Extramedullary Myxopapillary Ependymoma of the Filum Terminale Associated With Syringomyelia: A Case Report

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Abstract

Aim: Ependymomas compose the majority of all intradural intramedullary spinal lesions. Filum terminale lesions are rarely associated with syringomyelia. Here we describe a case of extramedullary myxopapillary ependymoma associated with extensive syringomyelia located in the filum terminale.

Material and Methods: A 34-year-old man admitted to our clinic with complaints of bilateral leg weakness, back pain and urinary dysfunction. The symptoms began three years before the admission and rapidly progressed in the last three months. Neurological examination revealed paraparesis (Grade 3/5) and diminished sensation below the Th 8 level. Deep tendon reflexes were reduced at the right lower extremity while they were increased at the right lower extremity. The anal sphincter tone was also decreased. The MRI of the patient showed us a contrast enhanced intraspinal mass lesion at the level of L 2 with a syringomyelia extending from L 2 level to Th7 level.

Results: The patient was operated via lumbar laminoplasty and tumor was totally resected.

Conclusion: The removal of tumor is also sufficient for the resolution of syringomyelia and no additional procedure is essential for the treatment of syringomyelia in such cases. This is the first case of extramedullary tumors of the cauda equine with extensive syringomyelia.

Key words: Ependymoma, Filum terminale, Syringomyelia.

Introduction and aim of the paper

Ependymomas compose the majority of all intradural intramedullary spinal lesions and consist 60% of all glial tumors in spinal cord (1, 2). Ependymomas are usually seen in cervical region (44%), followed by thoracic region (26%). Intradural intramedullary tumors are commonly related with syringomyelia in the cervicothoracic region and majority of these tumors are glial tumors. Filum terminale lesions are rarely associated with syringomyelia. Here we describe a case of extramedullary myxopapillary ependymoma associated with extensive syringomyelia located in the filum terminale.

Material and methods

A 34-year-old man was admitted to our hospital with the complaints of bilateral weakness in lower extremities, back pain and urinary dysfunction. The symptoms began three years before the admission and rapidly progressed in the last three months. Neurological examination revealed paraparesis (Grade 3/5) and diminished sensation below the Th 8 level. Deep tendon reflexes were reduced at the right lower extremity while they were increased at the right lower extremity. The anal sphincter tone was also decreased. The MRI of the patient showed us a contrast enhanced intraspinal mass lesion at the level of L 2 with a syringomyelia extending from L 2 level to Th 7 level (Figure 1, Figure 2).
A laminoplasty from L1 to L3 was performed. When the dura was opened, a dark brown extramedullary tumor 3 cm in length was encountered. The tumor arose from the filum terminale and displaced the nerve fibers of cauda equina anteriorly and laterally (Figure 3). The tumor was excised by microsurgical techniques. A thin membranous tissue covering syringomyelic cavity was opened during the tumor removal and slightly xanthochromic cystic fluid was drained.
Results

Postoperative course was uneventful and lower extremity weakness of the patient improved (Grade 4/5) immediately. Histopathological examination was reported as “myxopapillary ependymoma”. Three years after the operation, the patient was neurologically intact except an intermittent urinary dysfunction. Control MRI revealed total removal of the tumor and disappearance of syringomyelia (Figure 4).

Discussion

Syringomyelia associated with spinal tumors is common. In most cases, tumors are intramedullary glial tumors located in the cervicothoracic region (1, 3). However, the tumors which are located in the distal end of the spinal cord rarely cause syringomyelia (1). Barnett and Rewcastle reviewed seven cases of cauda equina and filum terminale tumors causing syringomyelia including five autopsy cases and two surgical cases (1). Other cases of intramedullary ependymoma of the conus medullaris associated with syringomyelia extending from cervical region were reported by McCormick and Sarikaya et al. (4, 5). However, in the case reported by Nagahiro et al. the tumor was an extramedullary ependymoma located in the cauda equina (6). In our case the tumor was purely extramedullary and arose from the filum terminale. This is the first case of extramedullary tumors of the cauda equina with extensive syringomyelia documented in the MRI era. The pathogenesis of syringomyelia associated with spinal cord tumors is still not well understood. It is usually believed that the effects of tumor including intramedullary softening due to a disturbed blood supply, stasis of the tissue fluid resulting from occlusion of drainage pathways, spread of edema, and hemorrhage into the cord from the tumor are responsible for the development of syringomyelia (1, 6, 7). However, the reason for the occurrence of syringomyelia associated with the tumors is uncertain. Probably the other entity known to be associated with syringomyelia such as Chiari I malformation, trauma, arachnoiditis, degenerative diseases of the spine, disturbance of the cerebrospinal fluid flow because of obliteration of the
subarachnoid space play an important role in the development of the syringomyelia. The subarachnoid space and extracellular space of the central nervous system act as a single fluid compartment without any barrier to prevent the fluid movements between them (7, 8). However, the removal of tumor is also sufficient for the resolution of syringomyelia and no additional procedure is essential for the treatment of syringomyelia in such cases.

Conclusion

A unique patient with extramedullary myxopapillary ependymoma located in the filum terminale associated with extensive syringomyelia has been reported. Serial MRI demonstrated the total removal of the lesion and spontaneous resolution of the syringomyelia.

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