Grisel’s Syndrome Induced by *Mycobacterium tuberculosis*

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Grisel’s syndrome is a non-traumatic subluxation of the atlantoaxial joints, which is caused by an inflammatory process in the upper neck. It is rare to find literary reports of Grisel’s syndrome with an evident pathogen in a lesion. For the first time in Korea, we report a 36-year-old female with Grisel’s syndrome having an atlantoaxial subluxation, which was caused by a retropharyngeal abscess secondary to pulmonary *Mycobacterium tuberculosis*. The patient was treated with an anti-tuberculosis regimen and was prescribed a Philadelphia collar for the control of torticollis. The result of magnetic resonance imaging (MRI) showed an improved atlantoaxial alignment, after drug treatment and immobilization. This patient was neurologically intact and free from symptomatic complaints at follow-up visit. Dynamic cervical radiograph confirmed that the atlantoaxial joints had been stable. The pathophysiology of Grisel’s syndrome, along with anatomical attributes, was explained on the basis of the patient’s clinical course.

Key Words: Grisel’s syndrome · *Mycobacterium tuberculosis* · Non-traumatic atlantoaxial rotatory subluxation · Torticollis

INTRODUCTION

Grisel’s syndrome or “La maladie de Grisel” is a rare non-traumatic atlantoaxial subluxation of the C1-C2 joint, characterized by a spontaneously-occurring unilateral or bilateral misalignment of the atlantoaxial joint. Grisel’s syndrome is sometimes associated with subluxation of the atlantooccipital joint. Sir Charles Bell first described a non-traumatic atlantoaxial subluxation in a patient with syphilis, pharyngitis and resultant lethal spinal compression in 1830. Subsequently in 1930, the syndrome was named after Pierre Grisel, who reported two cases of pharyngitis and atlantoaxial subluxation. It is a rare condition, usually seen in children with an upper cervical infection or after an otolaryngeal procedure. Grisel’s syndrome is not associated with a trauma or bone disease. Patients generally complain about neck stiffness, pain, and sometimes, difficulty swallowing. Diagnosis is established by clinical and radiological findings. Thus, demonstrable pathogens in cases with Grisel’s syndrome are rarely reported. It is rather rare to find reports of Grisel’s syndrome caused by *Mycobacterium tuberculosis*. This is the report of a 36-year-old female with Grisel’s syndrome, who developed atlantoaxial subluxation following a retropharyngeal abscess, secondary to pulmonary *Mycobacterium tuberculosis*.

CASE REPORT

A 36-year-old female patient was admitted to the outpatient department of this hospital with complaints of neck pain, stiffness, and twisted neck. She could not turn her head to the right, since having a bitter quarrel six months ago. There was no history of physical trauma. Nevertheless, she had suffered emotional stress by delivering a baby with fetal deformity. She had been diagnosed as having somatization disorder, neurologically, by consultants after review of several of her radiologic images at the time. A conservative management had not improved her symptoms, and she visited the department of neurosurgery of our hospital after several consultations. A neurological examination showed right ankle rigidity with markedly increased deep tendon reflex. She could not stand up by herself without the help of a crutch on her right side (Fig. 1). The range of motion in her neck was completely limited, and the patient suffered severe neck pain and stiffness. There was no fever and the routine blood tests were within normal limits. She had suffered pulmonary tuberculosis, which was completely resolved after she was treated with oral medication about 16 years ago. Her left foot cellulitis was also resolved after she was treated with intravenous antibiotics about a year ago.
Recently, this patient had no signs of retropharyngeal infection, but her body weight decreased about 12 kg during a four month period. Simple radiographs showed severe kyphoscoliotic cervical curvature (Fig. 2). The axial computed tomography (CT) scans (Fig. 3) of the cervical spine revealed a severe atlantoaxial subluxation with C1 rotated to the left on the odontoid process without anterior displacement. These are reliable findings, which correspond to type I in the Fielding classification of atlanto-axial rotatory subluxation. There were no evidences of bone injury, bone disease, or hemorrhagic sign. MRI verified the CT findings, which demonstrated that the transverse ligament was intact (Fig. 4). However, T2-weighted images revealed hyperintensity of the ligaments and edema of the atlantoaxial joint, and less of the atlanto-occipital joints, with thick peripheral enhancement from C1 to T2. This was interpreted as an infectious abscess pocket of the ligaments and joint. Indeed, multifocal nodular densities were observed in the simple chest radiographs. This suggested reactivation of pulmonary tuberculosis, compared to previous chest images.

