

Collateral blood vessels in acute ischemic stroke: a physiological window to predict future outcomes

Circulação colateral no acidente vascular cerebral isquêmico: uma janela fisiológica para prever resultados futuros

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ABSTRACT

Collateral circulation is a physiologic pathway that protects the brain against ischemic injury and can potentially bypass the effect of a blocked artery, thereby influencing ischemic lesion size and growth. Several recent stroke trials have provided information about the role of collaterals in stroke pathophysiology, and collateral perfusion has been recognized to influence arterial recanalization, reperfusion, hemorrhagic transformation, and neurological outcomes after stroke. Our current aim is to summarize the anatomy and physiology of the collateral circulation and to present and discuss a comprehensible review of the related knowledge, particularly the effects of collateral circulation on the time course of ischemic injury and stroke severity, as well as imaging findings and therapeutic implications.

Keywords: stroke; brain ischemia; collateral circulation; tomography; angiography; perfusion.

RESUMO

A circulação colateral é um circuito fisiológico de proteção contra alterações isquêmicas que, potencialmente, evita os efeitos de uma oclusão arterial e com isso pode influenciar nas dimensões e no crescimento de uma lesão isquêmica. Vários estudos recentes forneceram informações a respeito do papel das colaterais na fisiopatologia do acidente vascular encefálico isquêmico e demonstraram a capacidade da circulação colateral de influenciar as taxas de reperfusão, recanalização, transformação hemorrágica e com isso desfecho clínico dos pacientes. O objetivo desta revisão é sintetizar a anatomia e a fisiologia da circulação colateral encefálica, apresentando e discutindo, o que se conhece atualmente acerca do seu efeito na cronologia e gravidade da lesão isquêmica, além dos achados de imagens e implicações terapêuticas.

Palavras-chave: acidente vascular cerebral; isquemia encefálica; circulação colateral; tomografia; angiografia; perfusão.

Stroke is the second most common cause of death and was responsible for approximately 6.7 million deaths worldwide in 2012¹. Ischemia, or restricted blood flow, is the main cause of stroke and is typically due to abrupt occlusion of a cerebral artery as a result of progressive atherosclerosis or embolism². Acute ischemic stroke (AIS) can result in severe neurologic disability or death³.

Since the late 1990s intravenous thrombolysis has been a recommended treatment for AIS^{4,5}. Additionally, in the last decade, several clinical trials have investigated the effects of endovascular treatment (EVT) in the setting of an intracranial or extracranial large artery occlusion. Several studies, including MR CLEAN⁶, ESCAPE³, EXTEND-IA⁷ and SWIFT-PRIME⁸,

recently proved EVT to be more effective than standard medical care, with or without intravenous thrombolysis, using stentriever in the majority of the patients in the EVT arms.

However, although EVT has been shown to be generally effective, the trials have documented erratic individual and overall patient outcomes. These differences may not be solely the result of the various methods used but may also be related to patient-specific characteristics⁹. Among these patient-specific characteristics, collateral status has emerged as an independent factor that is associated with angiographic and clinical outcomes in AIS patients^{10,11}.

The collateral circulation is a physiologic pathway of specialized endogenous bypass vessels that is present in most

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tissues and protects against ischemic injury during initial oligemic status¹². In the setting of AIS, the extent of collateral circulation influences the size of the final infarct and the growth of the penumbra. Hence, the relationship between the collateralization grade and the predictability of infarct evolution has been a primary focus in recent years^{11,13,14}.

The refinement of diagnostic techniques for evaluation of collateral circulation may contribute to improved anatomical and pathophysiological characterization of this vascular network and its potential therapeutic and prognostic implications¹⁵. The ability to define physiologic parameters through diagnostic techniques is particularly useful when the time of the stroke is unknown or a wider nonconventional window for treatment is being considered.

In this review, we summarize the basic anatomy and physiology of the collateral circulation and its potential as an endogenous therapeutic target in AIS. The relevance of multidetector computed tomography (MDCT) in clinical settings to support medical decisions regarding AIS is highlighted, and recent evidence indicating that good collateral circulation can prevent or delay permanent neural damage is presented.

