Endovascular Management of Internal Carotid Artery Pseudoaneurysm Secondary to Pediatric Deep Neck Space Infection: A Case Report and Review of Literature

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Abstract

Pseudoaneurysms of extracranial internal carotid artery (ICA) are rare in children. Main causes include trauma, iatrogenic causes, and neck space infection. Prompt diagnosis and management is vital, in view of life-threatening complications like fatal airway hemorrhage and stroke. Endovascular management has currently become the preferred treatment strategy due to its minimally invasive nature and lower complication rates. We report a rare case of mycotic pseudoaneurysm of extracranial ICA in a 4-year-old child as a complication of neck space infection, which was successfully managed with endovascular parent artery occlusion.

Introduction

Pseudoaneurysms of extracranial internal carotid artery (ICA) are rare lesions in children with fewer than 100 cases reported.1,2 Main causes include trauma, iatrogenic causes, infections, and malignancies with infection and trauma being the most common causes in children.3,4 Prompt diagnosis and management is vital, in view of life-threatening complications like fatal airway hemorrhage and stroke.2,5 Endovascular management has become the preferred treatment strategy due to its minimally invasive nature and lower complication rates.6,7 We report a rare case of mycotic pseudoaneurysm of extracranial ICA in a 4-year-old child as a complication of neck space infection, which was successfully managed with endovascular parent artery occlusion (PAO).

Case

A 4-year male child presented to the emergency room with high-grade fever and tender right neck swelling for 2 days. The child had no neurological symptoms, prior history consistent with tonsillitis or surgery. Complete blood evaluation showed leukocytosis (35,000 cells/mm³) with neutrophilia. Neck ultrasound showed significant bilateral cervical lymphadenopathy with hypoechogenic lesion in the right submandibular region with internal swirling flow. Computed
tomography (CT) angiography showed a saccular pseudoaneurysm from the right extracranial cervical internal carotid artery (ICA) with peripheral thrombosis and enhancing walls (suggestive of mycotic pseudoaneurysm) close to the skull base (► Fig. 1). Adjacent airway was not compressed. Blood samples were sent for cultures. The child was admitted for intravenous antibiotics and rehydration. Within few hours of admission, the general status of the child deteriorated with purulent blood-streaked nasal discharge. With the clinical suspicion of rupture, the child was taken up for emergency endovascular PAO. Diagnostic angiograms confirmed the presence of saccular pseudoaneurysm from the right cervical ICA. On cerebral angiography, the right anterior cerebral artery and middle cerebral artery were shown to fill from left ICA and right vertebral artery with good-sized anterior communicating artery and right posterior communicating artery (► Fig. 2). Right ICA was occluded proximal to the pseudoaneurysm with coils using a vertebral diagnostic catheter (Interlock-35, 6 mm × 100 mm, Boston Scientific). Antibiotics (cultures isolated Staphylococcus sp.) were continued for up to 4 weeks in postprocedure period. No neurological deficits were seen at periprocedural period. Follow-up CT angiograms at 4 weeks showed thrombosed pseudoaneurysm with well-collateralized intracranial circulation (► Fig. 3). The child is well at 6 months of clinical follow-up.

**Discussion**

ICA pseudoaneurysms are an uncommon complication of deep neck space infections especially after advancements in antibiotic therapy. These are more common in children as compared with adults due to higher incidence of deep neck space infections in this age group. In a review by Ruff et al, only 31 pediatric cases of infectious pseudoaneurysms of extracranial carotid artery had been reported since 1968. Common organisms associated with this complication are *Staphylococcus aureus* and *Streptococcus pyogenes*. The mechanism of pseudoaneurysm formation is secondary to infectious arteritis of vessel wall with involvement of vasa vasorum, which results in ischemic vessel wall damage, subsequent erosion, and rupture.

Children with extracranial ICA pseudoaneurysms usually present with painful pulsatile neck mass. Though thromboembolic events like stroke, transient ischemic attacks, or amaurosis fugax have been reported in adults, they are less commonly seen in children. Other complications which may be seen with mycotic ICA pseudoaneurysms include hemorrhage into aerodigestive tract secondary to rupture, airway compromise, and stridor due to mass effect, cranial nerve palsies, and Horner syndrome. Risk of rupture is more
common in pseudoaneurysms in children, as compared with adults and in pseudoaneurysms occurring secondary to trauma and infections with high mortality rates in children (as high as 54%). Ultrasound is the initial imaging investigation of choice. Diagnosis is usually confirmed by CT angiography, though magnetic resonance angiography can be performed in older and cooperative children. Timely diagnosis and management is necessary to prevent morbidity and mortality from complications.

There are no consensus guidelines till date and the choice of treatment depends on factors like site and etiology of pseudoaneurysm, and status of intracranial collateral circulation. Historically, surgery (under antibiotic cover) has been the mainstay of treatment. It offers a definitive cure but has a high complication rate including perioperative hemorrhage, Horner Syndrome, and cranial nerve injury (rates as high as 40%) as well as mortality rates ranging from 9 to 20%. Surgery may also be technically difficult, especially in pseudoaneurysms near skull base and in small necks.

Endovascular strategies have a lower mortality rate (4.1%), lower risk of stroke (1.8%) and cranial nerve injury (0.5%), and shorter length of hospital stay. Endovascular strategies include PAO, coiling of pseudoaneurysm, using bare metal stents with/without coils, covered stents/stent grafts, and flow diverters to exclude the pseudoaneurysm from circulation. PAO using coils or detachable balloons is the preferred technique in children especially in emergencies, as they have good intracranial collateral circulation which is an essential prerequisite for PAO. Patency of stents and flow diverters with long-term antiplatelet medication is not well evaluated in children, hence making them less preferable in children. Bare metal stents, stent grafts, and flow diverters are not used in mycotic pseudoaneurysms, as they are prone to get infected and are not sufficient to prevent spread of septic emboli. In addition to lesion-specific management, the child should be kept on antimicrobial cover for 4 to 6 weeks. The probable complications of endovascular therapies include thromboembolism, arterial dissection, late stenosis, occlusion of stent graft, and recurrence. Immediate relief from compression-related symptoms may not be achieved with endovascular treatment. Long-term follow-up is essential to check for thrombosis and resolution of pseudoaneurysm, any recurrence, and patency of the stents.

**Conclusion**

Mycotic pseudoaneurysms of ICA in children are rare, but potentially life-threatening complications of deep neck space infections requiring prompt management. PAO with trapping of neck of the aneurysm using endovascular coils or balloons is a safe and effective method of treatment. Long-term antimicrobial therapy and follow-up is vital to achieve optimal outcomes.

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**References**


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**Fig. 3** Follow-up computed tomography (CT) angiography at 1 month showing complete thrombosis of pseudoaneurysm sac (white arrows in A and C) with the coil mass (seen in B).


