Pediatric Holohemispheric Developmental Venous Anomaly: Definitive characterization by 3D Susceptibility Weighted Magnetic Resonance Angiography

Michael A. Casey¹, Sourabh Lahoti², Ajeet Gordhan²*

¹. Advocate Bromenn Neurosurgery, Normal, Illinois, USA
². Department of Neurointerventional Radiology, St Joseph Medical Center, Bloomington, Illinois, USA

* Correspondence: Ajeet Gordhan MD, St Joseph Medical Center, Department of Neurointerventional Radiology, 2200 East Washington Street, Bloomington, IL, 61701, USA (agordhan@hotmail.com)

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ABSTRACT

We present a case of an incidentally discovered holohemispheric developmental venous anomaly (DVA) in a 12 year old, conclusively characterized by 3D T2* multi-echo sequence susceptibility weighted angiographic imaging (SWAN). For the evaluation of head trauma, abnormal right intraparenchymal and periventricular vascularity was identified by a non contrast head CT scan. Conventional MRI sequences revealed prominent veins with findings suspicious of a DVA. A definitive diagnosis was made by identifying angiographic features typical for DVA by augmented susceptibility weighted angiographic imaging. Using this sequence the entire hemispheric extent of the anomaly without complicating features was definitively characterized, negating the need for a catheter based angiographic study. A holohemispheric DVA in a child to our knowledge has not been previously described.

CASE REPORT

A 12 year old boy was kicked in the head at school and presented with a two day history of headache. His neurological examination was unremarkable. A non-enhanced CT (NECT) of the head showed an abnormal periventricular vascular lesion on the right cerebral hemisphere (Fig. 1). Conventional multi sequence gadolinium enhanced MRI of the brain study poorly identified a DVA (Fig. 2, 3, 4 and 5). Conventional time of flight MR angiography (MRA) and time resolved imaging of contrast kinetic (TRICKS) confirmed a vascular anomaly without differentiating an arteriovenous malformation (AVM) from a DVA (Fig. 6). A T2* weighted multi echo 3D volumetric susceptibility angiographic sequence (SWAN, Optima 450w, General Electric, Milwaukee, WI, USA) revealed the pathognomonic caput medusae venous structure and recipient veins of a DVA involving the entire right cerebral hemisphere (Fig. 4), definitively excluding other vascular malformations.

DISCUSSION

Developmental venous anomalies (DVAs) are low flow congenital brain vascular malformations that as the name describes are solely of venous origin. The anomaly drains normal brain tissue and has a distinctive imaging appearance, characterized by a linear branched configuration of veins, referred to as caput medusae. These veins converge into a dominant recipient vein that drains either into a dural sinus or ependymal veins with a varied transhemispheric brain parenchymal course [1]. They are the most common vascular brain malformation with an incidence of 2.6% of 4069...
autopsies performed and occur most commonly in the frontal lobes [2, 3]. Holohemispheric DVAs are uncommon and to our knowledge only one other case report of such an anomaly, diagnosed by catheter angiography in an adult has been reported [4].

Digital subtraction angiography (DSA) is the criterion standard for characterizing the micro and macro vascular structure as well as the hemodynamics of a DVA [2]. DVAs are visualized by CT and conventional MRI sequences. These non invasive methods are however limited in their ability to comprehensively determine the true extent and definitive angioarchitecture of the lesion for an accurate diagnosis. Conventional T2 spine echo sequences are sensitive to high flow vessels but are poor at defining low flow lesions. Gadolinium enhanced sequences opacify portions of the larger vessel components of a DVA and is acquired in relative thick sections. This limits a definitive diagnosis and may also miss small DVAs.