THE ANATOMY OF COLLATERAL CIRCULATION

Two main routes underlie collateral perfusion of the brain parenchyma. The anatomy of this arterial circulation includes extracranial sources of blood flow that can supply intracranial vessels, as well as intracranial routes that can supplement other intracranial areas when pathophysiologic mechanisms become activated¹⁵.

The extracranial sources consist of large connections between the extracranial and intracranial arteries. The external carotid artery gives rise to many branches in the neck that are potential sources of collateral blood flow, particularly when chronic stenosis or occlusion has developed in the internal carotid artery¹⁶. The facial, maxillary, middle meningeal, and occipital arteries are the main branches that can shunt flow via anastomoses to the intracranial arteries. Apart from these branches, common anastomotic routes include the ophthalmic artery, which may fill in a retrograde direction, as well as smaller and unnamed dural arteries¹⁷.

Intracranial collateral routes can be further subdivided into primary and secondary routes. The primary pathways include the permanently active components of the circle of Willis, and the secondary pathways include less direct routes that develop over time. The blood supply to the brain is unique because four major arteries coalesce to form an equalizing distributor, i.e., the circle of Willis, which, despite its variability and asymmetry, can redistribute blood flow in the event of sudden occlusion of a parent vessel.

The secondary pathway comprises leptomeningeal anastomoses that link distal sections of the major cerebral

arteries. It has been reported that some small arteriolar connections (~50–400 μm) allow retrograde perfusion of adjacent territories¹⁸. It is assumed that these connections are important routes for collateral blood flow, especially when an acute arterial occlusion occurs. These arteriolar anastomoses mimic the circle of arteries but connect a much larger extension of the microvasculature, joining territories of the middle cerebral artery (MCA) with both the anterior cerebral artery (ACA) and the posterior cerebral artery (PCA)¹⁹.

The development of native pial collateral circulation (collaterogenesis), which begins in the embryo, has been shown to determine the extent of the collaterals in adulthood²⁰. Acute obstructions induce blood flow across the collateral network (recruitment) followed by remodeling and, potentially, formation of additional collaterals in chronic obstructive disease (neocollateral formation)¹⁷.

There is wide variation in collateral status among healthy adults, and recent animal studies indicate that genetic background may be a major factor²¹. {Zhang, 2010 #110} A single polymorphic locus on chromosome 7 in mice, i.e., the determinant of collateral extent 1 (Dce1), has been shown to influence the extent of collateralization, blood flow and infarct volume after middle cerebral artery occlusion²². Whether human Dce1 or related loci are responsible for the wide variation in collateral status in humans is still under investigation.

A number of other factors, including environmental and clinical features, have also been shown to affect the quality and quantity of collaterals (rarefaction) at the time of presentation in AIS. Of these, the strongest predictor by far is age²³. Other clinical features include elevated glucose at the time of presentation, uric acid level, history of hypertension, and history of smoking^{24,25}.

PHYSIOLOGY OF COLLATERAL BLOOD FLOW REGULATION

The importance of the collateral circulation in brain physiology may be demonstrated with the concept of the collaterome, which is an extension of the connectome concept²⁶. The collaterome provides a physiologically relevant approach to the management of stroke and the influential balance of collateral perfusion that determines both stroke evolution and related clinical sequelae²⁷.

Beyond structural assessments of collateral circulation, advances in perfusion-based imaging have allowed for functional evaluations of the quality of collateral blood flow (effective parenchymal perfusion). Cerebral blood flow (CBF) is regulated by the metabolic demands of the brain itself, which vary regionally and with neuronal activity. Although the precise mechanisms underlying cerebral autoregulation are not fully understood, the process seems to be mediated at several levels and involves neurons, neuropil, and cerebral blood vessels²⁸.

Normal CBF ranges between 50 and 60 mL/100 g/min and is tightly controlled by cerebral autoregulation²⁹. The pace of cellular death in the brain after an arterial occlusion is closely linked to the severity of the decrease in blood flow within the local environment. When blood flow is less than 10 mL/100 g/min, damage is rapid, and most cells die within minutes of the insult^{30,31}. When CBF is between 10 and 20 mL/100 g/min (hypoperfusion), neurons cease to function but remain structurally intact and are potentially revivable if normal blood flow is restored³¹. Therefore, neuronal damage is not uniform when an intracranial artery is occluded, especially in the first few hours after an insult. Depending on the extent of collateral perfusion, infarction may not be complete for hours or even days³².

