Axis Metastasis, Atlantoaxial Dislocation and Cervical Syringomyelia: A Case Report

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Abstract
Syringomyelia associated with atlantoaxial dislocation is usually described in patients with complex craniovertebral junction anomaly. However, atlantoaxial dislocation without craniovertebral junction bony anomaly very rarely causes syringomyelia. In this report, we describe a case of syringomyelia associated with atlantoaxial dislocation secondary to metastatic tumour of the axis vertebra.

Keywords
Syringomyelia, Atlantoaxial dislocation, Craniovertebral junction, Tumour of the axis

Introduction
Syringomyelia is characterized by the presence of a fluid filled cystic cavity within the spinal cord. Cervical syringomyelia is usually associated with Chiari malformations, tumours or as a consequence of trauma [1,2]. Craniovertebral junction (CVJ) bony anomalies, including atlantoaxial vertical dislocation and basilar invagination with or without tonsillar herniation, can also cause cervical syringomyelia [3,4]. Simple or horizontal atlantoaxial dislocation-associated syringomyelia is very rarely reported [5-7]. We describe a very rare case of syringomyelia caused by atlantoaxial dislocation associated with metastatic tumour of the axis.

Case Report
A 65-year-old woman was admitted with a 6-month history of neck pain complaint of weakness in all extremities. Neurological examination revealed quadriparesis with brisk deep tendon reflexes in all four limbs. Pin prick sensation was reduced bilaterally below the C5 level; neck movement was reduced because of pain. The bilateral Babinski sign was positive. Plain skull and cervical X-rays showed an atlantoaxial dislocation. Magnetic resonance (MR) examination revealed an atlantoaxial dislocation with a tumoral mass involving the corpus of axis, with cervical syringomyelia (Figure 1). Computed tomography (CT) disclosed the destruction of the corpus of axis (Figure 2). In the first operation, C1 laminectomy and occipito-cervical fusion using C2 laminar and C3-C4 lateral mass screwing was performed. Following C1 laminectomy, dural pulsation was observed. Two days after the first operation, partial excision of the tumour mass was performed, and the bone defect was filled with acrylic to stabilize the segment. Histopathological examination resulted in a diagnosis of adenocarcinoma metastasis. Postoperative symptoms improved and control MR imaging showed that the syrinx was reduced (Figure 3). The patient was transferred to the oncology unit and died 6 months following surgery because of extensive metastasis of lung carcinoma.

Discussion
The most common cause of atlantoaxial dislocation is trauma. Rheumatoid arthritis, congenital CVJ anomalies, infection and tumours of the CVJ are other causes. Tumours affecting the osseous and ligamentous structures of CVJ are rare, and can be classified as primary, secondary or miscellaneous [8]. Primary tumours are bone tumours, such as osteoma, osteochondroma and chordoma. Metastatic tumours of the CVJ are very rare. Piper and Menezes reported 14 patients treated...
without basilar invagination [3]. In such cases, the abnormal position of atlantoaxial joint as a result of congenital bone anomaly causes slippage of the atlas over the axis and invagination of odontoid process [10]. That type of atlantoaxial dislocation is described as a ‘vertical’ dislocation. However, simple or horizontal atlantoaxial dislocations very rarely cause syringomyelia, and few cases have been reported [5-7].

Goel, et al. reported three infants with atlantoaxial dislocation-associated syringomyelia treated with atlantoaxial fixation [5]. In a case reported by Mizutani, et al., loosening of multistrand cables following C1-C2 Brooks-type fusion resulted in syringomyelia [6]. Yoong, et al. also reported a case of atlantoaxial dislocation-associated syringomyelia in a patient with rheumatoid arthritis [7].

Although the pathogenesis of syringomyelia is unclear, impairment of subarachnoid cerebrospinal fluid (CSF) flow is involved in cases of atlantoaxial dislocation with syringomyelia [4]. CSF flow obstruction at the dislocation level is similar to that of Chiari malformation, which prevents the return of spinal CSF and increases perivascular flow from the subarachnoid space, leading to the syrinx formation [2]. In horizontal atlantoaxial dislocation, as in our patient, dorsal compression by the posterior arch of the atlas is more prominent. In the case reported by Mizutani, loose cables of a Brooks-type fusion compressed subarachnoid pathways [6]. Decrease in the blood supply of the spinal cord resulting from atlantoaxial dislocation, tumours, or rheumatoid vascular compression can also cause ischemic necrosis and cavity formation in the spinal cord. Regardless of the pathology, treatment of atlantoaxial dislocation associated with syringomyelia includes atlantoaxial stabilization and restoration of the obstructed subarachnoid space.
Clinical and radiological improvement usually occurs soon after surgery. However, we chose occipitocervical stabilization and C1 laminectomy for decompression in our patient because of the malignant nature of the disease and the irreductable atlantoaxial dislocation.

**Conclusion**

Metastatic axis vertebrae tumours rarely cause atlantoaxial dislocation and syringomyelia. Occipitocervical stabilization and decompression of subarachnoid space are essential in the cases with irreducible atlantoaxial dislocation.

**References**