Giant cell arteritis related arteritic anterior ischemic optic neuropathy: Clinico-pathological correlation

A 63-year-old male presented with sudden painless vision loss in his right eye of 2 day duration. He had no perception of light, Relative afferent pupillary defect (RAPD), pale disc edema along with an inferotemporal branch retinal artery occlusion [Fig. 1a-d]. Clinical diagnosis of arteritic anterior ischemic optic neuropathy was made, and systemic examination revealed repeated bouts of fever and right sided scalp tenderness. [1,2] Right side temporal artery biopsy revealed giant cell arteritis. The patient was treated with intravenous pulse steroids and immunosuppressive therapy. [2,3] In cases of severe vision loss with pale disc edema, giant cell arteritis should be kept in mind. [4,5]

Declaration of patient consent

The authors certify that they have obtained all appropriate patient consent forms. In the form the patient(s) has/have given his/her/their consent for his/her/their images and other clinical information to be reported in the journal. The patients understand that their names and initials will not be published

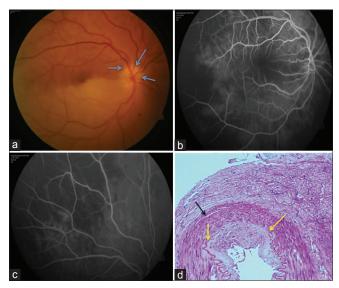


Figure 1: (a) showing pale disc edema (blue arrows) with an inferotemporal branch retinal artery occlusion, (b and c) showing "wedge shaped" equatorial choroidal infarcts on fluorescein angiogram, and (d) right side temporal artery biopsy at 200× magnification showing intimal proliferation with breaks in the internal elastic lamina (yellow arrows) with extravasation of red blood cells in the tunica media (black arrow) with minimal inflammatory cells, suggestive of vasculitis

and due efforts will be made to conceal their identity, but anonymity cannot be guaranteed.

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Conflicts of interest

There are no conflicts of interest.

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