A Step-by-step Sonographic Approach to Vascular Anomalies in the Pediatric Population: A Pictorial Essay

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Indian J Radiol Imaging 2021;31:157–171.

Abstract

Vascular anomalies are a common cause of soft-tissue masses in children and often referred for ultrasonographic (USG) evaluation. They are broadly classified as vascular tumors (hemangiomas, hemangioendotheliomas, and angiosarcomas) or vascular malformations (venous malformations, lymphatic malformations, and arteriovenous malformations). Findings on USG and Doppler imaging can be used to categorize vascular anomalies into high- or low-flow lesions, which forms the basis for further workup, diagnosis, and management. On careful evaluation of various sonographic features, in conjunction with clinical findings, an accurate clinicoradiological diagnosis can be made in most cases. Further imaging with magnetic resonance (MR) imaging or computed tomography (CT) helps in delineation of lesion extent, whereas MR or CT angiography is useful to map the vascular supply of high-flow lesions. We have illustrated and discussed a step-by-step approach to diagnose vascular anomalies using ultrasound and Doppler imaging.

Introduction

Vascular anomalies are commonly encountered in the pediatric population. They encompass a variety of disorders, from a simple small birthmark to life-threatening entities, which may cause complications like heart failure, Kasabach–Merritt syndrome, or significant airway narrowing. Traditionally, these anomalies have been diagnosed based on descriptive observations, including clinical appearance, location, and fluid contents. This has led to misdiagnosis and improper management of these lesions.¹,² The term vascular anomaly represents a broad spectrum of vascular pathology, including vasoproliferative tumors and vascular malformations. Vascular tumors demonstrate rapid growth, increased cellular turnover, and endothelial hypercellularity. Examples include infantile hemangioma, congenital hemangioma, Kaposiform hemangioendothelioma (KH), and angiosarcoma. Vascular malformations are not neoplasms; they result from errors in vascular morphogenesis. They usually grow at a rate proportionate to the growth of the child and comprise vascular spaces with normal endothelium. These include slow-flow vascular malformations (venous, lymphatic malformations [LMs]), high-flow malformations (arteriovenous malformations [AVMs]), and various combined malformations.

In 1996, the International Society for the Study of Vascular Anomalies (ISSVA) released a classification system for vascular anomalies, which was updated in 2014 and revised in 2018.³ The ISSVA classification (► Table 1) is widely accepted and used among clinicians managing vascular anomalies. It continues to evolve with our evolving knowledge of the biology and genetic background of these lesions.
The 1996 classification had fundamentally divided vascular lesions into vascular tumors and vascular malformations (malformations could be simple or combined; combined malformations are defined as more than two malformations in one lesion). The 2014 updated classification expanded the sections on vascular tumors and vascular malformations. Tumors were divided into three categories: benign, locally aggressive/borderline, and malignant. Vascular malformations, apart from simple or combined, were categorized as malformations of major named vessels and those associated with other anomalies like Klippel–Trenaunay syndrome, Sturge–Weber syndrome, etc.

With greater knowledge of the genetic basis of vasoproliferative lesions, the 2018 revised ISSVA classification included a list of causative genes (as an appendix).\(^3\) Most notable among these is the \(PIK3CA\) gene on chromosome 3, which is associated with several syndromes collectively called the \(PIK3CA\)-related overgrowth spectrum (PROS). A few vascular lesions—like fibroadipose vascular anomaly (FAVA)—still remain unclassified, because it is unclear whether they are tumors or malformations, or because their clinicopathological characteristics are as yet incompletely understood.

Vascular anomalies are further divided into "high-flow" or "low-flow" groups, based on their flow dynamics. High-flow lesions contain an arterial component. Examples include hemangiomas (infantile and congenital), other vascular tumors, AVMs, and arteriovenous fistulas (AVFs). Low-flow lesions are all other lesions that do not contain an arterial component, including capillary malformations, LMs, venous malformations (VMs), and also involving hemangiomas.\(^4\)

Sonography (along with Doppler imaging) is the first-line imaging modality employed for diagnosis of vascular anomalies. Usually, patients with these lesions are encountered in a busy routine ultrasound practice, sometimes even in emergencies, where a quick and accurate diagnosis is needed. A simple algorithmic approach on the basis of sonographic findings, in conjunction with clinical features, helps in making a confident diagnosis of these lesions, even by less experienced radiologists. Further imaging using magnetic resonance imaging (MRI) or computed tomography (CT) is helpful in assessment of lesion extent and vascular supply.

