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ABSTRACT

SUMMARY: Fetal vascular malperfusion includes a continuum of placental histologic abnormalities increasingly associated with perinatal brain injury, namely arterial ischemic stroke. Here, we describe the clinical-neuroimaging features of 5 neonates with arterial ischemic stroke and histologically proved fetal vascular malperfusion. All infarcts involved the anterior territories and were multiple in 2 patients. In 2 neonates, there were additional signs of marked dural sinus congestion, thrombosis, or both. A mixed pattern of chronic hypoxic-ischemic encephalopathy and acute infarcts was noted in 1 patient at birth. Systemic cardiac or thrombotic complications were present in 2 patients. These peculiar clinical-radiologic patterns may suggest fetal vascular malperfusion and should raise the suspicion of this rare, underdiagnosed condition carrying important implications in patient management, medicolegal actions, and future pregnancy counseling.

ABBREVIATIONS: FVM = fetal vascular malperfusion; PAIS = perinatal arterial ischemic stroke

Perinatal stroke is a group of cerebrovascular diseases occurring between 20 weeks of fetal life and 28 postnatal days caused by focal disruption of normal (arterial or venous) cerebral blood flow.^{1,2} Many patients are diagnosed several months after birth and therefore classified as having “presumed” perinatal stroke.^{1,2}

Perinatal arterial ischemic stroke (PAIS) and presumed PAIS are the most frequent subtypes of perinatal stroke,^{2,3} with an overall incidence up to 1 per 1600 live births.⁴ They represent an important cause of cerebral palsy and are frequently associated with epilepsy and abnormal cognitive, neurosensory, and behavioral outcomes.^{2,5} The pathomechanisms of PAIS and presumed PAIS are very likely similar but remain elusive, with multiple maternal, fetal, and placental risk factors proposed.^{2,6} Of these, placental abnormalities are the least investigated.⁶ Indeed, scant data about the placenta have been presented in the largest multicenter PAIS studies published so far.⁷

Fetal vascular malperfusion (FVM), previously known as fetal thrombotic vasculopathy,⁸ is an umbrella term introduced in 2015 by the Amsterdam International Consensus group of pathologists, encompassing a spectrum of histologic placental abnormalities, frequently at different stages compatible with reduced or absent perfusion in any location of the fetoplacental vasculature, such as occlusive thrombi, mainly in large-stem vessels, accompanied by downstream changes, including organization, septation, red cell extravasation, endothelial destruction, loss of vascularity (avascular villi), and ultimately fibrosis.⁹ According to the location or extension of placental abnormalities, it can be further classified into low- or high-grade FVM.⁹ The exact pathogenesis of FVM is still not clear. However, venous stasis, endothelial or vessel wall damage, and hypercoagulability are potentially involved (Virchow triad).^{9,10} Importantly, there is increasing evidence that FVM plays an important role in neonatal mortality, morbidity, and outcome, including brain injury.¹¹⁻¹⁴ In particular, the

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Ethics approval: All procedures performed in studies involving human participants were in accordance with ethical standards of the institutional and/or national research committee and with the 1964 Helsinki declaration and its later amendments or comparable ethical standards.

Informed consent was waived due to retrospective nature of the study.

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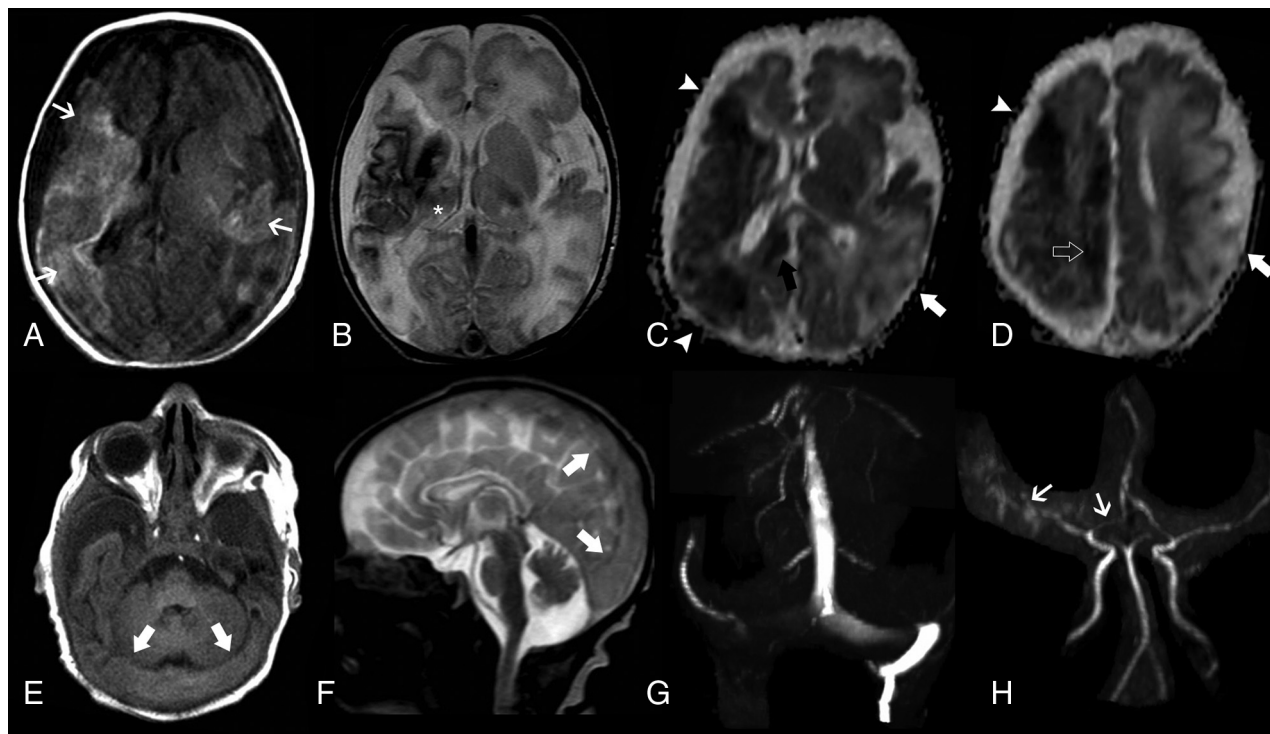


