idiopathic exudative outer retinopathy caused by abnormal retinal vascular development.

AIM

To evaluate the long-term outcomes of intravitreal conbercept injection with laser photocoagulation as a treatment for coats disease in adults.

METHODS

This retrospective case series study included patients diagnosed with Coats disease and treated with intravitreal conbercept injection and 532-nm laser photocoagulation at the Ophthalmology Department of Shenzhen People’s Hospital between January 2016 and January 2017. Best-corrected visual acuity (BCVA) measurements, noncontact tonometry, ophthalmoscopy, fundus photography, fundus fluorescein angiography and optical coherence tomography were performed before treatment and at 1 wk, 1 mo, 3 mo, 6 mo, 9 mo, 12 mo, 24 mo and 36 mo after therapy. Best-corrected visual acuity was measured using the early treatment of diabetic retinopathy study chart.

RESULTS

The study included eight eyes of 8 patients (7 men) aged 36.10 ± 6.65 years. The average BCVA of the affected eye before treatment was 51.17 ± 15.15 letters (range, 28–70 letters), and the average central macular thickness was 303.30 ± 107.87 µm (range, 221–673 µm). Four eyes were injected once, three were injected twice, and one was injected three times. Average follow-up duration was 37.33 ± 2.26 mo. Average BCVA of the affected eye was 51.17 ± 15.15 letters before treatment and was increased by 13.50 ± 3.20, 16.25 ± 7.73, 18.25 ± 8.96, 18.03 ± 5.27, 18.63 ± 3.35, 19.75 ± 6.96, 18.05 ± 5.36 and 17.88 ± 3.45 letters at 1 wk, 1 mo, 3 mo, 6 mo, 9 mo, 12 mo, 24 mo and 36 mo after treatment, respectively (*P* < 0.01). The patients showed varying degrees of subretinal fluid resorption after treatment. None of the patients had serious complications such as increased intraocular pressure, development/progression of cataracts, endophthalmitis or retinal detachment.

CONCLUSION

Intravitreal injection of conbercept combined with 532-nm laser photocoagulation may be a feasible treatment for coats disease in adult patients.

**Key Words:** Coats disease; Adults; Retinal telangiectasis; Conbercept; Laser therapy; Retrospective study

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**Core Tip:** Coats disease is an idiopathic exudative outer retinopathy caused by abnormal retinal vascular development. Intravitreal injection of conbercept combined with 532-nm laser photocoagulation may be a feasible treatment for coats disease in adult patients.

**INTRODUCTION**

Coats disease is an idiopathic exudative outer retinopathy caused by abnormal retinal vascular development[[1](#_ENREF_1),[2](#_ENREF_2)]. The main clinical features of Coats disease are dilated retinal capillaries, miliary or small aneurysms and retinal and subretinal exudation[[1](#_ENREF_1),[2](#_ENREF_2)]. The estimated incidence of Coats disease in the population is 0.09 per 100000 persons, and males are more commonly affected than females[[3](#_ENREF_3)]. The disorder usually affects adolescents with a mean age at presentation of around 12 years[[3](#_ENREF_3)]. However, a recent study reported that 7% of cases occur in adults (mainly males) with an onset age of 43–50 years, and the incidence of monocularity was 93.3%-100.0%[[4](#_ENREF_4)]. Standard treatments for Coats disease include laser therapy, cryotherapy and surgical techniques such as subretinal fluid drainage and vitrectomy[[5](#_ENREF_5)].

Recent studies have shown that patients with Coats disease have elevated levels of vascular endothelial growth factor (VEGF) in the aqueous humor, vitreous and subretinal fluid, suggesting that a disorder of VEGF-mediated vascular regeneration may be involved in the pathogenesis of Coats disease[[6-9](#_ENREF_6)]. It has been suggested that ischemia around a region of unperfused retinal capillaries may lead to VEGF upregulation that increases the permeability of dilated capillaries and causes exudation of lipoproteins below the retina, resulting in edema and exudative retinal detachment[[6-9](#_ENREF_6)]. There has been increasing interest in the use of anti-VEGF agents as an adjunct treatment for Coats disease[[1](#_ENREF_1),[2](#_ENREF_2),[10](#_ENREF_10)]. Several clinical studies have reported that the intravitreal injection of a VEGF-binding antibody (such as bevacizumab or ranibizumab) or aptamer (pegaptanib) exerts beneficial effects in patients with Coats disease, including a reduction in exudation and edema and an improvement in visual acuity[[11-15](#_ENREF_11)].

