Case Report

Dural sinus mechanical thrombectomy and continuous local rt-PA infusion in a child with refractory intracranial hypertension and progressive visual loss: A case report

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ABSTRACT

Background: Children with intracranial hypertension are at risk for visual loss and their visual function must be closely monitored. Surgery with the insertion of a ventriculoperitoneal shunt is imperative when vision is threatened.

Case Description: Herein, we report a case of a 5-year-old boy whose refractory intracranial hypertension and severe, progressive visual loss (secondary to a chronic, otogenic, right sigmoid sinus thrombosis, and a contralateral sinus tight stenosis) were resolved by a combination of continuous (6 h), locoregional, infusion of recombinant tissue plasminogen activator (rt-PA), and mechanical thrombectomy.

Conclusion: The association of in loco and continuous infusion of recombinant tissue plasminogen activator (rt-PA) with mechanical thrombectomy resulted in effective in partially reopening the occluded sinus and facilitating a good clinical recovery. This combined endovascular approach may represent an alternative, less invasive, therapeutic option to surgery in children with intracranial hypertension caused by chronic cerebral venous sinus thrombosis.

Keywords: Dural sinus thrombosis, Intracranial hypertension, Recombinant tissue plasminogen activator, Thrombectomy, Vision

INTRODUCTION

This paper would like to draw attention to a possible alternative therapeutic option to surgery in children with refractory intracranial hypertension secondary to cerebral venous sinus thrombosis (CVST) and progressive visual deterioration.

CASE DESCRIPTION

A 5-year-old boy with a more than 2 months history of otogenic CVST of the right sigmoid sinus was referred by pediatrician consultants. Despite adequate antibiotics and anticoagulant therapy, the child developed progressive bilateral papilledema and visual loss. An MR imaging (MRI) – MR venography (MRV) of the head showed a thrombotic occlusion of the right transverse-sigmoid
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sinus associated with a tight stenosis of the left transverse sinus [Figures 1a and b] and bilateral enlargement of the optic nerve sheath with bulging of the optic discs [Figure 1c]. Serial consecutive measurements of the visual acuity revealed a progressive and dramatic visual loss over 1 week. The case was discussed in a multidisciplinary meeting, and a decision was made to offer an endovascular reopening of the occluded right side, transverse-sigmoid sinus.

An informed consent signed by both parents was obtained, and the procedure was performed under general anesthetics on a biplanar angiography system (Philips Allura). A 7F guider was placed in the right jugular bulb and a diagnostic 4F catheter in the right internal carotid artery (ICA) through a femoral (venous 7F and arterial 4F) approach. After several attempts, we managed to cross the occluded segment of the sinus with a 0.16” guide and a 0.21” microcatheter. A 6 × 30 mm Solitaire (Medtronic) stent-retriever was then delivered across the thrombosed sinus, unsheathed and, after a few minutes, wait, trashed out. A second pass with the stent-retriever was repeated: each time only small and scanty clot fragments were retrieved. While attempting to cross the occlusion for the third time, the child presented a transient tachyarrhythmia (200 bpm) and hypotension requiring the anesthesiologist’s assistance with a quick resolution of the event. Once the child was hemodynamically back to normal, a dynaCT was obtained, and it resulted in normal appearances. Subsequent check angiograms showed an unsatisfactory partial reopening of the occluded sinus [Figure 2a]. A decision was made to stop the procedure after placement of two 0.21” microcatheters in the right transverse and sigmoid sinuses, respectively, their distal tips 2–3 cm apart. Then, a local infusion of rt-PA through the two microcatheters was started. The total dose of the rt-PA was titrated according to the child body weight and evenly split between the two microcatheters at a rate allowing a total 6 h infusion. The boy was transferred to the Intensive Care Unit (ICU), tubed, and sedated over the night. The following morning, a single pass with two unsheathed, and side-to-side 6 × 30 mm Solitaire stent-retrievers (“double stent-retrievers” technique), followed by aspiration through the 6F guider, resulted in a satisfactory reopening of the right sigmoid sinus [Figures 2b and c]. Final angiograms demonstrated an improved cerebral venous outflow [Figure 2d]. At the end of the procedure, a therapeutic dose of heparin was prescribed and the child was woken up in ICU. In the following days, the child vision progressively improved and he could play cards with his mother (recognizing the colors and numbers on the cards). A week later, the bilateral papilledema resolved with further visual acuity improvement. A follow-up MRI three weeks later showed a stable, though partial, reopening of the right sigmoid sinus with a resolution of the bilateral optic nerve sheath enlargement and disc bulging. The boy was discharged home on a full dose of fractionated heparin.