In thrombotic and embolic strokes, the intravascular pressure distal to the occlusion falls immediately. Concurrently, the pressure within the pial vessels is relatively well preserved, resulting in a gradient that is able to promote flow through anastomoses¹⁸. The effectiveness of collateral vessel flow can be assessed only with measurements of tissue perfusion, which reflect the statuses of both the microcirculation and the macrocirculation. Computed tomography (CT) and magnetic resonance imaging (MRI) perfusion techniques and other methods, such as positron emission tomography (PET) and single-emission computed tomography (SPECT), can provide insight into the collateral flow in patients with cerebrovascular disease^{17,33,34}. Physiologically effective collateral perfusion is evident when CBF and cerebral blood volume (CBV) are maintained within the territory of an occluded artery.

IMAGING OF COLLATERAL VESSELS

Digital subtraction angiography (DSA) remains the gold standard for the anatomic evaluation of the collateral circulation. This technique allows for the dynamic visualization of blood flow through pial collaterals or other secondary collaterals^{17,35}. The main limitations of DSA are its invasive nature, its reliance on iodinated contrast and ionizing radiation, and its inability to evaluate brain parenchyma. Furthermore, performing DSA in AIS when intra-arterial therapy is not considered may generate an additional delay to treatment³⁶.

Several noninvasive approaches have been proposed to evaluate intracranial collateral blood flow and the network, but none of these techniques have been shown to be as effective as a reference standard for quantifying collateral flow³⁷.

Computed tomography angiography (CTA) is fast, reproducible, and widely available, and its reasonable cost-to-effectiveness ratio makes this technique one of the most widely used methods of evaluating the locations of vascular occlusions and the collateral system. Analysis of CTA source images (CTA-SI) has a higher sensitivity for demonstrating the infarct core than non-contrast computed tomography (NCCT)³⁸. Post-processed CTA data involving maximal intensity projections (MIP) and multiplanar

reconstruction (MPR) allow for better visualization of the occluded vessel and the extent of leptomeningeal flow³⁹. The main limitation of CTA is that it is a snapshot of arterial contrast enhancement, providing limited information about flow dynamics. Some studies have attempted to circumvent this limitation with dual-phase CTA⁴⁰. Such variations in acquisition protocols and differences in classification have led to low levels of agreement among the relevant results even among experienced observers (K-alpha 0.3-0.6)⁴¹.

Dynamic CTA has become available for clinical practice in recent years and merges the noninvasive nature of CTA and the dynamic acquisition of DSA. This technique, also referred to as 4D-CTA, enables the noninvasive evaluation of the flow dynamics of the intracranial vasculature by multiple subsequent CT acquisitions or continuous volume CT acquisition over a period of time⁴².

Several protocols of acquisition have been proposed, including a toggling-table technique, shuttle mode scanning, and volume mode. The volume mode is considered the most versatile option and allows for complete or partial coverage of the whole brain during 1 rotation of the scanner⁴². Dynamic acquisitions in volume mode can be performed discontinuously or continuously, depending on the required temporal resolution. When collateral flow, as in the case of an arterial occlusion, needs to be evaluated, a lower temporal resolution is necessary. In the setting of AIS, 4D-CTA better estimates thrombus burden and the presence of collateral vessels than conventional CTA⁴². The challenge of the radiation dose level remains, although recently available noise-reduction filters have dramatically reduced radiation exposure⁴³.

Magnetic resonance angiography (MRA) using time-of-flight technique (TOF) is one of the most used MR techniques for accessing collateral circulation, but it remains controversial. MRA provides structural information based on flow-sensitive images but is less effective for collateral evaluations than CTA, especially when the objective is to estimate the occlusion of distal branches^{36,44}. Contrast-enhanced MRA (CE-MRA) allows better delineation of slow-moving blood in the distal branches and is a better predictor of infarct outcome, but it provides lower spatial resolution⁴⁴.

Transcranial Doppler (TCD) also provides information regarding cerebral autoregulation and cerebral circulation. Flow direction changes, such as those found in the ophthalmic artery, and increased velocity in vessels ipsilateral to a stenosis are correlated with the presence of leptomeningeal collaterals⁴⁵. However considerable variability has been found in TCD performance and interpretation^{35,45}.

