W J C C World Journal of Clinical Cases

Submit a Manuscript: https://www.f6publishing.com

World J Clin Cases 2020 December 26; 8(24): 6243-6251

DOI: 10.12998/wjcc.v8.i24.6243

Retrospective Study

ISSN 2307-8960 (online)

ORIGINAL ARTICLE

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

Li Jiang, Bo Qin, Xiao-Ling Luo, He Cao, Ting-Ming Deng, Ming-Ming Yang, Ting Meng, Hui-Qin Yang

ORCID number: Li Jiang 0000-0002-2478-620X; Bo Qin 0000-0002-7653-272X; Xiao-Ling Luo 0000-0001-7599-9822; He Cao 0000-0002-7540-0661; Ting-Ming Deng 0000-0001-8032-8281; Ming-Ming Yang 0000-0002-5550-729; Ting Meng 0000-0002-3390-5178; Hui-Qin Yang 0000-0001-9210-4648.

Author contributions: Jiang L was responsible for the conception of the article, case analysis, statistics and article writing and was the main contributor to the writing of the manuscript; Qin B and Luo XL were responsible for the guidance and revision of the article; Cao H was responsible for assisting with the query of literature and analysis of statistical results; Deng TM collected and reviewed follow-up records of major cases; Yang MM, Meng T and Yang HQ were responsible for the collection of case data and assisting with document search; All authors have read and approved the manuscript.

Supported by the Shenzhen Municipal Health and Family Planning System Research Project, No. SZFZ2017085.

Institutional review board

statement: The study was approved by the ethics committee of Shenzhen People's Hospital (No. LL-KY-2020238).

Li Jiang, Xiao-Ling Luo, He Cao, Ting-Ming Deng, Ming-Ming Yang, Ting Meng, Hui-Qin Yang, Department of Ophthalmology, Shenzhen People's Hospital, Second Clinical Medical College of Jinan University, Shenzhen 518020, Guangdong Province, China

Bo Qin, Department of Ophthalmology, Shenzhen Aier Eye Hospital Affiliated to Jinan University, Shenzhen 518032, Guangdong Province, China

Corresponding author: Bo Qin, MD, Doctor, Shenzhen Aier Eye Hospital Affiliated to Jinan University, No. 1017 Dongmen North Road, Luohu District, Shenzhen 518020, Guangdong Province, China. jiang.li@szhospital.com

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 $303.30 \pm$ 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 $37.33 \pm$ 2.26 mo. Average BCVA of the affected eye was 51.17 ± 15.15 letters before



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.

Open-Access: This article is an open-access article that was selected by an in-house editor and fully peer-reviewed by external reviewers. It is distributed in accordance with the Creative Commons Attribution NonCommercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: htt p://creativecommons.org/License s/by-nc/4.0/

Manuscript source: Unsolicited manuscript

Specialty type: Medicine, research and experimental

Country/Territory of origin: China

Peer-review report's scientific quality classification

Grade A (Excellent): 0 Grade B (Very good): B Grade C (Good): C Grade D (Fair): 0 Grade E (Poor): 0

Received: August 4, 2020 Peer-review started: August 4, 2020 First decision: August 21, 2020 Revised: September 22, 2020 Accepted: November 2, 2020 Article in press: November 2, 2020 Published online: December 26, 2020

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

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

©The Author(s) 2020. Published by Baishideng Publishing Group Inc. All rights reserved.

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.

Citation: Jiang L, Qin B, Luo XL, Cao H, Deng TM, Yang MM, Meng T, Yang HQ. Three-year follow-up of Coats disease treated with conbercept and 532-nm laser photocoagulation. World J Clin Cases 2020; 8(24): 6243-6251

URL: https://www.wjgnet.com/2307-8960/full/v8/i24/6243.htm DOI: https://dx.doi.org/10.12998/wjcc.v8.i24.6243

INTRODUCTION

Coats disease is an idiopathic exudative outer retinopathy caused by abnormal retinal vascular development^[1,2]. The main clinical features of Coats disease are dilated retinal capillaries, miliary or small aneurysms and retinal and subretinal exudation^[1,2]. The estimated incidence of Coats disease in the population is 0.09 per 100000 persons, and males are more commonly affected than females^[3]. The disorder usually affects adolescents with a mean age at presentation of around 12 years^[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]. Standard treatments for Coats disease include laser therapy, cryotherapy and surgical techniques such as subretinal fluid drainage and vitrectomy^[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]. 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]. There has been increasing interest in the use of anti-VEGF agents as an adjunct treatment for Coats disease^[1,2,10]. 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].

