Vein of Galen Malformations: Case Report & Review

Snigdha Goyal¹, Sunder Goyal², Isha Saini², Sarita Bishnoi³

¹Department of Pathology, Dr. RML Postgraduate Institute of Medical Sciences & Research, New Delhi, India
²Department of Surgery, Kalpana Chawla Government Medical College, Karnal, India
³Department of Radiology, Kalpana Chawla Government Medical College, Karnal, India

Email: *goyal.sunder@yahoo.in

Received 1 July 2015; accepted 18 July 2015; published 23 July 2015

Copyright © 2015 by authors and OALib. This work is licensed under the Creative Commons Attribution International License (CC BY).

Abstract

Vein of Galen aneurysmal malformations (VGAMs) are rare congenital abnormalities that can cause severe morbidity and mortality. VGAMs present in different way from vein of Galen aneurysmal dilations (VGADs). The VGAMS have a parenchymal arteriovenous malformation that drains through the vein of Galen whereas VGADs may present with intracranial hemorrhage. In neonates VGAMs result in high-output cardiac failure. Surgery offers little improvement, with fatal outcomes in 80% to 100% of cases.

Keywords

Vein of Galen Aneurysmal Malformation, Vein of Galen Aneurysmal Dilatation

Subject Areas: Clinical Trials, Public Health

1. Introduction

Vein of Galen aneurysmal malformations (VGAMs) and vein of Galen aneurysmal dilations (VGADs), are the most common arteriovenous malformations in infants and fetuses [1]. VGAM consisted of a jumbled mass of dilated vessels supplied by an enlarged artery [2]. The malformation increases greatly in size with age without any clear mechanism [2]. Dilation of the great cerebral vein of Galen is a secondary result of the force of arterial blood either directly from an artery via an arteriovenous fistula or by way of a tributary vein that receives the blood directly from an artery [1] [2]. There is usually a venous anomaly downstream from the draining vein that, together with the high blood flow into the great cerebral vein of Galen causes its dilation [3]. The right side cardiac chambers and pulmonary arteries also develop mild to severe dilation [4]. Mortality rate is about 80% to 100% of cases and open surgery offers little improvement [5]. Recently, endovascular management may im-

*Corresponding author.

prove results in infants and children. However, mortality ranges from 23% to 75% and morbidity from 21% to 88% in neonates [5]. We report a case of vein of Galen malformation in child of 2 years age.

2. Case Report

A 2 years old boy presented to the pediatrics outpatient department with a complaint of enlargement of head size. There was history delayed mile stones and also of seizure once. General and cardiovascular examinations, as well as routine biochemical analysis were within normal limits. There was no evidence of skin lesions to suggest capillary malformation neither there was any limb hypertrophy. There was no family history of skin lesions, limb hypertrophy or vascular malformations. Skull ultrasound (USG) done outside reported a heterogeneous, predominantly hypoechoic lesion in region posterior to third ventricle with dilated lateral and third ventricle and normal-sized fourth ventricle suggestive of obstructive hydrocephalus. Slow flow was also detected within the lesion. Computerized tomography skull done confirmed the findings of USG, and showed heterogeneous intensity lesion in the pineal region causing mass effect on aqueduct. CECT angiography and venography were also performed which well delineated the lesion (Figure 1). Patient was advised endovascular embolization. Patient lost the follow up.

![Figure 1. CECT showing vein of Galen malformation.](image)
3. Discussion

The vein of Galen (great cerebral vein or great vein of Galen) is a short vascular trunk formed by the union of the two internal cerebral veins and the basal veins of Rosenthal. Vein of Galen aneurysmal malformation (VGAM) occurs during 6 - 11 weeks of fetal life due to teratogenic effect. Aneurysmal dilatation of vein of Galen and arteriovenous shunting of blood occurs due to persistence of embryonic pros encephalic vein of Markowski. VGAM can be associated with capillary malformation-arteriovenous malformation (CM-AVM). It is an autosomal dominant disorder due to mutation in RASA1 gene [6].