FIG 1. Brain MR imaging of patient 1 performed on day 1. Axial T1-weighted image (A), T2-weighted image (B), and ADC maps (C and D) demonstrate acute ischemic infarcts in the territory of the right MCA (arrowhead) and right anterior cerebral artery (ACA) (open arrow) associated with mild diffuse atrophy of this cerebral hemisphere as well as a late subacute infarct in the territory of the left MCA (arrow). In addition, there are signs of early secondary network injury of the right thalamus (asterisk) and splenium of the corpus callosum (black arrow) as well as subcortical and deep white matter injury and laminar cortical necrosis (small arrows). Axial T1-weighted image (E) and sagittal T2-weighted image (F) depict marked distension of the transverse and superior sagittal sinuses (arrows) with signs of slowing of the venous blood flow, confirmed on the coronal view of MRV (G). MRA (H) shows reduced visualization of the right MCA and ACA branches compared with the contralateral side (small arrows).

association between FVM and perinatal arterial and venous stroke is increasingly acknowledged.^{10,11,13-17} On the other hand, FVM remains an underrecognized cause of PAIS because the onset of clinical manifestations is usually around 12–72 hours after birth or even months later in presumed PAIS, thus limiting the availability of placental tissue for histologic examination.^{2,18,19} Moreover, reports focusing on imaging data of patients with PAIS in the context of proved FVM are still limited, especially within the radiology literature.^{11,13,15-18} Nevertheless, failure to recognize this etiology may have important implications in patient management, as well as in medicolegal actions and future pregnancy counseling.¹⁹ Therefore, it is crucial to increase awareness of FVM-related PAIS in the neuroradiology community and to highlight the importance of knowing the results of placenta examination when reporting these cases.

Here, we describe the neuroradiologic features of 5 neonates with PAIS and histologically proved FVM and their correlation with histopathologic and clinical findings as well as neurologic outcome.

CASE SERIES

Regional review board approval was obtained for this single-center series (N.096/2019), waiving parental written informed consent. Five patients with PAIS and histologically proved FVM were identified from the institutional database of pediatric patients treated for stroke at the Gaslini Children's Hospital between 2012 and 2019. All patients were initially investigated by cranial sonography followed

by brain MR imaging in 4 patients (range: 1 day–2.3 months). Longitudinal brain MR imaging were obtained in 3 neonates (range: 14 days–2.4 years). In total, 11 brain MR imaging and 2 head CT studies were available for review. Images were analyzed in consensus by 2 pediatric neuroradiologists (M.S. and A.F.G., 10 and 5 years' experience, respectively). FVM histologic features were reviewed by 2 pathologists (E.F. and F.B., 30 and 5 years' experience, respectively) and classified into high or low grade.⁹ Clinical data at diagnosis and follow-up (range: 15 days–5 years) were obtained from electronic charts.

On-line Table 1 lists clinical and placental features and the outcomes of our cohort. Neuroimaging features are summarized in On-line Table 2. Parents were nonconsanguineous, and family history was unremarkable in all patients. All neonates were male and born by cesarean section (4/5 urgent procedures). Two neonates were born from dichorionic diamniotic twin pregnancies. Three pregnancies were complicated by threatened preterm labor, preterm premature rupture of membranes, and preeclampsia. Mean gestational age at birth was 34.2 weeks (range: 30–39 weeks). Intrauterine growth restriction was diagnosed in 3/5 patients. Mean birth weight was 1843 g (range: 910–3280 g), length was 42 cm (range: 35–51 cm), and head circumference was 30 cm (range: 26–34 cm). PAIS was symptomatic at birth in 3/5 neonates (clonic seizures and apneas) and was incidentally discovered in 2/5 neonates. In all patients, coagulopathies and congenital cardiac malformations were not detected. Associated clinical findings included a large