### Table 1

| Tumors                  | Benign                                   | Locally aggressive | Malignant       | Malformations          |
|-------------------------|------------------------------------------|--------------------|-----------------|-----------------------|
|                         | Infantile hemangioma, congenital hemangioma, tufted angioma, spindle cell hemangioma, epithelioid hemangioma, pyogenic granuloma, etc. | Kaposisform hemangioendothelioma, retiform hemangioendothelioma, papillary intralymphatic angioendothelioma, Dabska tumor, composite hemangioendothelioma, Kaposi sarcoma, etc. | Angiosarcoma, epithelioid hemangioendothelioma, others | Capillary (low flow) port wine stain, telangiectasia, cutis marmorata telangiectatica congenita, nevus simplex, etc. |
|                         | Lymphatic (low flow) common (cystic) LM, GLA, channel type LM, primary lymphedema (different types), etc. | Venous (low flow) common VM, familial VMMC, blue rubber bleb (bean) syndrome VM, GVM, CCM, etc. | Arteriovenous malformation (high flow): sporadic, in HHT, others | Lymphatic (low flow) common VM, familial VMMC, blue rubber bleb (bean) syndrome VM, GVM, CCM, etc. |
|                         | Arteriovenous malformation (high flow): sporadic, in HHT, others | Arteriovenous fistula (congenital) (high flow): sporadic, in HHT, others | Arteriovenous malformation (high flow): sporadic, in HHT, others | Arteriovenous malformation (high flow): sporadic, in HHT, others |
|                         | CVM, CLM, CAVM, LVM, CLVM, CLAVM, CAVVM, CLAVVM (C: capillary; V: venous; L: lymphatic; AV: arteriovenous) | Affects lymphatics, veins, arteries | Of major named vessels | Affects lymphatics, veins, arteries |
|                         | KT, PWS, Sturge–Weber syndrome, Maffucci syndrome, Proteus syndrome, etc. | Clinical Features | Associated with other anomalies | Associated with other anomalies |
|                         | Intramuscular hemangioma, angio-keratoma, sinusoidal hemangioma, acral arteriovenous "tumor," multifocal lymphangioendothelio-matosis with MLT/CT, PTEN (type) hamartoma of soft tissue/angiormatosis of soft tissue (PHOST), FAVA | Clinical Features | Provisionally unclassified vascular anomalies | Provisionally unclassified vascular anomalies |

Abbreviations: CCM, cerebral cavernous malformation; FAVA, fibroadipose vascular anomaly; GLA, generalized lymphatic anomaly; GVM, glomouavenous malformation; HHT, hereditary hemorrhagic telangiectasia; KTS, Klippel–Trenaunay syndrome; LMs, lymphatic malformations; MLT/CT, multifocal lymphangioendothelio-matosis with thrombo-cytopenia/cutananeous angiormatosis with thrombocytopenia; PTEN, phosphate and tensin homolog; PWS, Parkes–Weber syndrome; VMMC, venous malformation cutaneomucosal.

Clinical Features

The importance of detailed history and examination in a case of vascular anomaly cannot be emphasized enough, since they form the basis for differentiation. Lesions like superficial hemangiomas often do not need any imaging, being diagnosed solely on the basis of their age of onset, clinical appearance, and growth pattern.

These anomalies may involve any part of the body, most commonly the head and neck, followed by the extremities. Symptoms may range from skin discoloration, swelling, pain, and bleeding to systemic effects like high-output cardiac failure (which may be seen in high-flow lesions). The age of onset and pattern of growth (and involution) help diagnose infantile and congenital hemangiomas and differentiate them from vascular malformations.\(^5\-\(^15\)

On examination, deep-seated lesions may not produce any discoloration in the overlying skin. Superficial vascular anomalies are seen in the form of blush/violet/pink/red-colored lesions with varying morphologies. The important clinical features of common vascular anomalies have been summarized in Table 2.
Role of Various Imaging Modalities

Imaging is indicated in vascular anomalies to (1) confirm the clinical diagnosis; (2) characterize the lesion as high or low flow, and further diagnose the type of lesion; (3) know the complete extent of infiltrative, deep-seated, or widespread lesions; and (4) know the vascular supply in high-flow lesions where endovascular management is planned.

