Ischemia of the basal ganglia as an immediate consequence of minor head injury in children is rare (< 2% of all ischemic stroke in childhood) and is due to vasospasm of the lenticulostriate arteries. The clinical history of these lesions is particularly favourable because they are usually small, and also because the facial-brachial-crural hemiparesis typical of this pathology usually regresses after a period ranging from several weeks to several months, despite the persistence of an ischemic area on MRI. This is due to the well known neuronal plasticity of the CNS, in particular, of the primary motor cortex. The most effective therapeutic approach appears to be the conservative one, although the best treatment regimen is still not well defined.

Young patients should be closely monitored and treated conservatively with osmotic diuretics to reduce perilesional edema. At the same time, however, it is very important to exclude, by means of instrumental and laboratory studies, conditions that could favour the onset of ischemia, including emboligen heart disease, thrombophilia and acute traumatic arterial dissections. Generally speaking, the prognosis in these cases is good.

The authors describe their experience treating a 10-month old baby girl, with a left lenticular nucleus ischemia and report a literature review.

**Introduction**

Ischemia of the basal ganglia as an immediate consequence of minor head trauma in children under 18 months of age is a rare eventuality (< 2% of all ischemic stroke in childhood) [1], it is caused by vasospasm of the lenticulostriate arteries, frequent in childhood, which are disrupted by head injury. Generally, it manifests with nausea, vomiting, hemiparesis and drowsiness, a clinical picture known as JHTS syndrome (Juvenile Head Trauma Syndrome) [2]. We report the case of a 10 month old child who presented an acute ischemic lesion of the left lenticular nucleus, after a minor head trauma. The ensuing facio-brachio-crural hemiparesis improved over time until complete regression of symptoms 45 days later.

**Materials and methods**

**Case description**

A 10 month old girl with not clinical history of seizures was brought to our observation following a head injury involving the right temporo-occipital region: trauma had been caused by an accidental fall from a changing table at height of 1 m. She didn’t lose consciousness but developed a right hemiparesis after 15 minutes. At neurological examination the GCS score was 14, and the child was isochoric with isocyclic pupils: there was a right facio-brachio-crural hemiparesis with muscle strength of -3/5. A head CT scan, done in emergency, showed a right occipital bone fracture with moderate diastases of the right branch of the lambdoid suture, without any intra or extraxial blood collections. A brain MRI with Gd-DTPA was performed and documented the presence of a hyperacute ischemic lesion of about 2 cm in diameter, in the left lenticular nucleus extending to the internal capsule (Figure 1).
Since a suspicion of acute arterial dissection existed, the investigation was completed with angiography sequences showing anterior and posterior circulation and perviety of the vessels of the circle of Willis (Figure 2). To rule out the possibility of an embolic heart disease or thrombophilia, urgent haematological and cardiological videats were performed together with Doppler ultrasound evaluation of neck vessels and echocardiography to exclude pathological findings. The suspicion of congenital thrombophilia, prompted us to run a complete and specific blood workup: dosage of VIII, IX, XI, XIII factors, lipoprotein A and homocysteine which were all within normal range. We evaluated possible gene mutations of factor V Leiden and of the MTHFR which were heterozygous and
the CANAIA gene coding for a structural protein of Ca channel that was heterozygous. The prothrombin 20210 A did not show mutations. Platelet count, PT, PTT, Protein C, and D-dimer assays were negative. Anti-nuclear and anti ds-DNA antibody were evaluated and proved negative similarly to IgM and IgG antcardioliopin and IgM and IgG antiphospholipid dosage. We also executed thyroid hormone assays that were normal. The child immediately underwent treatment with oral anticoagulants (calciparine 80 U/kg until the exclusion of coagulopathy) and osmotic diuretics (mannitolo 0,25 g/kg, 4 times a day until the resolution of the perilesional edema). A new brain MRI was performed 12 hours later which showed a stable situation. A progressive improvement of the right hemiparesis was so observable from the first day of admission, but not of the facial paresis. After 45 days all symptoms had disappeared and a brain MRI with GD-DTPA only showed the sequelae of the left capsular lenticular ischemia (Figure 3).

