RECURRENT MENINGITIS AND HOLOCORD SUBDURAL ABSCESS CAUSED BY SPINAL DERMOID CYST

SPİNAL DERMÖİD KİSTE BAĞLI REKÜRENT MENENJİT VE HOLOKORD SUBDURAL ABSE

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INTRODUCTION
Spinal subdural abscess caused by spread of infection via the dermal sinus tract is rare in children (5). A child with a midline dimple or pit is one of the most common referrals to a pediatric neurosurgeon (10). Recurrent meningitis and hydrocephalus are also frequent consequences of infected spinal dermoid cyst and subdural abscess (2, 5, 6). Here, we present a child admitted to our clinic with the diagnosis of meningitis and hydrocephalus. Further evaluation of this child disclosed an infected intramedullary dermoid tumor and subdural abscess.

CASE
A 4 year-old-boy presented to the Department of Pediatrics with the complaints of fever, headache and decreased consciousness. On the examination, he was found apathic; his cooperation and orientation were decreased and he had meningeal irritation findings. Informed consent was obtained from his family after the nature of the procedure to be performed had been explained. Following an emergent cranial tomography (CT), demonstrating diffuse meningeal enhancement and ventricular dilatation, a ventricular puncture was performed. Evaluation of the cerebrospinal fluid (CSF) disclosed streptococcal meningitis. He had been treated with appropriate antibiotics. Then, a ventriculoperitoneal shunt system was inserted. He became conscious and cooperative. A control lumbar puncture at the end of the antibiotherapy revealed a purulent fluid and streptococci were cultured. Control cranial CT was normal and control ventricular puncture was clear. A spinal pathology was suspected and a spinal radiologic examination including X-ray, and spinal magnetic resonance imaging (MRI) was performed. On MRI, there was an enhanced
lesion of the conus region and subdural abscess all along the spinal cord (Figure 1a, b). He was operated on through a dorsal midline approach and an intramedullary dermoid tumor including hairy contents was removed totally. A fistula tract ending at the subcutaneous level was noticed during the surgery. After all, he was treated with intravenous antibiotics for subdural abscess. He did well postoperatively. On his last control examination 6 months after the surgery, he was normal and the shunt was doing well.

DISCUSSION
Spinal intramedullary tumors may be congenital or acquired. Sometimes lumbar puncture for any reason may also cause spinal tumor (4, 8). They may present with neurological motor and sensorial deficits (1, 6). Some tumors such as embryonic tumors-dermoid cysts, craniopharyngiomas, teratomas and malignant gliomas may present with the episodes of chemical meningitis, single or multiple in patients with no neurological symptoms (2, 7, 8, 9). In these cases, differential diagnosis from aseptic meningitis, especially from Mollaret aseptic meningitis, is essential (8). A careful physical and neurological examination is important. Especially skin examination of the dorsal midline and intergluteal regions for the dermal sinus tract is a must in meningitis and recurrent meningitis cases. A child with a midline dimple or pit is one of the most common referrals to a pediatric neurosurgeon. Regardless of their depth, those below the top of the intergluteal crease end blindly and never extend intraspinally. Families can be reassured that there is no infection or mechanical risk to the developing nervous system. A high index of suspicion must be maintained for all dimples above the intergluteal fold despite a normal examination or neuroradiologic studies. The midline must be carefully inspected when a child of any age suffers meningitis, especially when an unusual organism is cultured (3). Spinal congenital dermal sinus (CDS) is a rare entity which supposedly results from the failure of the neuroectoderm to separate from the cutaneous ectoderm during the process of neurulation. The lesions are most frequent at the lumbosacral followed by the occipital region. CDS of the thoracic region is very rare. The patients with spinal CDS present with meningitis and/or mass effect caused by the associated inclusion tumor. They are usually dermoid or epidermoid cysts. Teratoma is rarely associated with this condition (4, 8). Conservative management of dermal sinuses is not justified, and these lesions should be electively resected at the time of diagnosis. Dermal sinuses and inclusion tumors may lead to spinal cord tethering and progressive neurologic deterioration. Surgery in advance of deficits maintains normal neurologic function, and children can develop unencumbered by infection, motor, or bladder paralysis. Radiologic diagnosis of the dermoid tumors is generally dependent on MRI. In the literature, most of the lesions were iso- to hyperintense to CSF on T2-weighted images. Some demonstrates a mixed intensity pattern on all sequences; dominant component gives short T1 and short T2 values, consistent with fat (3). Dermal sinus tract may not be observed in all cases. In our case, we thought that there had been a dermal sinus associated with the intramedullary dermoid tumors. Then, the sinus had been infected and caused self obliteration and development of subdural abscess all along the spinal cord. At the end, when meningitis has developed the child’s family noticed the symptoms. Ventricular puncture drained a very clear CSF, but lumbar puncture drained a viscous infected fluid at the end of the antibiotic therapy for the meningitis.

CONCLUSION
This case indicates that meningitis can develop secondary to a congenital dermal sinus and dermoid tumors. So every child with central nervous system infection should be evaluated for dorsal midline defects. In cases of dermal sinus presence, exclusion of the intramedullary tumor and spinal abscess is a must to prevent recurrent meningitis and neurologic deficits caused by spinal abscess and tumors. In dermal sinus cases, an enhanced MRI evaluation of the spinal cord will help us to detail the pathology and to help for surgical management.

REFERENCES