Ultrasonography

Ultrasonography (USG) is the first-line imaging modality, with various advantages including lack of ionizing radiation, no need of sedation in small children, cost-effectiveness, portability, and wide availability. Most importantly, grayscale sonography along with color and spectral Doppler can accurately diagnose most of the cases, when interpreted in conjunction with clinical features. The categorization of vascular anomalies as high- or low-flow lesions is best achieved using Doppler-USG. Disadvantages include operator-dependence and inability to adequately assess deep lesions;16,17

Contrast-Enhanced Ultrasound

Contrast-enhanced ultrasound combines all the advantages of USG with the ability to evaluate the microvasculature of the lesion and quantify perfusion parameters using a time-intensity curve analysis. Various parameters like time to peak and area under the curve can be studied in different regions of interest in the center and at the periphery of the lesion, and comparison of pretreatment and posttreatment values has a role in monitoring response to therapy.18

Magnetic Resonance Imaging

MRI is very frequently the next step after USG for assessment of the extent of a deep-seated or large vascular anomaly, its relation with surrounding vital structures, and the tissue planes involved. Magnetic resonance (MR) angiography provides information about the vascular supply in high-flow lesions. The need for sedation, long scan times, high cost, and limited availability are some disadvantages of MRI.

Computed Tomography

The presence of ionizing radiation, lack of soft-tissue contrast, and contrast-related side effects are some important drawbacks of CT. It is performed in select cases where an urgent evaluation needs to be done, for example, in evaluation of airway compromise caused by a lesion, and when MRI is not available or the child cannot be sedated for MRI.19

Step-by-Step Doppler-USG Approach to the Diagnosis of Vascular Anomalies

A step-wise approach to sonographic diagnosis of vascular anomalies has been summarized in – Fig. 1.

First step: Assess the presence/absence of arterial flow

The presence of arterial flow suggests a high-flow lesion, whereas low-flow lesions do not show arterial flow.

- When arterial flow is detected within the lesion on color Doppler imaging, the following features should be evaluated:
  - Vessel density: A semiquantitative assessment of vessel density is possible using color Doppler imaging. The area of greatest vascularity should be identified within the lesion, and the number of vessels counted in an area of 1 cm². Less than 2 vessels/cm² is described as low vessel density, 2 to 4 vessels/cm² as moderate, and ≥5 vessels/cm² as high vessel density. High vessel density is typically seen in “high-flow” vascular anomalies, like hemangiomas, other vascular tumors, and high-flow vascular malformations (AVMs, AVFs).
  - Peak arterial systolic velocity: Peak systolic velocity (PSV) of the arterial vessels within the lesion should be measured, after appropriate angle correction. High values of arterial PSV are seen in high-flow lesions. A study by Paltiel et al found no significant difference in mean arterial PSV values between hemangiomas and AVMs.20

### Table 2  Clinical features of common vascular anomalies

| Vascular malformations | Age of onset                  | Growth                  | Involution (spontaneous) | Morphology and clinical examination |
|------------------------|-------------------------------|-------------------------|--------------------------|------------------------------------|
| Infantile hemangioma   | 2–6 wk after birth            | Rapid till 18 mo        | Usually by 9 y           | Superficial lesion—bright red      |
| Congenital hemangioma  | At birth                      | Variable                | RICH by 1 y              | NICH—pink/violet with central telangiectasias |
| High-flow (AVM)        | Variable, usually discovered in childhood | In proportion to the growth of child | Absent | Bruit and thrill over the lesion or a pulsatile lesion |
| Low-flow VM            |                               |                         |                          | Superficial lesion: blue/purple, fluctuant, compressible |
| Low-flow LM            |                               |                         |                          | Soft lesions showing transillumination |

Abbreviations: LM, lymphatic malformation; NICH, noninvoluting congenital hemangioma; PICH, partially involuting congenital hemangioma; RICH, rapidly involuting congenital hemangioma; VM, venous malformation.
Step-by-step Sonographic Approach to Vascular Anomalies in Children

Mittal et al.

**Resistive index:** The arteries within high-flow vascular anomalies show low-resistance flow, with low values of resistive index (RI). In some instances, a single or a few areas of arterial flow may also be seen within low-flow vascular malformations—for instance, venous and LM often infiltrate around normal structures, and a few arterial vessels may be seen passing through these lesions. Also, LMs may show areas of arterial flow within the septa/walls of their constituent cysts. These foci of arterial flow in low-flow lesions show high RI values, in contrast to arteries within high-flow lesions.

**Imaging Approach if Arterial Flow Is Present (High-Flow Lesion)**

**Second Step: Is the Lesion Typical for Hemangioma or Not**

As the name suggests, congenital hemangiomas are present at birth, whereas infantile hemangiomas develop soon after birth. An initial phase of rapid growth followed by involution is typically seen in infantile hemangiomas or rapidly involuting congenital hemangiomas (RICH).

