

The clinical variability of midbrain lesions

The midbrain is a crucial part of the central nervous system. In a small volume, it not only contains the nuclei of the oculomotor nerves^[1] (III and IV cranial nerves) but it is also crossed by many structures coming from the cerebral hemispheres, the cerebellum, and from the periphery. The midbrain also contains an important structure such as the red nucleus, the substantia nigra, and the superior cerebellar peduncle among others. Hence, it is easy to understand how different lesions in this area can lead to such a great variability in clinical presentation.^[2]

The midbrain can be involved in many different pathological process such as vascular accident (Weber's syndrome, Parinaud's syndrome with the typical upward gaze palsy), tumors (one of the first patients described as affected by supranuclear progressive palsy was instead affected by brainstem tumor), demyelinating processes (multiple sclerosis), infections, and neurodegenerative diseases (Parkinson's disease and supranuclear progressive palsy). Each one of these conditions can present with a broad clinical spectrum, from movement disorder to focal signs. One of the key symptoms is definitely the involvement of the third cranial nerve and often there is also the involvement of the conjugate upward gaze. The classic syndrome with this type of symptom is the supranuclear progressive palsy where the upward gaze palsy is one of the strongest symptoms for the diagnosis.^[3]

In the paper "Bilateral ptosis without upward gaze palsy-unusual presentation of midbrain tuberculoma,"^[4] Sarkar *et al.* describe a very interesting case of a young patient affected by midbrain tuberculoma with ataxia and ptosis. Neurotuberculosis is found in more or less 5–15% cases of extrapulmonary tuberculosis and common locations are cerebral and cerebellar hemisphere, instead brainstem locations are very rare. Brainstem tuberculomas were already described in literature with a wide range of clinical presentations such as "one and half syndrome," "eight and half syndrome," and "bilateral ptosis" as the readers will be able to find in reading the paper of

the authors. In many of these case reports, the lesions were located in the midbrain showing how much can be variable the clinical presentation of a lesion affecting the midbrain.

The paper is very interesting because the authors highlight the structure of the midbrain, especially the area where the oculomotor nucleus lies and its architecture. As well described in the text, due to the particular structure of the affected area of the midbrain with tuberculoma, it is rare that there is a bilateral ptosis without upward gaze palsy. Moreover, they show how the complexity of the midbrain can lead to a heterogeneous combination of symptom in this case with ataxia.

They showed the importance of the magnetic resonance spectroscopy in the diagnosis of tuberculoma as this condition has a particular spectroscopy pattern, and it is very helpful in the differential diagnosis between this and other pathological conditions.

The authors demonstrate the importance of the right therapy. In fact, after few days, the therapy was started; the patient got better till a complete regression of his neurological deficits.

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