Conbercept is a recombinant fusion protein comprising extracellular domain 2 of VEGF receptor-1, extracellular domains 3 and 4 of VEGF receptor-2 and the Fc region of human IgG1[[16](#_ENREF_16)]. This fusion protein binds VEGF with high affinity, has a long half-life in the vitreousand blocks all isoforms of VEGF-A, VEGF-B, VEGF-C and placental growth factor[[16](#_ENREF_16)]. Conbercept has been shown to be an effective treatment for retinal diseases such as diabetic retinopathy-induced macular edema[[17](#_ENREF_17),[18](#_ENREF_18)] and age-related macular degeneration[[19-22](#_ENREF_19)]. However, to the best of our knowledge, only one previous study has investigated the use of conbercept as an adjunct treatment to laser therapy in the management of Coats disease[[23](#_ENREF_23)].

The aim of this retrospective case series study was to summarize the clinical outcomes of adult patients who were diagnosed with Coats disease at our ophthalmology department and treated with intravitreal injection of conbercept in combination with laser photocoagulation.

**MATERIALS AND METHODS**

***Patients***

This retrospective case series study included patients diagnosed with Coats disease and treated with intravitreal injection of conbercept and laser photocoagulation at the Department of Ophthalmology of Shenzhen People’s Hospital between January 2016 and January 2017. The inclusion criteria were: (1) age > 18 years; (2) newly diagnosed with Coats disease; and (3) treated with intravitreal injection of conbercept and laser photocoagulation. Coats disease was diagnosed according to the results of the fundus examination (the presence of abnormal dilated retinal blood vessels and retinal interstitial or subretinal exudation with or without retinal detachment), fluorescein fundus angiography (FFA; typical dilatation of retinal capillaries and small blood vessels) and optical coherence tomography (OCT; characteristic features such as retinal thickening, exudation and subretinal fluid)[[24](#_ENREF_24)]. The exclusion criteria were: (1) other eye diseases such as age-related macular degeneration; and (2) complications such as severe refractive interstitial opacity or vitreous hemorrhage that impaired observation of the fundus.

The severity of Coats disease in each patient was classified as stage 1 (retinal capillary dilation only), stage 2A (retinal capillary dilation and subretinal exudation outside the central fovea), stage 2B (retinal capillary dilation and exudation inside the central fovea), stage 3A (incomplete retinal detachment), stage 3B (complete retinal detachment), stage 4 (total retinal detachment with secondary glaucoma) or stage 5 (the terminal stage of eyeball atrophy)[[24](#_ENREF_24)].

The study was approved by the ethics committee of Shenzhen People’s Hospital, No. LL-KY-2020238.

***Treatment***

Written consent was obtained from each patient after they and their families had been fully informed about the treatment strategy (intravitreal injection of conbercept combined with laser photocoagulation) and potential adverse effects. A standard method was used for intravitreal injection of conbercept into the affected eye. Before treatment, 0.5% levofloxacin eye drops were administered as a local anti-inflammatory therapy. The same doctor, who had a third-level surgery qualification, performed all the procedures. The patient was placed in a supine position in the operating room. Tropicamide was used for mydriasis, and oxybuprocaine hydrochloride was applied for surface anesthesia. Disinfection and surgical draping were performed routinely according to the requirements of internal eye surgery. A protective film was pasted, an eyelid opener was used to open the eyelids, and the conjunctival sac was washed with diluted povidone iodine. A 30-gauge needle was inserted into the flat part of the ciliary body at a distance of 3.5–4 mm from the limbus. The intravitreal cavity was injected with 0.5 mg conbercept (trade name, Langmu; Chengdu Kanghong Biological Co., Ltd, China) administered as 0.05 mL of a 10 mg/mL solution. Gentle pressure was applied with a cotton swab for 10 s to prevent reflux. At the end of the procedure, light sensation was checked, tobramycin-dexamethasone eye ointment was applied, and the eye was covered with a blindfold. Postoperatively, 0.5% levofloxacin eye drops were given for 3 d as prophylaxis against bacterial infection. Subsequently, the anterior segment of the eye was examined for signs of an inflammatory reaction, intraocular pressure was measured, and fundus examination and FFA were performed. Based on the FFA results, 532-nm laser photocoagulation was carried out after conbercept injection to block retinal aneurysms, abnormally dilated capillaries or capillaries without perfusion. The laser equipment used in this study was Lumenis Vision One. Retinal laser photocoagulation was operated routinely with a 532-nm light. The size of the spot was 200 µm, the energy was 150 MW, and the duration was 200 ms.

