Vein of Galen Aneurysmal Malformation: An Updated Review

Li-Rong Cao¹ Chun-Quan Cai²

¹ Graduate College of Tianjin Medical University, Tianjin, China

² Department of Neurosurgery, Tianjin Children's Hospital, Tianjin, China

Address for correspondence Chun-Quan Cai, PhD, Department of Neurosurgery, Tianjin Children's Hospital, Beichen District 300134, Tianjin, China (e-mail: cqcns6@126.com).

J Pediatr Neurol 2019;17:45–56.

Abstract

Vein of Galen aneurysmal malformation (VGAM) is a rare intracranial vascular anomaly in the choroidal arteries characterized by the persistence of the embryonic median prosencephalic vein of Markowski (MProsV). It is common in children with high mortality and disability rates. VGAM-associated clinical presentations include heart failure, hydrocephalus, and neurological symptoms, which make it difficult for diagnosis. Simultaneously, the complex VGAM vascular architecture, characterized by a wide interindividual variability, makes the management of this condition challenging to clinicians. Thus, timely diagnosis and treatment of VGAM are crucial. In this review article, we elucidated the latest research progresses of VGAM in the aspects of the pathogenesis, classification, clinical features, imaging, management, and prognosis based on recent relevant literatures. It is beneficial to raise the awareness of VGAM and improve the level of management and treatment.

Keywords

- ▶ vein of Galen aneurysmal malformation
- ▶ vein of Galen
- ▶ review

Introduction

Vein of Galen aneurysmal malformation (VGAM) is a rare congenital intracranial arteriovenous deformation resulting from an anomalous fistulous connection between the primitive choroidal vessels and the most proximal part of the median prosencephalic vein of Markowski (MProsV). The incidence has been reported as 1:25,000.¹ It accounts for 1% of cerebral vascular malformations and 30% of pediatric vascular malformations.² Clinically, VGAM is commonly diagnosed during the neonatal period and early childhood according to the symptoms of congestive heart failure and intracranial hypertension.

Steinheil first described this abnormal intracranial arteriovenous connection as “varix aneurysm” in 1895.³ Since then, there were various terms used to depict this condition, including “vein of Galen malformation,” “vein of Galen aneurysmal malformation,” “arteriovenous aneurysms of the vein of Galen,” and “aneurysms of the vein of Galen.” However, these medical terms are misnomers because the malformation is actually caused by MProsV, rather than the vein of Galen. The true VGAM was reported by Boldrey and Miller in 1949.⁴ The term

of vein of Galen “aneurysms” was first proposed by Jaeger et al in 1937.⁵ In current understanding, the word, “aneurysms,” is also an description error. This tendency is arbitrary since the word is derived from the Greek *aneurynein*, that is, to dilate. Therefore, the use of a “venous aneurysms” is etymologically legitimate rather than a misnomer.⁶

Endovascular embolization is generally considered to be the first-line therapeutic option. A meta-analysis performed by Yan et al⁷ revealed that the mortality rate of embolized patients decreased from 17 to 12% from 1980s to 2000s. But the management is still challenging. This article reviewed the pathogenesis, classification, clinical manifestations, imaging of VGAM, and summarized the key points of endovascular embolization to improve the rate of treatment and prognosis for VGAM.

Pathogenesis

The persistent arteriovenous shunting from primitive choroidal vessels into the MProsV is the predominant pathogenesis of VGAM, which occurs at the 6th to 11th week of gestation, after development of the circle of Willis (▶ Fig. 1). The posterior part

received

October 16, 2017

accepted after revision

January 23, 2018

published online

April 8, 2018

Copyright © 2019 by Georg Thieme
Verlag KG, Stuttgart · New York

DOI <https://doi.org/10.1055/s-0038-1635108>.
ISSN 1304-2580.

Fig. 1 The formation of VGAM.²⁵ (A) Normal vessel structure between the pericallosal arteries and the PCA is present in embryonic period. (B) The abnormal high-output arteriovenous connection contributes to aneurysmal dilation of the MProsV, which leads to the formation of VGAM. This connection drains into the SS and subsequently into T. In addition, it also bloods into the FS and then into the AT when the SS is absent, hypogenetic, or stenotic. Reproduced with permission from Mortazavi et al.²⁵ AT, accessory torcula; FS, falcine sinus; MProsV, median prosencephalic vein; MRI, magnetic resonance imaging; PCA, posterior cerebral arteries; SS, straight sinus; T, trocula; VGAM, vein of Galen aneurysmal malformation.

of the MProsV persists as the vein of Galen, while the anterior portion regresses in parallel with the formation of the internal cerebral veins. The remnants of the MProsV then join the internal cerebral veins to form the vein of Galen, which may cause the existence of connection between the VGAM and the deep venous system. Raybaud and Strother⁸ identified the dilated venous structure as the MProsV at first. The arteriovenous connection between primitive choroidal vessels and MProsV leads to hemodynamic abnormalities, which results in the obstruction of embryologic vein regression and the development of vein of Galen. Subsequently, the high-output arteriovenous connection contributes to venous hypertension and aneurysmal dilation of MProsV. Besides, stenosis of dural sinuses is another cause of the dilatation, resulting from the rapid and turbulent flow caused by the arteriovenous connection.

The drainage of the malformation is toward the straight sinus, and it also drains into sagittal sinus through the

persistent falcine sinus, in this case the straight sinus is absent, hypogenetic or stenotic.⁹ The cases of VGAM with persistent falcine sinus have been reported previously by us¹⁰ (► Fig. 2). The arteriovenous deformation is supplied by a variety of arterial feeding vessels including subependymal network originating from the posterior circle of Willis, thalamoperforating arteries, choroidal arteries, and the limbic arterial arch.⁴ It is located in the cistern of velum interpositum as well as the quadrigeminal cistern, and there is no connection between the malformation and the deep venous system, which has been considered traditionally. However, this theory has been challenged in recent literature reports.^{11–13} In these studies, the drainage of the malformation into the deep venous system has been confirmed by follow-up imaging, including internal cerebral veins, straight sinus, falcine sinus, basal vein of Rosenthal, and inferior sagittal sinus.

