A Rare Complication Following Adenotonsillectomy: Grisel’s Syndrome

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The nontraumatic subluxation of the atlantoaxial joint, also called Grisel’s syndrome, is a rarely seen phenomenon in the pediatric population after nasopharyngeal inflammation or otolaryngological surgery. The patients are usually between 5 and 12 years old, and there is no gender difference in occurrence. Grisel’s syndrome should be considered in patients who have painful torticollis after an upper airway infection or otolaryngological surgery. The most important factor affecting the prognosis is early diagnosis and treatment. Early diagnosis not only allows for conservative treatment, but it also prevents permanent neck deformity and the development of severe neurological deficit and morbidity due to extensive surgical procedures. Otolaryngologists should be aware of this rare, but potentially serious, condition.

ABSTRACT
The nontraumatic subluxation of the atlantoaxial joint, also called Grisel’s syndrome, is a rarely seen phenomenon in the pediatric population after nasopharyngeal inflammation or otolaryngological surgery. The patients are usually between 5 and 12 years old, and there is no gender difference in occurrence. Grisel’s syndrome should be considered in patients who have painful torticollis after an upper airway infection or otolaryngological surgery. The most important factor affecting the prognosis is early diagnosis and treatment. Early diagnosis not only allows for conservative treatment, but it also prevents permanent neck deformity and the development of severe neurological deficit and morbidity due to extensive surgical procedures. Otolaryngologists should be aware of this rare, but potentially serious, condition.

INTRODUCTION
Nontraumatic atlantoaxial rotatory subluxation was first described by Charles Bell in 1830 in a patient with syphilis and pharyngitis. Grisel defined it as a syndrome in 1951. Grisel’s syndrome is atlantoaxial rotatory subluxation that occurs after a previous infection or otolaryngological surgery other than trauma.[1]

Children are more often affected; patients are usually between 5 and 12 years old. There is no gender difference in occurrence. Patients typically have torticollis with neck stiffness or painful neck movements. Torticollis may occur related to pharyngitis or after a slight trauma to the neck. This syndrome has been reported to be associated with surgical procedures, such as mastoidectomy, adenotonsillectomy, choanal atresia repair, and other situations, like rhinopharyngitis, cervical osteomyelitis, and rheumatological diseases. Tension or abnormal laxity of the ligaments around the atlantoaxial joint resulting from direct expansion of pharyngeal or nasopharyngeal inflammation can cause neurological damage or cervical spine instability.[2]

CASE REPORT
An 8-year-old boy was brought by his parents to the clinic with complaints of recurrent tonsillitis, nasal obstruction, and snoring for three years. An otolaryngological examination revealed that both palatine tonsils were quite large with adenoid tissue closely covering the choana, while the bilateral tympanic membranes were normal in appearance. Other systemic examinations were within the normal limits. The patient was diagnosed as having chronic adenotonsillitis with adenoid vegetation. An adenotonsillectomy operation was planned. After a successful surgical intervention, on the third postoperative day, the patient was admitted to the otorhinolaryngology clinic with the
complaints of the neck leaning to one side and painful neck movements. Oropharyngeal examination revealed no evidence of infection. A severe spasm of the right sternocleidomastoid muscle was observed. A neurology consultation was requested, and the results of a neurological examination were normal. There was no history of trauma after the operation. For further investigation, a cervical vertebrae X-ray was obtained in the anterior-posterior (open-mouth) and lateral projections. The X-ray images demonstrated a widened atlantodental interval and anterior displacement of the cervical C1 and C2 vertebrae bodies (Fig. 1a). According to these findings, the radiologists recommended cervical computed tomography (CT). A cervical CT scan confirmed the widening of the atlanto-dental interval and anterior displacement of the C1 and C2 vertebrae bodies (Fig. 1b). CT images also showed a widened space between the dens and left C1 lateral mass, and rotation of the atlas on the lateral articular process with 5.2 mm of anterior displacement consistent with type II atlantoaxial rotatory fixation (AARF) (Fig. 2a). The C1 vertebrae were turned to the left, and subluxation on the occipital condyle of 14° was seen (Fig. 2b). Following a consultation with a neurosurgeon, a Philadelphia cervical collar, arms, antibiotherapy, and anti-inflammatory treatment were applied for 2 weeks. At the end of 2 weeks, the torticollis was resolved and the painful neck movements were healed.

**DISCUSSION**

The nontraumatic subluxation of the atlantoaxial joint is only rarely seen in the pediatric population after nasopharyngeal inflammation or otolaryngological surgery, and its pathogenesis is not fully understood.\(^3\)

Grisel’s syndrome is graded according to the Fielding classification, based on the degree of displacement of the atlas on the axis determined by radiological examination.\(^5\) The Fielding classification of atlantoaxial rotatory fixation includes 4 types: Type 1, simple rotatory displacement without anterior shift, the transverse ligament is intact and the dens acts as a pivot point; type 2, rotatory and anterior displacement between 3 and 5 mm, the transverse ligament is injured and the opposite facet acts as a pivot point; type 3, rotatory and anterior displacement greater than 5 mm with both lateral atlantoaxial joints anteriorly subluxated, and the transverse and alar ligaments are injured; and type 4, subluxation of the both lateral atlantoaxial joints.\(^2\)

Although Grisel’s syndrome was first described in the 1830s, there has been long-standing and unresolved controversy regarding the pathogenesis, diagnosis, and the best treatment modalities for the syndrome. There is no universally accepted treatment model. Some authors\(^6\) suggest conservative treatment (immobilization, antibiotics, resting, and analgesics) for types 1 and 2 (cervical traction and muscle relaxants in addition to type 1) and more invasive methods (halo immobilization arthrodesis and C1-C2 cervical fusion) for types 3 and 4. Wetzel and Lorracca\(^7\) proposed another treatment algorithm: a soft
cervical collar for type 1, rigid arms for type 2, close fixation with halo for type 3, and open fixation with halo for type 4.

The goal of treatment of Grisel's syndrome is to prevent bone deformities and to treat infectious disease and prevent neurological sequelae. All of these treatment modalities can be personalized to the patient. In our case, we applied antibiotherapy for 10 days, a soft cervical collar, and 2 weeks of resting as analgesic treatment for type 2 subluxation. After 2 weeks, we observed complete recovery.

Grisel's syndrome is a rare condition. In patients who have painful torticollis after an upper airway infection or otolaryngological surgery, Grisel's syndrome should be suspected. The most important factor affecting the prognosis is early diagnosis and treatment. Early diagnosis not only allows for conservative treatment, but also prevents permanent neck deformity and the development of severe neurological deficits and morbidity due to extensive surgical procedures. Otolaryngologists should be aware of this rare, but potentially serious, condition.

Informed Consent
Written informed consent was obtained from the patient for the publication of the case report and the accompanying images.

Peer-review
Internally peer-reviewed.

REFERENCES