In the literature and by MRI machine manufacturers, the term susceptibility weighted imaging (SWI) has been used to describe varied phenomenon. Some authors have used this term to describe any sequences that allow for the detection of tissue related T2* signal susceptibility effect, such as conventional 2D T2* gradient recall echo (GRE) sequences. Susceptibility weighted imaging (SWI) has also been designated as a method by which the tissue related T2* susceptibility effect is enhanced by 3D velocity compensated gradient recalled echo sequences, using concomitantly acquired phase imaging information [5]. Although conventional 2D T2* GRE sequences are sensitive for identifying small venous vessels with low flow, they are limited by slice thickness, poor signal to noise ratio and an increased acquisition time that may be up to 8 minutes [2,5]. SWI minimal intensity projection images provide high resolution images sensitive to T2* effect, inclusive of low venous flow as seen in DVAs.

T2* weighted susceptibility angiography (SWAN, Optima 450, General Electric, Milwaukee, WI, USA) is a unique 3D-gradient echo T2*-based multi-echo sequence with varied TE times within one TR time period that allows for imaging of tissues with varying degrees of T2* contrast and is technically distinct from conventional 2D T2*GRE and SWI. A combined weighted average of all the echoes is collated by a specialized post-processing algorithm to obtain whole brain sub-millimeter-resolution 3D images [6]. The varied TE times reduces chemical shift artifact that contributes to image blurring on conventional 2D T2* GRE sequences. This also increases susceptibility signal and doubles the signal-to-noise ratio as compared to conventional 2D T2* GRE sequences. With this method large and particularly small venous structures containing deoxygenated hemoglobin as in a DVA are better characterized with greater spatial resolution for a more accurate imaging diagnosis. Although SWAN and SWI imaging are acquired differently and are vendor specific, their indications and image quality are similar.

The majority of DVAs found incidentally have a benign course [7]. The retrospective risk for hemorrhage is 0.22% per year and the prospective risk 0.68 % per year [1, 2]. They are symptomatic when spontaneous thrombosis of the draining vein with subsequent venous infarction or hemorrhage occurs. A review of symptomatic thrombosed DVA's by Ruiz et al identified venous infarction in 53%, parenchymal hemorrhage in 37%, subarachnoid hemorrhage in 5% and no lesions in 5% of 21 patients [1]. DVAs may be additionally symptomatic if associated with other vascular malformation, most commonly cavernous malformations (CMs), which are found in 13 to 40% of the cases [2]. Parenchymal bleeds from these malformations are considered to account for the majority of symptomatic DVAs.

Varied arterIALIZATION associated with DVAs is referred to as atypical types and occur very rarely [1, 2]. In particular, SWI may also be helpful in identifying high flow within arterialized vessels and has been quoted as being 100% sensitive and 96% specific for arteriovenous shunts [9]. This has not been documented for SWAN imaging in the literature. Arteriovenous shunting may also be identified by time resolved imaging of contrast kinetics (TRICKS) and is a key factor in differentiating DVA's from AVM's. MRI sequences that allow for detection of paramagnetic substances allows for identification of CMs associated with DVAs [8]. The presence of clinical symptoms, CM, hemorrhage, and arterIALIZATION with and without an AVM nidus requires further diagnostic imaging with DSA [1]. Improved characterization of DVAs by SWAN and SWI imaging could allow for differentiation between typical and types associated with CMs and arterIALIZATION.

The current understanding of DVAs is based primarily on adult studies. There are no published studies for the evaluation DVAs in the pediatric population [2]. The type and frequency of associated features in the pediatric population may be different [2]. Vigilance for accurate diagnosis of DVAs using advanced MRI techniques and to additionally evaluate for associated features are important considerations for both the radiologist and clinician in the work up and management of such conditions in the pediatric population.