Pulmonary tuberculosis was observed in both upper lobe and left lower lobe from the chest CT. Tuberculosis was immunologically confirmed by utilizing tuberculosis polyme-
Grisel’s syndrome involves a subluxation of the atlantoaxial joint and sometimes lead to a partial dislocation of the atlanto-occipital joint. It usually affects children between 6 and 12 years of age, but may also occur in adults. Grisel’s syndrome was first described in 1930. However, it is a rare disease with an apparently low index of suspicion. History taking and clinical signs and symptoms are important clues in the diagnosis of Grisel’s syndrome.

Patients generally seek treatment for progressive throat and neck pain, and pain on attempted motion. One of the outstanding symptoms of Grisel’s syndrome is a spontaneously arising, which is the symptomatic manifestation of unilateral or bilateral atlantoaxial subluxation. Neurological complications occur in approximately 15% of these cases and can range from radiculopathy to myelopathy, and may even result in death. A subluxation may not yet be evident at the onset of the disease. Early recognition of the infectious disease and a prompt attempt of appropriate conservative treatment are essential in the prevention of progression from such infectious disease to Grisel’s syndrome. Several theories have been proposed to explain the etiology and pathogenesis of atlantoaxial subluxation in Grisel’s syndrome. There is a general agreement today that the disorder may occur in association with any condition or surgical procedure that results in hyperemia and pathologic laxity of the transverse ligament and of the atlantoaxial joint. Therefore, the prospective pathophysiology of subluxation of the atlantoaxial joint may be inflammatory ligamentous laxity caused by an infection. It is largely associated with an overt or occult head and neck infection, especially, an otolaryngologic infection or cervical lymphadenitis. Furthermore, Grisel’s syndrome has been reported postoperatively after tonsillectomy, adenoidectomy, repair of choanal atresia, surgery for the otitis media and surgery for a neck disease. It is well known that patients with clefts are prone to upperairway infections, even more so after surgery. It is also postulated that infection may spread directly into the small atlantoaxial joints and thus into the surrounding ligaments. This infection may increase the laxity of the ligaments of the atlantoaxial joint. This in turn may explain the gap between the dens and the front arch of atlas. Several authors support these assumptions, but pathogens of Grisel’s syndrome are rarely identified. Paping et al. found Fusobacterium necrophorum, an anaerobic bacterium, in the blood and ear cultures of a child with Grisel’s syndrome after the child had been inflicted with bilateral fulminant otitis media. Clark et al. described a case of infection-related atlantoaxial subluxation without torticollis, from which a retropharyngeal beta-hemolytic streptococcal abscess was found. Those pathogens described by another study included: Pseudomonas aeruginosa (1 case), Staphylococcus aureus (1 case) or Methicillin-resistant Staphylococcus epidermidis (1 case) (isolated variably from the neck and blood cultures), and Epstein-Barr virus (of infectious mononucleosis). However, there was no case associated with Mycobacterium tuberculosis.

