Giant Virchow-Robin Spaces Presenting as Pseudo-Tumour of the Brainstem

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Abstract

We report a rare case of giant Virchow Robin spaces in the mesencephalon in a middle aged male suffering from ataxia, incomplete occulo-motor paresis, neurogenic urinary incontinence and cognitive impairment. A well documented case with its MRI findings and clinical relevance are discussed.

Keywords: Virchow Robin Spaces; Perivascular Spaces; Magnet Resonance Imaging; Ataxia; Tumour

Abbreviations: VRSs: Virchow-Robin Spaces; MRI: Magnetic Resonance Imaging; CT: Computed Tomography; CSF: Cerebrospinal Fluid; MRS: Magnetic Resonance Spectroscopy; NAA: N-Acetyl-Aspartate.

Introduction

Virchow-Robin spaces (VRSs) are perivascular spaces of the brain which are pial-lined interstitial fluid-filled areas that accompany the penetrating arteries and arterioles. They penetrate the cerebral substance for a variable depth and appear as cystic structures, round or tubular in form. VRSs are a complex entity and are commonly seen in MRI scans in all age groups. Giant forms of VRSs are a rare entity and may mimic cystic neoplasia and may lead to mass effect. We present a well-documented case of giant VRSs of the mesencephalon with its MRI features and the relevant clinical manifestations.

Case Report

A middle aged male presented to the emergency department of our hospital with symptoms of progressive ataxia and loss of memory of recent onset. He also had episodes of vertigo and urinary incontinence for a year. The clinical examination revealed incomplete paresis of the right third cranial nerve and left sided ataxia. The initial CT scan showed no evidence of acute stroke or intracranial haemorrhages. Cystic changes were observed around the right sided Thalamus and in the midbrain with secondary extensively dilated third and lateral ventricle systems suggestive of an obstructive hydrocephalus (Figure 1).

Figure 1: NECT showing cystic mass in the mesencephalon extending on the right sided thalamus with mass effect and secondary dilatation of the third and lateral ventricle.
MRI scan of the brain showed bizarre, purely cystic mass of the mesencephalon extending to the right thalamus (Figures 2a-2d) with no significant oedema of the adjacent brain parenchyma. No diffusion restriction or accompanying haemorrhages within the lesion were seen. MRI also showed significant dilatation of the third and lateral ventricles. No enhancement was noted (Figures 3a & 3b). The initial CT scan and MRI scan were suggestive of a probable tumour such as a Germinoma or an epidermoid cyst, or an Arteriovenous malformation. But the absence of diffusion restriction and contrast enhancement led to a probable diagnosis of unusual giant VRSs.

Results of Cerebrospinal fluid examinations, the EEG and the Duplex examination were within normal ranges. Funduscopic examinations of the eyes show no abnormal findings suggestive of raised intracranial pressure. The patient was treated conservatively without ventricle shunting/drainage which were reserved as an optional
surgical management in case of persistence or deterioration of the symptoms. On follow-up after 3 months on conservative medical treatment, the patient showed considerable improvement of symptoms. Follow-up CT and MRI scans showed no changes in the findings.

Discussion

Virchow-Robin spaces are named after Rudolf Virchow, a German pathologist and Charles Philippe Robin, a French anatomist [1,2]. Perivascular spaces or Virchow-Robin spaces are visualized as small CSF equivalent foci in all the sequences in MRI and follow the pathway of penetrating vessels from the subarachnoid space to various depths of penetration in the underlying brain substance. Small VRSs of 2-3 mm size are a usual finding on routine MRI scans in any age group [3]. There is evidence suggesting that the VRSs may represent a lymphatic system draining the deep substance of the brain [4].

Several studies have shown that the increase in number of VRSs has a strong correlation with age, hypertension, dementia or other causes of brain atrophy [4,5]. It is also postulated that VRSs have a close association with neuropsychiatric disorders [5], multiple sclerosis [6] and mild traumatic brain injury [7]. Technically heavy weight T2 weighted sequence or MRI imaging with higher Tesla will delineate more VRSs. They should neither show diffusion-restrictions nor enhance on contrast sequences [4,8]. There are three types of VRSs as per their location [6,9,10]. Type 1 VRSs are seen along the course of lenticulostriate arteries located in basal ganglia whereas type 2 VRSs track towards the white matter from the convexity along the path of medullary arteries and veins and are located in both the hemispheres of the brain as well as in the cerebellum. The third types of VRSs are located along the course of the collicular and accessory collicular arteries in the ponto-mesencephalic junctions.

Giant Type I lesions should be differentiated from lacunar infarctions. These are unilateral compared to VRSs and show diffusion restriction in early stages as well as contrast enhancement [10]. Cystic periventricular leukomalacia in children and multiple sclerosis can mimic type II VRSs. Leukomalacia shows loss of white matter and ventricular dilatation whereas multiple sclerosis are oriented perpendicular to the corpus callosum which helps to differentiate them from VRSs [11]. Type II giant VRSs can mimic mucopolysaccharidosis which are inherited metabolic disorders of metabolism of glycosaminoglycans, leading to accumulation of toxic intracellular substrate [12]. In mucopolysaccharidosis, the brain changes are dominantly seen in white matter, corpus callosum and basal ganglia with increase in signal intensities in Flair and T2 weighted MRI sequences due to dysmyelination. In contrast, giant VRSs are signal hypointense in Flair sequences.

When VRSs become enlarged, they are known as giant VRSs or “cavernous dilatation”. Giant VRSs type 3 are a rare presentation as in our case [6,10]. They are located in the mesencephalothalamic region in the territory of mesencephalothalamo-thalamic (collicular) artery as mentioned above [6,13]. Obstruction of CSF drainage pathways could be a major factor in the pathogenesis of giant VRSs, although the exact nature of the obstruction is
In conclusion, VRSs should be considered in differential diagnosis in patients with large cystic lesions without enhancement located mainly in the mesencephalon. This can, as in our patient, help avoid unnecessary neurosurgical interventions.

References