IMAGING STUDIES OF COLLATERAL CIRCULATION IN THE CLINICAL SETTING OF ACUTE ISCHEMIA

The concept that a vascular network can potentially bypass the effects of a blocked cerebral artery and influence

ischemic lesion size and growth^{37,46} has recently had an increased impact on the management of stroke patients⁴.

Assessments of collaterals in angiographic studies have proven be widely variable. A recent meta-analysis by Leng et al.⁹ found 12 studies that used the American Society of Interventional and Therapeutic Neuroradiology/Society of Interventional Radiology (ASITN/SIR) collateral flow grading system by DSA and primarily defined grades 3-4 and 0-2 as good and poor collateral statuses, respectively. Eleven studies have used other grading methods for DSA, 9 studies have used different grading methods for CTA, and others have used CTP or combined grading methods with different imaging modalities.

Two DSA classifications stand out: Higashida⁴⁷ used the ASITN/SIR assessment, which was based on the extent and delay of retrograde filling, whereas Christoforidis⁴⁸ proposed a distinct model that is based solely on the extent of the feedback.

The ASITN/SIR classification⁴⁷ seems to be the most appropriate because it has higher reproducibility and accounts for both the parameters of the extent and the delay of the feedback via collaterals. The ASITN/SIR classification involves a five-point system (Table 1) that has been used in several endovascular trials^{9,10,49,50,51,52}. Grades 0 and 1 indicate only marginal flow, grade 2 indicates only partial filling of the ischemic territory, and grades 3 and 4 indicate varying rates of complete filling of the occluded arterial territory.

Additional non-invasive grading systems for assessing collateral blood flow circulation have also been described⁵³. Several strategies have been used to describe CTA-visualized vessels by either comparing them with the contralateral brain hemisphere or estimating the percentage of MCA branches that become filled by contrast media during examination^{54,55}.

Souza et al.⁵⁶ proposed a simple and reliable grading system that correlates collateral scores and diffusion-weighted imaging (DWI) lesion volumes on admission.

The best method of evaluating and grading collateral flow remains controversial. The currently proposed methods of assessing collaterals are largely qualitative or semiquantitative, and there are no clear indications of the superiority of any of the available techniques⁵⁷. This lack of consensus may contribute to an overall under appreciation of the fundamental role of collateral circulation in outcomes following AIS.

Nevertheless, Shet and Liebeskind⁵⁸, when revising the status of collateral circulation in endovascular therapy for stroke, concluded that collateral blood supply is pivotal in determining clinical outcomes. In the setting of vascular occlusion, patients with more robust collaterals have smaller infarcts (Figure 1)^{17,59}. Regardless of the method used to determine the collateral score,

Table 1. American Society of Interventional and Therapeutic Neuroradiology / Society of interventional Radiology (ASITN/SIR) collateral grade scale.

Grade	Angiographic collaterals (Digital subtraction angiography)
0	No collaterals visible to the ischemic site
1	Slow collaterals to the periphery of the ischemic site with persistence of some of the defect
2	Rapid collaterals to the periphery of ischemic site with persistence of some of the defect and to only a portion of the ischemic territory
3	Collaterals with slow but complete angiographic blood flow of the ischemic bed by the late venous phase
4	Complete and rapid collateral blood flow to the vascular bed in the entire ischemic territory by retrograde perfusion