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]. This fusion protein binds VEGF with high affinity, has a long halflife in the vitreous and blocks all isoforms of VEGF-A, VEGF-B, VEGF-C and placental growth factor^[16]. Conbercept has been shown to be an effective treatment for retinal diseases such as diabetic retinopathy-induced macular edema^[17,18] and age-related





macular degeneration^[19-22]. 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].

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]. 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].

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 μ m, the energy was 150 MW, and the duration was 200 ms.

Zaishidena® WJCC | https://www.wjgnet.com

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 of 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].

Statistical analysis

SPSS 23.0 (IBM Corp., Armonk, NY, United States) was used for the statistical analyses. Measurement data are expressed as mean \pm 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 $303.30 \pm 107.87 \mu m$ (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*-trend_{linear} < 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 μ m before treatment but decreased to 587 μ m, 372 μ m, 322 μ m, 267 μ m, 221 μ m and 208 μ m 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).

WJCC | https://www.wjgnet.com

Table	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	HTN.	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; HTN.: Hypertension; os: Oculus sinister; od: Oculus dexter.

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]; 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

Jiang L et al. Conbercept and laser for Coats disease

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						
	<i>P</i> -trend _{linear} < 0.001							

Data are presented as mean ± standard deviation.



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

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], 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] 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]. 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]. 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]: The combination of intravitreal injection of conbercept with laser



WJCC | https://www.wjgnet.com



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.

> 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.