DISCUSSION

The aforementioned case opens up new therapeutic options in children with refractory intracranial hypertension from CVST. In children, a systematic review of the literature identified nine children with acute CVST, treated using several endovascular methods, including local rt-PA, microguidewire and catheter disruption, balloon angioplasty, and thromboaspiration. A series of 15 children with acute CVST reported 11 patients treated with intrasinus urokinase or rt-PA (6 and 5 patients, respectively), 1 using the AngioJet device (Possis, now Medrad Inc, Warrendale, Pennsylvania) and 1 with the penumbra aspiration device. In all the reported cases, the indication for endovascular treatment was

Figure 1: MR imaging (a and c) and MR venogram in the sagittal view (b). An abnormal signal in the right transverse-sigmoid sinus (white arrowhead), associated with inflammatory changes in the right mastoid is noted in a. Absence of flow signal of a sigmoid sinus and a tight narrowing of the contralateral transverse sinus (white arrowhead) are depicted in b. Bilateral enlargement of the optic nerve sheaths (white waved arrowheads) and bulging of the optic discs (black arrowheads) are depicted in c. Appearances are consistent with an otogenic, right transverse-sigmoid sinus thrombotic occlusion with contralateral transverse sinus tight stenosis and indirect signs of an increased intracranial pressure.
It is known that the dural sinus narrowing is structural or secondary to the increased intracranial pressure. Finally, in refractory intracranial hypertension, it is still an issue of hot debate whether the dural sinus narrowing is structural or secondary to the increased intracranial pressure. In our case, ventriculo/ lumbo peritoneal shunting was deemed challenging by the neurosurgeons, due to the slit-like ventricles and large body habitus. The consensus view of the Multidisciplinary Team (MDT) meeting opted for endovascular reopening of the sinus (a reconstructive procedure targeted to the primary cause of the intracranial hypertension) because it represented a viable, less invasive/expensive procedure, compared to surgery. Furthermore, it was considered to be a better guarantee of a long-term outcome.

In children, there are only scant reports of intrasinus rt-PA infusion, the rates, and total dose of the fibrinolytic differing greatly as well as the outcomes. In Waugh series of six patients, a continuous infusion of rt-PA was administered over a mean of 49 hours, with a maximum dose of 60 mg. In their series, 5/6 patients demonstrated hemorrhagic complications. Another series of three patients demonstrated a favorable outcome in 2/3 patients. It is known that the efficacy of rt-PA is greatly hampered by a large clot burden, not allowing the lytic agent to seep through the clot and to exert its action on a wider clot surface. In our case, the total rt-PA dose was titrated to the body weight and administered in a relatively short time window (6 h) to limit both the risk of a bleed and complications related to long-term intravenous lodging of (micro) catheters. The following day, it was much easier to track the microwires past the clot and the “double” stent-retrievers with thromboaspiration resulted in much more effective in reopening the sinus. It is possible that both results may have been favored by the pre-emptive fibrinolytic action of the continuous local infusion of rt-PA. Our case confirms that it is often unnecessary to achieve a complete thrombus resolution; even a partial recanalization is often sufficient to relieve the intracranial hypertension, possibly allowing the intrinsic fibrinolytic system to lyse the remaining thrombus.