Several proposed classification systems have been used to describe malformations of the vein of Galen. Five patterns of Galenic arteriovenous malformations have been described:

Pattern 1—Many vessels, including anterior cerebral arteries, thalamic perforating arteries, and superior cerebellar arteries discharge into the vein of Galen [7].

Pattern 2—A single posterior choroidal artery drains into the vein of Galen [7].

Pattern 3—One or both posterior choroidal and one or both anterior cerebral arteries drain directly into the Galenic system [7].

Pattern 4—An angiomatous network of posterior choroidal and thalamic perforating arteries enters the vein of Galen directly [7].

Pattern 5—A high flow arteriovenous malformation in the right inferior frontal lobe drains via the inferior sagittal sinus and pericallosal vein into the Vein of Galen [3].

Small arteriovenous shunts causes progressive neurological impairment where as larger arteriovenous shunts are linked with greater hemodynamic effects and thus result in earlier symptom [8].

Malformations frequently cause cardiac failure, cranial bruits (pattern 1), hydrocephaly, and subarachnoid hemorrhage in neonates. The heart failure is due to the size of the arteriovenous shunt that can steal 80% or more of the cardiac output, with large volumes of blood under high pressure returning to the right heart and pulmonary circulation and sinus venous atrialseptal defects [3] [4]. It is the most common cause of death in such patients [9].

There can be extremely rare non-developmental syndromes like superior vena cava syndrome (SVCS), and thrombosis of the lateral sinus, superior sagittal sinus, internal jugular vein, or of the Great Cerebral Vein of Galen itself.

In a child with unexplained heart failure, malformed vein of Galen should be suspected. On examination there can be dilated facial veins along with cranial bruits. Ultrasound, Doppler, Computerized tomography (CECT) and magnetic imaging resonance (MRI) are useful radiological diagnostic tools for vein of Galen malformations [3]. To improve outcomes in survivors, transcranial sonography and fetal MR imaging can help in antenatal diagnosis [10]-[12]. Many cases are diagnosed only during autopsy as congestive heart failure occurs very early [7].

VGAM should be differentiated from arachnoid cyst, cavum vergae and porencephalic cyst. There is no evidence of intralesional flow in these lesions.

Color-flow imaging and pulsed Doppler ultrasonography are noninvasive tools to assess anatomical and pathophysiological information about cardiac hemodynamics and intracranial blood flow [13].

Treatment depends on the structure of the malformation [3]. A ventriculoperitoneal shunt may be required if there is hydrocephalous. To reduce the blood flow into the vein, the feeding fistulous arteries into the Vein of Galen must be blocked [7].

As structure of malformation is complicated, so, open surgery is very difficult. Mostly endovascular procedures are very useful [14]. In the highest-risk neonates with VGAMs and cardiac failure, endovascular therapy can reduce the mortality nearly 100% [15]. With the help of catheters drugs, balloons, or coils can be delivered at the site of the malformation to stop blood flow through the vein [14]. During catheterization of vein, guide wire can damage the wall of the vein can be damaged and, in some cases, may dislodge the emboli [4]. Endovascular procedures offer safe, successful treatment if cardiac failure cannot be controlled with medical therapy [15].

Another treatment option is radio surgery in which blood vessel is damaged with the help of focused beam but it is very slow process and may takes months to years [16]. Seizures usually are managed with antiepileptic medications [16].

Most noted complication is intracranial hemorrhages [17]. In about half of the patients malformations are untreatable and these patients mostly die in the neonatal period or in early infancy [4]. Incidence of mortality is about 77% in untreated cases [18]. The mortality rate is about 39.4% even after surgery [18].
4. Conclusion
Vein of Galen malformations are rare malformations. Clinicians should be given awareness about this condition for proper management and for better prognosis. With proper selection of cases, good results can be acquired.

References