On clinical examination, superficial hemangiomas are characteristic in appearance—red/blue/violet in color without any bruit/thrill/transillumination.

Hemangiomas, which are generally high flow in nature (Fig. 2), may not show arterial flow in their involuting phase/posttreatment (Fig. 3). Such hemangiomas are, thus, low-flow lesions.

**Third Step: Assess the Presence of Soft-tissue Component within the High-Flow Lesion**

Intralesional soft tissue is seen in vascular tumors, whereas lesions without a soft-tissue component are high-flow malformations (AVM/AVF). Presence of soft-tissue mass is the most reliable predictor for differentiating hemangiomas (or other vascular tumors) from AVMs.

The most common vascular tumors are infantile hemangiomas. Lesions with atypical imaging/clinical findings may be vascular tumors other than hemangiomas. These usually require biopsy for diagnosis.

**Fourth Step: Does the Lesion Show Arterialization of Venous Flow and Low-Resistance Arterial Waveform?**

Such a lesion, which lacks a soft-tissue component, is likely to be an AVM/AVF. The term arterialization of venous flow refers to high venous peak velocities seen within these lesions. Mean venous peak velocities in AVMs are significantly higher than those seen in hemangiomas and VMs.

**Imaging Findings of Various High-Flow Lesions**

**Vascular Tumors**

**Infantile Hemangioma**

Infantile hemangiomas are the most common vascular tumors, may be single or multiple, and are seen in ~4 to 10% of infants. They express the glucose transporter isoform 1 (GLUT1) protein, which is not expressed by any other normal...
tissue or vascular tumor. Skin is the most common location, with multiple cutaneous hemangiomas in 10 to 25% cases. These lesions are most commonly located in the subcutaneous plane and may infiltrate into the deeper tissue planes when large. Liver is the most common extracutaneous site (Fig. 4). The typical Doppler-USG appearance is of a well-circumscribed, homogenous, hypo-/hyperechoic lesion with internal arterial as well as venous flow on color Doppler (Fig. 5). In the proliferative phase, hemangiomas typically show a high vessel density (>5 vessels/cm²) and high Doppler shift (>2 kHz). Involuting lesions may show mixed echogenicity due to the intermixing of fat and may not show arterial flow within. Small, superficial lesions with typical clinical as well as USG features do not usually require further imaging, unless there is suspicion of associated visceral involvement. Uncommonly, large hemangiomas may be infiltrative in appearance (Fig. 6). Such lesions require further imaging evaluation using MRI/CT. On MRI, hemangiomas typically show multiple flow voids and intense enhancement on postcontrast images. In large facial hemangiomas, MRI of the brain is indicated to rule out PHACES syndrome (P—posterior fossa malformations, H—hemangiomas, A—arterial anomalies, C—coarctation of aorta and cardiac anomalies, E—eye/ocular anomalies, S—sternal defects). Although the majority of hemangiomas are self-limiting, treatment is required in the presence of complications like heart failure or Kasabach-Merritt syndrome.

**Congenital Hemangioma**

Congenital hemangiomas are much less common than infantile hemangiomas, usually solitary, and test negative for...
the immunohistochemical marker GLUT1. They are of two major types—RICH and noninvoluting congenital hemangiomas (NICH). Imaging findings are broadly similar to those of infantile hemangiomas. Presence of calcification, which is not a feature of infantile hemangiomas, may be seen in congenital hemangiomas (17% NICH, 37.5% RICH vs. none in infantile hemangiomas). A partially involuting congenital hemangioma has also been recently described, which shows initial rapid involution like RICH, but does not completely involute and persists instead, like a NICH.

Pyogenic Granuloma/Lobular Capillary Hemangioma
These are common, benign vascular tumors diagnosed clinically in most cases. They are seen as small growths, usually blood red in color, with a “minced meat”–like surface. However, they have an increased tendency to bleed, which may sometimes make it difficult to differentiate them from AVMs. In such cases, imaging may help by demonstrating a well-defined, homogenous, vascular mass (Fig. 7), without the flow voids or the nidus typical of an AVM. Final diagnosis is established on histopathology.