The genetic susceptibility to VGAM is still unclear, while there are some potentially causative genes reported in

Fig. 2 Cranial sagittal MRI in a patient with VGAM reveals a dilated MProsV (short arrow) connected with a persistent falcine sinus (long arrow) blooded into the sagittal sinus and the absence of straight sinus on T1WI (A) and T2WI (B). MProsV, median prosencephalic vein; MRI, magnetic resonance imaging; VGAM, vein of Galen aneurysmal malformation; WI, weighted imaging.

association with this condition. Revencu et al¹⁴ identified *RASA1* gene mutation in two patients with VGAM. Chida et al¹⁵ reported one variant of *ACVRL1* in a case with VGAM. Tsutsumi et al¹⁶ published a patient with VGAM carrying a mutation in the *ENG* gene encoding endoglin; this patient presented with a family history of hereditary hemorrhagic telangiectasia and the authors hypothesized the gene might be etiologically related to VGAM. Additionally, Xu et al¹⁷ reported a familial VGAM in an adult female patient with VGAM, who had a history of a stillbirth pregnancy caused by VGAM. However, the relationship between genetic factors and VGAM needs to be further investigated.

Classification

Vein of Galen Aneurysmal Malformation

The first classification of VGAM was proposed by Litvak et al¹⁸ in 1960. There are two most widely accepted classifications described by Lasjaunias and Yasargil, respectively (►Table 1). Based on the age at presentation and angioarchitecture of the fistulae, Lasjaunias et al¹⁹ divided VGAM into two types: choroidal type and mural type (►Fig. 3).²⁰ The choroidal type is the most common type accounting for 56 to 76% of all VGAMs and is common in males.²¹ It mainly occurs

in neonates,²² characterized by congestive heart failure due to a high flow fistula. In choroidal type, multiple arterial feeders like a nidus shunt into the anterior aspect of MProSV. The feeder vessels mostly come from the choroidal arteries, involving the anterior and posterior choroidal, anterior cerebral, as well as quadrigeminal and thalamoperforating arteries occasionally.²³ In mural type, one or more arteriovenous fistulae (AVF) shunt into the inferolateral margin in the wall of MProSV with lower flow. The arterial feeders arise from unilateral or bilateral quadrigeminal arteries and posterior choroidal arteries. Clinically, it is common in infants and children associated with the symptoms of macrocephaly, hydrocephalus, and developmental delay. Due to slow flow and few fistulae, it has a better prognosis, and thrombosis is more common in this type compared with choroidal type.

Yasargil²⁴ dichotomized VGAM into four types according to whether the malformation is a pure AVF (types I–III) or an arteriovenous malformation (AVM) with or without associated AVF (types IVA–C), and the exact origin of the feeding arteries. Type I is a pure AVF between posterior cerebral artery (P4 and its branches), anterior or posterior pericallosal arteries, and the vein of Galen. Type II is the fistulous connection between the thalamoperforators (basilar and P1 segment) and the vein of Galen. Type III is a mixture of type I and II, which is the commonest type. Type IV comprises three subtypes. Type IVA is similar to type IVB. Both types IVA and IVB are aneurysmal dilation of the vein of Galen caused by shunting from an adjacent thalamic and mesencephalic AVM, respectively. Type IVC is a thalamomesencephalic or mesodiencephalic plexiform malformation along with an adjacent and separate cisternal AVF to the vein of Galen.^{25,26} Though the Yasargil's classification is classic and widely recognized, there are several deficiencies. The clinical presentation and outcome are not involved in the Yasargil's classification.

Table 1 Classification of VGAM

Lasjaunias' classification	Yasargil's classification
Mural type	Type I
Choroidal type	Type II
	Type III
	Type IV (type IVA, type IVB, type IVC)

Abbreviation: VGAM, vein of Galen aneurysmal malformation.

Fig. 3 (A) Mural type of VGAM. MR angiography displays an AVF directly into the wall of the venous pouch. Note the drainage of the malformation into the stenosis straight sinus; (B) Choroidal type of VGAM. In comparison with mural type, MR angiography of choroidal type shows an extensive arterial network like a nidus centered in the region of the quadrigeminal plate cistern (closed arrow), flowing into the malformation and primitive falxine sinus (open arrow).²⁰ Reproduced with permission from Triffo et al.²⁰ AVF, arteriovenous fistula; MR, magnetic resonance; VGAM, vein of Galen aneurysmal malformation.

Vein of Galen Aneurysmal Dilatation

Vein of Galen aneurysmal dilatation (VGAD) refers to the dilated vein of Galen arising from the drainage of an AVM into the true vein of Galen rather than the precursor of the vein of Galen. It belongs to the group of dura AVM. Different from VGAM, it is a real malformation of Galen vein and contacted with the deep venous system. The arterial supply comprises the internal carotid artery, meningeal branches of internal and external carotid artery, normal sinus, the drainage from Galen vein into brain parenchyma, and mesencephalic or diencephalic AVM.²⁷ Based on the source of abnormal blood, VGAD can be differentiated into (1) solid type, (2) varicose type, and (3) dura mater type. According to the shunt location of VGAD, Iizuka et al²⁸ proposed a new anatomical classification, involving (1) ventricular type, (2) cisternal type, and (3) parenchymal type.

Vein of Galen Varices

Vein of Galen varices is the varicose dilatation of the Galen vein without an arteriovenous shunt. There are two types of the vein of Galen varices in children. The first type is a transient dilatation of the Galen vein accompanied with heart failure caused by other origins. This type is asymptomatic and cardiac function is improved along with the disappearance of the dilatation. The second type occurs when flowing a complete hemisphere into the deep venous system. It is also asymptomatic, but the venous ischemia disorders may occur as a result of insufficient venous drainage.²⁵

Clinical Manifestations

Clinical presentations vary with the age of onset and vascular architecture. The main manifestations include congestive heart failure, hydrocephalus, and neurological symptoms. In 1964, Gold et al²⁹ proposed a clinical classification based on the age of onset, including the group of neonates, infants as well as children and adults. In neonates, the predominant clinical manifestation is represented by serious congestive heart failure. The prognosis is extremely poor with high mortality, and choroidal type is common in this group. During infancy, clinical presentations mainly consist of hydrocephalus, macrocephaly, seizures, developmental delay, and mild heart failure. The mechanisms causing hydrocephalus include hypertension, impaired cerebrospinal fluid reabsorption, and compression of mesencephalic aqueduct. Mural type usually occurs in this group. In children and young adults, the symptoms consist of headache, hydrocephalus, seizure, neural dysfunction, and subarachnoid hemorrhage. Both choroidal and mural types are observed in this group. In addition, there are some rare clinical manifestations reported in the literature, which include galactosemia,³⁰ abnormal movement (e.g., chronic asymmetric tremor),³¹ a pulsatile right-sided neck swelling due to tortuous carotid artery and bilateral dilatation of neck veins,³² jugular bulb stenosis,³³ dilated scalp veins, and tonsillar prolapse. But the relationship between these clinical manifestation and VGAM needs to be further confirmed.

