Multimodal images of chronic branch retinal vein occlusion

Abstract: Two illustrative cases of chronic branch retinal vein occlusion are presented with multimodal imaging, including commercially available optical coherence tomography angiography. In these two patients, retinal ischemia and collateral vessels were well imaged without the need to use traditional fluorescein angiography. Optical coherence tomography angiography provides useful information for the diagnosis and management of patients with branch retinal vein occlusion and other retinal vascular diseases.

Keywords: branch retinal vein occlusion, optical coherence tomography angiography, spectral domain optical coherence tomography, multimodal imaging, BRVO management

Introduction
Branch retinal vein occlusion (BRVO) is a common retinal vascular disease that may lead to cystoid macular edema (CME), retinal ischemia, retinal neovascularization, and other complications. Spectral domain optical coherence tomography (SD-OCT) is important in the management of this entity. More recently, OCT angiography (OCT-A) has become available and has provided new insights for clinicians managing this disease.

Using multimodal imaging, including OCT-A imaging with the commercially available Cirrus 5000 with AngioPlex (Zeiss, Jena, Germany), the pathologic features of chronic BRVO in two patients are demonstrated.

Report of two cases
Case 1
A 56-year-old female with a history of chronic BRVO left eye returned for scheduled follow-up, noting long-standing but stable blurred vision. She had been treated with intravitreal Avastin® (bevacizumab) (Avastin®; Genentech, South San Francisco, CA, USA) and triamcinolone acetonide in the past, but no treatments for several years. On examination, best-corrected visual acuity was 20/40 in the left eye. Fundus examination revealed sclerotic vessels in the superotemporal quadrant with collaterals near the optic disc, as well as CME (Figure 1A). Humphrey visual field 24-2 testing revealed an inferior arcuate defect encroaching fixation (Figure 1B). SD-OCT revealed mild thickening centrally and atrophy superotemporally (Figure 1C). OCT-A through the center of the macula revealed a prominent central cyst as well as temporal thinning (Figure 1D). The OCT-A retina slab (6×6 mm) demonstrated marked capillary
nonperfusion in the superotemporal macula, involving the foveal avascular zone (FAZ) (Figure 1E).

Case 2
A 52-year-old female with a history of chronic hemiretinal vein occlusion right eye presented with recurrent episodes of floaters and visual loss. She had been treated with scatter photocoagulation elsewhere about six months previously. On examination, best-corrected visual acuity was 20/25. Fundus examination revealed sclerotic veins inferotemporally and inferonasally. Scatter photocoagulation was evident inferotemporally. Collateral vessels temporal to the center of the macula and crossing the horizontal raphe were present (Figure 2A). Fluorescein angiography revealed areas of ischemia temporally and inferonasally, with collateral vessels temporal to the center of the macula (Figure 2B). Late frames revealed mild macular leakage and more prominent leakage inferiorly, consistent with retinal neovascularization (Figure 2C). SD-OCT through the center of the macula showed no frank CME (Figure 2D). The OCT angiography retina slab (6x6 mm) demonstrated prominent collateral vessels temporal (and some nasal) to the foveal avascular zone, as well as capillary nonperfusion involving the inferior FAZ as well as inferotemporal to the center of the macula (Figure 2E).

Discussion
These two cases illustrate the utility of OCT-A in providing additional information in the management of patients with chronic BRVO. In case 1, OCT-A demonstrates capillary nonperfusion involving the FAZ, which correlates with the visual field defect encroaching fixation. This suggests that visual loss is due to both CME and macular ischemia and that further pharmacologic treatment of the CME might not benefit the patient. In case 2, OCT-A demonstrates capillary nonperfusion and highlights the collateral vessels, which were poorly visualized on fundus photography but were visualized using fluorescein angiography. Case 2 also provides a comparison between OCT-A and fluorescein angiography, demonstrating concordance between the collateral vessels and the macular ischemia on each test.

The role of OCT-A in the management of BRVO is evolving. OCT-A may offer similar information to fluorescein angiography and can be obtained rapidly and without the need for dye injection. OCT-A is reported to image the FAZ well in patients with BRVO. However, the technology is currently expensive, and image quality depends on the patient’s ability to fixate.

As additional patients with acute and chronic BRVO, and other retinal vascular diseases, are imaged with OCT-A, the utility of this noninvasive imaging technology will become...
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better understood and may help in screening or predicting visual outcomes.

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Figure 2 Hemiretinal vein occlusion, right eye.

Notes: (A) Fundus photography demonstrated the site of the occlusion at the disc (upward vertical arrow) and sclerotic veins in the inferotemporal and inferonasal quadrants (downward vertical arrows). Collateral vessels temporal to the center of the macula did not photograph well. Photocoagulation burns (horizontal arrow) are evident. (B) Fluorescein angiography demonstrated areas of capillary nonperfusion temporally and inferonasally (vertical arrows). Collateral vessels temporal to the macula were noted (horizontal arrow). (C) Fluorescein angiography late frames demonstrated mild macular leakage (horizontal arrow) and more prominent leakage inferiorly (vertical arrow). (D) SD-OCT demonstrated no frank cystoid macular edema. (E) OCT-A retina slab (6x6 mm) demonstrated prominent collateral vessels temporal to (and some nasal to) the center of the macula, as well as areas of capillary nonperfusion inferior to the fovea and inferotemporal to the fovea.

Abbreviations: OCT-A, optical coherence tomography-angiography; SD-OCT, spectral domain optical coherence tomography.

Written informed consent has been provided by the patients to have the case details and any accompanying images published.
References


