

Complete Bilateral Horizontal Gaze Palsy as Initial Manifestation of Multiple Sclerosis

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Abstract

Multiple sclerosis is a chronic inflammatory disease of the central nervous system that commonly affects young adults. Although ocular movement disturbances are commonly encountered in multiple sclerosis, bilateral horizontal gaze palsy is quite rare. We present a case of young woman presented with dizziness, intractable vomiting and visual disturbances for one week, on examination found to have complete bilateral horizontal gaze palsy while vertical gaze was intact. Brain MRI showed a small lesion in posterior part of medial pontine tegmentum bilaterally which was responsible for patient`s symptoms. She received five doses of one-gram methylprednisolone, her symptoms gradually improved over one week after treatment.

Keywords

Multiple sclerosis, bilateral horizontal gaze palsy, paramedian pontine reticular formation, medial longitudinal fasciculus

Case report

A 26-year-old woman, a known case of migraine and sinusitis otherwise healthy, referred from emergency to neurology for evaluation as she presented with acute onset and progressive course of severe dizziness,

frequent vomiting, imbalance and visual disturbances for 1 week. She was seen by ophthalmology 5 days prior to presentation and her examination was normal. Also she was seen by otolaryngology and given vestibular suppressants but without improvement.

On examination she was fully conscious, looked ill and pale, bilateral horizontal gaze was totally impaired with both eyes fixed in midline. Convergence and vertical eye movements were intact. No appreciated horizontal saccades with oculocephalic maneuver. She denied diplopia but there was visual blurring which disappears on closing either eye. She had bilateral peripheral facial weakness, other cranial nerves were intact, she had normal motor power, sensation and coordination in upper and lower limbs but she could not tandem walk and had an unsteady gait.

Brain MRI showed a small lesion in the brain stem involving the posterior part of medial pontine tegmentum bilaterally appeared hyperintense in T2WI and FLAIR sequences, with subtle restricted signal on DWI (Figure 1). Several foci of altered signal in T2WI and FLAIR noted at the periventricular white matter, centrum semiovale bilaterally and left temporal sub-cortical region, some of the lesions are oriented

perpendicular to lateral ventricles (Figure 2). No appreciable post contrast enhancement.

Cerebrospinal fluid was within normal limits but oligoclonal IgG bands were detected in CSF but not in serum. Anti-MOG and Anti-AQP4 antibodies were negative. Other investigations included cervical and thoracic spine MRI, routine blood tests and immunology blood tests were normal.

After MRI results, methylprednisolone 1000 mg/day for 5 days was started. On 2nd day of treatment dizziness and gait improved. On 3rd day, a mild improvement in right eye abduction observed as well as facial weakness. One week after completion of pulse steroid therapy, her eye movements totally returned to normal.

Diagnosis of multiple sclerosis was made based on revised McDonald criteria 2017 (one clinical attack + dissemination in space in MRI+ demonstration of specific oligoclonal bands in CSF).¹ On follow up, she was started on interferon B-1a weekly intramuscular injections.

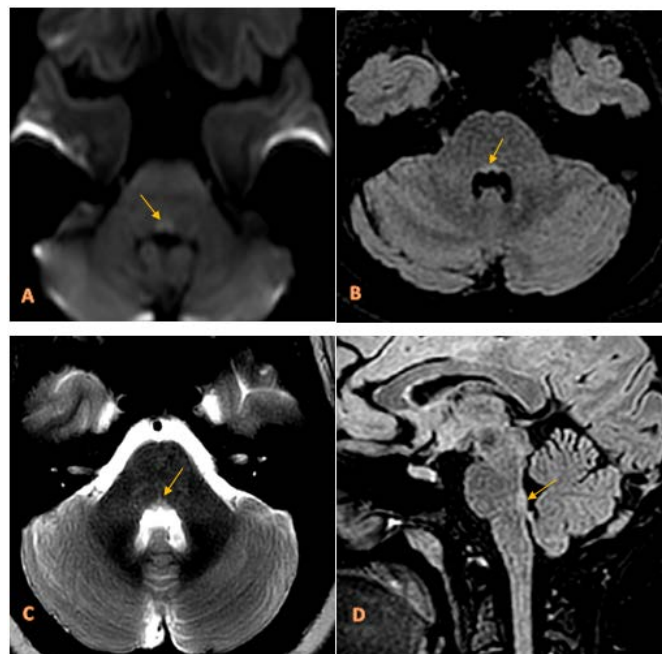


Figure 1: MRI brain shows a lesion of restricted diffusion (arrow) in the posterior part of pons involving

PPRF region bilaterally (A). Same lesion appears hyperintense in FLAIR and T2WI axial cuts passing through the lower pons (B) & (C) respectively and FLAIR sagittal cut (D).