Additionally, Lasjaunias et al²² recommended the Bicêtre neonatal evaluation score system, which takes into account neurological, cardiac, respiratory, renal, and hepatic functions (►Table 2). It helps to estimate the comprehensive symptoms of patients with VGAM and assists with the therapeutic decision. A score of less than 8 indicates a poor outcome needing expectant treatment. An emergency endovascular treatment is advised relating to a score of 8 to 12. The embolization can be delayed until 5 months of age associated with a score of 13 to 21. Besides, the evaluation of the neurocognitive status could be performed using the Denver test and the Brunet-Lézine test.³⁴

Imaging

VGAM is one of the diseases mostly diagnosed prenatally with the proportion of approximately 40%.³⁵ However, a retrospective analysis of 25 patients with VGAM conducted by Geibprasert et al³⁶ showed there was no relevance between early prenatal diagnosis and the follow-up outcome. But it assisted in the decision of therapy plan and facilities. Prenatal diagnosis is usually made during the third trimester of pregnancy by ultrasonography according to the visualization of an anechogenic or hypoechogenic round or ovoid structure with regular borders located in the midline of the posterior part of the third ventricle. Enough dilatation of the malformation is required if it can be detected on the first- or even second-trimester ultrasound³⁶ and the conformation of the antenatal diagnosis is usually performed by color Doppler imaging and magnetic resonance imaging (MRI). There is an antenatal VGAM classification including isolated VGAM, when VGAM is the only manifestation and associated VGAM, when VGAM is associated with cardiac and neurological abnormalities.³⁷ However, it might be challenged in the future research. Termination of pregnancy should be performed when fetal hydrops and serious heart failure occur.

The chest radiograph shows cardiomegaly resulting from congestive heart failure. Brain radiograph may display a rim of calcification within the wall of the aneurysmal sac, which is considered to be a poor indicator associated with the tendency of subsequent thrombosis.²⁵ Cranial computed tomography (CT) usually displays a spherical mass located in the quadrigeminal cistern or the cistern of velum interpositum. Dilated ventricular system, periventricular white matter hypodensities, and diffuse cerebral atrophy are usually associated MRI findings (►Fig. 4). MRI has the advantage to accurately demonstrate the location of the lesion, vascular architecture, and status of vessel drainage in the VGAM malformation (►Fig. 5); furthermore, the position and identity of major arterial trunks, primary branches as well as secondary branches feeding the fistula are better recognized on brain MRI than on cranial CT.³⁸ Additionally, parenchymal changes including focal encephalomalacia and diffuse brain volume loss can also be visible in CT and MRI, which are considered to be predictors of poor outcome.³⁶ Angiography (including MR angiography and CT angiography) remains the gold standard for the diagnosis. It has advantages in revealing the site, size,

Table 2 Bicêtre neonatal evaluation score²²

Points	Cardiac function	Cerebral function	Respiratory function	Hepatic function	Renal function
5	Normal	Normal	Normal	—	—
4	Overload, no medical treatment	Subclinical, isolated EEG abnormalities	Tachypnea, finishes bottle	—	—
3	Failure, stable with medical treatment	Nonconvulsive intermittent neurological signs	Tachypnea, does not finish bottle	No hepatomegaly, normal hepatic function	Normal
2	Failure, not stable with medical treatment	Isolated convulsion	Assisted ventilation, normal saturation $\text{FIO}_2 < 25\%$	Hepatomegaly, normal hepatic function	Transient anuria
1	Ventilation necessary	Seizures	Assisted ventilation, normal saturation $\text{FIO}_2 > 25\%$	Moderate or transient hepatic insufficiency	Unstable diuresis with treatment
0	Resistant to medical therapy	Permanent neurological signs	Assisted ventilation, desaturation	Abnormal coagulation, elevated enzymes	Anuria

Abbreviations: EEG, electroencephalogram; FIO_2 , fractional inspired oxygen.

Maximal score = 5 (cardiac) + 5 (cerebral) + 5 (respiratory) + 3 (hepatic) + 3 (renal) = 21.

Source: Reproduced with permission from Lasjaunias et al.²²

Fig. 4 Axial CT scan of brain in a patient with VGAM reveals a large homogeneous, well-defined spherical mass associated with dilated third and lateral ventricles and hydrocephalus because of the compression of the aqueduct, as well as periventricular white matter hypodensities (arrows). CT, computed tomography; VGAM, vein of Galen aneurysmal malformation.

morphology, and abnormal flow dynamics of vessels (► Fig. 6). Similarly, it better shows the small feeders supplying the fistula, as well as the dynamic aspects of the venous drainage into the normal brain and hemodynamics of the arteriovenous shunt.^{26,35} Strategies for the decision of treatment can be made according to the evidence on angiography. In future applications, four-dimensional flow MRI is probably useful in the management and monitoring intracranial hemodynamic alteration during embolizations, to estimate the efficacy and

risk of reperfusion complications.³⁹ Wu et al⁴⁰ indicated that it assisted in detecting the arteriovenous shunt flow, cerebral vessel inflow, and cerebral flow redistribution after embolizations during staged embolizations of six patients with VGAM.

The spontaneous thrombosis of VGAM is rare and the underlying causal mechanism is not entirely understood. There are multifactors described including the alteration of hemodynamics (slow flow, increased pressure, or turbulence), vascular spasm, vascular proliferation, and regressive atherosclerotic changes in the vessels.^{41,42} The ventricular shunting and the use of contrast media for angiography are also considered to be correlated with the thrombosis of VGAM.⁴¹ The imaging investigation can well demonstrate the features of thrombosis. On CT, it reveals heterogeneous signal associated with calcified capsule occasionally. Contrast-enhanced CT shows a central thrombus and peripheral circulating blood along the wall of the sac, called “target-sign.”²⁶ MRI of the thrombosis VGAM displays extracellular methemoglobin in the central part and hemosiderin in the peripheral part of lesion.⁴³

Management

The purpose of therapy is various according to the age of onset. In newborns, the goal of therapy is mainly the correction of heart failure, while in older children the aim is to improve the neurological symptoms.⁴ It is advised that the therapy outcome is to reverse the pathophysiologic effects caused by the arteriovenous deformation rather than to obtain a complete normal morphologic appearance.⁴⁴ The management of VGAM includes a combination of surgical approaches, endovascular treatments, radiosurgery, and conservative treatments. Endovascular embolization is the preferred treatment and others are considered to be the complementary options, when embolization is limited.