**Literature review**

A review of the literature was conducted using Pubmed. The search was undertaken using the terms “mild head trauma”, “child”, “basal ganglia ischemia”. References from the retrieved reports were checked to identify other possible reports. We selected manuscripts reporting young patients from the age of 0 to 18 months presenting lenticulo-capsular ischemia due to a mild head injury. The data extracted included, number of patients, male/female ratio, age distribution, location of the lesions, treatment, and clinical outcomes (Table 1) [1,3-5].

**Results**

From 2000 to 2010, nine patients were found in the literature, 7 males and 2 females. All of them were studied by brain MRI that showed ischemia involving the left internal capsule and corona radiate ischemia in 3 cases, the right lentiform nucleus and the corona radiate in 3, the bilateral basal ganglia, internal capsule and periven- tricular white matter in 1, the internal capsule in 1 and the lentiform nucleus in 1 (the side in the last 2 cases was not specified). Hemiparesis occurred on the side opposite to the ischemia. All patients were treated conservatively, but the type of drugs used were not described in detail in any of the reports. All of them made a clinical improvement in less than 13 months (Figure 4). All of the patients underwent radiological
follow up with Gd-DTPA MRI that always showed radiological evidence of the lesion. Hence, 7 patients were symptom-free and in 2 cases there was an improvement of the clinical status without complete resolution. None of the patients had associated risk factors.

Discussion
To classify the mild head injury in the case described here, we took into account the guidelines of the Mild Traumatic Brain Injury Interdisciplinary Special Interest Group of the American Congress of Rehabilitation Medicine. These state that mild head trauma is a condition in which there is:

1. loss of consciousness not exceeding 30 minutes;
2. GCS between 13 and 15 after 30 minutes from the trauma;
3. post-traumatic amnesia not exceeding 24 hours [6].

In childhood, cerebral infarction due to a mild head trauma is a rare eventuality, with an estimated incidence of about 1.9% [1]. Despite the rarity of the event, in children less than 18 months old the occurrence of this condition causes ischemia of the basal ganglia and, therefore, facio-brachio-crural hemiparesis, which frequently undergoes remission in a period ranging from one week to several months (usually 3 months) [4]. Secondary causes responsible for cerebral ischemic lesions in children below 18 months are attributable to traumatic dissection of the common carotid, internal carotid arteries or of the vessels of circle of Willis, congenital predisposition to rupture of cervical or intracranial arteries, emboligen heart disease and congenital thrombophilia [7,8]. In this case, before classifying a cerebral infarction in children...
below 18 months of age as idiopathic, it is imperative to exclude all possible secondary causes.

In cases similar to the one we describe here, it is necessary to exclude a possibility of surgical treatment of the lesions primarily responsible for symptoms such as bleeding, so the first investigation to perform is a CT scan. Because children are more sensitive to radiation, we prefer not to have a CTA study, but CT scan alone does not show hyperacute ischemic lesions. In this case young patients have to be submitted to:

2 - Brain MRI with Gd-DTPA which shows, when there is a cerebral infarction, hypointense image on T1 and hyperintense on T2, FLAIR and DWI corresponding to the ischemic area. 3 - Angio-MRI to rule out any congenital or anatomical abnormalities due to the trauma of the circle of Willis, responsible for ischemia. Although the gold standard for this purpose remains angiography, we think Angio-MRI should be the first study to be performed after CT scan.

4 - echocardiogram to exclude emboligen heart disease and echocolor doppler to exclude a traumatic carotid dissection.

5 - complete blood work-up in suspected congenital thrombophilia.