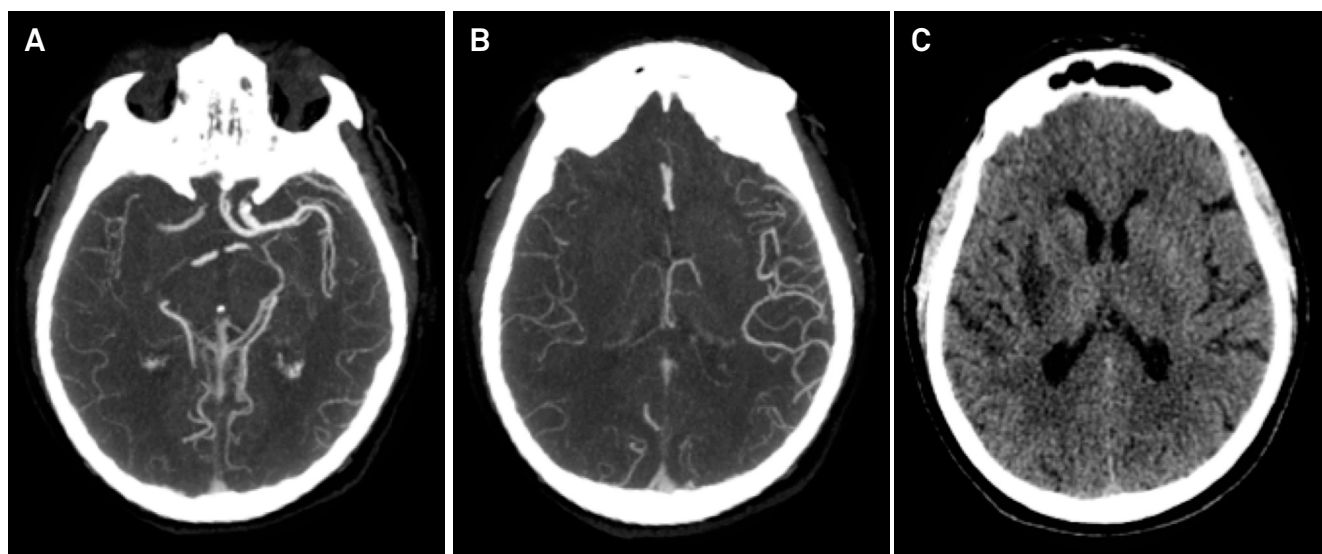


Figure 1. Proximal occlusion with suitable reconstitution of the distal middle cerebral artery (MCA) branches (Mittelf grade 3). (A) Computed tomography angiography (CTA) identified right MCA occlusion. (B) Note collateral circulation throughout the MCA territory, filled from leptomeningeal branches. (C) Follow-up non-contrast compute tomography (NCCT) 24 hours later demonstrating a small final infarct volume confined to the lenticulostriate territory (proximal MCA); this patient exhibited a good clinical outcome.

extremely poor outcomes are predicted when collateral blood flow is reduced or absent (Figures 2 and 3)^{60,61,62,63}.

Physiology is now considered to be more relevant than time in AIS because it is the degree of collateral flow and not simply the time elapsed since the stroke that is the main factor determining core infarct volume within the first 6 hours of stroke onset⁴⁶. Collateral status is recognized as having an influence on stroke prognosis, particularly in terms of recanalization, reperfusion, hemorrhagic transformation and subsequent neurological outcomes⁵⁹. Collateral grading (Table 2) may represent a currently available opportunity to predict possible future outcomes, whereas elapsed time solely reflects the past and does not have a causal connection with the future of infarcted brain tissue.

RELATIONSHIP WITH PARENCHYMAL PERFUSION

Collaterals are usually evaluated by examining arterial flow with angiography techniques, and parenchymal perfusion is profoundly influenced by downstream microcirculation⁵⁹. The microcirculation is crucial to the restoration of the blood supply to the brain, and collateral circulation may increase ischemic tolerance by enhancing microvascular perfusion³⁵.

A relatively simple method of evaluating brain perfusion and viable tissue is the assessment of capillary blush using DSA. The capillary index score (CIS) can define an ischemic area in two ways: either by a lack of anterograde flow (in an area that receives blood supply in a retrograde fashion through pial collaterals) or by a significant delay in anterograde flow due to a proximal partially recanalized clot⁶⁴. This angiographic index has been shown to be a good predictor of outcomes and a powerful strategy for improving outcomes in endovascular treatment⁶⁵.

Beyond structural assessments, perfusion-based images have allowed functional evaluations to differentiate critically hypoperfused areas (infarct cores), penumbral areas (potentially savable areas) and benign oligemic tissues. Current evidence indicates that the goals of acute stroke treatment should be to determine tissue viability by noninvasive techniques, use this information to individualize thrombolytic therapy, extend the therapeutic time window and rescue penumbral tissue⁴⁶. Therefore, knowing the factors that influence the loss of penumbral tissue is crucial, and collaterals have emerged as a major feature that is relevant to this knowledge.