Fig. 5 Cerebral MRI in axial of a patient with VGAM shows a well-defined midline round mass within the pineal region accompanied with the dilatation of lateral and third ventricles on T1WI (A) and T2WI (B). MRI, magnetic resonance imaging; VGAM, vein of Galen aneurysmal malformation; WI, weighted imaging.

Fig. 6 (A) MR angiography shows the dilated aneurysmal mass with some feeding vessels which drains into the straight sinus. (B) Vertebral artery angiography displays the dilated median prosencephalic vein bleeding into the straight sinus, which belongs to the mural type of VGAM. (C) Lateral view reveals the stenosis of straight sinus (arrow) draining directly into the superior sagittal sinus. MR, magnetic resonance; VGAM, vein of Galen aneurysmal malformation.

Surgery and Radiosurgery

Though the method of surgery is widely used, the complete resection and surgical clipping of the vessels are still challenging with higher morbidity and mortality. The problem is associated with surgery including the deep location of the lesion, high-flow shunt, complex vascular architec-

ture, and poor tolerance in infants. Therefore, surgery is usually considered as a complementary approach when embolization is failed. Additionally, ventricular shunting may exacerbate the cerebral venous hypertension and brain ischemia due to the alteration of venous drainage. Hence, ventricular shunting should be performed after

Fig. 7 Carotid angiogram of a patient with VGAM displays a dilated aneurysmal malformation (A). Post-embolization angiogram shows the complete occlusion of the malformation using Guglielmi coils in anteroposterior (B) and lateral (C) views. VGAM, vein of Galen aneurysmal malformation.

embolization.^{22,26} Similarly, the approach of Gamma knife surgery is also a complementary alternative since it does not reduce the size of the lesion immediately. Stereotactic radiotherapy may have an effect on some patients with VGAM, but its effect is still not clear. Besides, there is controversy about the lesion to the development of brain.

Endovascular Therapy

The first case treated by embolization was reported in 1950 by Davis.⁴⁵ With the development of endovascular treatment, the mortality rate has dropped from 60–82% to 11%.⁴⁶ The goal of endovascular therapy is to reduce intracranial shunting or even occlude all the fistulae (→Fig. 7). In fact, total obliteration is not necessary because a reduction in one-third to half of the shunt is enough to correct the clinical symptoms.⁴⁷ Recently, a meta-analysis showed the predominant complications of endovascular treatment consisted of cerebral vessel perforation, hemorrhage, cerebral ischemia, hydrocephalus, and leg ischemia.⁷ The prevention of complications depends on proper super selective embolization, selection of embolic material and exact technique, as well as preservation of normal cerebral deep venous drainage. Additionally, the existence of normal drainage between deep venous system and VGAM should be considered before embolization in case of the occurrence of adverse effects to the normal cerebral vessel after endovascular management. Although it is difficult to be seen through angiograms or noninvasive imaging presumably due to technical factors and hemodynamics, the drainage is a critical element of the treatment planning.¹¹

There are two options of embolization including transarterial and transvenous approach, of which the transarterial route is preferred. Umbilical artery approach can be used for embolization in the early neonatal period, and the femoral artery is an alternative route when the umbilical artery is not available. As for transvenous approach, there is a controversy regarding the outcome of transvenous embolization. In previous literature, it was considered to

be not feasible due to increased morbidity. In recent article, Orlov et al⁴⁸ considered it was a safe and effective salvage method for VGAM with restricted transarterial access. Besides, Meila et al⁴⁹ described a combined transvenous and transarterial approach, called kissing microcatheter technique. It was considered to be a safe method with lower complication rate and the overall outcome can be improved, particularly in the neonate with congestive heart failure. The embolic agents comprise coils, microballoons, N-butyl-cyanoacrylate (NBCA), Onyx liquid, and ethylene vinyl alcohol copolymer. In choroidal type of VGAM, Onyx is recommended because of the advantage of more definitive and stable occlusion. In mural type, NBCA is a preferred embolic material due to the properties of quick precipitation and adhesion to vessel.⁴⁸ In addition, Joo et al⁴² first reported a patient with a mural-type VGAM who was successfully treated using the Micro Vascular Plug system, which may be a potential available option.

The staged embolization is recommended to minimize the occurrence of complications such as a normal perfusion pressure breakthrough phenomenon or a multiple venous thrombosis.¹¹ The interval is usually 3 to 6 months.^{4,23} There is a new classification scoring system for the decision of treatment proposed by Mortazavi et al,²⁵ which combines arterial feeders with clinical symptoms and age (→Table 3). Embolization is not urgent and one stage of embolization is suggested for patients with a score of 0 to 1. Urgent embolization and multiple stages of treatment are recommended when the score is 2 points. Embolization in multiple stages or palliative treatment is advised for patients with a score of 3.

The time of endovascular management depends on the clinical symptom and is preferably at the age of 5 to 6 months for maximum safety and efficacy of treatment.²⁶ However, symptoms related to the possible progression of the heart failure and the potential development of hydrocephalus need to be closely followed up.⁵⁰ The decision of management can be evaluated by the Bicêtre scoring system proposed by

Table 3 New classification system proposed by Mortazavi et al²⁵

	Score	
Parameter	0	1
Arterial feeders	Any feeders other than P1–2, thalamoperforators, choroidal, basilar	Any of the following feeders: P1–2, thalamoperforators, choroidal, basilar
Clinical symptoms	No heart failure	Heart failure
Age	≥ 5 months	< 5 months

Source: Reproduced with permission from Mortazavi et al.²⁵

Alvarez et al⁴ (► **Table 2**). A score of more than 12 suggests the clinical symptom is stable and it is better to prolong therapy until the age of 5 to 6 months. Endovascular embolization is recommended when a score is between 8 and 12. There is no indication for embolization associated with a score of less than 8.