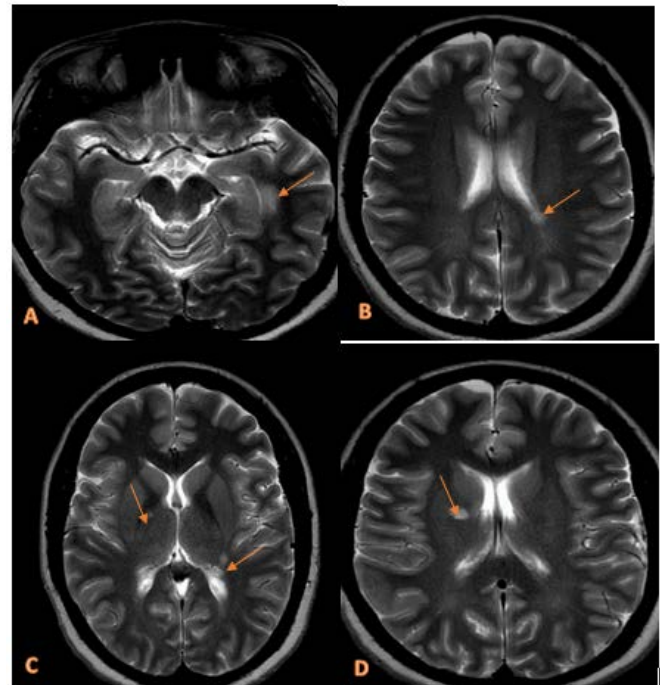


Figure 2: MRI brain T2WI axial cuts show hyperintense lesions (arrows) in left temporal subcortical region (A) and bilateral periventricular white matter (B, C & D)

Discussion

Complete bilateral horizontal gaze paralysis is a rare disorder which has been reported in bilateral pontine lesions due to infarction, hemorrhage, metastasis or demyelinating lesions.² Bilateral horizontal gaze palsy may be caused by bilateral involvement of the paramedian pontine reticular formation (PPRF) and/or bilateral abducens nucleus, or the raphe which is affecting the descending horizontal gaze pathways.³ Lateral gaze is controlled by specific pathways in medial longitudinal fasciculus and nuclei of oculomotor and abducens nerves. The abducens nucleus contains two types of neurons: motor neurons and internuclear neurons. The axons of the internuclear neurons cross to

the contralateral side in the lower pons and, after ascending in the medial longitudinal fasciculus (MLF), synapse in the portion of the oculomotor subnucleus that innervates the medial rectus muscle. Thus, horizontal gaze involves synchronous activation of the lateral rectus muscle of one eye and the medial rectus muscle of the other, via the MLF. This final common pathway for horizontal gaze is controlled by the vestibular, smooth pursuit, optokinetic, and saccadic systems.⁴ The oculocephalic or doll's eye maneuver can be used at bedside to differentiate between pontine lesions from supranuclear lesions in the frontal lobe. Rotating the head passively directly stimulates the abducens nucleus through the vestibuloocular reflex. This will overcome gaze palsies caused by supranuclear lesions but not pontine lesions.⁵

Thus, bilateral horizontal gaze palsy in our patient can be explained by involvement of abducens nuclei and either the MLF or PPRF bilaterally. Bilateral lower motor neuron facial weakness indicated additional involvement of facial nerve nuclei or fasciculi which may label this syndrome as a variant of eight and half syndrome. Classic eight-and-a-half syndrome is caused by a unilateral lesion in the posterior tegmentum of the lower pons involving the paramedian pontine reticular formation and the medial longitudinal fasciculus, as well as the nucleus and/or the fasciculus of the facial nerve. It is characterized by one-and-a-half syndrome (internuclear ophthalmoplegia in addition to horizontal gaze palsy) and an ipsilateral lower motor neuron facial nerve palsy. The affected nuclei may include either a combination of the ipsilateral facial or abducens nerve nuclei with the ipsilateral MLF or a combination of the ipsilateral facial nerve nucleus and ipsilateral PPRF along with the ipsilateral MLF.⁶

An important differential diagnosis needed to rule out was neuromyelitis optica (NMO). In the light of presenting symptoms of nausea and persistent vomiting and the lower pontine lesion which close to the area postrema in medulla, NMO should be suspected. However, absence of anti AQP-4 and anti-MOG antibodies in serum, absence of optic neuritis and absence of long extensive transverse myelitis in cervical and thoracic spine MRI as well as presence of IgG oligoclonal bands in CSF made the possibility of NMO less likely.⁷ Lastly, our patient was diagnosed with multiple sclerosis based on characteristic brain MRI demyelinating plaques, presence of oligoclonal bands in CSF (which were not simultaneously found in serum) and her clinical syndrome that was justified by a lesion in the brain MRI^{1,8}. She showed rapid improvement on methylprednisolone therapy and was started on a DMT on follow-up.

Conclusion

Although ocular movement disorders are common in multiple sclerosis, bilateral horizontal gaze palsy is exceptionally rare. There are few case reports describe patients with similar clinical presentation finally diagnosed with MS. Understanding anatomical distribution of gaze control centers in the brain stem help localizing the lesion which further guide clinicians to suspect the underlying pathology and to order appropriate investigations that eventually lead to proper diagnosis and management.

Abbreviations

MRI: Magnetic Resonance Imaging

FLAIR: Fluid Attenuated Inversion Recovery

DWI: Diffusion Weighted Image

CSF: Cerebrospinal Fluid

MOG: Myelin-Oligodendrocyte Glycoprotein

AQP: Aquaporin

PPRF: Paramedian Pontine Reticular Formation

MLF: Medial Longitudinal Fasciculus

NMO: Neuromyelitis Optica

MS: Multiple Sclerosis

DMT: Disease Modifying Therapy

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