Solitary Fibrous Tumor/Hemangiopericytoma
A solitary fibrous tumor (previously referred to as a hemangiopericytoma) is a mesenchymal tumor, extremely rare in children. Most tumors are benign, but some may show variable malignant potential. On USG, these are solid or rarely cystic masses, well demarcated, with homogeneous or heterogeneous echogenicity. On Doppler, the presence of intratumoral arteriovenous shunting can be diagnosed based on the presence of low-resistance flow in the feeding arteries, arterialization of the venous flow, and large internal vessels with low vascular impedance (Fig. 8). Presence of soft tissue helps differentiate from an AVM. Further imaging with MRI/CT may demonstrate a locally aggressive soft-tissue lesion with marked contrast enhancement (Fig. 8). Biopsy is needed for diagnosis.

Kaposiform Hemangioendothelioma
KH is a rare, locally aggressive, vasoproliferative tumor, which has a low malignant potential and usually presents shortly after birth. It may be seen in the head and neck, trunk, extremities, retroperitoneum, and rarely in other locations. KH has both vascular and lymphatic components and may be associated with Kasabach–Merritt phenomenon, a term that describes profound thrombocytopenia caused by platelet sequestration resulting from a consumptive coagulopathy. The imaging appearance of various vasoproliferative tumors overlaps with each other. The diagnosis of hemangiomas can be usually made in the presence of typical clinical features. If the history, clinical examination, and/or Doppler-USG findings are atypical, then further imaging is indicated. If a vascular tumor other than hemangioma is suspected, biopsy is required for diagnosis.

Vascular Malformations: AVM/AVF
An AVM typically shows an abnormal cluster of arterial channels communicating with venous channels via a nidus. An AVF is similar to an AVM except for the lack of a nidus.
Histologically, AVMs and AVFs consist of dysplastic arteries that drain into arterialized veins, bypassing capillary beds. Unlike other high-flow lesions, no soft-tissue component or lesion matrix is seen in these lesions. Typically, most AVMs are congenital, whereas most AVFs are acquired. A congenital AVF may be sporadic or occur as a part of hereditary hemorrhagic telangiectasia.

Doppler-USG demonstrates low-resistance flow in the involved arteries, arterIALIZATION of flow in the venous system, and high peak velocities in both the arterial and venous channels (►Fig. 9). It may show the site of arteriovenous communication in an AVF (►Fig. 10). CT or MR angiography helps by demonstration of the vascular anatomy of the lesion (►Fig. 11), which helps in planning its management.

Imaging Approach if Arterial Flow Is Absent (Low-Flow Lesion)

Second Step: Is the Lesion Typical for Hemangioma or Not?
It is important to remember that hemangiomas may not show arterial flow in the involuting phase, or on treatment. These lesions will have a typical history and should not be misdiagnosed as VMs. In typical cases, no further imaging is necessary and the patient can be managed conservatively.

Third Step: Look for the Presence or Absence of Venous Flow
If venous flow is seen, the lesion is a VM. However, not all VMs will show flow on Doppler-USG. In case no flow is seen, the lesion may be either a lymphatic or a VM (with very slow flow or thrombosis). Approximately 16% of all VMs show no detectable flow on Doppler-USG.

Fourth Step: Look for Phleboliths
Calcified phleboliths, if seen, are highly specific for the diagnosis of VMs, even in the absence of flow (►Fig. 12). Phleboliths are seen in ~16% of all VMs.

Fifth Step: Is the Lesion Cystic with Peripheral Septal Vascularity?
LMs are cystic, usually multilocular cystic lesions, lack phleboliths, and show no flow or only peripheral/ septal flow. Internal echoes may be seen. VMs, on the other hand, are often hypoechoic and not multicystic; they may comprise venous spaces or ectatic/dysplastic veins.

Imaging Findings of Various Low-Flow Lesions
Common low-flow lesions are vascular malformations including capillary, venous, and LMs, and their combinations. Capillary malformations typically involve superficial layers of skin—like the facial malformations seen in Sturge–Weber syndrome and are clinically diagnosed. Venous, lymphatic, and venolymphatic malformations are the most common...
Fig. 9 (A) An arteriovenous malformation (AVM) in an 18-year-old man, who presented with a vascular swelling over his foot. (B–D) Ultrasonography (USG) showed multiple vascular channels with both arterial and venous flow. (E) There was arterialization of venous flow, whereas the arteries showed low-resistance flow with (F) a resistive index (RI) of 0.41. (G–I) Computed tomography (CT) angiography showed multiple enlarged vessels, and early filling of venous channels in the arterial phase.

