

Short Report

Open Access, Volume 6

Cerebellopontine angle schwannoma with obstructive hydrocephalus causing posterior disc bulge of spinal cord: A rare clinical imageVishal Telrandhe¹; Ruchika Zade²; Sharath Hullumani V^{3*}¹UG Scholar, Department of Paediatric Physiotherapy, Ravi Nair Physiotherapy college, Datta Meghe Institute of Higher Education and Research (DU), Sawangi (Meghe), Wardha, Maharashtra, India.²Resident, Department of Neurophysiotherapy, Ravi Nair Physiotherapy College, Datta Meghe Institute of Higher Education and Research (DU), Sawangi (Meghe), Wardha, India.³Assistant Professor, Department of Paediatric Physiotherapy, Ravi Nair Physiotherapy College, Datta Meghe Institute Higher Education and Research (DU), Sawangi (Meghe), Wardha, India.***Corresponding Author: Sharath Hullumani V**

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Abstract

A 14-year-old girl complained of weakness in both lower limbs and balance issues with hypotonic tone in both lower limbs. Knee and ankle jerk reflexes were absent in both lower limbs, although plantar response extensor bilaterally grade 1 clonus was present in the right lower limb. Investigation were done like Magnetic Resonance Imaging (MRI) of brain which revealed that right CP angle schwannoma size 4.5 x 4 x 3.5 cm causing obstructive hydrocephalus brainstem rotated, fourth ventricle is effaced and the lesion shows perilesional edema with neurofibroma 6 x 5 x 4 cm medial to right parotid, multiple plaque like nodular intraspinal meningiomas seen at anterior epidural space at C5 level. Nodular pachymeningeal enhancement is seen in the cervical level. Nodular enhancement is seen in the right parasellar and suprasellar region. Magnetic Resonance Imaging (MRI) whole spine revealed that posterior disc bulge causing thecal sac indentation in seen involving C3-C4, C4-C5, and C5-C6 levels. Lobulated intensely enhancing Dural based masses are seen in the spinal canal at D3-D4 (32 x 15 mm), D5-D6 (24 x 16 mm), and at the D8-D9 (23 x 6 mm) level.

Received: Dec 16, 2024

Accepted: Jan 10, 2025

Published: Jan 17, 2025

Archived: www.jcimcr.org

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DOI: www.doi.org/10.52768/2766-7820/3428

Introduction

A 14-year-old girl complained of weakness in both lower limbs and balance issues with hypotonic tone in both lower limbs. Knee and ankle jerk reflexes were absent in both lower limbs, although plantar response extensor bilaterally grade 1 clonus was present in the right lower limb. Investigation was done like Magnetic Resonance Imaging (MRI) of brain which revealed that right CP angle schwannoma size 4.5 x 4 x 3.5cm causing obstructive hydrocephalus brainstem rotated, fourth ventricle is effaced and the lesion shows perilesional edema

with neurofibroma 6 x 5 x 4 cm medial to right parotid, multiple plaque like nodular intraspinal meningiomas seen at anterior epidural space at C5 level. Nodular pachymeningeal enhancement is seen in the cervical level. Nodular enhancement is seen in the right parasellar and suprasellar region. Magnetic Resonance Imaging (MRI) whole spine revealed that posterior disc bulge causing thecal sac indentation in seen involving C3-C4, C4-C5, and C5-C6 levels. Lobulated intensely enhancing Dural based masses are seen in the spinal canal at D3-D4 (32 x 15 mm), D5-D6 (24 x 16 mm), and at the D8-D9 (23 x 6 mm) level. The symptomatic accumulation of Cerebrospinal Fluid (CSF) in-

Citation: Telrandhe V, Zade R, Hullumani VS. Cerebellopontine angle schwannoma with obstructive hydrocephalus causing posterior disc bulge of spinal cord: A rare clinical image. *J Clin Images Med Case Rep.* 2025; 6(1): 3428.

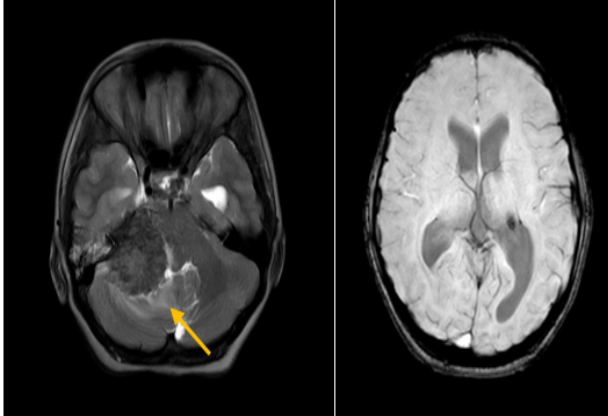


Figure 1: Right CP angle schwannoma size 4.5 x 4 x 3.5 cm causing obstructive hydrocephalus brainstem rotated, fourth ventricle is effaced.

