

Comment on: Diagnostic positron emission tomography–computed tomography in clinically elusive giant cell arteritis

Sir,

Giant cell arteritis (GCA) can be categorized into cranial GCA and large vessel giant cell arteritis (LV-GCA).^[1] Cranial GCA frequently presents with headache, jaw claudication, and visual disturbances due to involvement of external carotid artery, whereas LV-GCA usually involves the aorta and its main branches and is often subclinical.^[2] The frequency of inflammatory aortic involvement varies from 22% to 85% of GCA cases.^[3]

Temporal artery biopsy (TAB) remains the gold standard for diagnosis of cranial GCA with hypoechoic halo on Doppler being similarly useful.^[2] The LV-GCA usually spares the temporal arteries, and hence, TAB has a low diagnostic yield for it. Conversely, positron emission tomography–computed tomography (PET-CT) of aorta is

a good diagnostic tool for LV-GCA, which presents with constitutional symptoms and has very low risk of ocular involvement.^[4]

Mohamed *et al.* in their article on ‘Diagnostic positron emission tomography–computed tomography in clinically elusive giant cell arteritis’ describe the utility of PET-CT for diagnosing a patient with headaches and raised erythrocyte sedimentation rate (ESR).^[5] We would like to ask the authors why PET-CT of aorta was done as the first investigation for a patient with signs of only cranial GCA. A negative aortic PET-CT cannot rule out cranial GCA. Not just the high cost and limited availability, but the low diagnostic yield of PET-CT in cranial GCA makes it an unlikely choice.

To conclude, PET-CT is of value in LV-GCA presenting with unexplained constitutional symptoms, raised inflammatory markers with negative TAB or Doppler. It is usually not recommended as first line in a patient with headaches or visual disturbances.

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Conflicts of interest

There are no conflicts of interest.

Sonali Gupta, Shreyans Jain¹

Birmingham and Midland Eye Centre, Birmingham, UK, ¹Dr. Rajendra Prasad Centre for Ophthalmic Sciences, All India Institute of Medical Sciences, New Delhi, India

Correspondence to: Dr. Sonali Gupta,
Birmingham and Midland Eye Centre, Dudley Road,
Birmingham B18 7QH, UK.
E-mail: sonali1286@yahoo.com

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