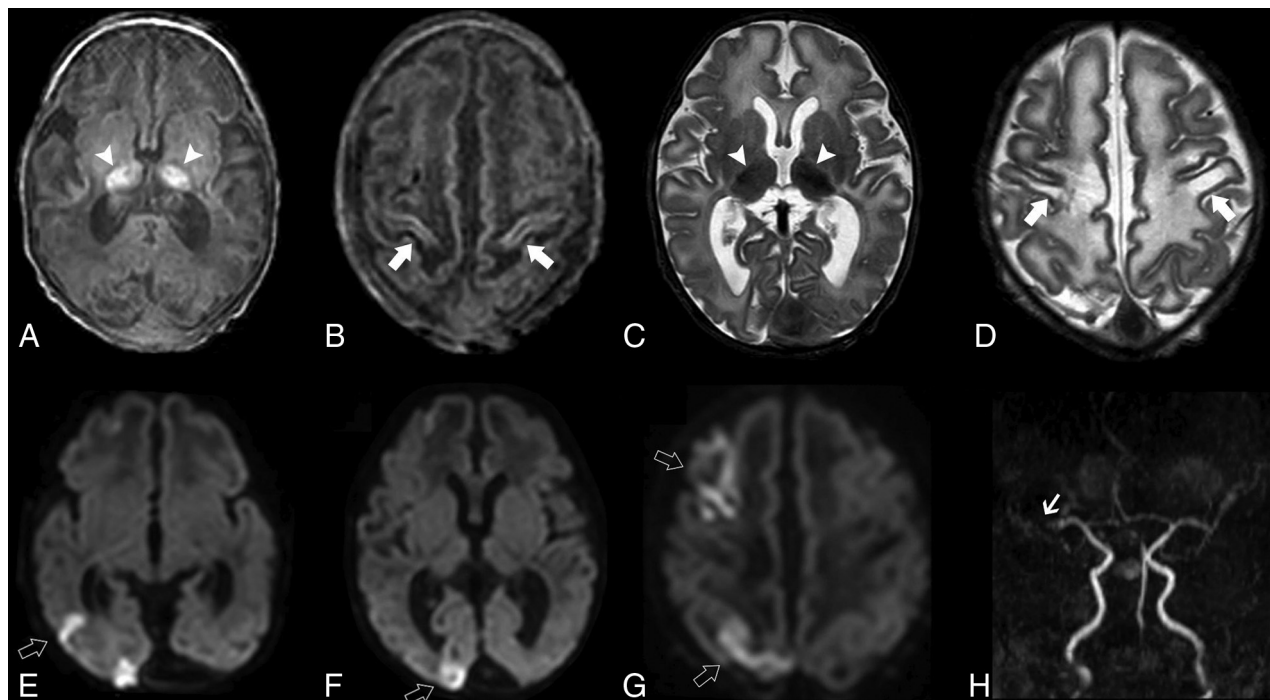


FIG 2. Brain MR imaging of patient 2 performed on day 1. Axial T1-weighted image (A and B), T2-weighted (C and D) and diffusion-weighted (E–G) images show bilateral atrophy and abnormal T1 and T2 signal intensity of the thalami (*arrowheads*) and perirolandic regions (*arrows*) in the context of hypoxic-ischemic lesions. Also note multiple areas of acute ischemic infarct distributed in the right MCA territory and watershed region (*open arrows*). MRA (H) demonstrates reduced flow-related signal of the right MCA branches (*small arrow*).

deep skin ulcer related to distal thromboembolism ($n=1$, On-line Fig 1), transient myocardial ischemia ($n=1$), respiratory distress syndrome of prematurity ($n=3$), and necrotizing enterocolitis ($n=1$).

Cranial sonography findings were abnormal in 4/5 patients, showing unilateral or bilateral WM hyperechogenicity ($n=3$) and a focal irregular cavitation in the parietal subcortical WM ($n=1$, On-line Fig 2).

On imaging, PAISs were multiple in 2/5 patients and involved the anterior circulation in all patients (MCA in 4 neonates, with additional anterior cerebral artery involvement in 1 patient), mostly with a superficial involvement (4/5 patients, On-line Fig 3). In 2/5 patients, PAIS was detected in different phases (acute, subacute, or chronic) at birth (Figs 1 and 2). Signs of early corticobasal diaschisis were noted in 3/4 patients, and early Wallerian degeneration of the corticospinal tract was present in 2/4 patients (Fig 3). Concomitant dural sinus congestion, thrombosis, or both was depicted in 2/5 patients, 1 of whom also displayed signs of chronic hypoxic-ischemic encephalopathy at birth (Figs 1, 2 and On-line Fig 4). MRA at diagnosis was available in 2 patients, showing reduced flow-signal in the involved arteries. Follow-up brain MR imaging and MRA studies demonstrated chronic evolutions of PAIS, without new arterial infarcts, in 2/3 patients. In 1 patient, we observed the phenomenon of DWI reversal of the cortical ischemic lesions, in keeping with spontaneous blood flow restoration in the affected distal MCA branches.²⁰ Resolution of dural sinus congestion and thrombosis was noted in 2/2 patients (On-line Figs 4 and 5).