TEACHING POINT

Holohemispheric DVAs in the pediatric population are rarely diagnosed, although the entity is the most common brain vascular anomaly. Susceptibility weighted angiography (SWAN) is a unique T2 star weighted sequence that definitively characterizes the angioarchitecture and associated features of a DVA for an accurate imaging diagnosis. Acquiring this or similar type of sequences such as SWI may negate the need for catheter based angiography if no atypical features of a DVA are identified.
REFERENCES


FIGURES

Figure 1: 12 year old male with right holohemispheric Developmental Venous anomaly (DVA). Select axial non-enhanced CT of the head. (Protocol: kV 120, mA 250, 2.5mm slice thickness). Multiple serpiginous hyperattenuating vascular structures within the right cerebral hemisphere and marginating the right occipital horn (small white arrows).
Figure 2: 12 year old male with right holohemispheric developmental venous anomaly (DVA). Select axial T1 MRI brain images. (Protocol: Pulse sequence; 2D Spin Echo, TR: 516.7, TE: 13, slice thickness: 5mm, Nex 1.0). Vascular flow voids within the right cerebral hemisphere and marginating the right lateral ventricle (white arrows). Note no caput medusae. Post traumatic left parietal scalp hematoma (arrow head).

Figure 3: 12 year old male with right holohemispheric developmental venous anomaly (DVA). Select axial MRI T2 brain images. (Protocol: Pulse sequence; 2D Spin Echo, TR: 4053.9, TE: 81.5, slice thickness: 5mm, Nex 1.0). Vascular flow voids within the right cerebral hemisphere and marginating the right lateral ventricle (white arrows). Note no caput medusae. Post traumatic left parietal scalp hematoma (arrow head).
Figure 4: 12 year old male with right holohemispheric developmental venous anomaly (DVA). Select axial FLAIR MRI brain images. (Protocol: Pulse sequence; 2D FLAIR, TR: 516.7, TE 13, slice thickness 5mm, Nex 1.0). Vascular flow voids within the right cerebral hemisphere and marginating the right lateral ventricle (white arrows). Note no caput medusae. Post traumatic left parietal scalp hematoma (arrow head).

Figure 5: 12 year old male with right holohemispheric developmental venous anomaly (DVA). Select axial T1 post contrast MRI brain images. (Protocol: Pulse Sequence; 2D Spin Echo, TR 846, TE 11.4, slice thickness 5mm, Nex 1.0, Multihance (Bracco Diagnostics, Princeton,NJ), 13ml injected through right forearm). Enhancement of the draining vein (large arrows) with poor visualization of radicular veins of the caput medusae (small arrows). Post traumatic scalp hematoma (arrow head).
Figure 6: 12 year old male with right holohemispheric developmental venous anomaly (DVA). Select axial TRICKS (Time resolved imaging of contrast kinetics, Optima 450, General Electric, Milwaukee, WI, USA) contrast enhanced MR angiography image of the brain. (Protocol: Pulse Sequence; 3D Gradient Recall Echo, TR: 3.6, TE: 1.4, Slice Thickness 2.4mm Nex: 0.5). Tortuous draining vein terminating in straight sinus (white arrow). Cluster of vascularity (small arrow) not typical in appearance for a caput medusae. B. Select axial Time of Flight (TOF) MRA image of the brain. (Protocol: Pulse sequence; 3D Gradient Recall Echo, Maximum intensity projection TR: 25, TE: 3.4, Slice Thickness 1.4mm, Nex 1.0), No prominent right hemispheric arterial feeder vessel or vascular nidus (arrow).

Figure 7: 12 year old male with right holohemispheric developmental venous anomaly (DVA). Serial minimum projection intensity sequential axial susceptibility weighted angiography (SWAN) images of the brain. (Protocol: Pulse sequence; 3D Gradient Recall Echo, TR: 76.6, TE: 48.1, slice thickness 8mm Nex: 0.70). Distinct characterization of the diminutive veins comprising multiple venous radicals constituting the medusa head within all the right hemispheric lobes (small black arrows). Singular recipient vein coursing along the right lateral ventricle with entry to the straight sinus through the occipital horn (white arrows).
Neuroradiology: Pediatric Holohemispheric Developmental Venous Anomaly: Definitive characterization by 3D Susceptibility Weighted Magnetic Resonance Angiography