The presence of robust collaterals both markedly reduces and slows down penumbra loss¹⁴. This can be seen in patients without significant reperfusion after treatment, reinforcing the fact that poor collaterals alone are able to predict larger infarcts. In contrast, smaller infarcts, at least in part, result from good collaterals. Reperfusion can occur not only via successful recanalization of the primary occlusion and restoration of downstream flow but also via viable collateral blood flow⁶⁶. However, it is important to note that mismatch (the difference between the penumbra and the infarct core) is also an independent prognostic factor with a strong association with better outcomes in target mismatch patients, adding information to the study of collateral profiles⁶⁷.

Some studies have demonstrated the use of ischemia-induced vascular damage estimates in AIS in combination with collateral scores and brain perfusion analysis to predict hemorrhagic complications^{17,68,69}. In the setting of poor collaterals, a finding of hyperperfusion may indicate a higher risk of hemorrhagic transformation. When revascularization is achieved, symptomatic hemorrhagic transformation may occur more frequently in patients who have presented with poor collaterals⁴⁹. Higher frequencies of infarct growth and symptomatic hemorrhagic

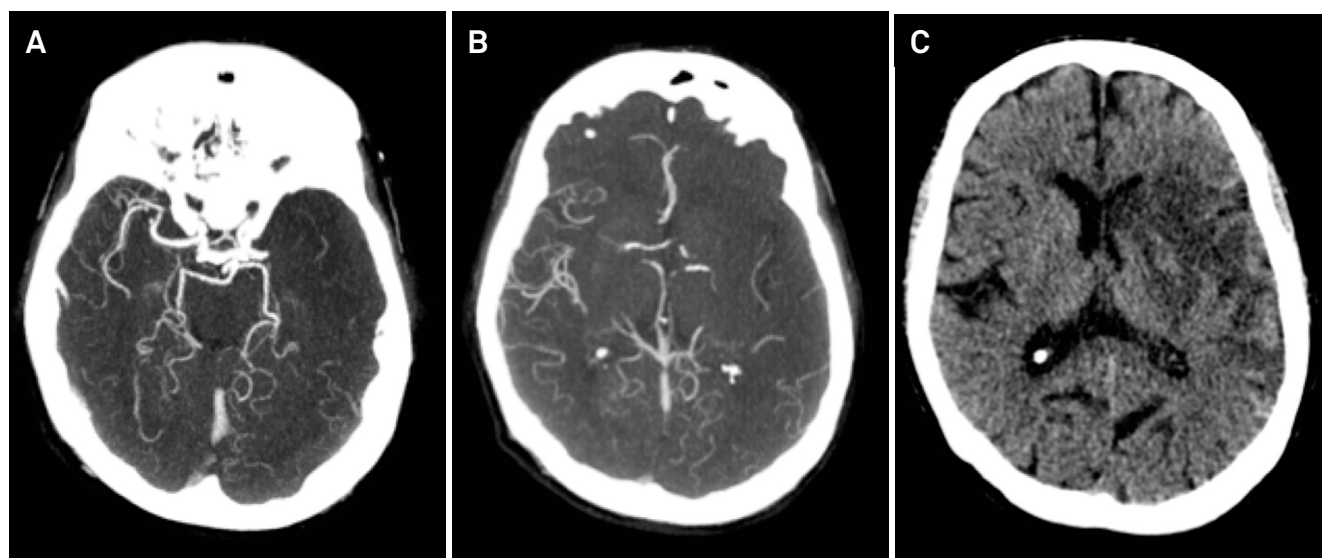


Figure 2. Proximal occlusion with vessel filling restricted to the Sylvian fissure (Mittelf grade 2). (A) A left Mittef grade 1 (M1) occlusion was demonstrated on computed tomography angiography (CTA), whereas only minimal collateral filling was noted in the middle cerebral artery (MCA) territory (B). (C) Follow-up non-contrast compute tomography (NCCT) 48 hours later illustrating a large infarct volume involving the lenticulostriate territory, insula and temporal lobe.