Fig. 10 Intrahepatic arterioportal fistula in a 6-month-old infant with Down’s syndrome and abdominal distension. (A,B) Ultrasonography (USG) shows an aneurysmally dilated left portal vein, with an abnormal intrahepatic arterioportal communication (arrow in B). (C,D) Color Doppler shows turbulent intralesional flow. Spectral Doppler confirms (E) a high-velocity, low-resistance flow in the feeding artery and (F) chaotic, arterIALIZED flow in the dilated vein.

Fig. 11 (A) An arteriovenous malformation (AVM) of the pinna in a 16-year-old adolescent boy who had a swollen pinna showing multiple dilated vascular channels over it. (B–D) Ultrasonography (USG) showed multiple dilated vessels showing arterialization of venous flow, diagnostic of AVM. (E–G) Computed tomography (CT) angiography well depicted the tangle of vessels, with arterial supply from branches of the left external carotid artery. The patient was referred for embolization.

Fig. 12 (A) A venous malformation in an 11-year-old boy with a long-standing neck swelling. Ultrasonography (USG) reveals (B) a circumscribed lesion with cystic and hyperechoic areas and (C) a phlebolith, but (D) no flow even on power Doppler. (E–G) Computed tomography (CT) confirms intralesional phleboliths with only minimal enhancement.
types of vascular malformations; their prevalence is ~1% in the general population.4

**Simple Low-flow Vascular Malformations**

**Venous Malformations**

VMs are usually present at birth but may not be apparent. They grow in proportion to the growth of the child. They are composed of vascular channels, sometimes containing intraluminal thrombi, lined by thin endothelium. Based on the pattern of venous drainage, these may be of different types—isolated malformations without venous drainage; those draining into normal veins, or into dysplastic veins; or malformations comprising primarily of venous ectasia.19

On Doppler-USG, VMs are often hypoechoic, may be infiltrative lesions involving multiple tissue planes (►Fig. 13), or may be well marginated (►Fig. 14). They may comprise venous spaces or ectatic/dysplastic veins and show venous flow or no flow. Phleboliths (well defined, round, calcific foci), if seen, are characteristic (►Fig. 15). These represent areas of spontaneous thrombosis.

MRI is the preferred examination to evaluate the extent of large or infiltrative lesions. Fat-suppressed T2-weighted or short tau inversion recovery images can depict the lesions, which are hyperintense, well. On postcontrast CT or MRI, VMs have a wide range of contrast enhancement patterns: from homogeneous to heterogeneous, faint to vivid, and rapid to delayed.13–34 Phleboliths within the lesion may be missed on MRI and are best depicted on CT images.

**Lymphatic Malformations**

LMs comprise dilated lymphatic channels, forming cystlike structures, isolated from the normal lymphatic system. They are congenital, and the majority (80–90%) of these lesions are discovered by the age of 2 years. They are subclassified on the basis of the size of the lymphatic spaces into macrocystic (cysts >1 cm in diameter), microcystic (cysts <1 cm in diameter), and combined (►Fig. 16).20,35,36 They commonly involve the subcutaneous tissues, and most often occur in the head and neck region. They are often trans-spatial and appear infiltrative, involving multiple tissue planes (►Fig. 17).

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**Fig. 13** (A) An infiltrative venous malformation of the neck in a 12-year-old boy. (B–D) Ultrasonography (USG) showed an infiltrative lesion with multiple abnormal vessels showing (E) venous flow. (F) Magnetic resonance imaging (MRI) coronal short tau inversion recovery (STIR) and (G,H) axial T1-weighted (T1W) and fat-saturated T2W images clearly demonstrated an infiltrative lesion involving the right buccal, parotid, masticator, carotid, prevertebral, and perivertebral spaces.

**Fig. 14** (A) Intramuscular venous malformation of the left thigh in a 6-year-old girl. Ultrasonography (USG) shows a well-circumscribed lesion with (B) phleboliths and (C) venous flow. (D) An arterial branch is seen within the lesion, with high resistive index (RI) and biphasic flow, suggestive of a muscular arterial branch. Computed tomography (CT) confirms (E) phleboliths (non–contrast-enhanced CT (NCCT)) and (F) enhancement within the lesion.
LMs can show periods of sudden growth, in response to immunological stimuli like the common cold, or in the presence of hemorrhage or infection. On USG, LMs appear as multilocular cystic masses with increased through-transmission. They may show low-level internal floating echoes (►Fig. 18) or a fluid–fluid level representing chyle, blood, or pus. On Doppler study, no internal vascularity is noted (►Fig. 19); however, peripheral vascularity may be visible within the fibrous septations (►Fig. 18) and capsule.\textsuperscript{20,35,36} If arterial flow is seen within the septa or cyst wall, it shows high-resistance flow and high RI values.