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<table>
<thead>
<tr>
<th>Etiology</th>
<th>Uncertain, possibly arrested intrauterine loss of venous structures with compensatory recruitment of local veins</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gender Ratio</td>
<td>Slight male dominance</td>
</tr>
<tr>
<td>Treatment</td>
<td>None unless symptomatic or with atypical types or association with other vascular lesions</td>
</tr>
<tr>
<td>Age</td>
<td>Adult and pediatric</td>
</tr>
</tbody>
</table>
| Associated Entities                                                     | 1. Cavernous Hemangiomas  
2. Arterialized DVA  
3. Blue rubber bleb nevus syndrome  
4. Superficial venous malformation of the neck |
| Prognosis                                                                | 1. The retrospective risk for hemorrhage is 0.22% per year and the prospective risk 0.68% per year.  
2. Hemorrhagic infarction if draining vein surgically removed or spontaneously occluded.  
3. Hemorrhage from associated vascular malformations |

Table 1. Summary table for developmental venous anomaly (DVA)

<table>
<thead>
<tr>
<th>CT without and or with iodinated contrast</th>
<th>MRI T1 and T2</th>
<th>Gadolinium enhanced MRI</th>
<th>Hemosiderin and deoxyhemoglobin sensitive MRI sequence (SWI and SWAN)</th>
<th>Catheter Angiography</th>
</tr>
</thead>
</table>
| 1. Prominent brain parenchymal and or periventricular iso to hyper attenuating veins.  
2. Hyperdense vein if associated thrombotic occlusion.  
3. May identify associated parenchymal hemorrhage, calcification or brain atrophy.  
4. Variable enhancement of veins post iodinated contrast.  
5. Thin section CT venography for identification of the caput medusae and draining veins. | 1 Hypointense signal voids and or phase shift artifact within large veins of the caput medusae and draining veins.  
2. Superior to CT in identifying white matter changes and localized atrophy. | 1. Varied visualization of veins of the caput medusae and draining veins. More conspicuous visualization than MRI sequences without contrast. | 1. Most sensitive for detection of associated cavernous malformation  
2. Recognize associated arterialized components. | 1. Caput medusa identified during venous phase simultaneously with other vein  
2. May have simultaneous or delay of contrast clearance from draining vein.  
3. No shunt vascularity.  
4. No prominent arterial feeder vessels.  
5. No arteriovenous malformation vascular nidus. |

Table 2. Imaging Findings of developmental venous anomalies (DVA)
### CT without or with iodinated contrast

1. Prominent brain parenchymal and or periventricular iso to hyperattenuating veins.
2. Hyperdense vein if associated thrombotic occlusion.
3. May identify associated parenchymal hemorrhage, calcification or brain atrophy.
4. Variable enhancement of veins post iodinated contrast.
5. Thin section CT venography for identification of the caput medusae and draining veins.

### Dural sinus thrombosis with collateral venous drainage

1. Hyperattenuating dural and or cortical veins.
2. Enhancement of dura surrounding the thrombus. Referred to as the “Empty delta” sign.

### Sturge-Weber Syndrome

1. Tram track gyriform calcification in affected region.
2. Cortical atrophy in affected area.
3. Ipsilateral choroid plexus enlargement.
4. Leptomeningeal enhancement after contrast administration in affected area.

### Arteriovenous Malformation

1. Prominent iso and/or hyperattenuating vascularity.
2. Heterogenous attenuation of a vascular nidus with or without calcification.

### MRI T1

1. Hypointense signal voids and/or phase shift artifact within large veins of the caput medusae and draining veins.
2. Superior to CT in identifying white matter changes and localized atrophy.

### MRI T2

1. Hypointense signal voids and or phase shift artifact within large veins of the caput medusae and draining veins.
2. Superior to CT in identifying white matter changes and localized atrophy.

### Gadolinium enhanced MRI

1. Varied visualization of veins of the caput medusae and draining veins. More conspicuous visualization than MRI sequences without contrast.

### Developmental Venous anomaly

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### Arteriovenous Malformation

1. Prominent iso and/or hyperattenuating vascularity.
2. Heterogenous attenuation of a vascular nidus with or without calcification.

