Susceptibility Weighted Imaging: A Novel Method to Determine the Etiology of Aqueduct Stenosis

Chanabasappa Chavadi1  Keerthiraj Bele1  Anand Venugopal1  Santosh Rai1

1Department of Radiodiagnosis, Kasturba Medical College, Manipal University, Mangalore, India

Address for correspondence  Chanabasappa Chavadi, DNB, Flat No. C-1-13, 3rd Floor, K.M.C Staff Quarters, Light House Hill Road, Mangalore 575001, India (e-mail: chavadidoc@gmail.com).


Abstract

The stenosis of aqueduct of Sylvius (AS) is a very common cause of obstruction to cerebrospinal fluid. Multiple etiologies are proposed for this condition. Because treatment is specific for correctable disorder, assessment of etiology gains importance. A case of pediatric hydrocephalus was diagnosed with stenosis of AS on magnetic resonance imaging (MRI). Susceptibility-weighted imaging (SWI) demonstrated blooming in the distal aqueduct and lateral ventricle, which was not seen on routine MRI sequences. The findings suggest that old hemorrhage is a cause of chemical arachnoiditis and adhesions causing aqueduct stenosis and hydrocephalus. To our knowledge literature is very scarce, wherein SWI is being used to confirm blood products as a cause of aqueduct stenosis; hence SWI should be routine protocol in imaging of pediatric hydrocephalus. Etiology, clinical presentation, role of imaging, and, in particular, SWI in evaluation of aqueductal stenosis is discussed.

Keywords

► susceptibility weighted imaging
► aqueductal stenosis
► magnetic resonance imaging
► hydrocephalus

Introduction

Aqueduct of Sylvius (AS) is the narrowest segment of the cerebrospinal fluid (CSF) pathway and is the most common site of intraventricular obstruction to the flow of CSF. Stenosis of the AS is responsible for 6 to 66% of cases of hydrocephalus in children and 5 to 49% in adults.1,2 There are two peaks of distribution for age: one in the first year of life, the other in the adolescence. Susceptibility-weighted imaging (SWI) is a unique gradient-echo magnetic resonance imaging (MRI), which is flow-compensated and uses magnitude and phase information. Phase postprocessing accentuates the magnetic properties of extravascular blood products and produces high contrast, when compared with other conventional MRI sequences.3 SWI is being extensively used in trauma, vascular pathologies and tumors causing microbleeds, multiple sclerosis, and stroke. In our case, SWI confirmed the cause of stenosis of AS and the case is being discussed. Etiology, pathogenesis, and imaging of aqueduct stenosis are summarized.

Case Report

An 8-month-old child presented with increased head size, developmental delay, and an episode of seizure. No history of birth asphyxia was found. Laboratory investigations were normal. MRI revealed enlargement of the lateral and third ventricle (Fig. 1a, b) and dilation of proximal aqueduct with narrowing at the distal segment secondary to adhesions (Fig. 1c, d). The temporal horn diameter was 28 mm and ventricle size index (VSI) was 50% suggestive of significant hydrocephalus. There was narrow angle of the frontal horn and effacement of the sulci at vertex (Fig. 1b). SWI demonstrated blooming at the distal AS (Fig. 1f, g), confirming the etiology as adhesions due to extravascular blood products. Blooming due to hemorrhage was also noted occipital horn (Fig. 1h) with dilation of the suprapineal recess and absence of normal flow void in the fourth ventricle (Fig. 1e).