Histologic examination of the placenta revealed long-standing high-grade FVM in different evolutionary stages in all affected patients (Fig 4). In both dichorionic diamniotic twin pregnancies, FVM was detected in the chorionic disk of each twin with brain injury, but no signs of FVM were found in the chorionic disk of the unaffected co-twin. The placentas of the co-twins presented minor signs of long-standing hypoxic damage and asynchronous patchy ischemic lesions in 1 patient and focal congestion with recent intravillous hemorrhages in the other. Signs of maternal vascular malperfusion were present in 4/5 patients. Umbilical cord abnormalities were present in all affected patients: nuchal cord ($n=2$), hypercoiled ($n=1$), hypocoiled ($n=1$), and false knots with torsion ($n=1$). No signs of inflammation of the umbilical cord, membranes, or amniochorial vessels were depicted in any of the patients.

The preterm neonate who developed necrotizing enterocolitis died at 15 days of life. Mean age at last clinical follow-up of the remaining patients was 28 months (range: 6–60). Moderate to severe developmental delay was diagnosed in 3 children, associated with microcephaly in 2 patients. Motor deficits were present in 3/5 children; 1 developed drug-resistant epilepsy, required noninvasive respiratory support and enteral artificial nutrition through gastrostomy at 9 months of age, and died at 10 months after a respiratory infection. Both co-twins of the 2 affected patients were born with appropriate birth weight, length, and head circumference for gestational age and presented with normal psychomotor development at last follow-up (24 and 36 months, respectively).

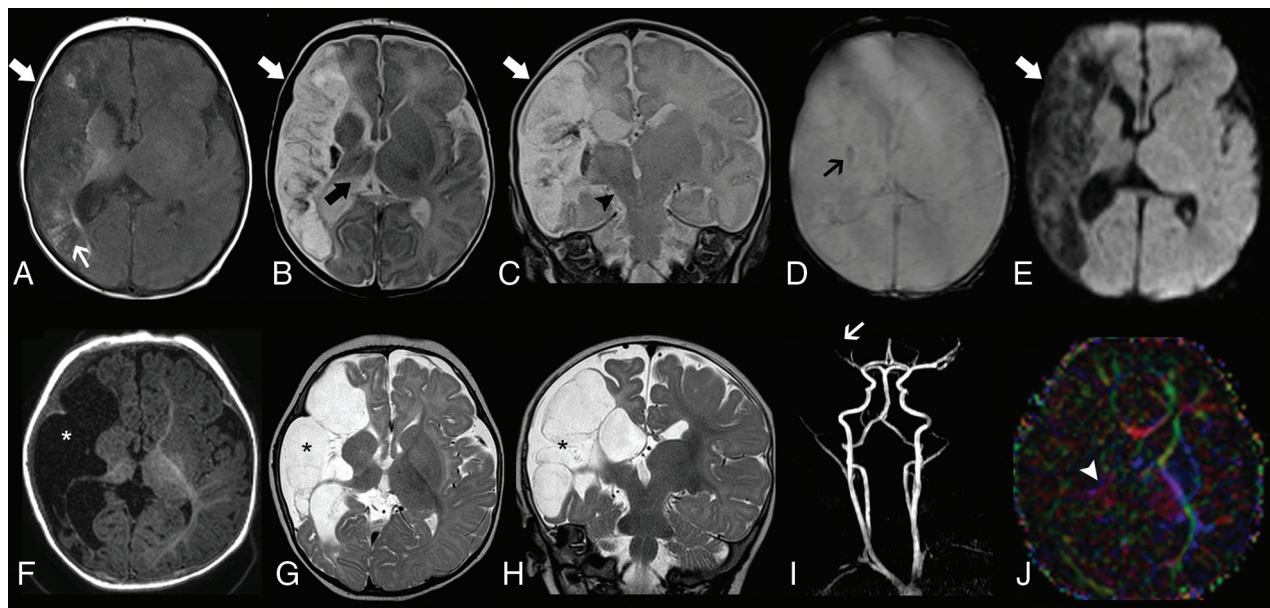


FIG 3. Longitudinal neuroimaging evaluation of patient 5. Initial brain MR imaging obtained at 18 days of life, including axial T1-weighted image (A), axial (B) and coronal (C) T2-weighted images, axial T2* image (D) and diffusion-weighted image (E), demonstrate a large ischemic infarct in the superficial and deep territory of the right MCA in the late subacute or early chronic phase (arrows). There are associated areas of cortical laminar necrosis (thin white arrow) and a focus of hemorrhagic transformation (thin black arrow). Also note signs of ipsilateral thalamic diaschisis (thick black arrow) and Wallerian degeneration of the corticospinal tract (arrowhead). Follow-up brain MR imaging performed at 6 months (F–H). Axial T1-weighted image (A) and axial (B) and coronal (C) T2-weighted images depict progressive cystic encephalomalacia and atrophy in the infarcted area with ex vacuo enlargement of the adjacent CSF spaces (asterisks). MRA reveals reduced visibility of right MCA branches (arrow). Axial directionally encoded color fractional anisotropy (FA) map at the level of the basal ganglia shows markedly reduced FA of the posterior limb of the right internal capsule (arrowhead).