Any one treatment method for Grisel’s syndrome would not be universally accepted. Several authors would consider a conservative treatment for type I (immobilization, antibiotics, rest and analgesia) and type II (includingcervical traction or muscle relaxants) while they may utilize a more invasive approach for types III and IV (halo immobilization, arthrodesis and C1-C2 cervical fusion). Nevertheless, these various therapeutic approaches may be applicable in each case, and patients should be individually managed. A proper management of Grisel’s syndrome should be based on three aspects: (1) a specific treatment of the infectious disease, (2) correction of the bone deformity and (3) prevention of neurological damage. Further...
Moreover, a neurosurgical consultation is essential for all cases, even mild ones7). This case had no anterior displacement and had less than 3 mm of atlantoaxial space, which corresponds to Type 1 subluxation in the Fielding’s Classification. Concurring to the recommendations for the management of type 1 subluxation of the Fielding’s Classification, she was prescribed with a soft collar to wear for cervical immobilization, antibiotics, and botulinum toxin as a supplementary muscle relaxant. In an attempt to correct severe subluxation with a kyphoscoliotic curvature, anterior cervical disectomy and fusion (ACDF) were performed at the C4-5 level in this patient. Also, she was put on an oral medication for a long time. Her symptoms of torticollis and pain improved. Furthermore, radiographs and MR images showed an improved atlantoaxial alignment and decreased posterior epidural abscess after administration of oral drugs or after surgery.

**CONCLUSION**

Proper and effective treatment leads to excellent prognosis. Mild subluxation types (Fielding types I and II) are not typically associated with a neurologic impairment. Nevertheless, it was reported that cases with Fielding type III and IV lesions were associated with 15% incidence of neurologic impairment, ranging from radiculopathy to paralysis and death10. Most of these cases reported in current literature have a successful outcome with a conservative treatment, including this case. Without treatment, an atlantoaxial subluxation becomes permanent, and will result in chronic torticollis, with a possibility of facial flattening and neurological complications.

**REFERENCES**

1. Battiata AP, Pazos G: Grisel’s syndrome: the two-hit hypothesis - a case report and literature review. *Ear Nose Throat J* 83:553-555, 2004
2. Bredenkamp JK, Maceri DR: Inflammatory torticollis in children. *Arch Otolaryngol Head Neck Surg* 116:310-313, 1990
3. Clark WC, Coscia M, Acker JD, Wajnschtit K, Robertson JT: Infection-related spontaneous atlantoaxial dislocation in an adult. Case report. *J Neurosurg* 69:455-458, 1988
4. Fielding JW, Hawkins RJ. Ratzan SA: Spine fusion for atlantoaxial instability. *J Bone Joint Surg Am* 58:400-407, 1976
5. Fielding JW, Hawkins RJ, Hersinger RN, Francis WR: Atlantoaxial rotary deformities. *Orthop Clin North Am* 9:955-967, 1978
6. Galer C, Holbrook E, Treves J, Leopold D: Grisel’s syndrome: a case report and review of the literature. *Int J Pediatr Otorhinolaryngol* 69:1689-1692, 2005
7. Karkos PD, Benton J, Leong SC, Mushi E, Sivaji N, Assimakopoulos DA: Grisel’s syndrome in otolaryngology: a systematic review. *Int J Pediatr Otorhinolaryngol* 71:1823-1827, 2007
8. Konttinen YT, Bergroth V, Santavirta S, Sandelin J: Inflammatory involvement of cervical spine ligaments in patients with rheumatoid arthritis and atlantoaxial subluxation. *J Rheumatol* 14:531-534, 1987
9. Kraft M, Tschopp K: Evaluation of persistent torticollis following adenoidectomy. *J Laryngol Otol* 115:669-672, 2001
10. Meek MF, Hermens RA, Robinson PH: La maladie de Grisel: spontaneous atlantoaxial subluxation. *Cleft Palate Craniofac J* 38:268-270, 2001
11. Mezue WC, Taha ZM, Bashir EM: Fever and acquired torticollis in hospitalized children. *J Laryngol Otol* 116:280-284, 2002
12. Papin RH, Annyas AA, Mooij JJ, Penning L: Nasopharyngeal torticollis (Grisel’s syndrome). *Ned Tijdschr Geneesk* 129:1787-1789, 1985
13. Welinder NR, Hoffmann P, Hakansson S: Pathogenesis of non-traumatic atlanto-axial subluxation (Grisel’s syndrome). *Eur Arch Otorhinolaryngol* 254:251-254, 1997
14. Wurm G, Aichholzer M, Nussbacher K: Acquired torticollis due to Grisel’s syndrome: case report and follow-up of non-traumatic atlantoaxial rotatory subluxation. *Neuropediatrics* 35:134-138, 2004