***Follow-up***

Periodic reviews were conducted postoperatively at 1 wk, 1 mo, 3 mo, 6 mo, 9 mo, 12 mo and annually thereafter. Changes in the condition of the eye were used to determine whether the treatment needed to be repeated.

***Data collection***

Clinical data were extracted from the medical records. Best-corrected visual acuity (BCVA) was measured using the early treatment of diabetic retinopathy study chart. Intraocular pressure was measured with a noncontact tonometer (NT-530; Nidek Co., Ltd, Japan). Fundus color photography and FFA examinations were performed using a TRC-50DX retinal camera (Topcon Medical Systems Inc., Japan). Spectral-domain OCT was carried out with a Spectralis system (Heidelberg Engineering GmbH, Germany). The central macular thickness was measured in high-definition scanning mode. The change in vision at each follow-up was evaluated as follows: an increase of ≥ 15 in the number of letters in the chart was considered a substantial improvement in vision; an increase of 5–14 was considered an improvement in vision; an increase or decrease of < 5 was considered stable vision; a decrease of 5–14 was considered a worsening of vision; and a decrease of ≥ 15 was considered a substantial worsening of vision[[25](#_ENREF_25)].

***Statistical analysis***

SPSS 23.0 (IBM Corp., Armonk, NY, United States) was used for the statistical analyses. Measurement data are expressed as mean ± standard deviation values. BCVA was compared between time points using repeated-measures analysis of variance and Fisher’s LSD post-hoc test. *P* < 0.05 was considered statistically significant.

**RESULTS**

***Clinical characteristics of the study participants***

The study included eight eyes of 8 patients (7 men, 87.5%) aged 36.10 ± 6.65 years (range, 27-51 years). The clinical characteristics of the study participants are summarized in Table 1. The average BCVA of the affected eye before treatment was 51.17 ± 15.15 letters (range, 28-70 letters), and the average central macular thickness was 303.30 ± 107.87 µm (range, 221–673 µm). Intraocular pressure was normal in all affected eyes. The average number of injections in the affected eyes was 1.37 ± 0.92: four eyes were injected once, three eyes were injected twice, and one eye was injected thrice. The average follow-up duration was 37.33 ± 2.26 mo (range, 34–39 mo).

***Comparison of BCVA before and after treatment***

There was a substantial improvement in vision in four eyes (50.0%), an improvement in vision in three eyes (37.5%) and stable vision in one eye (12.5%). Table 2 presents the BCVA before treatment and at 1 wk, 1 mo, 3 mo, 6 mo, 9 mo, 1 year, 2 years and 3 years after treatment. There was a significant upward trend in the BCVA (*P*-trendlinear < 0.001). Compared to the pretreatment value, the average BCVA at 1 wk, 1 mo, 3 mo, 6 mo, 9 mo, 1 year, 2 years and 3 years after treatment increased by 13.50 ± 3.20, 16.25 ± 7.73, 18.25 ± 8.96, 18.03 ± 5.27, 18.63 ± 3.35, 19.75 ± 6.96, 18.05 ± 5.36 and 17.88 ± 3.45 letters, respectively (*P* < 0.01 for all pairwise comparisons; Figure 1). The number of letters by which the average BCVA increased at 1 mo after the treatment showed no significant difference compared to that at 3 mo, 12 mo and 24 mo post-treatment and at the final follow-up (*P* > 0.05), suggesting that 1 mo after treatment was the critical period for recovery of visual acuity.

***Comparison of subretinal edema before and after treatment***

Only one of the patients in this study had macular lesions and developed macular edema. The central macular thickness in this patient was 673 µm before treatment but decreased to 587 µm, 372 µm, 322 µm, 267 µm, 221 µm and 208 µm at 1 wk, 1 mo, 3 mo, 1 year, 2 years and 3 years after treatment, respectively. Macular edema was substantially improved at the final follow-up. In the other patients, treatment led to varying degrees of subretinal fluid resorption and a decrease in retinal thickness (a representative case is shown in Figure 2).

***Complications***

No patients had serious complications such as increased intraocular pressure, development/progression of cataracts, endophthalmitis or retinal detachment.

