**Letter-to-the Editor: Focal cerebral arteriopathy and acute ischaemic stroke in children: A diagnostic-therapeutical conundrum**

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Acute stroke and mechanical thrombectomy in children: “where is the conundrum?”

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ABSTRACT

At difference with the adults, acute ischaemic stroke (AIS) in children presents significant differences mainly regarding aetiology and physiological response to an ischaemic insult (collateral vessels). Herein we report a small series of children (<16 yo) with AIS who underwent mechanical thrombectomy (MT). We speculate on the peculiar findings on the preliminary subtraction angiography (DSA), suggesting a possible aetiology of AIS, and on indications and limits of the currently available treatments.

PAPER

Background: MT is the standard care of patients with large vessel occlusions (LVO) causing acute ischaemic stroke (AIS). On the preliminary DSA it is usually not possible to demonstrate the underlying cause of the vessel’s occlusion. In the majority of patients, the vessel’s occlusion is caused by an embolus in otherwise normal vessels and diagnosis is usually made on hindsight. However, in a minority of patients the occlusion is caused by a local vessel wall lesion, such as an atheromatic plaque in adults, or a focal arteriopathy in children. MT devices, such as stentrievers and large-bore catheters, are designed to efficiently remove emboli, but they are not necessarily as efficient in different lesions. One of the Authors (MC), observed a peculiar angiographic pattern in two children with AIS submitted to MT. Subsequently, a revision was started to pull out similar cases in different centres. Herein we report a small series of AIS cases in children whose diagnostic appearances, endovascular treatment strategies and outcome pose more questions than answers. The paper focuses on the peculiar findings of the preliminary DSA.

Methods: we retrospectively reviewed the pre-procedural DSA of patients <16 yo, submitted to MT because of AIS of the anterior circulation, from January 2018 to December 2019, in three large volume centres. Patients were classified as having either a vessel’s occlusion or a stenosis. Only patients with a stenosis were included. Procedural imaging and clinical notes were then extracted and reviewed.
Results:
Out of 777 patients with AIS submitted to MT 4 patients (\(<16\) yo, range 13-16 years) with no history of cardiopathy nor cervical dissection nor coagulopaties had similar appearances on the preliminary DSA. Appearances were characterized by a tight narrowing of the vessel lumen at the carotid terminus caused by a “minus” lump of the vessel’s lumen. (fig 1A,2A,3A,4A).

Case description

Case #1 torino
Thirteen yo boy, previously fit and well. He presented with a sudden onset right hemiparesis, while skying. He was admitted to a local hospital one hour later with a NIHSS=14 and a CT ASPECT score of 10. On MRI appearances were in keeping with an acute ischemic stroke in the left MCA territory. No i.v. lysis was performed. He was transferred to a comprehensive hospital for further treatment. At arrival (9 hours since symptoms onset), DSA demonstrated a diffuse filling defect of the left carotid terminus extending to the A1 and the M1 tracts of the anterior and middle cerebral arteries (fig.1A). Delayed albeit complete filling of the distal MCA branches was noted. MT with an ADAPT technique (3 passes) resulted in a complete occlusion of the distal M1 and A1 segments (fig.1B). The child had an unfavourable outcome with right MCA infarct ending with a dense right hemiplegia and mRS of 4 at discharge (1 mo later)

Case #2 bologna
Fifteen yo girl previously fit and well. She presented with sudden onset of a right hemiparesis and transitory loss of consciousness. She was tubed at the scene. On A&E arrival 1hrs and a half later she was submitted to a CT that resulted in normal appearances. A subsequent MRI-MRA study showed an area of restricted diffusion in the left lenticulostriate nuclei with no flow signal in the distal left carotid siphon and proximal M1 arteries. On DSA appearances were consistent with a severe stenosis of the supraclinoid left carotid siphon and proximal M1 with a delayed, albeit complete and orthograde filling of the distal left middle cerebral artery (MCA) branches (fig 2A) and good collateral circulation from pial branches of the anterior cerebral arteries (ACA). Two consecutive stent-triever passes under aspiration (Solumbra technique) resulted in an overall improvement of orthograde flow to the
left hemisphere with an area of vascular void in the left temporoparietal region (TICI=2B) and a residual stenosis
the origin of the left M1 segment (fig.2B). Follow-up MRI-MRA (7 days and 3 mo later) showed an established
left lenticulostriate arteries infarct with persistent flow signal abnormalities in the left distal carotid siphon and
M1. The child recovered gradually to a 3 month mRS of 2

Case #3 torino

Sixteen yo female previously fit and well. She presented at the local hospital with high temperature (39° C) and a
sudden onset left hemiparesis (NIHSS=14). She was transferred to the comprehensive hospital. At arrival 4 hrs1/2
later a CT/CTA scan showed early ischemic signs in the right basal ganglia (ASPECTS=7 ) and a severe stenosis
of the distal right ICA She was transferred to the angiosuite where the preliminary DSA confirmed the diagnosis
doing severe carotid terminus stenosis (fig.3A). A self-expandable stent was deployed in the right supraclinoid ICA,
achieving an immediate and complete reopening of the vessel lumen (fig.3B). The patient was administered
Tirofibam (i.v.bolus + 12 hrs infusion) followed by Clopidogrel loading and maintenance doses. She was
discharged 2 weeks later with a mild left hemiparesis. An MRI scan obtained one month later (mRS =2) was
consistent with a partial infarction in the right ACA and anterior MCA.