DISCUSSION

Data on neuroimaging features of neonatal stroke associated with thrombotic or inflammatory placental disorders are very rare.^{15,17} In the largest study, which was by Bernson-Leung et al,¹⁷ collecting 46 children with neonatal arterial or venous ischemic stroke from 3 academic medical centers, 13/28 (46.4%) PAISs and 10/18 (55.5%) venous strokes were linked to FVM. Regarding the imaging pattern, PAIS mostly involved the left hemisphere in the MCA territory and was frequently multiple or superficial, but the right hemisphere or perforator territories were affected only in a minority of patients.^{15,17} Similarly, we found that the anterior territories were always involved with sparing of the posterior circulation. In addition, multiple recurrent infarcts and/or small corticosubcortical infarcts were prevalent, suggesting a thromboembolic phenomenon occurring at different phases as the most plausible pathomechanism linking FVM to PAIS. Indeed, because the rate of thrombophilia is similar in placentas with and without FVM,^{12,13} the main hypothesis is the spread of FVM-related thromboemboli into the venous fetal circulation, in turn reaching the aorta and then the intracranial vessels through a patent foramen ovale and a right-to-left direction of blood flow in the fetal ductus arteriosus.^{21,22} Predominant involvement of the MCA territory is also described in most studies including patients with PAIS of mixed or unknown cause,^{3,7,23} though multiple infarcts are usually less frequent (<20%–30%).²³ Interestingly, some of the recurrent infarcts seen in our cohort appeared to be chronic with cortical T1 hyperintensity indicative of laminar necrosis despite having been detected in neonates on day 1; such observation favors an intrauterine onset, as expected to occur in FVM.

Reduced flow signal involving distal arterial branches ipsilateral to the ischemic strokes was present in 2 patients in this series. Intracranial arterial abnormalities have been reported in a subset of neonates with PAIS of variable causes,^{24–26} likely reflecting reduced arterial blood supply to the infarcted territory and/or persistently occluded or near-occluded vessels in the context of a proximal thromboembolism.^{24,25} We did not identify large-vessel occlusions in any of our patients, but these have only been exceptionally described in FVM-related PAIS,^{19,21} probably caused by rapid vessel recanalization in neonates and/or more distal embolization.^{19,27}

Interestingly, 2 of our neonates presented additional features related to severe venous congestion and/or thrombosis involving the superior sagittal and transverse sinuses. The co-occurrence of both arterial and venous complications in the setting of FVM has not been reported before but is theoretically plausible because FVM has been described in association with both PAIS and isolated neonatal cerebral sinovenous thrombosis.^{5,17} In particular, this peculiar combination might be caused by common risk factors underlying FVM, activating coagulation and inflammatory pathways in both the arterial and venous system²⁸ or by secondary marked reduction of venous flow in the dural sinuses caused by diffuse fetoplacental venous stasis.¹⁷ On the other hand, additional risk factors previously described in the literature for neonatal dural sinovenous thrombosis^{29,30} and not necessarily related to FVM, such as emergent cesarean section, might have contributed to venous complications in the present patients. Therefore, future studies on larger series are needed to elucidate the role of FVM in perinatal venous stroke and to estimate the exact prevalence of both arterial and venous abnormalities in neonates with documented FVM.