### Differential diagnosis table for developmental venous anomaly (DVA)

<table>
<thead>
<tr>
<th>CT without or with iodinated contrast</th>
<th>MRI T1</th>
<th>MRI T2</th>
<th>Gadolinium enhanced MRI</th>
</tr>
</thead>
<tbody>
<tr>
<td>1. Prominent brain parenchymal and or periventricular iso to hyperattenuating veins.</td>
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<td>1. Varied visualization of veins of the caput medusae and draining veins. More conspicuous visualization than MRI sequences without contrast.</td>
</tr>
<tr>
<td>2. Hyperdense vein if associated thrombotic occlusion.</td>
<td>2. Superior to CT in identifying white matter changes and localized atrophy.</td>
<td>2. Superior to CT in identifying white matter changes and localized atrophy.</td>
<td></td>
</tr>
<tr>
<td>3. May identify associated parenchymal hemorrhage, calcification or brain atrophy.</td>
<td>1. Acute thrombus appears isointense.</td>
<td>1. Acute thrombus appears hypointense.</td>
<td>1. Peripherally enhancement around clot, chronic thrombus may enhance.</td>
</tr>
<tr>
<td>5. Thin section CT venography for identification of the caput medusae and draining veins.</td>
<td></td>
<td>3. Chronically occluded sinus becomes isointense.</td>
<td>1. Prominent leptomeningeal enhancement in region affected.</td>
</tr>
</tbody>
</table>

**Table 3.** Differential diagnosis table for developmental venous anomaly (DVA) – continued on next page
### Table 3 (continued). Differential diagnosis table for developmental venous anomaly (DVA)

<table>
<thead>
<tr>
<th>MRI diffusion</th>
<th>Dural sinus thrombosis with collateral venous drainage</th>
<th>Sturge-Weber Syndrome</th>
<th>Arteriovenous Malformation</th>
</tr>
</thead>
<tbody>
<tr>
<td>1. May identify venous infarct if associated thrombotic occlusion.</td>
<td>1. There is mixture of vasogenic and cytotoxic edema.</td>
<td>1. Restricted diffusion, if there is ischemic injury.</td>
<td>1. Usually normal.</td>
</tr>
<tr>
<td>1. Caput medusa identified during venous phase simultaneously with other veins.</td>
<td>1. Occluded sinus with slowed flow of adjacent veins, development of collateral venous drainage.</td>
<td></td>
<td></td>
</tr>
<tr>
<td>2. May have simultaneous or delay of contrast clearance from draining vein.</td>
<td>2. Abnormal multiple enlarged collateralized deep venous drainage.</td>
<td></td>
<td></td>
</tr>
<tr>
<td>3. No shunt vascularity.</td>
<td></td>
<td>1. Solitary or multiple prominent arterial feeder vessels.</td>
<td></td>
</tr>
<tr>
<td>4. No prominent arterial feeder vessels.</td>
<td></td>
<td>2. An arteriovenous nidus of varied size.</td>
<td></td>
</tr>
<tr>
<td>5. No arteriovenous malformation vascular nidus.</td>
<td></td>
<td>3. Singular or multiple draining veins of varied size.</td>
<td></td>
</tr>
<tr>
<td></td>
<td>5. May have aneurysms associated with or remote to nidus.</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

### Abbreviations

AVM = Ateriovenous malformation  
CM = Cavernous malformation  
CT = Computed tomography  
DSA = Digital subtraction angiography  
DVA = Developmental venous anomaly  
FLAIR = Fluid attenuated inversion recovery  
GRE = Gradient recall echo  
MR = Magnetic resonance  
MRI = Magnetic resonance imaging  
NECT = Non-enhanced computed tomography  
SWAN = Susceptibility weighted angiography  
SWI = Susceptibility weighted imaging  
TE = Echo time  
TOF = Time of flight  
TR = Repetition time  
TRICKS = Time resolved imaging of contrast kinetics  
T2* = Susceptibility Weighted

### Keywords

Developmental Venous Anomaly; DVA; Susceptibility weighted imaging; MRI

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