**DISCUSSION**

An important finding of this study was that adult patients with Coats disease exhibited significant improvements in BCVA that were maintained for at least 3 years after treatment with intravitreal injection of conbercept and laser photocoagulation. Furthermore, the treatment resulted in the resorption of subretinal fluid. Notably, no patients had serious complications such as increased intraocular pressure, development/progression of cataracts, endophthalmitis or retinal detachment. Our findings suggest that intravitreal injection of conbercept combined with 532-nm laser photocoagulation may be a feasible treatment for Coats disease in adult patients.

Coats disease is more common in men and is usually monocular. The average age of onset is 8-16 years[[26](#_ENREF_26)]; hence, Coats disease is rarely diagnosed in adults. It can be challenging to distinguish Coats disease from other ocular diseases such as exudative age-related macular degeneration, diabetic retinopathy and retinal vasculitis. We observed that Coats disease in adulthood has similar fundal manifestations to Coats disease of childhood onset, although FFA showed that the abnormal expansion of the retinal capillaries was limited and mostly confined to the temporal retina. Only one patient in this study had an affected macula, which resulted in local superficial retinal detachment. Our observations are similar to those reported previously[[27](#_ENREF_27)], suggesting that adult patients with Coats disease have a limited disease type with slow progression, mild macular damage and a good visual prognosis. Otani *et al*[[28](#_ENREF_28)] followed two adult patients with Coats disease for more than 10 years: 1 patient showed a reduction in subretinal exudation within 1 year after retinal laser photocoagulation and an improvement in vision that was maintained for 11 years; the other patient exhibited increased exudation at 1 mo after photocoagulation and retinal detachment at 2 mo, but these subsequently resolved to result in visual improvement that was maintained for 10 years.

There are various management strategies for Coats disease, and the decision whether to use a single therapy or combination of treatments is made according to the disease stage. The pathogenesis of Coats disease involves abnormal dilatation and enhanced permeability of retinal vessels leading to vascular leakage. It is thought that VEGF may play an important role in these pathogenetic mechanisms, and several studies have reported that VEGF concentration is significantly increased in the affected eyes of patients with Coats disease[[6-9](#_ENREF_6)]. Intravitreal injection of anti-VEGF drugs can reduce VEGF concentration, stabilize abnormal blood vessels and promote the resorption of subretinal effusion and exudation. Numerous clinical investigations have found that the intravitreal injection of a VEGF-binding agent (such as bevacizumab, ranibizumab or pegaptanib) can reduce exudation and edema and improve visual acuity in patients with Coats disease[[29-31](#_ENREF_29)]. However, there is no current consensus on the choice, dosage and timing of anti-VEGF drug in the treatment of Coats disease, and research in this area is still in the exploratory stage.

Only one previous study has described the use of conbercept in the management of Coats disease[[23](#_ENREF_23)]: The combination of intravitreal injection of conbercept with laser therapy resulted in a significant improvement in BCVA after 2 years with no serious adverse effects. The above observations are consistent with our findings, which showed a reduction in subretinal fluid and an improvement in BCVA that was sustained for 3 years without any major adverse effects.

This study has some limitations. First, the results may be prone to information bias or selection bias due to the retrospective study design. Second, the generalizability of the findings is not known because this was a single center study with a small sample size. Third, a comparator group was not included. Prospective randomized clinical trials are needed to confirm our findings and establish whether conbercept is a useful adjunct therapy for Coats disease.

**CONCLUSION**

In conclusion, intravitreal injection of conbercept combined with 532-nm laser photocoagulation may be a feasible treatment for Coats disease in adult patients.

**ARTICLE HIGHLIGHTS**

***Research background***

Coats disease is an idiopathic exudative outer retinopathy caused by abnormal retinal vascular development

***Research motivation***

The disorder usually affects adolescents. However, a recent study reported that 7% of cases occur in adults.

***Research objectives***

We investigated the long-term outcomes of intravitreal conbercept injection with laser photocoagulation to treat Coats disease in adults.

***Research methods***

This is a retrospective case series on 8 adult patients (eight eyes) diagnosed with Coats disease. Patients were treated with intravitreal conbercept injection and 532-nm laser photocoagulation.

***Research results***

There was a significant upward trend in the best-corrected visual acuity after treatment, which was maintained for at least 3 years.

***Research conclusions***

Intravitreal conbercept injection plus 532-nm laser photocoagulation may be a feasible treatment for Coats disease in adult patients.

***Research perspectives***

Prospective randomized clinical trials are needed to confirm our findings and establish whether conbercept is a useful adjunct therapy for Coats disease.

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**Footnotes**

**Institutional review board statement:** The study was approved by the ethics committee of Shenzhen People’s Hospital (No. LL-KY-2020238).