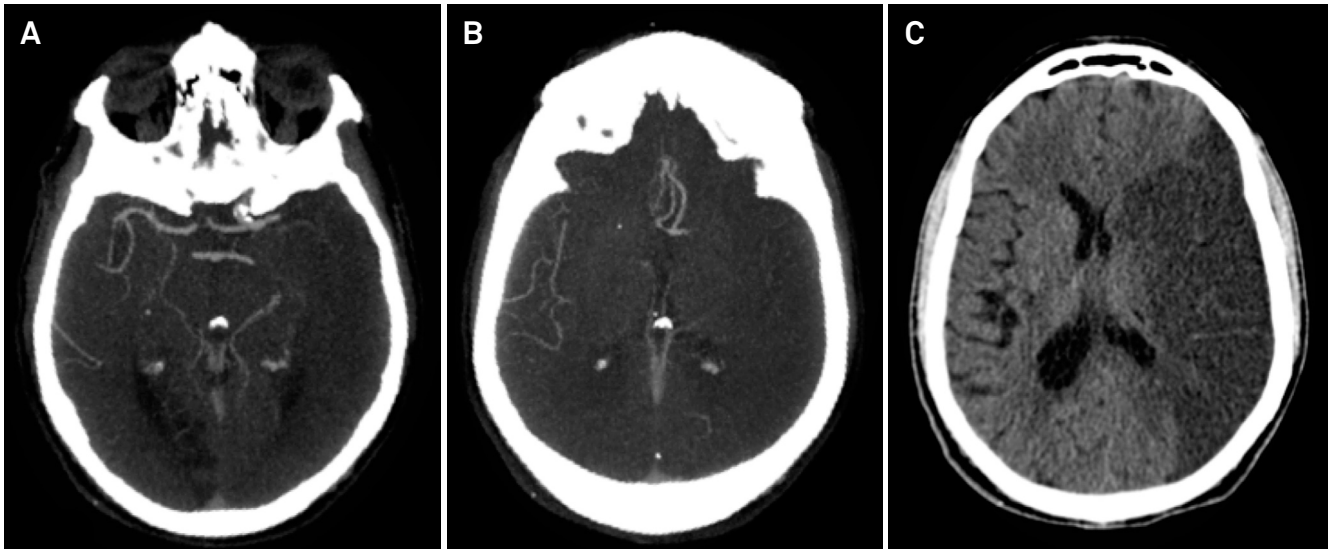


Figure 3. Proximal occlusion with faint contrast opacification restricted to the distal superficial branches (Miteff grade 1 – M1). (A) A left M1 occlusion was found on computed tomography angiography (CTA), and the collateral circulation in the middle cerebral artery (MCA) territory was considered absent (B). (C) Follow-up non-contrast computed tomography (NCCT) 24 hours later demonstrating a malignant infarct. This patient died 3 days later.

Table 2. Collateral grade by Miteff system.

Grade	Computed tomography angiography collaterals
1	The contrast opacification is merely seen in the distal superficial branches
2	Vessels can be seen at the Sylvian fissure
3	If the vessels are reconstituted distal to the occlusion

transformation in patients with poor collaterals in whom therapeutic recanalization has been achieved may support the concept of reperfusion injury⁷⁰.

TEACHING POINTS FOR CLINICAL USE

Larger recent trials have established the use of intracranial vascular studies in the setting of AIS to detect proximal obstructions, define the treatment subtype and select an intraarterial approach^{3,6,7}. Therefore, the collateral profile should be determined in all of these patients. Some critical findings may be established with the knowledge of collateral status.

Abundant native (preexisting) collateral circulation is directly correlated with better clinical status and smaller volumes of infarcted brain¹⁰. Improved collateral circulation also

predicts higher rates of recanalization, favorable outcomes and lower rates of mortality^{9,10,50}.

The absence or relative paucity of a collateral network is a major predictor of extensive infarct on admission⁵⁶, and a proximal thrombus associated with such an absence has been termed a “malignant profile”⁷¹. Recent studies have also demonstrated that worse collaterals are associated with increased hemorrhagic complications, reinforcing the relevance of specific approaches for these patients^{49,70}.

FINAL REMARKS

This literature review supports the view that noninvasive vascular studies should be used to identify proximal arterial occlusions and to estimate collateral grading in the setting of AIS. A personalized approach that is not solely restricted by time should be provided to maximize the effect of therapy, including appropriate patient selection for EVT.

The currently available knowledge has increased the pathophysiologic understanding of intrinsic compensatory vascular mechanisms, supports the use of MDCT techniques to rapidly evaluate hyperacute AIS, and provides evidence for therapeutic decisions. Modern techniques for reducing radiation exposure should be employed to ensure that diagnostic tests preserve patient safety.

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