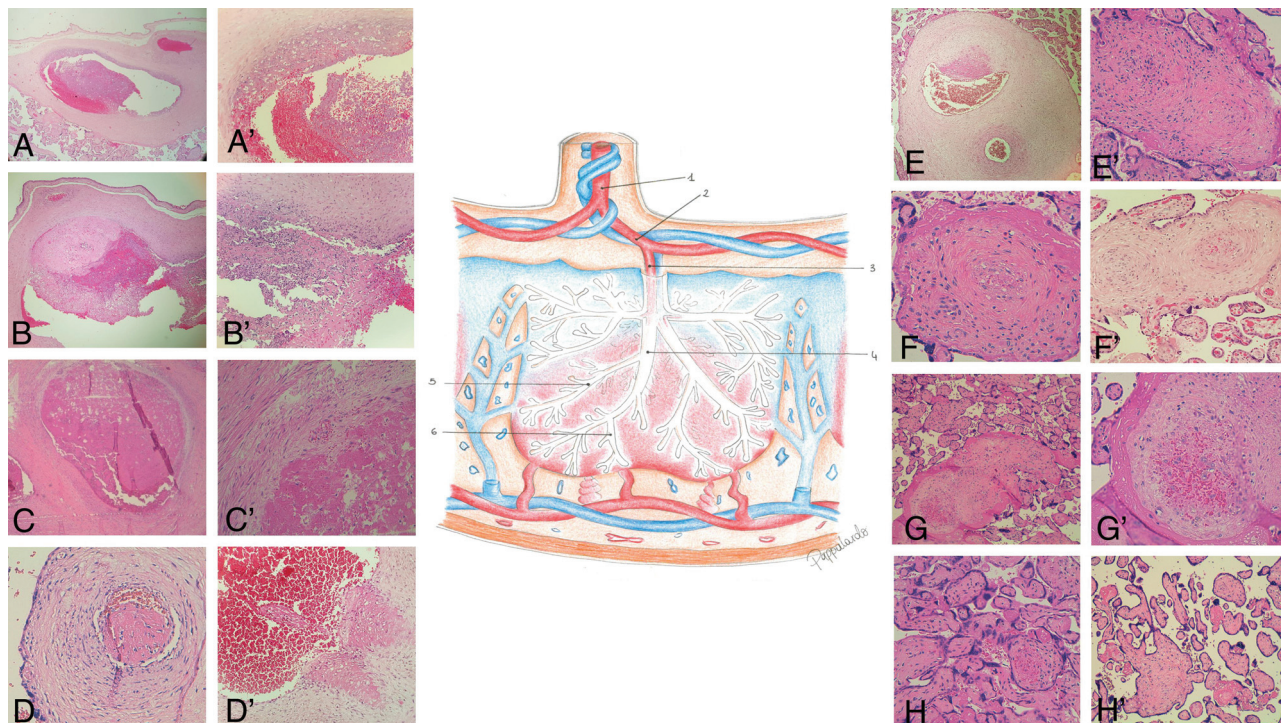


FIG 4. Schematic representation of the placental fetal vascularization and histologic features of fetal vascular malperfusion. 1. Vessel of umbilical cord. 2. Amniochorial vessels (A, A', B, B'). 3. First-order stem vessels (C, C'). 4. Second-order stem vessels (D, D'). 5. Third-order stem vessels (E, E', F). 6. Fourth-order stem vessels (F', G, G', H). A. Amniochorial vessel with dilated lumen occupied by a floating thrombus adhering to the wall by means of coagulation (patient 5). A'. Detail of the previous figure: evidence of wall damage in the form of a necrotic fibrinoid band (patient 5). B. Amniochorial vessel with dilated lumen in which a thrombus in various phases of evolution adheres to the wall; the thrombus has a fibrous nucleus, at the margins of which a recent coagulation and a fibrous thrombus in a medium stage of evolution are observed (patient 5). B'. Detail of B: granulocytic infiltrate of the wall indicating vasculitis as the cause of thrombosis (patient 5). C. First-order stem vessel totally occupied by a long-standing thrombus adherent to the wall (patient 1). C'. Detail of C, a segment of the destructured wall shows evolving lesions after adhesion of the thrombus (patient 1). D. Second-order stem vessel with a thrombotic formation occluding the lumen without adhering to the wall (patient 3). D'. Second-order stem vessel with wall lesion and initial detachment of a floating thrombus into the bloodstream. E. Third-order stem vessel displaying wall damage and initial adhesion of a fibrinoid thrombus. E'. Third-order stem vessel with the lumen completely occluded by a long-standing thrombotic formation hampering the recognition of vessel wall structure (patient 1). F. Third-order stem vessel with a long-standing partly recanalized occlusive thrombus (patient 1). F'. Fourth-order stem vessel: occluded lumen with revascularization; adjacent to this, a totally occluded vessel whose wall structure is no longer recognizable (patient 4). G. Fourth-order stem vessel displaying initial recanalization and endothelial proliferation promoting endoluminal neoangiogenesis (patient 1). G'. Detail of G showing the intermixing of nonagglutinated erythrocytes and incomplete neo-formed vessels (patient 1). H. Fourth-order stem vessel: the lumen is occupied by a long-standing thrombotic formation but displays recanalization; villi undergoing ischemic necrosis caused by perfusion block (patient 1). H'. Group of villi undergoing ischemic necrosis; long-standing lesion integrates the picture of so-called avascularized villi (patient 3).

Another peculiar finding in our series was the concurrence of clinical and imaging signs of chronic hypoxic-ischemic encephalopathy and multiple acute infarcts in 1 neonate at birth. Although it is accepted that FVM is a risk factor for both hypoxic-ischemic encephalopathy and PAIS^{31,32} and both neonatal hypoxic-ischemic encephalopathy and PAIS involve hypoxia-ischemia, patients with a mixed pattern on neuroimaging have not been described in the setting of proven FVM to the best of our knowledge. Such combination, although rare, may be a potential alerting sign of FVM in neonates with PAIS. In addition, as documented in our series and previously described in the literature, the presence of intrauterine growth retardation, thromboembolism in multiple organs, or cardiac complications in neonates with PAIS are also associated with FVM and should raise suspicion of this underestimated condition.^{12,19,33}

Previous studies reported an extremely low rate of recurrence of stroke after PAIS even in the context of FVM,¹⁰ supporting the

concept that risk factors associated with both FVM and PAIS completely subside outside the perinatal period in most patients. Similarly, we did not detect additional infarcts on follow-up studies nor the appearance of clinical manifestations related to new arterial ischemic strokes. Independent of the subjacent etiology, PAIS frequently results in epilepsy, cerebral palsy, and cognitive deficits.^{2,5} In our cohort, 1 preterm neonate died during the neonatal period because of severe necrotizing colitis. Adverse neurologic outcomes were detected in most of the other patients, representing a higher proportion than that described in most series of PAIS.²² However, these data should be considered with caution because they might be related to selection bias caused by nonsystematic evaluation of placenta histology.

