Isolated cotton-wool spots of unknown etiology: management and sequential spectral domain optical coherence tomography documentation

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Abstract: Cotton-wool spots (CWSs) are common retinal manifestations of many diseases including diabetes mellitus, systemic hypertension, and acquired immunodeficiency syndrome. Clinically they appear as whitish, fluffy patches on the retina and eventually fade with time. In this study, spectral domain optical coherence tomography (SD-OCT) with mapping was used to demonstrate in vivo the characteristics of an isolated CWS in a 59-year-old patient as well as its appearance immediately after ophthalmoscopic resolution. Presented here is the work-up and management of this clinical problem for the ophthalmologist. The authors propose that SD-OCT could be a valuable research tool in characterizing and following the dynamic CWS changes at individual retinal layer level, with potential clinical applications as a screening or diagnostic tool in CWS-related diseases.

Keywords: SD-OCT, spectral domain optical coherence tomography, retinal lesion

Introduction
Cotton-wool spots (CWSs) are retinal lesions, most commonly seen as manifestations of diabetes mellitus and systemic hypertension. They are also associated with a number of other etiologies including ischemic, embolic, connective tissue, neoplastic, and infectious, but occasionally no underlying cause can be identified.

CWSs are thought to represent axoplasmic debris at the level of retinal ganglion cell axons resulting from axoplasmic flow interruption due to vascular or mechanical causes. Ophthalmoscopically, CWSs appear as whitish, fluffy retinal patches that eventually fade with time. Described in this paper is the use of spectral domain optical coherence tomography (SD-OCT) to document the progression of an isolated cotton-wool spot, from presentation to fundoscopic resolution.

Case report
A 59-year-old chemist presented to the ophthalmology department with a 3-day history of acute onset, painless reduction of the paracentral vision of his left eye. He used spectacles for reading and was previously fit and well with no history of ophthalmic or systemic disease. He wasn’t taking any medication, had never smoked, and there was no family history of note.

His uncorrected visual acuities were 20/16 right and 20/20 left. Ocular examination showed an isolated CWS located between his left optic disc and macula (Figure 1A) but no other pathology. His blood pressure was 138/82 mmHg and systemic examination was unremarkable.
A full hematological work-up was performed that included fasting glucose, complete blood count, urea, electrolytes and calcium, lipid profile, inflammatory markers, coagulation, connective tissue disease markers, human immunodeficiency virus (HIV), and serum protein electrophoresis. In addition, 24-hour urine was collected for quantification. Results were negative. Carotid Doppler ultrasound and echocardiography were also unremarkable.

The patient’s fundus fluorescein angiogram images at presentation were consistent with the ophthalmoscopic findings and did not reveal any evidence of other abnormalities than the isolated CWS in his left retina.

SD-OCT with Cirrus HD-OCT (Carl Zeiss Meditec Inc, Dublin, CA), also performed at presentation, showed marked retinal thickening corresponding to the area of the CWS (Figure 2A). The thickening was confined to the level of the retinal nerve fiber layer (Figure 3A).

The patient was reviewed 9 weeks later. He reported that the vision in his left eye had returned to normal. Fundoscopically, there was resolution of the CWS (Figure 1B) without any intervention. His blood pressure was still within normal limits at 135/79 mmHg. Based on the whole clinical picture, the authors of this paper felt that further exhaustive investigations to establish an underlying cause for the CWS were inappropriate, and a diagnosis of isolated cotton-wool spot of unknown etiology was made.

At 9 weeks from presentation, the area of the resolved CWS was scanned again with the Cirrus HD-OCT using software (version 4.0.1.3) that allowed real-time automatic registration for thickness change analysis as well as side-by-side visual inspection. There was significant intraretinal thinning at the area of the lesion compared with adjacent healthy retina, evident on the retinal thickness map (Figure 2B) as well as the longitudinal image (Figure 3B), although it was not clear which individual retinal layer levels were affected.

**Discussion**

CWSs are useful fundoscopic signs for grading hypertensive retinopathy and diabetic retinopathy. They signal a declining CD4 count in HIV disease and are commonly seen in the course of central and branch retinal vein occlusions. CWSs can be a presenting sign of multiple myeloma and post-radiation retinopathy and are associated with a wide spectrum of diseases including cardiac valvular disease, carotid artery obstruction, dermatomyositis, systemic lupus erythematosus, polyarteritis nodosa, leukemia, metastatic carcinoma, Purtscher retinopathy, and giant cell arteritis.

Patients may experience relative or absolute scotomas corresponding to the areas of CWSs. Following ophthalmoscopic resolution, visual function does not usually recover, although recovery was reported by the patient in this study and has also been documented by other authors. The mechanism of visual recovery is unknown, but a plausible explanation might be survival of enough retinal ganglion cell axons during the initial “assault” by the axoplasmic debris. If the number of surviving retinal ganglion cells is above a certain threshold, signal transmission – or at least part of it – is reestablished following resolution of the axoplasmic debris.

In recent years, SD-OCT has been established as a powerful weapon in the armamentarium of the retina specialist. But until now, there has been limited research on the SD-OCT
Characteristics of ophthalmoscopically visible CWSs and practically no clinical applications. Using SD-OCT as well as software mapping to more accurately identify, scan, and measure the same anatomical area of the retina in the patient, the structural changes associated with a CWS episode were able to be demonstrated and characterized in vivo.

Firstly, in the acute phase, 3 days from the onset of symptoms, there was marked thickening of the retina (Figure 2A) confined to the retinal nerve fiber layer with no apparent thickness change in the other layers (Figure 3A). This is consistent with the theory of accumulation of axoplasmic debris at the level of retinal ganglion cell axons14 that is visible ophthalmoscopically as a CWS.

Secondly, following ophthalmoscopic resolution at 9 weeks, there was overall retinal thinning at the area of the lesion compared with adjacent healthy retina, evident on the retinal thickness map (Figure 2B) as well as the longitudinal image (Figure 3B). Although not all individual retinal layers were distinguishable, a previous case series17 has showed marked ganglion cell layer thinning and moderate retinal nerve fiber layer, inner nuclear layer, inner plexiform layer, outer plexiform layer thinning with moderate outer nuclear layer thickening following resolution (2–11 years) of CWS in HIV disease. This is the first OCT report demonstrating retinal thinning, just 9 weeks from presentation of a CWS.

Conclusion
SD-OCT imaging has provided a new insight into the dynamic changes of the diseased retina. This image study suggests that SD-OCT can be a valuable research tool in following, in vivo, the anatomical and histological changes at individual retinal layer level of CWSs acutely and after apparent resolution.

As the use of retinal OCT imaging in ophthalmology clinics becomes more common and resolution of images improves further, it is possible that high-resolution OCT could also prove to be a useful screening or diagnostic tool in systemic disease for establishing areas of resolved CWS, in fundoscopically healthy looking retinas.

Additionally it can be used to determine the stage of disease progression or disease regression depending on the nature of retinal vascular pathology and severity in conjunction with the clinical symptomatology. The advances in high-resolution OCT technology allow histological analysis of the neurosensory retina in health and disease with multiple clinical applications.

Disclosure
The authors declare no conflicts of interest in this work.

References