VGAD is the pathological dilatation of Galen vein due to the stenosis or obstruction of the outflow. The degree of dilatation is varied with the extent of stenosis. So, the treatment of VGAD with high flow is mainly planned embolization or stereotactic radiotherapy. For low level VGAD, intravascular embolization is the major approach of management.²⁷ It is associated with a good prognosis in comparison with VGAM because VGAM has high morbidity and mortality due to refractory high-output heart failure and neurological damage.⁵¹ As for the vein of Galen varices, it has lower hemorrhage risk and better outcome. The adverse consequence may occur as a result of an inappropriate surgical or interventional management. Conservative treatment is recommended for patients with stable symptoms,⁵² and close follow-up with imaging investigations is essential.

Prognosis

Without treatment, more than 90% of patients with VGAM die in the neonatal and infant period. A meta-analysis about the outcome and complications of endovascular embolization for VGAM revealed the proportion of the good outcome increased from 49 to 70% from 1980s to 2000s.⁷ But the morbidity and mortality are still very high. There are some factors that correlated with the clinical outcome involving embolization technology, clinical presentation (multiorgan failure and neurological symptoms), and imaging findings (extensive encephalomalacia, arterial steal, parenchymal calcification, and high-flow multifeeder shunts).³⁶ Additionally, the choroidal type of VGAM has worse prognosis than mural type. Agarwal et al⁵³ retrospectively analyzed 36 patients with VGAM and concluded that jugular atresia was significantly associated with poor outcome. A systematic review conducted by Yan et al⁵⁴ showed the outcome was good in the patient with an early spontaneous thrombosis, while the outcome was poor in the late stages of the VGAM with spontaneous thrombosis.

The main sequela of VGAM is represented by permanent neurological deficit with intellectual disability and epilepsy. The evaluation of therapy outcome in patients with VGAM can be performed by the system of Bicêtre admission and outcome score proposed by Lasjaunias et al²² (► **Table 4**). This scoring system can help to estimate the effect of clinical treatment, contribute to correct the therapy approaches, and improve the level of management of VGAM.

In this study, we performed a literature retrospective of relevant articles in the recent 15 years using the database of PubMed (<https://www.ncbi.nlm.nih.gov/pubmed>) with the key words “vein of Galen aneurysmal malformation” or “VGAM.” The detail information was collected, including age, sex, types of VGAM, clinical feature, treatment, and outcome (► **Tables 5 and 6**). A total of 43 patients (36 children and 7 adults) were enrolled. We found that the mortality of VGAM was still high. The prognosis of VGAM in adult was better than children, and the mural type was associated with a good outcome compared with choroid type.

Table 4 Bicetre admission and outcome score²²

Score	Condition
5	Normal (N)
4	Minimal non-neurological symptoms, not treated (MS), and/or asymptomatic enlargement of the cardiac silhouette
3	Transient neurological symptoms, not treated (TNS), and/or asymptomatic cardiac overload under treatment
2	Permanent minor neurological symptoms, mental retardation of up to 20%, nonpermanent neurological symptoms under treatment (MNS), normal school with support, and/or cardiac failure stabilized with treatment
1	Severe neurological symptoms, mental retardation of more than 20% (SNS), specialized school and/or cardiac failure unstable despite treatment
0	Death(D)

Note: Does not apply to neonates.
Source: Reproduced with permission from Lasjaunias et al.²²

Table 5 Literature review of clinical outcomes after treatment of VGAM in children

Authors	Age/sex	Type	Clinical features	Treatment	Outcome
Pop et al ⁵⁵	11 mo/M	Mural	IHC, DD	Femoral TA Embolization	Normal
Gupta et al ²⁶	1y/M	Mural	MC, ASD occipital bruit	Embolization (TA)	Normal
Triffo et al ²⁰	Neonate/M	Mural	PD, CHF, RD	Embolization (TA), GKRS	Normal
Jones et al ⁵⁶	Neonate/F	Mural	PD, Mild CHF	Embolization (TA)	Normal
Ellis et al ⁵⁷	5 mo/M	Mural	MC, cranial bruit	Embolization	Normal
Ellis et al ⁵⁷	Neonate/F	Mural	PD, CHF	Embolization	Normal
Joo et al ⁴²	4 mo/F	Mural	IHC, DFV, NSC, lethargy	Embolization (TA)	Normal
Gupta et al ²⁶	3 y/M	Mural	H	Embolization (TV and TA)	Died
Filippi et al ³⁰	Neonate/M	Mural	PD, H, CD, HF, HM	Embolization (TA)	Died
Orlov et al ⁴⁸	Neonate/M	Mural	PD, CHF	Embolization (TA)	Sz, mild DD
Ellis et al ⁵⁷	21 mo/F	Mural	MC, cranial bruit	Embolization	LSH, ANP, DVA
Heuer et al ⁵⁰	Neonate/M	Mural	RVH, RVD	None	Improved
Cai et al ⁵⁸	15 mo/M	Mural	Sz, DFV, IHC	Embolization (TV)	Improved
Kong we al ⁵⁹	Neonate/M	Mural	PD, H, IVH, V	Embolization (TA)	Improved
Orlov et al ⁴⁸	Neonate/M	Choroidal	PD, no symptom	Embolization (TA)	Normal
Tsutsumi et al ¹⁶	Neonate/M	Choroidal	CD, HM, HF, cardiac bruit,	Embolization (TA)	Normal
Ellis et al ⁵⁷	Neonate/M	Choroidal	CHF, cranial bruit	Embolization	Normal
Heuer et al ⁵⁰	5 d/M	Choroidal	HF	Embolization	Normal
Heuer et al ⁵⁰	7 d/F	Choroidal	HF	Embolization	Normal
Heuer et al ⁵⁰	31 mo/M	Choroidal	Mild LVH hemorrhage	Embolization	Normal
Heuer et al ⁵⁰	6 mo/M	Choroidal	None	Embolization	Normal
Heuer et al ⁵⁰	Neonate/M	Choroidal	Mild HF	Embolization	Normal
Heuer et al ⁵⁰	Neonate/F	Choroidal	HF	Embolization	Died
Heuer et al ⁵⁰	6 d/M	Choroidal	HF, RVH, RVD	Embolization	Died
Heuer et al ⁵⁰	Neonate/M	Choroidal	HF, PDA, Sz, ischemia	Embolization	Died
Jones et al ⁵⁶	Neonate/M	Choroidal	PD, CHF, ASD	Embolization (TA)	Died
Komiyama et al ⁶⁰	Neonate/M	Choroidal	CHF	Femoral TA Embolization	Died
Orlov et al ⁴⁸	Neonate/M	Choroidal	H	Embolization (TA)	Mild DD
Heuer et al ⁵⁰	Neonate/M	Choroidal	RVD, PFO, Sz, ischemia,	Embolization	Severe MR
Heuer et al ⁵⁰	Neonate/F	Choroidal	HF, Sz, ischemia,	Embolization	Hemiparesis, mild MR
Heuer et al ⁵⁰	14 mo/M	Choroidal	Sz, ischemia	Embolization	Hemiparesis, autistic, epilepsy in remission
Heuer et al ⁵⁰	45 d/M	Choroidal	PFO, hemorrhage	Embolization	H, mild motor delay
Demartini et al ⁴⁷	Neonate/M	Choroidal	HF, PC, DD, V	Embolization (TA and TV)	Improved
Fourie et al ⁶¹	Neonate/M	Choroidal	PC, CHF, cerebral bruit	Embolization (TA and TV)	Improved
Jagadeesan et al ⁶²	Neonate/M	Mixed type	ASD, PFO	Embolization (TV and TA)	Dysconjugate gaze
Levrier et al ¹³	11 y/F	Mixed type	MC, MR, H	Embolization (TV and TA)	Improved