In the event of an idiopathic ictus cerebri in a child < 18 months of age after a mild head trauma, the aetiology is due to the particular anatomical characteristics of the arteries and of the brain parenchyma at this age, especially the lenticulostriate arteries, terminal branches of ACM responsible for the flow to caudate, putamen, internal capsule and pallidus. These vessels create an acute angle with the middle cerebral artery, which is more acute in childhood than in the adulthood [9,10]. Other vessels that may be involved are the thalamoperforating and choroidal arteries. In addition, the lateral perforators make a more acute angle than the medial perforators, which more readily undergo stretching during head trauma. The lateral perforators are shorter in children, encouraging them to stretch. Anatomically, therefore, between the intraparenchymal and the extraparenchymal segment of the lenticulostriate arteries, which are relatively fixed (the first connected to the brain parenchyma and the second to the middle cerebral artery), there is a mobile segment which, if stretched by trauma, results in vasospasm and/or thrombosis, causing ischemia of the tributary territory. Moreover, the sphenoid bone in children is not yet fully developed and does not therefore fully cover the temporal lobes,
meaning that the brain has greater mobility than the skull base in case of violent deceleration. This would facilitate stretching of the lenticulostriate arteries by traumatic forces, causing vasospasm and thrombosis [11-13]. Another rare condition which may cause stroke in children is a rare syndrome characterized by diffuse cerebral edema and coma as a result of a mutation of the CACNA1A gene, coding for a structural protein for the calcium channel [14] not present in the case we described. Clearly, the occurrence of a minor head trauma in patients with this syndrome can cause an acute ischemic stroke due to a massive edema. Another pathogenetic theory views that the friction generated between the lenticulostriate arteries and the brain parenchyma that occurs during the separation of the gray and white matter following a brain trauma in children may cause vasospasm and thrombosis. This would lead to a shutdown of the lenticulostriate flow causing ischemia in the internal capsule with a subsequent facio-brachio-crural hemiparesis [9]. It has also been shown that certain viral infections, especially varicella zoster [15], can cause secondary vascular disease, thus increasing susceptibility to develop ment of thrombosis or post-traumatic arterial vasospasm. This parameter was negative in our case because the patient had no clinical history of varicella or herpes.

The clinical history of these lesions is particularly favorable because they are usually small, and also because in a period ranging from several weeks to several months, there is usually a complete remission of facial-brachial-crural hemiparesis, typical of this pathology, despite persistence of the ischemic area on MRI. This is due to the well-known neuronal plasticity of the CNS and in particular of the primary motor cortex [16,17]. The most effective therapeutic approach thus appears to be the conservative one although a review of the pertinent literature over the last 10 years does not clearly indicate which type of treatment is better. In our experience, and in the light of the different pathologies responsible for this type of injury, administration of osmotic diuretics (such as mannitol 0.25-0.50 g/kg 4-6 times a day), to reduce perilesional edema and oral anticoagulants (such as calciparine 80-100 U/kg), to prevent acute stroke and secondary pathologies (emboligen heart disease, thrombophilia and acute traumatic arterial dissections) [18] represent an optimal treatment option in childhood.

Conclusion

Stroke after a minor head trauma in children under 18 months of age is a very rare condition (< 2% of all ischemic stroke in childhood), and is closely related to the anatomic peculiarities of the brain and skull base in childhood. Young patients should be closely monitored and treated conservatively with osmotic diuretics to reduce perilesional edema and oral anticoagulants to prevent acute stroke. Thanks to neuronal plasticity, deficits generally resolve in a period ranging from 1 week to 3 months. At the same time, however, it is very important to exclude, by means of instrumental and laboratory studies, conditions that could favor the onset of ischemia, including emboligen heart disease, thrombophilia and acute traumatic arterial dissections. Generally, prognosis in these cases is good. In our experience, administration of osmotic diuretics and oral anticoagulant drugs represents an optimal form of treatment.

Consent

Written informed consent was obtained from the parents of the patient for publication of this case report and accompanying images.

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Authors’ contributions
AL performed the neurosurgical management. NM participated to the neurosurgical management. CM participated in the literature review and in the article’s translation. MS participated in the design of the study. RD conceived of the study, and participated in its design and coordination. All authors read and approved the final manuscript.

Competing interests
The authors declare that they have no competing interests.

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