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## Central Retinal Vein Occlusion in a 13-Year-Old COVID-19 Patient: A Case Report

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Dear Editor.

The recent COVID-19 pandemic, caused by infection from the SARS-COV-2, has affected more than 328 million people and is responsible for more than 5 million deaths worldwide from 2019 to January 2022 [1]. COVID-19 patients may have a number of complex and varied coagulation abnormalities that induce a hypercoagulable state and provoke venous thromboembolism, a condition called COVID-19-associated coagulopathy [2]. In fact, COVD-19 appears to also affect the retina, with the most common retinal manifestations involving microvascular changes such as cotton wool spots and retinal microhemorrhages [3]. Thus, retinal vein occlusions (RVO) in COVID-19 patients can be associated with this hypercoagulable state. Herein, we report a case of central RVO (CRVO) in a young teenage girl with COVID-19. Written informed consent for publication of the research details and clinical images was obtained from the patient.

A 13-year-old girl presented with decreased vision in her left eye for 13 days. Her past medical history was unremarkable, except for COVID-19 infection, with which she was diagnosed 16 days before the onset of ocular symptoms. Her best-corrected visual acuity was 20 / 25 in the

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right eye and 20 / 63 in the left eye. Her intraocular pressures were 20 and 17 mmHg in the right and left eyes, respectively. Anterior segment findings were within normal limits in both eyes. Fundus findings in her right eye was unremarkable. The left eve fundus showed multiple retinal hemorrhages with optic disc edema and dilated tortuous vessels in all quadrants (Fig. 1A). Optical coherence tomography (OCT) showed macula edema with subretinal fluid and a central foveal thickness of 397 µm (Fig. 1B). Fluorescein angiography showed marked delay in arteriovenous transit time and staining of the dilated tortuous veins (Fig. 1C, 1D). Based on these clinical findings, the patient was diagnosed with nonischemic CRVO, with possible association with COVID-19. A full diagnostic workup, including blood pressure measurement, complete blood count and hypercoagulability tests with peripheral blood smears, tests for homocysteine, anti-DNA, ELISA, anti-β 2 glycoprotein 1 (anti-β2GP1) immunoglobulin G (IgG), anti-β2GP1 IgM, anti-cyclic citrullinated peptide antibody, cholesterol, and anticardiolipin antibodies IgM and IgG, were unremarkable, except for positive antinuclear antibody. After 2 monthly intravitreal anti-vascular endothelial growth factor injections in the left eye, her visual acuity improved to 20 / 40 and OCT showed resolution of macula edema (Fig. 1E, 1F), with resolution of flame-shaped hemorrhage and decreased vascular tortuosity on fundus examination.

CRVO typically occurs in elderlies and far less commonly in healthy teenagers. Our 13-year-old patient did not have major risk factors of CRVO, including diabetes mellitus, hypertension, hypercoagulability, atherosclerosis, and

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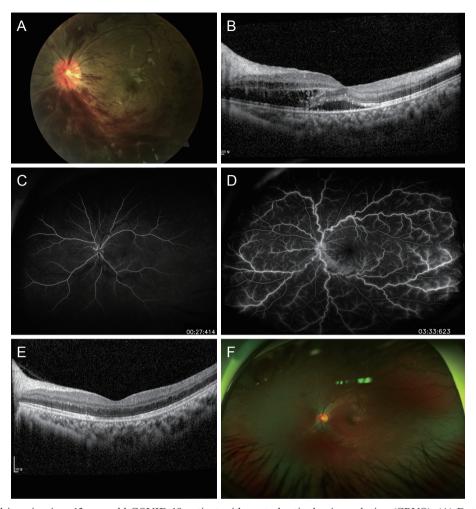


Fig. 1. Multimodal imaging in a 13-year-old COVID-19 patient with central retinal vein occlusion (CRVO). (A) Dot-, blot-, and flame-shaped hemorrhages throughout all four retinal quadrants can be observed, with tortuous and dilated veins predominantly in the inferior retina and optic disc edema. (B) Macular edema can be observed on optical coherence tomography. (C) Early phase fluorescein angiography shows marked delay in the arteriovenous transit time, while (D) late phase shows staining of the dilated tortuous veins. There is no evidence of widespread capillary nonperfusion, providing evidence for the diagnosis of nonischemic CRVO. (E,F) After a second intravitreal injection of anti–vascular endothelial growth factor, there was resolution of the secondary macular edema, and marked improvement in the retinal hemorrhage, although there remains venous dilation.

vasculitis secondary to systemic inflammation, except that she had recently been diagnosed with COVID-19 approximately 2 weeks before the onset of visual symptoms. Although she had tested positive for antinuclear antibody (ANA), she did not have other symptoms or signs indicating autoimmune diseases including systemic lupus erythematosus, scleroderma, rheumatoid arthritis, Sjögren's syndrome or other rheumatic systemic diseases. Additionally, there was no family history related to autoimmune diseases, and a positive ANA can be seen in healthy individuals.

COVID-19-associated coagulopathy with venous thromboembolic disease have been described in many studies, including cerebral venous thrombosis in three young COVID-19 patients [4], and autopsy findings of diffuse small vessel thrombosis in patients with severe COVID-19 [5]. Excessive inflammation, platelet activation, endothelial dysfunction, and venous stasis have been suggested as the cause of arterial and venous circulation problems in COVID-19 patients. It is yet to be determined whether there is a direct causal relationship between CRVO and COVID-19, but the occurrence of the disease shortly after COVID-19 infection at such a young age in the absence of other systemic risk factors suggests that the prothrombotic state created by COVID-19 may have contributed to CRVO

development in our patient. This case adds to the existing literature that there may be possible association in the occurrence of CRVO in the setting of COVID-19. As COVID-19 patients are at risk for vascular occlusive disease, patient education for retinal vascular complications and early detection with prompt treatment are imperative for visual prognosis.

Conflicts of Interest: None.
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