REFERENCES

- Sen M, Shields CL, Honavar SG, Shields JA. Coats disease: An overview of classification, 1 management and outcomes. Indian J Ophthalmol 2019; 67: 763-771 [PMID: 31124484 DOI: 10.4103/ijo.IJO_841_19]
- 2 Yang X, Wang C, Su G. Recent advances in the diagnosis and treatment of Coats' disease. Int Ophthalmol 2019; 39: 957-970 [PMID: 30895419 DOI: 10.1007/s10792-019-01095-8]
- 3 Morris B, Foot B, Mulvihill A. A population-based study of Coats disease in the United Kingdom I: epidemiology and clinical features at diagnosis. Eye (Lond) 2010; 24: 1797-1801 [PMID: 20865031 DOI: 10.1038/eye.2010.126]
- Rishi E, Rishi P, Appukuttan B, Uparkar M, Sharma T, Gopal L. Coats' disease of adult-onset in 48 4 eyes. Indian J Ophthalmol 2016; 64: 518-523 [PMID: 27609165 DOI: 10.4103/0301-4738.190141]
- Kusaka S. Surgical Management of Coats Disease. Asia Pac J Ophthalmol (Phila) 2018; 7: 156-159 5 [PMID: 29664241 DOI: 10.22608/APO.201867]
- Kase S, Rao NA, Yoshikawa H, Fukuhara J, Noda K, Kanda A, Ishida S. Expression of vascular 6 endothelial growth factor in eyes with Coats' disease. Invest Ophthalmol Vis Sci 2013; 54: 57-62 [PMID: 23221067 DOI: 10.1167/iovs.12-10613]
- 7 He YG, Wang H, Zhao B, Lee J, Bahl D, McCluskey J. Elevated vascular endothelial growth factor level in Coats' disease and possible therapeutic role of bevacizumab. Graefes Arch Clin Exp Ophthalmol 2010; 248: 1519-1521 [PMID: 20379736 DOI: 10.1007/s00417-010-1366-1]
- Sun Y, Jain A, Moshfeghi DM. Elevated vascular endothelial growth factor levels in Coats disease: 8 rapid response to pegaptanib sodium. Graefes Arch Clin Exp Ophthalmol 2007; 245: 1387-1388 [PMID: 17342503 DOI: 10.1007/s00417-007-0559-8]
- Zhao Q, Peng XY, Chen FH, Zhang YP, Wang L, You QS, Jonas JB. Vascular endothelial growth factor in Coats' disease. Acta Ophthalmol 2014; 92: e225-e228 [PMID: 23764089 DOI: 10.1111/aos.12158
- Shields CL, Udyaver S, Dalvin LA, Lim LS, Atalay HT, L Khoo CT, Mazloumi M, Shields JA. Coats 10 disease in 351 eyes: Analysis of features and outcomes over 45 years (by decade) at a single center. Indian J Ophthalmol 2019; 67: 772-783 [PMID: 31124485 DOI: 10.4103/ijo.IJO 449 19]
- 11 Zheng XX, Jiang YR. The effect of intravitreal bevacizumab injection as the initial treatment for Coats' disease. Graefes Arch Clin Exp Ophthalmol 2014; 252: 35-42 [PMID: 23873253 DOI: 10.1007/s00417-013-2409-1]
- 12 Li S, Deng G, Liu J, Ma Y, Lu H. The effects of a treatment combination of anti-VEGF injections, laser coagulation and cryotherapy on patients with type 3 Coat's disease. BMC Ophthalmol 2017; 17: 76 [PMID: 28532448 DOI: 10.1186/s12886-017-0469-4]
- 13 Gaillard MC, Mataftsi A, Balmer A, Houghton S, Munier FL. ranibizumab in the management of advanced Coats disease Stages 3B and 4: long-term outcomes. Retina 2014; 34: 2275-2281 [PMID: 25075562 DOI: 10.1097/IAE.000000000000248]
- 14 Raoof N, Quhill F. Successful use of intravitreal bevacizumab and pascal laser photocoagulation in the management of adult Coats' disease. Middle East Afr J Ophthalmol 2013; 20: 256-258 [PMID: 24014992 DOI: 10.4103/0974-9233.114805]
- 15 Villegas VM, Gold AS, Berrocal AM, Murray TG. Advanced Coats' disease treated with intravitreal bevacizumab combined with laser vascular ablation. Clin Ophthalmol 2014; 8: 973-976 [PMID: 24876764 DOI: 10.2147/OPTH.S62816]
- 16 de Oliveira Dias JR, de Andrade GC, Novais EA, Farah ME, Rodrigues EB. Fusion proteins for treatment of retinal diseases: aflibercept, ziv-aflibercept, and conbercept. Int J Retina Vitreous 2016; 2: 3 [PMID: 27847621 DOI: 10.1186/s40942-016-0026-y]
- 17 Zhou Q, Guo C, You A, Wang D, Wang W, Zhang X. One-year outcomes of novel VEGF decoy receptor therapy with intravitreal conbercept in diabetic retinopathy-induced macular edema. Mol Vis 2019; 25: 636-644 [PMID: 31700228]
- Li F, Zhang L, Wang Y, Xu W, Jiao W, Ma A, Zhao B. One-Year Outcome of Conbercept Therapy 18 for Diabetic Macular Edema. Curr Eye Res 2018; 43: 218-223 [PMID: 29265939 DOI: 10.1080/02713683.2017.1379542
- 19 Wu BH, Wang B, Wu HQ, Chang Q, Lu HQ. Intravitreal conbercept injection for neovascular age-



related macular degeneration. Int J Ophthalmol 2019; 12: 252-257 [PMID: 30809481 DOI: 10.18240/ijo.2019.02.11]