**Informed consent statement:** Patients were not required to give informed consent to the study because the analysis used anonymous clinical data that were obtained after each patient agreed to treatment by written consent.

**Conflict-of-interest statement:** The authors declare that they have no competing interests.

**Data sharing statement:** Not applicable.

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Grade A (Excellent): 0

Grade B (Very good): B

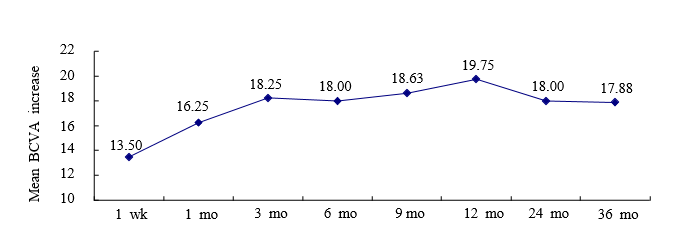
Grade C (Good): C

Grade D (Fair): 0

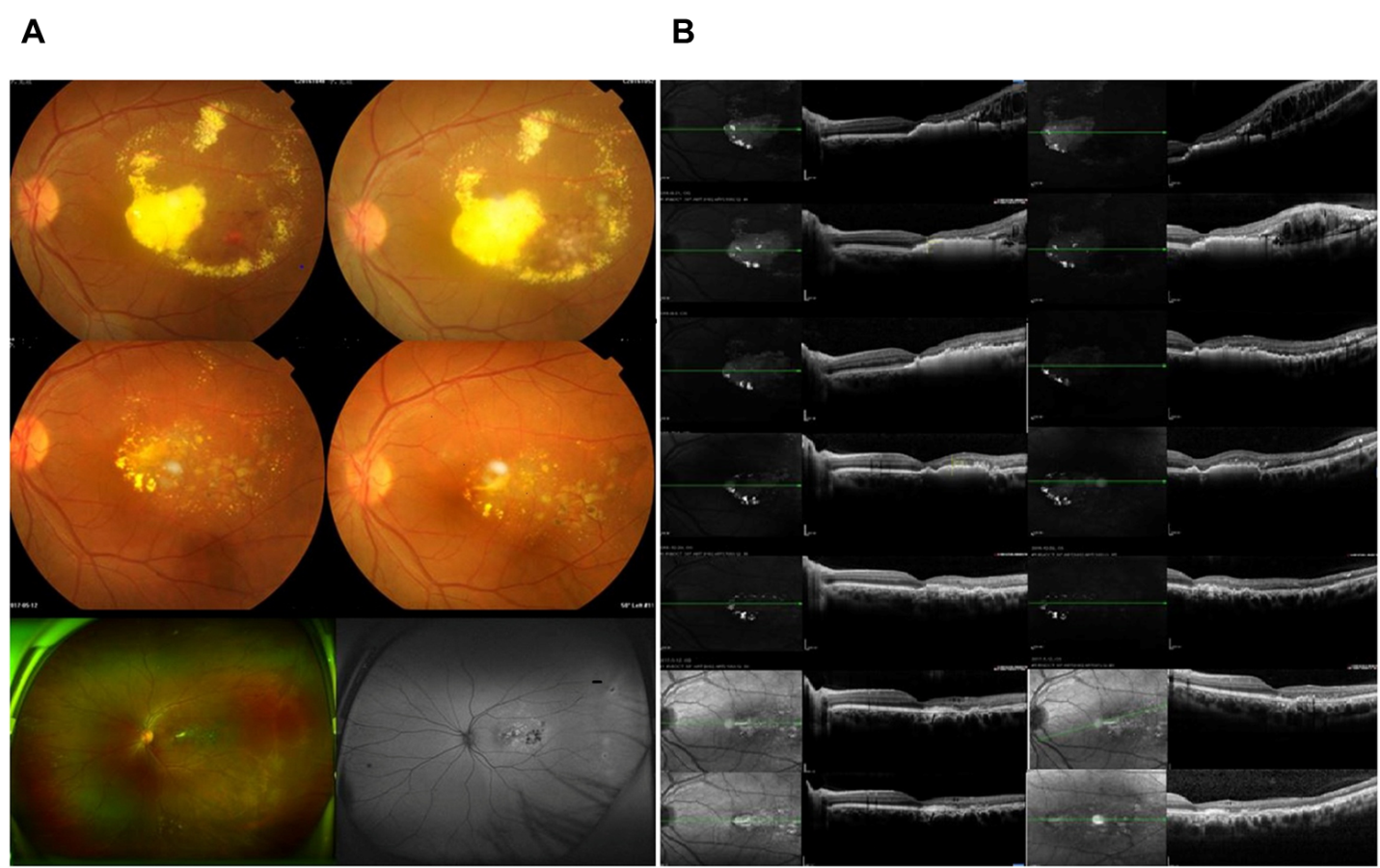
Grade E (Poor): 0

**P-Reviewer:** Buchaim RL, Mehdipour P **S-Editor:** Zhang H **L-Editor:** Filipodia **P-Editor:** Liu JH

**Figure Legends**



**Figure 1 Mean increase in the** **best-corrected visual acuity from the pretreatment value.** BCVA: Best-corrected visual acuity.

****

**Figure 2 Coats disease of the left eye treated with conbercept combined with 532-laser photocoagulation.** A: Fundus photographs of the patient’s eye before treatment and at 1 mo, 1 yr and 2 yrs after treatment as well as at the last follow-up; B: Changes in central retinal thickness and subretinal fluid before treatment and at 1 wk, 1 mo, 3 mo, 1 yr and 2 yrs after treatment as well as at the final follow-up. The retinal interstitial layer and subretinal regions showed gradual reductions in edema and exudation.

**Table 1 Clinical characteristics of the study participants**

|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- |
| **Case no.** | **Sex/age in yr/eye** | **Stage** | **Comorbidity** | **Alcohol/smoking status** | **Quadrants of telangiectasia** | **No. of injections** | **No. of laser therapies** | **BCVA**  **baseline** | **BCVA 1 wk** | **BCVA 1 mo** | **BCVA 3 mo** | **BCVA 6 mo** | **BCVA 9 mo** | **BCVA 12 mo** | **BCVA 24 mo** | **BCVA 36 mo** | **Termination** |