Abbreviations: ANP, abducens nerve palsy; ASD, atrial septal defect; CD, cardiomegaly; CHF, congestive heart failure; DFV, dilate scalp and facial veins; DVA, decreased visual acuity; F, female; GKRS, Gamma Knife radiosurgery; H, hydrocephalus; HF, heart failure; HM, hepatomegaly; IHC, increasing head circumference; IVH, intraventricular hemorrhage; LSH, left sided hemiparesis; LVH, left ventricular hypertrophy; M, male; MC, macrocrania; MR, mental retardation; NSC, nasal and sinus congestion; PC, perioral cyanosis; DD, delay development; PD, prenatal diagnosis; PDA, patent ductus arteriosus; PFO, patent foramen ovale; RD, respiratory distress; RVD, right ventricular dilation; RVH, right ventricular hypertrophy; Sz, seizure; TA, transarterial; TV, transvenous; V, vomiting; VAGM, vein of Galen aneurysmal malformation.

Table 6 Literature review of clinical outcomes after treatment of VGAM in adult

Authors	Age/sex	Type	Clinical features	Treatment	Outcome
Gupta et al ²⁶	21 y/F	Mural	H, SAH, neck stiffness	Embolization (TA)	Normal
Levrier et al ¹³	40 y/M	Mural	Pituitary dysfunction	Femoral TA Embolization	Normal
Ellis et al ⁵⁷	24 y/M	Mural	Face pain, dementia	Embolization	Normal
Xu et al ¹⁷	44 y/F	Choroidal	Dizziness, vertigo	None	Normal
Gupta et al ²⁶	28 y/M	Choroidal	HA, IVH	Embolization (TV and TA)	Died
Porzionato et al ⁶³	36 y/M	Choroidal	IVH, coma	VS	Died
Triffo et al ²⁰	27 y/M	Choroidal	RTH, VM	GKRS	DFNT

Abbreviations: DFNT, dysmetria on finger-to-nose test; F, female; GKRS, Gamma Knife radiosurgery; H, hydrocephalus; HA, headache; IVH, intraventricular hemorrhage; M, male; RTH, right thalamic hemorrhage; SAH, subarachnoid hemorrhage; TA, transarterial; TV, transvenous; VGAM, vein of Galen aneurysmal malformation; VM, ventriculomegaly; VS, ventricle-peritoneal shunt.

Conclusion

VGAM is a rare congenital intracranial arteriovenous deformation associated with high mortality. To increase and ameliorate the VGAM diagnosis and therapy, it is crucial to fully understand the pathogenesis, clinical classification, and features of this condition. Imaging plays an important role in the visualization of vascular architecture and also in assisting the treatment with embolization procedures. The evaluation of the time and stages for endovascular treatment based on the score systems will be beneficial to decrease the complication rate, morbidity, and mortality.

Conflict of Interest
None declared.

Acknowledgment
This work was supported by the key project of the Tianjin Health and Family Planning Commission Natural Science Foundation (grant number: 2015KR12); the key project of the Tianjin Health Care Professionals (grant number: 16KG166); and the National Natural Science Foundation of China (grant number: 81771589).

References

- Stephan S, Rodesch G, Elolf E, Wiemann D, Jorch G. Vein of Galen aneurysmal malformations: an ultrasonographic incidental finding—a case report. *Case Rep Pediatr* 2012;2012:824284
- Kośla K, Majos M, Polgaj M, Antosik-Biernacka A, Stefańczyk L, Majos A. Prenatal diagnosis of a vein of Galen aneurysmal malformation with MR imaging – report of two cases. *Pol J Radiol* 2013;78(04):88–92
- Dandy WE. Arteriovenous aneurysm of the brain. *Arch Surg* 1982;117(02):190–243
- Boldrey E, Miller ER. Arteriovenous fistula (aneurysm) of the great cerebral vein (of Galen) and the circle of Willis; report on two patients treated by ligation. *Arch Neur Psych* 1949;62(06):778–783
- Jaeger JR, Forbes RP, Dandy WE. Bilateral congenital cerebral arteriovenous communication aneurysm. *Trans Am Neurol Assoc* 1937;63:173–176

