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An 18-year-old patient presenting right holocranial headache and convulsive seizures, with progressive hemiparesis at left.

*Figure 1.* Skull computed tomography.

*Figure 2.* Skull magnetic resonance imaging.

*Figure 3.* Arteriogram.
Images description

Figure 1. Non-contrast-enhanced (A) and contrast-enhanced (B) skull computed tomography demonstrating a diffuse meshwork of tortuous, dilated vessels in the right cerebral hemisphere.

Figure 2. Skull magnetic resonance imaging: contrast-enhanced axial, T1-weighted image demonstrating multiple abnormal vessels intermingled with healthy cerebral parenchyma (A,B), and MRI angiogram of intracranial vessels, venous phase (C), demonstrating anterior, middle and posterior right afferent cerebral arteries, and capillary ectasias with cortical reflux.

Figure 3. Digital angiogram of right external carotid artery (lateral) demonstrating transdural shunts (A) and right internal carotid artery (lateral), arterial and venous phases demonstrating capillary ectasias and drainage through cortical and deep veins (B,C).

Diagnosis: Proliferative cerebral angiopathy.

COMMENTS

Notwithstanding its uncertain etiology, proliferative cerebral angiopathy is characterized by endothelial proliferation and angiogenesis, corresponding to 2–4% of cerebral arteriovenous malformations.

Such condition has a predilection for young, female patients, presenting progressive neurologic deficit, transient ischemic attack, convulsions and disabling headaches as most common symptoms. Hemorrhage episodes are less frequent than in classic arteriovenous malformations.

The natural history, clinical presentation and the recognition of radiological findings allow the differentiation of proliferative cerebral angiopathy from classic arteriovenous malformations, determining different therapeutic strategies.

There is a progressive, vicious circle cascade where the arterial supply associated with venous ectasia creates an environment of local increase in the blood volume and perinidal areas of severe hyperperfusion, triggering an uncontrolled, progressive angiogenic response to this healthy brain with an abnormal blood demand.

The name proliferative cerebral angiopathy was strictly based on the angiographic evidence of non-focal angiogenic activity, i.e., the presence of transdural supply and afferent arteries stenosis. Other distinct characteristics include absence of dominant feeder vessels to the nidus (generally lobar or hemispheric), draining veins moderately dilated in relation to the size of the arteriovenous shunting zone, and presence of healthy cerebral tissue intermingled with vascular spaces, as demonstrated by magnetic resonance imaging (Figure 2). Such findings are determining factors in the diagnostic differentiation between proliferative cerebral angiopathy and classic arteriovenous malformations.

Differential diagnoses include hereditary hemorrhagic telangiectasia (Rendu-Osler-Weber syndrome), Sturge-Weber syndrome, Wyburn-Mason syndrome and Dirvy-Van Bogaert angiomatosis.

The therapeutic strategy to be adopted should consider the presence of healthy cerebral tissue intermingled with vascular spaces. Arterial embolization of pial afferents in extensive areas determines high risk for permanent neurological deficits. The symptoms can be reduced by means of arterial embolization limited to non-eloquent areas and to transdural shunts.

Chemica is the main pathological mechanism of the disease, leading to decelerated angiogenesis, convulsions, headaches and theft syndrome, similar to Moyamoya disease, such signs and symptoms can be reduced by trepanation in the calvarium, which enhances the cortical blood supply by recruitment of additional blood supply.

As a function of the disease development stage and symptoms, a conservative treatment, with symptoms management and rehabilitation of deficits already present, may be adopted in cases where the risk for occlusion of healthy structures is greater than the arterial embolization benefits. In the present case, such a conservative approach was adopted.

REFERENCES