Case #4 parma

12yo boy admitted to hospital with a left dense hemiplegia (NIHSS=12) 5hrs and 40 min since symptoms onset.
An immediate plain CT scan did not demonstrate early ischemic signs (ASPECTS=10) and a subsequent MRI-
MRA study revealed an area of restricted diffusion in the right insula and frontoparietal region with a distal right
ICA and MCA occlusion. No i.v. lysis was performed. The boy was transferred in the angiosuite and a
preliminary DSA showed an eccentric, severe narrowing of the right distal ICA –M1 junction with good pial
collaterals (fig. 4A). One pass with a stent retrieve (fig.4B,C) resulted in a complete reopening of the vessel lumen
(TICI=3) and a good 3 month clinical outcome (mRS=0 )
<table>
<thead>
<tr>
<th>Case#</th>
<th>Age</th>
<th>Sex</th>
<th>Clinical onset</th>
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<td>13yo M</td>
<td>R hemi.</td>
<td>NIHSS= 12</td>
<td>CT ASPECTs=10</td>
<td>Distal ICA, proximal M1-A1 stenosis</td>
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<td></td>
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<td></td>
<td>No i.v. rT-PA</td>
<td>MRI L.MCA territory diffusion hyperintensity</td>
<td>Good collaterals</td>
<td>L.MCA</td>
<td>TICI=0</td>
<td>L.MCA occlusion</td>
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<td></td>
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<td></td>
<td>No i.v. rT-PA</td>
<td>MRA distal L ica - M1CA occlusion</td>
<td>O-to-G: 9 hrs</td>
<td></td>
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<td>15yo F</td>
<td>R. hemi, LOC</td>
<td>No NIHSS (tubed at the scene)</td>
<td>CT ASPECTs=10</td>
<td>Distal ICA proximal M1-A1 stenosis</td>
<td>2 passes ADAPT</td>
<td>Improved perfusion but persistent severe stenosis at M1 CA origin</td>
<td>MRI L.basal ganglia infarct</td>
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<td></td>
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<td></td>
<td>No i.v. rT-PA</td>
<td>MRI L. basal ganglia lesion</td>
<td>Good collaterals</td>
<td>1 pass Solumbra stent-triever</td>
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<td>MRA abnormal flow signal in L. carotid siphon and M1</td>
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<td></td>
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<td>MRA Distal ICA – M1 occlusion</td>
<td>O-to-G::3hrs</td>
<td></td>
<td>TICI=2b</td>
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</tr>
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<td>3</td>
<td>16yo F</td>
<td>L. hemi</td>
<td>NIHSS= 10</td>
<td>CT ASPECTs=7</td>
<td>Distal ICA proximal M1-A1 stenosis</td>
<td>Primary stenting of the distal segment of R carotid siphon</td>
<td>Improved complete vessel lumen reopening</td>
<td>MRI R.partial ischaemic infarct in ACA and anterior MCA territories</td>
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<td></td>
<td></td>
<td></td>
<td>No i.v. rT-PA</td>
<td>CTA preocclusive stenosis of R. distal carotid siphon</td>
<td>Good collaterals</td>
<td>TICI=3</td>
<td></td>
<td>mRS (1 mo)= 2</td>
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<td></td>
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<td></td>
<td></td>
<td>R sigmoid sinus occlusion</td>
<td>O-to-G : 2 dd</td>
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<tr>
<td>4</td>
<td>12yo M</td>
<td>L hemi, NIHSS 12</td>
<td>No i.v. rT-PA</td>
<td>MR R insula and frontoparietal cortex lesion</td>
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<td>1 pass with stent triever</td>
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<td>MRI basal ganglia and insula and frontoparietal cortex infarct ; mRS (3 mo)</td>
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<tr>
<td></td>
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<td></td>
<td></td>
<td>MRA distal R ICA, M1, A1 occlusion</td>
<td>poor collaterals , O-to-G 5 hrs</td>
<td></td>
<td>TICI=3</td>
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DISCUSSION

Age apart, the patients in our series are not in any way different from all other stroke patients, as far as the clinical presentation and cross sectional imaging are concerned. However, they are indeed different. On DSA, we found a peculiar and constant pattern characterized by a focal “minus” defect of the vessel’s lumen at the carotid terminus causing a severe stenosis. To our knowledge this pattern has not been described yet, and it deserves some thoughts.