MRI is the preferred imaging modality for further evaluation of the extent of the lesion. These lesions are typically hyperintense on T2-weighted images and may show variable signal intensity on T1-weighted images, depending on cyst content. CT may be useful if MRI cannot be done. No enhancement is seen within LMs (►Fig. 20), and this is very useful to differentiate them from VMs.

Fig. 15 (A) A venous malformation along the thenar eminence of palm in a 4-year-old boy. Ultrasonography (USG) showed (B) a phlebolith and (C) low flow within. (D) Radiograph also confirmed intralobesional phlebolith. (E,F) Magnetic resonance imaging (MRI) coronal T1-weighted (T1W) and (G) short tau inversion recovery (STIR) images well depicted the lesion extent.

Fig. 16 (A) Lymphatic malformation of the abdominal wall in a 9-month-old infant. (B–D) Ultrasonography (USG) revealed a well-defined lesion with multiple macrocysts (large arrow) and few microcysts (small arrow) and (E) septal flow. (F) Contrast-enhanced magnetic resonance imaging (MRI) also showed peripheral and septal enhancement.

Fig. 17 (A) Multispatial lymphatic malformation of the neck in an 11-month-old infant. (B,C) Ultrasonography (USG) showed a multicystic lesion with internal echoes and (D,E) no color flow. (F-H) Computed tomography (CT) depicted the extent of the lesion both anterior and posterior to the left sternocleidomastoid muscle (arrow), with a component in the prevertebral space.
Complex Lymphatic Anomalies

Rare systemic lymphatic anomalies that can involve multiple organs and have significant morbidity and mortality include generalized lymphatic anomaly, Kaposiform lymphangiomatosis, Gorham–Stout disease, and central conducting lymphatic anomaly (channel type LM, lymphangiectasia). USG has a limited role in evaluation of these lesions, and MRI or CT evaluation is indicated, depending on the site(s) of involvement.

Combined Vascular Malformations

Combined vascular malformations comprise two or more malformations in a single lesion. They may occur in sporadic or syndrome forms and are named by hyphenation of the names of the types of component tissues—for example, CLM for capillary lymphatic malformations. Combined venolymphatic malformations may occur, which show imaging features of both VMs and LMs (Fig. 21).

Malformations of Major Named Vessels

The malformations of major named vessels are also known as the “channel-type” or “truncal” vascular malformations and include a wide range of anomalies of origin, course, number, and diameter of vessels as well as residual embryonal vessels.
Vascular Malformations Associated with Other Anomalies

PIK3CA-Related Overgrowth Spectrum
The PIK3CA gene on chromosome 3 is important for growth regulation and mutations in this gene are associated with several cancers. This gene is also implicated in a spectrum of overgrowth syndromes/phenotypes, many of which have vascular anomalies as a part. PROS lesions include congenital lipomatous overgrowth, vascular malformations, epidermal nevi, scoliosis/skeletal and spinal syndrome, and CLAPO syndrome (which stands for—Capillary malformations of the lower lip, cervicofacial LM, Asymmetry and Partial or generalized Overgrowth), fibroadipose hyperplasia or overgrowth, macrodactyly, and Klippel–Trenaunay syndrome.38

Klippel–Trenaunay Syndrome
Klippel–Trenaunay syndrome classically comprises a triad of capillary malformations, VMs, and soft-tissue/bone overgrowth, most commonly involving (one) lower limb. A wide spectrum of venous anomalies may be present, including ectasia, aplasia/hypoplasia of deep veins, and VMs involving the superficial and deep venous systems. LMs may also be present. USG is useful as an initial investigation, whereas MRI is helpful for delineating the extent of abnormality (►Fig. 22).

Provisionally Unclassified Vascular Anomalies: Fibroadipose Vascular Anomaly
The 2018 ISSVA classification has expanded the list of provisionally unclassified entities. FAVA is a recently described, complex vascular malformation that has certain typical clinical and imaging findings.39

Adolescent females are most often affected and present with constant pain. FAVA lesions occur within the muscle, most often of the lower limb (gastrocnemius, soleus). On imaging, they may be focal mass–like, focal infiltrative, or diffuse infiltrative. Variable amounts of fat and fibrous tissue are present within the lesion. Phlebectasia is a characteristic imaging finding (►Fig. 23) that may be seen on USG, MRI, or venography. On USG, presence of an echogenic soft-tissue component helps differentiate FAVA from VM. Macroscopic
venous channels and dilated veins may be seen. On MRI, hyperintense areas are seen on T1-weighted images (due to intrallesional fat), and these lesions are less hyperintense on T2-weighted images compared with conventional VMs. Postcontrast enhancement is seen. FAVA lesions may be mistaken for VMs on imaging, but it is important to diagnose them correctly as the management is different. They respond poorly to sclerotherapy and may require surgical resection or ablative/cryotherapy for pain relief.

