CYCLOPHOSPHAMIDE TREATMENT FOR UNRELENTING CNS VASCULITIS SECONDARY TO TUBERCULOUS MENINGITIS

Gregory Youngnam Chang, Phoenix: Gonzalez-Duarte et al.\(^1\) presented a 19-year-old woman with delayed recurrent unilateral ischemic strokes due to left middle cerebral arteritis while being treated over several months. The following case demonstrates that multiple ischemic strokes may be the initial presenting clinical feature of TB meningitis.

A 58-year-old Filipino woman complained of headache for several days and was found stuporous with left-sided weakness. MRI revealed bilateral ischemic strokes predominantly in the deep gray matter (figure). CSF examination revealed 52 leukocytes/mm\(^3\), protein of 237 mg/dL, glucose of 32 mg/dL, and later culturing out *Mycobacterium tuberculosis*. No further strokes occurred after starting anti-TB drugs and dexamethasone. Four weeks later, after discontinuing ventriculostomy and following steroid taper, the patient remained conversant yet clumsy in all limbs with residual right oculomotor nerve paresis.

These cases highlight the range of cerebral infarctions with different pathophysiologic mechanisms in the setting of TB meningitis. Early predominantly bilateral diencephalic ischemic strokes result from small perforator involvement from the inflammatory basilar meningitis surrounding the circle of Willis.\(^2\) Large vessel vasculitis, as Gonzalez-Duarte et al. describe, occurred due to a delayed-type hypersensitivity reaction.

Author Response: Alejandra Gonzalez-Duarte, Carlos Cantú-Brito, Mexico City, Mexico: Dr. Chang describes a case of TB meningitis with ischemic strokes that arose as the initial presentation. We understand that the MRI showed bilateral cerebral ischemia localized in the basal ganglia known as the “tubercular zone” that has been typically attributed to the involvement of the perforating branches at the base of the brain. Cerebrovascular complications of tuberculous meningitis are common.\(^3\) The controversy over the pathogenesis of vasculitis is far from resolved. Small vessel pathology may be a consequence of its immersion in the local inflammatory exudate or of direct luminal thrombosis. However, cortical strokes can also occur due to the involvement of the proximal portions of the middle, anterior, and posterior cerebral arteries, as well as the supraclinoid portion of the internal carotid. Moreover, other mechanisms such as changes in microvascular reactivity to cytokines may play a role. Dr. Chang’s case illustrates the concept that small vessel involvement may occur early in the course of the infection. However, the mechanisms of stroke in the later phases of the disease are less clear, possibly involving proliferation of the vascular intima, pointing toward a cascade of immune mechanisms.\(^4\)\(^5\)

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Figure Axial fluid-attenuated inversion recovery sequences reveal subacute strokes involving bilateral basal ganglia and thalamus (A, B), extending out to the right temporal lobe cortex and upper midbrain parenchyma (C)
ROLE OF DATSCAN AND CLINICAL DIAGNOSIS IN PARKINSON DISEASE
Tao Xie, Peter Warnke, Un Jung Kang, Chicago: We read with interest the article by Dr. de la Fuente-Fernandez1 and the editorial by Drs. Perlmutter and Eidelberg.2 We agree with the authors that the overall accuracy of a clinical diagnosis is numerically identical to the accuracy of DaTSCAN, which questions the DaTSCAN as a diagnostic tool.3 However, DaTSCAN has important and unique clinical value. One example is the diagnosis of early-onset Parkinson disease (PD), often with dystonia in addition to parkinsonism,4 which mimics dopa-responsive dystonia (DRD). The empirical levodopa/carbidopa trial may not differentiate one from the other because they both respond very well and both may develop dyskinesia. DaTSCAN is normal in DRD,5 while it shows decreased DAT binding in PD. Another scenario is to confirm the diagnosis of subtle early PD in a patient with severe essential tremor (ET) warranting deep brain stimulation (DBS) treatment, as 7% of patients with PD have prior ET.6 An abnormal DaTSCAN would lead us to select the subthalamic nucleus instead of ventral intermediate nucleus (VIM) as the DBS target given the lack of effect of VIM on anticipated progression of PD symptoms such as rigidity and bradykinesia.

Author Response: Raul de la Fuente-Fernandez, Ferrol, Spain: I appreciate the comments by Xie et al. on the potential diagnostic utility of DaTSCAN in 2 specific clinical scenarios. In the case of early-onset PD vs DRD, it might be best to use gene testing. Risks associated with radiation exposure are particularly high in young individuals.7 In severe ET plus subtle early PD, it should be decided if the treatment is directed to symptom severity (tremor) or the possibility of later occurrence of new symptoms (rigidity and bradykinesia). Then, as recent evidence suggests, the subthalamic nucleus could be considered a better target for neurosurgical treatment of ET than the VIM.7,8

CORRECTION
Caffeine for treatment of Parkinson disease: A randomized controlled trial
In the article “Caffeine for treatment of Parkinson disease: A randomized controlled trial” by R. Postuma et al. (Neurology® 2012;79:651–658), there is an omission in the study funding at the bottom of page 651. The statement should have read: “Supported by grants from the Canadian Institute of Health Research, the Webster Foundation, and the Drummond Foundation.” The authors regret the omission.
Cyclophosphamide treatment for unrelenting CNS vasculitis secondary to tuberculous meningitis

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