| 1 | M/31/os | 2A | None | Smoker | 3 | 2 | 1 | 60 | 70 | 68 | 75 | 75 | 74 | 72 | 66 | 65 | 65 |
| 2 | M/35/os | 2A | None | Drinker | 2 | 1 | 1 | 55 | 70 | 75 | 70 | 70 | 70 | 75 | 80 | 76 | 76 |
| 3 | M/45/od | 2A | HT | Drinker | 2 | 1 | 1 | 28 | 40 | 45 | 50 | 53 | 55 | 55 | 53 | 55 | 55 |
| 4 | M/29/os | 2A | None | None | 3 | 1 | 2 | 39 | 55 | 65 | 60 | 58 | 60 | 60 | 58 | 62 | 62 |
| 5 | M/33/od | 2B | None | None | 3 | 2 | 1 | 55 | 70 | 70 | 75 | 75 | 75 | 77 | 75 | 75 | 75 |
| 6 | F/32/od | 2B | None | None | 3 | 2 | 2 | 70 | 78 | 75 | 78 | 78 | 80 | 80 | 77 | 75 | 75 |
| 7 | M/35/os | 2B | None | Drinker | 3 | 3 | 2 | 56 | 70 | 75 | 78 | 75 | 75 | 78 | 75 | 75 | 75 |
| 8 | M/37/od | 2A | Cervical spondylosis | Smoker | 2 | 1 | 1 | 54 | 72 | 74 | 77 | 77 | 77 | 78 | 77 | 77 | 77 |

BCVA: Best-corrected visual acuity; HT: Hypertension; os: Oculus sinister; od: Oculus dexter.

**Table 2 Best-corrected visual acuity**

| **Time of measurement** | **Best-corrected visual acuity** | ***P* (*vs* pretreatment value)** |
| --- | --- | --- |
| Pretreatment | 52.12 ± 12.93 |  |
| 1 wk after treatment | 65.63 ± 12.19 | < 0.001 |
| 1 mo after treatment | 68.38 ± 10.17 | < 0.001 |
| 3 mo after treatment | 70.38 ± 10.18 | < 0.001 |
| 6 mo after treatment | 70.13 ± 9.42 | < 0.001 |
| 9 mo after treatment | 70.75 ± 8.75 | < 0.001 |
| 12 mo after treatment | 71.88 ± 9.28 | < 0.001 |
| 24 mo after treatment | 70.13 ± 9.98 | < 0.001 |
| 36 mo after treatment | 70.01 ± 8.23 | < 0.001 |
|  | *P*-trendlinear < 0.001 |  |

Data are presented as mean ± standard deviation.