High-grade FVM is associated with a higher risk of brain injury.¹⁶ Indeed, all our patients were classified as having high-grade FVM, showing occlusive thrombi, mainly in large-stem vessels, accompanied by downstream changes, including organization,

separation, red cell extravasation, endothelial destruction, loss of vascularity and, ultimately, fibrosis. These changes are similar to those seen in intrauterine fetal demise but are focal rather than diffuse and, especially, in different stages of evolution.⁹ However, no differences in terms of severity and extension of FVM-related lesions were noted among patients with severe or mild imaging patterns in our series. This is in agreement with prior literature showing no stringent correlation between MR imaging and histologic data.¹⁰

Although the likelihood of FVM and PAIS is thought to be increased by endothelial damage caused by fetal inflammatory response,⁹ these lesions were not detected in the present series. On the other hand, as documented in almost all our patients, FVM is frequently associated with other placental abnormalities, such as maternal vascular malperfusion.^{10,34} Interestingly, we had 2 cases of twin dichorionic diamniotic pregnancies, but no co-twin had a stroke or evidence of FVM features in the corresponding placenta, as previously demonstrated.¹⁷ Of note, umbilical cord abnormalities (ie, potentially obstructing anatomic lesions and/or clinical conditions) are considered major risk factors of FVM, probably through promotion of venous obstruction, congestion, and stasis.^{10,11} Accordingly, 4 neonates presented cord abnormalities associated with FVM, including entanglements, hypercoiling, and/or torsion. Unfortunately, placentas are generally discarded soon after delivery, but the clinical presentation of PAIS is often mild and nonspecific, and the diagnosis is most often made a few days or even months after birth, as in the present series.^{2,6} These are major limitations in the diagnosis of FVM in PAIS, and the extent to which different FVM histologic patterns and umbilical cord anomalies may impact on the type of PAIS or brain injury at birth remains an intriguing issue awaiting further exploration.

In conclusion, we confirm the prevalent involvement of the anterior circulation in FVM-related PAISs that were frequently multiple, asynchronous, and distributed superficially, thus suggesting an embolic origin. Moreover, we describe for the first time the co-occurrence of imaging signs of hypoxic-ischemic lesions and/or dural sinus congestion or thrombosis in these neonates, and the association of these features with FVM is worth further investigation. Together with clinical manifestations such as intrauterine growth retardation, thromboembolism in other organs, and cardiac complications, these MR imaging features may raise the suspicion of a subjacent FVM, a rare but likely underdiagnosed condition. Larger prospective studies including placental examination in all neonates are needed to clarify the pathomechanisms of FVM-related PAIS and the correlation between clinical, imaging, and histologic findings.

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REFERENCES

1. Raju TNK, Nelson KB, Ferriero D, et al. Ischemic perinatal stroke: summary of a workshop sponsored by the National Institute of Child Health and Human Development and the National Institute of Neurological Disorders and Stroke. *Pediatrics* 2007;120:609–16 [CrossRef Medline](#)
2. Dunbar M, Kirton A. Perinatal stroke: mechanisms, management, and outcomes of early cerebrovascular brain injury. *Lancet Child Adolesc Health* 2018;2:666–76 [CrossRef Medline](#)
3. Wagenaar N, Martinez-Biarge M, van der Aa NE, et al. Neurodevelopment after perinatal arterial ischemic stroke. *Pediatrics* 2018;142:e20174164 [CrossRef Medline](#)
4. Laugesaar R, Kolk A, Tomberg T, et al. Acutely and retrospectively diagnosed perinatal stroke: a population-based study. *Stroke* 2007;38:2234–40 [CrossRef Medline](#)
5. Kirton A, deVeber G. Life after perinatal stroke. *Stroke* 2013;44:3265–71 [CrossRef Medline](#)
6. Mineyko A, Kirton A. The black box of perinatal ischemic stroke pathogenesis. *J Child Neurol* 2011;26:1154–62 [CrossRef Medline](#)
7. Kirton A, Armstrong-Wells J, Chang T, et al. Symptomatic neonatal arterial ischemic stroke: the International Pediatric Stroke Study. *Pediatrics* 2011;128:e1402–10 [CrossRef Medline](#)
8. Redline RW, Pappin A. Fetal thrombotic vasculopathy: the clinical significance of extensive avascular villi. *Hum Pathol* 1995;26:80–85 [CrossRef Medline](#)
9. Khong TY, Mooney EE, Ariel I, et al. Sampling and definitions of placental lesions: Amsterdam Placental Workshop Group Consensus Statement. *Arch Pathol Lab Med* 2016;140:698–713 [CrossRef Medline](#)
10. Redline RW, Ravishanker S. Fetal vascular malperfusion, an update. *APMIS* 2018;126:561–69 [CrossRef Medline](#)
11. Redline RW. Severe fetal placental vascular lesions in term infants with neurologic impairment. *Am J Obstet Gynecol* 2005;192:452–57 [CrossRef Medline](#)
12. Saleemuddin A, Tantbirojn P, Sirois K, et al. Obstetric and perinatal complications in placentas with fetal thrombotic vasculopathy. *Pediatr Dev Pathol* 2010;13:459–64 [CrossRef Medline](#)
13. Roescher AM, Timmer A, Erwich JJHM, et al. Placental pathology, perinatal death, neonatal outcome, and neurological development: a systematic review. *PLoS One* 2014;9:e89419 [CrossRef Medline](#)
14. Kraus FT, Acheen VI. Fetal thrombotic vasculopathy in the placenta: cerebral thrombi and infarcts, coagulopathies, and cerebral palsy. *Hum Pathol* 1999;30:759–69 [CrossRef Medline](#)
15. Elbers J, Viero S, MacGregor D, et al. Placental pathology in neonatal stroke. *Pediatrics* 2011;127:e722–29 [CrossRef Medline](#)
16. Chisholm KM, Heerema-McKenney A. Fetal thrombotic vasculopathy. *Am J Surg Pathol* 2015;39:274–80 [CrossRef Medline](#)