- Gailloud P, O’Riordan DP, Burger I, et al. Diagnosis and management of vein of Galen aneurysmal malformations. *J Perinatol* 2005;25(08):542–551
- Yan J, Wen J, Gopaul R, Zhang CY, Xiao SW. Outcome and complications of endovascular embolization for vein of Galen malformations: a systematic review and meta-analysis. *J Neurosurg* 2015;123(04):872–890
- Raybaud CA, Strother CM. Persisting abnormal embryonic vessels in intracranial arteriovenous malformations. *Acta Radiol Suppl* 1986;369(369):136–138
- Chauhan U, Tullu M, Muranjan M, Lahiri K. Thalamic haemorrhage: a rare presentation of vein of Galen aneurysmal malformation in infancy. *N Z Med J* 2003;116(1186):U687
- Cai CQ, Zhang QJ, Yang WD, Wang CX, Shen CH. Neuroimages of persistent falcine sinus in children. *World J Pediatr* 2009;5(01):63–64
- Iizuka Y, Kakihara T, Suzuki M, Komura S, Azusawa H. Endovascular remodeling technique for vein of Galen aneurysmal malformations—angiographic confirmation of a connection between the median prosencephalic vein and the deep venous system. *J Neurosurg Pediatr* 2008;1(01):75–78
- Gailloud P, O’rordan DP, Burger I, Lehmann CU. Confirmation of communication between deep venous drainage and the vein of Galen after treatment of a vein of Galen aneurysmal malformation in an infant presenting with severe pulmonary hypertension. *Am J Neuroradiol* 2006;27(02):317–320
- Levrier O, Gailloud PH, Souei M, Manera L, Brunel H, Raybaud C. Normal galenic drainage of the deep cerebral venous system in two cases of vein of Galen aneurysmal malformation. *Childs Nerv Syst* 2004;20(02):91–97, discussion 98–99
- Revenu N, Boon LM, Mulliken JB, et al. Parkes Weber syndrome, vein of Galen aneurysmal malformation, and other fast-flow vascular anomalies are caused by RASA1 mutations. *Hum Mutat* 2008;29(07):959–965
- Chida A, Shintani M, Wakamatsu H, et al. ACVRL1 gene variant in a patient with vein of Galen aneurysmal malformation. *J Pediatr Genet* 2013;2(04):181–189
- Tsutsumi Y, Kosaki R, Itoh Y, et al. Vein of Galen aneurysmal malformation associated with an endoglin gene mutation. *Pediatrics* 2011;128(05):e1307–e1310
- Xu DS, Usman AA, Hurley MC, Eddleman CS, Bendok BR. Adult presentation of a familial-associated vein of Galen aneurysmal malformation: case report. *Neurosurgery* 2010;67(06):E1845–E1851, discussion 1851
- Litvak J, Yahr MD, Ransohoff J. Aneurysms of the great vein of Galen and midline cerebral arteriovenous anomalies. *J Neurosurg* 1960;17:945–954