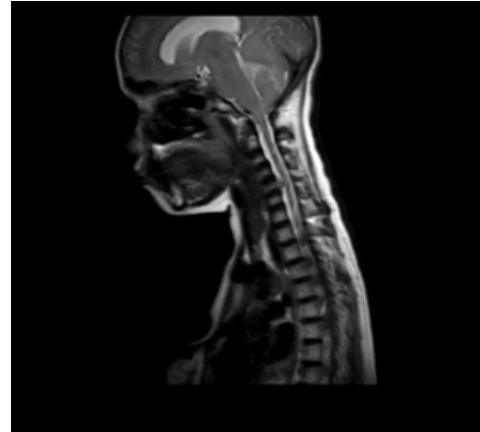


Figure 4: Posterior disc bulge causing thecal sac indentation in seen involving C3-C4, C4-C5, and C5-C6 levels.

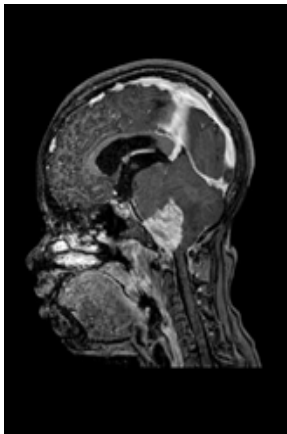


Figure 2: Perilesional edema with neurofibroma 6 x 5 x 4 cm medial to right parotid.



Figure 5: Lobulated intensely enhancing Dural based masses are seen in the spinal canal at D3-D4 (32 x 15 mm), D5-D6 (24 x 16 mm), and at the D8-D9 (23 x 6 mm) level.

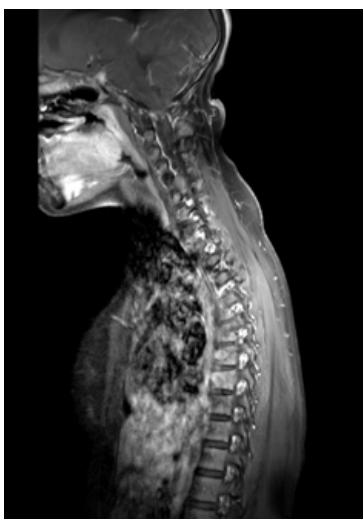


Figure 3: Multiple plaque like nodular intraspinal meningiomas seen at anterior epidural space at C5 level.

side the brain ventricles is known as hydrocephalus [1]. This buildup could be caused by a blockage in the CSF's normal flow, a difficulty with the Pacchionian arachnoid granulations' ability to absorb CSF into the venous system, or an excessive amount of CSF being produced. Early in 1913, Dandy offered the first classifications for hydrocephalus: communicative and non-communicative (obstructive) [2,3]. Since then, numerous additional categories have been presented. Obstructive, communicative, hypersecretory, and Normal Pressure Hydrocephalus (NPH) are the four forms that affect adults. A genetic condition or spinal dysraphism is frequently associated with congenital or developmental hydrocephalus, which is frequently evident at birth [4]. An obstruction in the CSF channels leads to obstructive hydrocephalus [5]. The foramina Monro, the aqueduct of Sylvius, the fourth ventricle, and the foramen magnum are the locations where the obstruction occurs most frequently, however most tumors of a size that is significant can obstruct at any point of the CSF pathways. Ependymoma, subependymal giant cell astrocytoma, choroid plexus papilloma, craniopharyngioma, pituitary adenoma, hypothalamic or optic nerve glioma, hamartoma, and metastatic cancers are a few of the most common

malignancies linked to hydrocephalus. Hydrocephalus development is frequently linked to posterior fossa tumors [6]. Impaired CSF absorption leads to communicating hydrocephalus. Post-hemorrhagic or post-inflammatory alterations are the most frequent causes. One-third of these instances are attributable to subarachnoid bleeding, which prevents CSF absorption at the arachnoid granulations [7].

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