First, regarding the patients. Our review was focused on patients less than 16 y.o. because our first cases were children. However we do not have any recollection of adults with AIS with a similar pattern. This may be due both to a pathogenetic mechanism selectively affecting young people. Alternatively, appearances may be caused by a focal artheriopathy (inflammatory or non-imflammatory) more common in children than adults since the relative rarity of AIS and the absence of atherosclerosis in the paediatric age.

Second, regarding the nature of the lesion. A relevant element is that the ICA was not completely occluded. Indeed in all patients the affected MCA territories were supplied by a satisfactory amount of orthograde and collateral flow. It is unlikely that such a limited flow reduction especially in children and young adults, may cause symptoms as relevant as those that affected our patients. A possible explanation is that a partial resolution of the ICA-MCA occlusion may have occurred within the time window from symptom onset to DSA. Such behaviour, (occlusion and partial spontaneous reopening) although not exclusive, is in keeping with a subintimal dissection with a moving flap. Although a partially recanalized embolus may not be completely excluded, the angiographic appearances are not consistent at all with this hypothesis if the location (ICA instead of MCA) the eccentricity of the lesion (a recanalized embolus most likely appears as a concentric recanalization) and its profile (smooth versus irregular) are taken into account. A different aetiology to be considered is an inflammatory
focal arteriopathy. This is a leading cause of stroke in children in the absence of other risk factors. The angiographic pattern has been described (lasjaunias) and clinical features have been recently revised (1). Although this disease may not be completely excluded, focal arteriopathies have different angiographic appearances, and they usually involve the first portion of MCA, while in our patients the disease was located in terminal ICA.

Third, regarding treatment. The stentriever and direct stenting techniques were efficient in 3 cases, while aspiration resulted harmful in a single patient. Due to the limited size of the sample, we obviously cannot generalize these results. Anyway, while aspiration and stentriever are both efficient in embolic occlusions, (responsible for the vast majority of LVO in AIS), this may not necessarily be the case in patients with a dissective arteriopathy. In the hypothesis of a focal subintimal dissection, the retrieval of a stentriever moving retrogradely to flow direction (from distal-to- proximal) may have a very different effect on a flap than a large bore aspiration catheter moving the other way round (from proximal- to-distal). The presence of a subintimal dissection with a flap may also explain the optimal and immediate vessel reopening in case #3 following the deployment of a self-expandable stent in the right terminal ICA.

As a final consideration, it is possible that in these patients there is a yet-unrecognized benefit in relying on collateral circulation rather than attempting to aggressively reopen the vessel. Aggressive management of acute stroke in children involves general anaesthesia, possibly lowering blood pressure and disrupting normal compensatory mechanism for an efficient collateral circulation.

**Conclusions:** we identified a peculiar angiographic pattern of lesions in four children with anterior circulation AIS. Although a definitive diagnosis could not be made, a focal subintimal dissective pathology was suspected based on the preliminary DSA appearances and the procedural behaviour during MT. This pattern should suggest a cautious attitude for MT and/ or different strategies for revascularization.
REFERENCES


FIGURES AND CAPTIONS

FIGURE 1 (A-B)

Fig. 1 (A-B) DSA selective left ICA injection pre and post MT, frontal view (case #1) On the preliminary angiogram there is a scant filling of the carotid terminus and the M1 segment with persistent orthograde flow. The ACAs are not opacified because of a wash out phenomenon from the right ICA. After MT (B) there is a dramatic worsening of the angiogram with complete occlusion of the carotid terminus.

FIGURE 2(A-B)

Fig. 1 (A-B) DSA selective left ICA injection pre and post MT, frontal view (case #2) On the preliminary angiogram (A) there is a severe narrowing of the carotid terminus-M1 junction with an eccentric “minus” image (“lump”) protruding into the vessel’s lumen with a smooth profile. After MT (B) there is a satisfactory reopening of the vessel with a focal vasospasm in the proximal M1 segment.
FIGURE 3 (A-B)

Fig. 3 (A-B) DSA selective right ICA injection pre and post MT, frontal view (case #3) On the preliminary angiogram (A) there is a smooth narrowing of the carotid terminus-M1 junction with persistent orthograde flow. The ACAs are not opacified because of a left dominant A1 segment. After MT (B) there is a successful reopening of the vessel lumen.

FIGURE 4 (A-C)

Fig. 4 (A-C) DSA selective right ICA injection pre , during and post MT, frontal view (case #4) On the preliminary angiogram (A) there is a smooth eccentric narrowing of the carotid terminus-M1 junction with delayed orthograde filling of the distal MCA branches. After placement and unsheathing of the stentriever (B) there is a satisfactory reopening of the vessel with restoration of a full orthograde flow. After retrieval of the stentriever (C) there is a complete vessel’s patency restoration with filling of a hypoplastic A1 segment as well.