Finally, in refractory intracranial hypertension, it is still an issue of hot debate whether the dural sinus narrowing is structural or secondary to the increased intracranial pressure. In our case, the stenosis of the contralateral transverse sinus was still present on the follow-up MRI 3 weeks later, when the child was discharged home with the incompletely understood long-term outcomes (what if after PTA and stenting the intracranial hypertension failed to improve?). In summary, the associated risk of the endovascular procedure and the antiplatelet therapy was considered to exceed the possible benefit.
a substantial clinical recovery and resolution of the radiological signs of intracranial hypertension. Although no invasive measurement of the CSF pressure was obtained, follow-up MRI appearances were consistent with a significant decrease of the intracranial pressure, suggesting a possible structural cause of the left transverse sinus stenosis. Further, clinical and MRI follow-ups are planned to check for the recovery of the visual deficits and the possible development of idiopathic intracranial hypertension or dural AV fistulas.

CONCLUSION

Mechanical thrombectomy with a double stent-retrievers technique following a continuous, local, and 6 h infusion of rt-PA resolved intracranial hypertension and progressive visual loss in a 5-year-old child with a chronic otogenic transverse-sigmoid sinus occlusion. Mechanical thrombectomy with a double stent-retrievers with/without local infusion of rt-PA may represent a viable, less invasive and expensive, and alternative therapeutic option to surgery. However, due to the intra-periprocedural hemorrhagic risks, this technique should be reserved for only the most carefully selected cases of idiopathic intracranial hypertension secondary to chronic CVST.

Declaration of patient consent

The authors certify that they have obtained all appropriate patient consent.

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Nil.

Conflicts of interest

There are no conflicts of interest.

REFERENCES

1. Bynke G, Zemack G, Bynke H, Romner B. Ventriculoperitoneal shunting for idiopathic intracranial hypertension. Neurology 2004;63:1314-6.
2. Gebara BM, Goetting MG, Wang AM. Dural sinus thrombosis complicating subclavian vein catheterization: Treatment with local thrombolysis. Pediatrics 1995;95:138-40.
3. Griesemer DA, Theodorou AA, Berg RA, Spera TD. Local fibrinolysis in cerebral venous thrombosis. Pediatr Neurol 1994;10:78-80.
4. Klisch J, Sychara V, Strasilla C, Taschner CA, Reinhard M, Urbach H, et al. Double solitaire mechanical thrombectomy in acute stroke: Effective rescue strategy for refractory artery occlusions? AJNR Am J Neuroradiol 2015;36:552-6.
5. Kulcsár Z, Marosfoi M, Berentei Z, Szikora I. Continuous thrombolysis and repeated thrombectomy with the Penumbra System" in a child with hemorrhagic sinus thrombosis: Technical note. Acta Neurochir (Wien) 2010;152:911-6.
6. Mortimer AM, Bradley MD, O'Leary S, Renowden SA. Endovascular treatment of children with cerebral venous sinus thrombosis: A case series. Pediatric Neurol 2013;49:305-12.
7. Ramos SK, Phillips J, Lin J, Moresco K, Meagher S. Successful rheolytic mechanical thrombectomy of cerebral venous thrombosis in a pediatric patient. J Neurosurg Pediatr 2013;11:140-3.
8. Wasay M, Bakshi R, Dai A, Roach S. Local thrombolytic treatment of cerebral venous thrombosis in three paediatric patients. J Pak Med Assoc 2006;56:555-6.
9. Waugh J, Plumb P, Rollins N, Dowling MM. Prolonged direct catheter thrombolysis of cerebral venous sinus thrombosis in children: A case series. J Child Neurol 2012;27:337-45.
10. Zhang A, Collinson RL, Hurst RW, Weigele JB. Rheolytic thrombectomy for cerebral sinus thrombosis. Neurocrit Care 2008;9:17-26.

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