- 19 Lasjaunias P, Ter Brugge K, Lopez Ibor L, et al. The role of dural anomalies in vein of Galen aneurysms: report of six cases and review of the literature. *Am J Neuroradiol* 1987;8(02):185–192
- 20 Triffo WJ, Bourland JD, Couture DE, McMullen KP, Tatter SB, Morris PP. Definitive treatment of vein of Galen aneurysmal malformation with stereotactic radiosurgery. *J Neurosurg* 2014;120(01):120–125
- 21 Pearl M, Gomez J, Gregg L, Gailloud P. Endovascular management of vein of Galen aneurysmal malformations. Influence of the normal venous drainage on the choice of a treatment strategy. *Childs Nerv Syst* 2010;26(10):1367–1379
- 22 Lasjaunias PL, Chng SM, Sachet M, Alvarez H, Rodesch G, Garcia-Monaco R. The management of vein of Galen aneurysmal malformations. *Neurosurgery* 2006;59(05, Suppl 3):S184–S194, discussion S3–S13
- 23 Puvabanditsin S, Mehta R, Palomares K, et al. Vein of Galen malformation in a neonate: a case report and review of endovascular management. *World J Clin Pediatr* 2017;6(01):103–109
- 24 Yasargil MG. *Microneurosurgery* IIIB. New York: Thieme Medical Publishers; 1988:323–357
- 25 Mortazavi MM, Griessenauer CJ, Foreman P, et al. Vein of Galen aneurysmal malformations: critical analysis of the literature with proposal of a new classification system. *J Neurosurg Pediatr* 2013;12(03):293–306
- 26 Gupta AK, Varma DR. Vein of Galen malformations: review. *reviewNeurol India* 2004;52(01):43–53
- 27 Tang F, Zhang L, Huang X, Li XW, Zhang Y, Zhai XW. Effects of intravascular embolization operation on adult VGAD. *Eur Rev Med Pharmacol Sci* 2016;20(09):1829–1833
- 28 Iizuka Y, Kakihara T, Yoshimura N, Kimizuka T, Sumi Y, Kaneko K. Anatomical subcategorization for pediatric vein of Galen aneurysmal dilatation. A report of three cases. *Neuroradiol J* 2007;20(05):551–561
- 29 Gold A, Ransohoff J, Carter S. Vein of Galen malformation. *Acta Neurol Scand Suppl* 1964;40(Suppl 11):1–31
- 30 Filippi L, Pezzati M, Poggi C, Pasquini E. Vein of Galen aneurysmal malformation and galactosemia in a neonate: a previously unreported association. *Paediatr Anaesth* 2007;17(12):1221–1223
- 31 de Souza A, Fernandes YS, Bhatkar SR, Bhonsle SK. Chronic asymmetric tremor and levodopa-responsive parkinsonism due to a vein of Galen aneurysmal malformation. *Parkinsonism Relat Disord* 2017;43:122–123
- 32 Agarwal A, Firdouse M, De Almeida CL, Mondal T. Distinctive neck swelling in a patient with a vein of Galen arteriovenous malformation. *J Ultrasound* 2014;18(02):197–200
- 33 Saliou G, Dirks P, Slater LA, Krings T. Is jugular bulb stenosis in vein of Galen aneurysmal malformation associated with bony remodeling of the jugular foramina? *J Neurosurg Pediatr* 2016;18(01):92–96
- 34 Frankenburg WK, Dodds J, Archer P, Shapiro H, Bresnick B. The Denver II: a major revision and restandardization of the Denver Developmental Screening Test. *Pediatrics* 1992;89(01):91–97
- 35 Bohiltea RE, Turcan N, Mihalea C, et al. Ultrasound prenatal diagnosis and emergency interventional radiologic therapy of Galen aneurysmal malformation in a newborn. *Maedica (Buchar)* 2016;11(04):334–340
- 36 Geibprasert S, Krings T, Armstrong D, Terbrugge KG, Raybaud CA. Predicting factors for the follow-up outcome and management decisions in vein of Galen aneurysmal malformations. *Childs Nerv Syst* 2010;26(01):35–46
- 37 Deloison B, Chalouhi GE, Sonigo P, et al. Hidden mortality of prenatally diagnosed vein of Galen aneurysmal malformation: retrospective study and review of the literature. *Ultrasound Obstet Gynecol* 2012;40(06):652–658
- 38 Cai C, Zhang Q, Shen C. Vein of Galen malformations-report of 2 cases and literature review. *Front Med China* 2008;2(03):317–322
- 39 Schnell S, Wu C, Ansari SA. Four-dimensional MRI flow examinations in cerebral and extracerebral vessels - ready for clinical routine? *Curr Opin Neurol* 2016;29(04):419–428
- 40 Wu C, Schoeneman SE, Kuhn R, et al. Complex alterations of intracranial 4D hemodynamics in vein of Galen aneurysmal malformations during staged endovascular embolization. *Operative Neurosurgery* 2016;12(03):1
- 41 Irfan M, Lohman B, McKinney AM. Confirmation of T1-bright vein of Galen aneurysm spontaneous thrombosis by subtraction magnetic resonance venography: a case report. *Acta Radiol* 2009;50(07):812–815
- 42 Joo W, Mercier P, Kheradmand S, et al. Vein of Galen malformation treated with the Micro Vascular Plug system: case report. *J Neurosurg Pediatr* 2017;19(06):729–733
- 43 Mahmoodi R, Habibi Z, Heidari V, Nejat F. Spontaneous regression and complete disappearance of the vein of Galen aneurysmal malformation. *Childs Nerv Syst* 2016;32(04):593–598
- 44 Pearl M, Gregg L, Gandhi D. Cerebral venous development in relation to developmental venous anomalies and vein of Galen aneurysmal malformations. *Semin Ultrasound CT MR* 2011;32(03):252–263
- 45 Davis PB. Thrombosis of the internal cerebral veins of Galen; report of two cases involving adults. *J Am Osteopath Assoc* 1950;50(01):21–22
- 46 Berenstein A, Fifi JT, Niimi Y, et al. Vein of Galen malformations in neonates: new management paradigms for improving outcomes. *Neurosurgery* 2012;70(05):1207–1213, discussion 1213–1214
- 47 Demartini Z Jr, Dos Santos ML, Koppe GL, Cardoso-Demartini AA. Sinus thrombosis after endovascular treatment of vein of Galen aneurysmal malformation. *Pediatr Neurosurg* 2017;52(02):136–139
- 48 Orlov K, Gorbatykh A, Berestov V, et al. Superselective transvenous embolization with Onyx and n-BCA for vein of Galen aneurysmal malformations with restricted transarterial access: safety, efficacy, and technical aspects. *Childs Nerv Syst* 2017;33(11):2003–2010
- 49 Meila D, Hannak R, Feldkamp A, et al. Vein of Galen aneurysmal malformation: combined transvenous and transarterial method using a “kissing microcatheter technique”. *Neuroradiology* 2012;54(01):51–59
- 50 Heuer GG, Gabel B, Beslow LA, et al. Diagnosis and treatment of vein of Galen aneurysmal malformations. *Childs Nerv Syst* 2010;26(07):879–887
- 51 Sharma J, Toubas P, Sklar T, Pole-Spellman J. Spontaneous resolution of marked dilatation of cerebral dural-venous system in newborn presenting with fetal hydrops. *Indian J Pediatr* 2010;77(03):313–315
- 52 Xu H, Zheng J, Chen G, Chen J. Hemispheric developmental venous anomaly draining into a vein of Galen varix: a case report and literature review. *Chin J Neurosurg* 2017;3(02):1
- 53 Agarwal H, Sebastian LJ, Gaikwad SB, Garg A, Mishra NK. Vein of Galen aneurysmal malformation-clinical and angiographic spectrum with management perspective: an institutional experience. *J Neurointerv Surg* 2017;9(02):159–164
- 54 Yan J, Gopaul R, Wen J, Li XS, Tang JF. The natural progression of VGAMs and the need for urgent medical attention: a systematic review and meta-analysis. *J Neurointerv Surg* 2017;9(06):564–570
- 55 Pop R, Manisor M, Wolff V, Kehrli P, Marescaux C, Beaujeux R. Flow control using Scepter™ balloons for Onyx embolization of a vein of Galen aneurysmal malformation. *Childs Nerv Syst* 2015;31(01):135–140
- 56 Jones BV, Ball WS, Tomsick TA, Millard J, Crone KR. Vein of Galen aneurysmal malformation: diagnosis and treatment of 13 children with extended clinical follow-up. *Am J Neuroradiol* 2002;23(10):1717–1724
- 57 Ellis JA, Orr L, Li PC, Anderson RC, Feldstein NA, Meyers PM. Cognitive and functional status after vein of Galen aneurysmal malformation endovascular occlusion. *World J Radiol* 2012;4(03):83–89

- 58 Cai MJ, Ma LT, Yang M, Pan L. Vein of Galen aneurysmal malformation: strategy transarterial embolization using coils alone. *Neurol India* 2012;60(05):559–560
- 59 Kong JC, Cheng KM, Cheung YL, Chan CM. Transarterial embolisation with Guglielmi detachable coils in an infant with a vein of Galen aneurysmal malformation. *Hong Kong Med J* 2012;18(05):435–438
- 60 Komiya M, Honnda Y, Matsusaka Y, Morikawa T, Kitano S, Sakamoto H. Cerebral diagnostic and therapeutic angiography for neonatal arteriovenous fistulas. *Interv Neuroradiol* 2004;10 (Suppl 1):39–42
- 61 Fourie PA, Potze FP, Hay N, Du Toit E, Lippert MM, Mennen J. Transcranial placement of an amplatzer device to control intractable cardiac failure in an infant with a vein of Galen anomaly. A case report. *Interv Neuroradiol* 2010;16(02):191–197
- 62 Jagadeesan BD, Zacharatos H, Nascene DR, Grande AW, Guillaume DJ, Tummala RP. Endovascular management of a vein of Galen aneurysmal malformation in an infant with challenging femoral arterial access. *J Neurosurg Pediatr* 2016;18(02):231–234
- 63 Porzionato A, Macchi V, Parenti A, De Caro R. Vein of Galen aneurysm: anatomical study of an adult autopsy case. *Clin Anat* 2004;17(06):458–462