Management of Various Vascular Anomalies

The management of most vascular anomalies requires a multidisciplinary approach and close cooperation between clinicians and radiologists. Increasingly, interventional radiology techniques have an important role to play in the management of these lesions. Uncomplicated infantile hemangiommas may be managed conservatively or with oral propranolol, whereas complicated lesions can be managed surgically with laser treatment or with resection. RICH presenting with acute cardiac failure may require emergency embolization to reduce the risk of arteriovenous shunting. NICH, usually small, do not require treatment but can be surgically treated. In most cases of congenital hemangiomas, conservative management with a “watch-and-wait” approach is adopted. However, cases complicated by hemorrhage, ulceration, airway obstruction or ophthalmic involvement, heart failure, and those causing severe psychological distress or disfigurement warrant further discussion and consideration for treatment. Most common treatment for solitary fibrous tumor/hemangiopericytoma is wide surgical excision, whereas for AVM, it is embolization focused on obliteration of the nidus, which can be achieved percutaneously, via an endovascular approach, or surgically. The mainstay of treatment for VMs or LMs is percutaneous sclerotherapy, or surgery when feasible.

Pitfalls in USG Evaluation/Diagnosis of Vascular Anomalies

- There are several pitfalls that must be kept in mind while using sonography and Doppler imaging for the evaluation of vascular anomalies. Some of these are the following:  
  - Optimal Doppler settings are essential for the evaluation of any vascular anomaly. For evaluation of low-flow lesions, it is imperative to utilize low-flow settings on color Doppler. Power Doppler imaging is also a valuable tool, as it has greater flow sensitivity. Sometimes, presence of low flow may be seen as moving echoes on real-time grayscale USG.
• All hemangiomas do not show arterial flow. Lesions in the involuting phase may show venous flow or no flow on Doppler study. These patients need to be diagnosed based on typical clinical appearance and response to treatment.

• In LMs/VMs, sometimes a few arterial channels may be seen traversing through the lesion. Such arteries on spectral evaluation show high-resistance and low-velocity flow, in contrast to low-resistance flow within the arterial component of a high-flow vascular anomaly.

• Some LMs may not appear cystic and instead appear hypoechoic or heteroechoic on sonography. This may be due to the presence of multiple internal echoes or due to their microcystic nature. These should not be confused with VMs or solid tumors. The presence of increased through-transmission establishes their cystic nature.

• Certain cystic tumors (like teratomas) and tumorlike lesions (like epidermoid/dermoid cysts) may mimic LMs on ultrasound, especially in the head and neck region. The presence of a solid component, along with demonstration of calcification and/or fat within the lesion, suggests the diagnosis of a teratoma rather than an LM.

• VMs, in some cases, show a very slow flow that is not detected by Doppler-USG. Absence of flow does not preclude the diagnosis of a VM. VMs with thrombosis may show absence of flow on Doppler and may even show absence of enhancement on postcontrast MRI or CT.

• Certain entities can mimic VMs on imaging; a common example is a plexiform neurofibroma, seen in patients with neurofibromatosis type 1 (►Fig. 24). On USG, these are hypoechoic, trans-spatial lesions, which may mimic infiltrative VMs. On MRI, they are T2-hyperintense lesions, typically showing avid postcontrast enhancement.

Conclusion

Sonography can accurately diagnose most vascular anomalies if examined in a systematic manner and should be the initial imaging modality of choice, especially in the pediatric population. Doppler sonography helps in categorizing vascular anomalies into high- and low-flow lesions, which forms the basis of imaging diagnosis and guides further management. Sonographic findings must be interpreted in conjunction with history and clinical examination findings. Further imaging evaluation using MRI or CT has a role in assessment of lesion extent, and in angiographic evaluation of high-flow lesions.

Financial Support and Sponsorship
Nil.

Conflict of Interest
There are no conflicts of interest.

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Step-by-step Sonographic Approach to Vascular Anomalies in Children  Mittal et al.

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