17. Bernson-Leung ME, Boyd TK, Meserve EE, et al. **Placental pathology in neonatal stroke: a retrospective case-control study.** *J Pediatr* 2018;195:39–47.e5 [CrossRef Medline](#)
18. Fluss J, Garcia-Tarodo S, Granier M, et al. **Perinatal arterial ischemic stroke related to carotid artery occlusion.** *Eur J Paediatr Neurol* 2016;20:639–48 [CrossRef Medline](#)
19. Heider A. **Fetal vascular malperfusion.** *Arch Pathol Lab Med* 2017;141:1484–89 [CrossRef Medline](#)
20. Nagaraja N, Forder JR, Warach S, et al. **Reversible diffusion-weighted imaging lesions in acute ischemic stroke: a systematic review.** *Neurology* 2020;94:571–87 [CrossRef Medline](#)
21. Cheong JLY, Cowan FM. **Neonatal arterial ischaemic stroke: obstetric issues.** *Semin Fetal Neonatal Med* 2009;14:267–71 [CrossRef Medline](#)
22. Chabrier S, Saliba E, Nguyen The Tich S, et al. **Obstetrical and neonatal characteristics vary with birthweight in a cohort of 100 term newborns with symptomatic arterial ischemic stroke.** *Eur J Paediatr Neurol* 2010;14:206–13 [CrossRef Medline](#)
23. Husson B, Hertz-Pannier L, Renaud C, et al. **Motor outcomes after neonatal arterial ischemic stroke related to early MRI data in a prospective study.** *Pediatrics* 2010;126:e912 [CrossRef Medline](#)
24. Husson B, Hertz-Pannier L, Adamsbaum C, et al. **MR angiography findings in infants with neonatal arterial ischemic stroke in the middle cerebral artery territory: a prospective study using circle of Willis MR angiography.** *Eur J Radiol* 2016;85:1329–35 [CrossRef Medline](#)
25. Siddiq I, Armstrong D, Surmava A-M, et al. **Utility of neurovascular imaging in acute neonatal arterial ischemic stroke.** *J Pediatr* 2017;188:110–14 [CrossRef](#)
26. Wintermark M, Hills NK, DeVeber GA, et al. **Clinical and imaging characteristics of arteriopathy subtypes in children with arterial ischemic stroke: results of the VIPS study.** *AJNR Am J Neuroradiol* 2017;38:2172–79 [CrossRef Medline](#)
27. van der Aa NE, Porsius ED, Hendrikse J, et al. **Changes in carotid blood flow after unilateral perinatal arterial ischemic stroke.** *Pediatr Res* 2012;72:50–56 [CrossRef Medline](#)
28. Prandoni P. **Venous and arterial thrombosis: is there a link?** *Adv Exp Med Biol* 2017;906:273–83 [CrossRef Medline](#)
29. Rutherford MA, Ramenghi LA, Cowan FM. **Neonatal stroke.** *Arch Dis Child Fetal Neonatal Ed* 2013;98:F180 [CrossRef Medline](#)
30. Ramenghi LA, Cardiello V, Rossi A. **Neonatal cerebral sinovenous thrombosis.** *Handb Clin Neurol* 2019;162:267–80 [CrossRef Medline](#)
31. Mir IN, Johnson-Welch SF, Nelson DB, et al. **Placental pathology is associated with severity of neonatal encephalopathy and adverse developmental outcomes following hypothermia.** *Am J Obstet Gynecol* 2015;213:849.e1–7–849.e7 [CrossRef Medline](#)
32. Vik T, Redline R, Nelson KB, et al. **The placenta in neonatal encephalopathy: a case-control study.** *J Pediatr* 2018;202:77–85.e3 [CrossRef Medline](#)
33. Gluck O, Schreiber L, Marciano A, et al. **Pregnancy outcome and placental pathology in small for gestational age neonates in relation to the severity of their growth restriction.** *J Matern Fetal Neonatal Med* 2019;32:1468–73 [CrossRef](#)
34. Zur RL, Kingdom JC, Parks WT, et al. **The placental basis of fetal growth restriction.** *Obstet Gynecol Clin North Am* 2020;47:81–98 [CrossRef Medline](#)