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Discussion

The AS is a tubular channel situated in the dorsal midbrain, connecting the third and the fourth ventricles. Stenosis of AS may be tumoral or may be nontumoral which is referred to as intrinsic aqueductal stenosis. In about three-quarters of patients with aqueductal stenosis, the etiology of the disorder is not known. In the remaining cases, it can be attributed to different causes, namely genetic, infection–inflammation, prematurity-related intraventricular hemorrhage, in which case aqueductal stenosis may develop in the acute phase, secondary to direct obstruction of aqueduct by blood, or in a chronic phase, secondary to organization of clots. Other causes include intoxications and deficiencies, congenital malformations, and vascular malformations. Russell classified nontumoral aqueduct stenosis by histopathologic studies into four subtypes: (1) stenosis without gliosis of the surrounding tissue, proposed to be due to “developmental errors”; (2) forking caused by incomplete fusion of the median fissure; (3) membranous obstruction; (4) gliosis usually a reaction to irritant agents, such as hemorrhage, infection, or toxic agents, and is often part of a widespread ependymitis of the ventricles.

In the first year of life, the clinical presentation usually consists of rapid enlargement of the circumference of the head. Other features include retarded psychomotor development, cognitive dysfunction, temporary headache, urinary incontinence, seizures, and growth retardation. Sometimes, decompensation may follow minor head injury, febrile affections, or subarachnoid hemorrhage. Visual disturbances are related to papilledema, and chronic compression of the optic chiasm is exerted by an enlarged third ventricle. Endocrine manifestations occur in 10% of adolescents and adults, due to chronic compression of the hypothalamo-hypophyseal axis by the enlarged anterior third ventricle.

Imaging is the cornerstone for diagnosis, intrauterine diagnosis by ultrasonography primarily by excluding other causes of hydrocephalus in presence of triventriculomegaly with macrocephaly. Antenatal MRI may allow diagnosis of aqueductal stenosis by showing hydrocephalus in association with the normal fourth ventricle and absence of aqueductal lumen. MRI is investigation of choice for evaluation of hydrocephalus and aqueductal stenosis. Anatomical deformations of bulging of the third ventricle are better evaluated. The optimal view for the evaluation of anatomical details is a midline sagittal section. On T1-weighted image (WI) sequences, the aqueduct has the same signal intensity as CSF, whereas on T2 WI, a hypointense signal is seen as a result of CSF flow. This “flow void” extending inferiorly into the superior aspect of the fourth ventricle has been considered as an important sign for diagnosis of patency of aqueduct, with a few limitations. If the aqueduct remains visible, the obstruction is...
presumably caused by a membrane. Ultra-thin three-dimensional constructive steady-state sequences (CISS) sequence due to inherent high CSF-to-brain contrast provides exquisite detail of the ventricular system and permits better identification of aqueductal obstruction by membranes, webs, hypoplasia or cysts, and tumors. The absence of CSF flow at the aqueductal level in flow-sensitive phase contrast MRI is a reliable, reproducible, and rapid method, which supports the diagnosis of stenosis of AS in patients with clinical and/or radiologic suggestion of obstructive hydrocephalus.

In cases of long-standing hydrocephalus, CSF pulsations against the thinnest segment of the ventricular walls may determine focal enlargement of some portions of the ventricular system. MRI is helpful by detecting and correctly explaining pulsion diverticula, third ventricle bulging in the chiasmatic and interpeduncular cisterns, and cystic expansion of the suprapineal recess, subependymal dissection, and spontaneous ventriculocisternostomies.

**Conclusion**

Triventriculomegaly is usually accepted as a sign of stenosis of AS. The etiology of stenosis of AS may be obscure on routine MRI. Our observation suggests that detection of blooming on SWI in distal aqueduct with flaring of the proximal aqueduct confirms the causative factor of stenosis of AS, being secondary to intraventricular hemorrhage. To our knowledge literature is very scarce, wherein SWI is being used to confirm blood products as a cause of aqueduct stenosis. With varied etiology and limitation of routine MRI sequence, SWI is a novel method, and should be a part of routine MRI protocol for imaging in congenital and pediatric hydrocephalus.

**Conflict of Interest**

The authors declare that there is no actual or potential conflict of interest in relation to this article. The manuscript has been read and approved by all the authors, the requirements for authorship as stated earlier in this document have been met, and that each author believes that the manuscript represents honest work.

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