- Liu K, Song Y, Xu G, Ye J, Wu Z, Liu X, Dong X, Zhang M, Xing Y, Zhu S, Chen X, Shen Y, 20 Huang H, Yu L, Ke Z, Rosenfeld PJ, Kaiser PK, Ying G, Sun X, Xu X; PHOENIX Study Group. Conbercept for Treatment of Neovascular Age-related Macular Degeneration: Results of the Randomized Phase 3 PHOENIX Study. Am J Ophthalmol 2019; 197: 156-167 [PMID: 30148987 DOI: 10.1016/j.ajo.2018.08.026]
- Cui C, Lu H. Clinical observations on the use of new anti-VEGF drug, conbercept, in age-related 21 macular degeneration therapy: a meta-analysis. Clin Interv Aging 2018; 13: 51-62 [PMID: 29343949 DOI: 10.2147/CIA.S151225]
- 22 Cui J, Sun D, Lu H, Dai R, Xing L, Dong H, Wang L, Wei D, Jiang B, Jiao Y, Jablonski MM, Charles S, Gu W, Chen H. Comparison of effectiveness and safety between conbercept and ranibizumab for treatment of neovascular age-related macular degeneration. A retrospective casecontrolled non-inferiority multiple center study. Eye (Lond) 2018; 32: 391-399 [PMID: 28937147 DOI: 10.1038/eye.2017.187]
- Zhang L, Ke Y, Wang W, Shi X, Hei K, Li X. The efficacy of conbercept or ranibizumab intravitreal 23 injection combined with laser therapy for Coats' disease. Graefes Arch Clin Exp Ophthalmol 2018; 256: 1339-1346 [PMID: 29549425 DOI: 10.1007/s00417-018-3949-1]
- Shields JA, Shields CL, Honavar SG, Demirci H. Clinical variations and complications of Coats 24 disease in 150 cases: the 2000 Sanford Gifford Memorial Lecture. Am J Ophthalmol 2001; 131: 561-571 [PMID: 11336930 DOI: 10.1016/s0002-9394(00)00883-7]
- Holz FG, Amoaku W, Donate J, Guymer RH, Kellner U, Schlingemann RO, Weichselberger A, 25 Staurenghi G; SUSTAIN Study Group. Safety and efficacy of a flexible dosing regimen of ranibizumab in neovascular age-related macular degeneration: the SUSTAIN study. Ophthalmology 2011; 118: 663-671 [PMID: 21459217 DOI: 10.1016/j.ophtha.2010.12.019]
- 26 Wang KY, Cheng CK. A combination of intravitreal bevacizumab injection with tunable argon yellow laser photocoagulation as a treatment for adult-onset Coats' disease. J Ocul Pharmacol Ther 2011; 27: 525-530 [PMID: 21951171 DOI: 10.1089/jop.2011.0088]
- 27 Zhang C, Dong F, Wang R. Clinical characteristics and diagnosis and treatment of Coats disease in adults. Zhonghua Yandibing Zazhi 2008; 24: 279-282
- Otani T, Yasuda K, Aizawa N, Sakai F, Nakazawa T, Shimura M. Over 10 years follow-up of Coats' 28 disease in adulthood. Clin Ophthalmol 2011; 5: 1729-1732 [PMID: 22205832 DOI: 10.2147/OPTH.S27938]
- Sigler EJ, Randolph JC, Calzada JI, Wilson MW, Haik BG. Current management of Coats disease. 29 Surv Ophthalmol 2014; 59: 30-46 [PMID: 24138893 DOI: 10.1016/j.survophthal.2013.03.007]
- Kaul S, Uparkar M, Mody K, Walinjkar J, Kothari M, Natarajan S. Intravitreal anti-vascular 30 endothelial growth factor agents as an adjunct in the management of Coats' disease in children. Indian J Ophthalmol 2010; 58: 76-78 [PMID: 20029154 DOI: 10.4103/0301-4738.58480]
- Lin CJ, Hwang JF, Chen YT, Chen SN. The effect of intravitreal bevacizumab in the treatment of 31 Coats disease in children. Retina 2010; 30: 617-622 [PMID: 19996822 DOI: 10.1097/IAE.0b013e3181c2e0b7]



WJCC | https://www.wjgnet.com



Published by Baishideng Publishing Group Inc 7041 Koll Center Parkway, Suite 160, Pleasanton, CA 94566, USA Telephone: +1-925-3991568 E-mail: bpgoffice@wjgnet.com Help Desk: https://www.f6publishing.com/helpdesk https://www.wjgnet.com

