A case of combined congenital anterior and posterior arch anomaly of C1 vertebra

C1 vertebrenin kombine konjenital anterior ve posterior arkus anomalisi olgusu

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ABSTRACT

A very rare case of the craniovertebral junction (CVJ) anomaly is reported owing to its rarity and clinical significance. Computed tomography (CT) and magnetic resonance imaging (MRI) studies of the cervical spine of a 56-year-old female patient presented with head trauma, revealed an anterior midline arch cleft of atlas with totally aplastic posterior neural arch, as well as anteriorly subluxed right atlantoaxial joint. The coexistence of a total aplasia of the posterior neural arch and anterior arch cleft of atlas is very rare. Congenital anomalies affecting the craniovertebral junction (CVJ) may mimic traumatic injury in the setting of a previous trauma. In this presented case, the absence of an associated bone and soft-tissue edema on the MR imagings helped us to make a differentiation between the acute traumatic injury and congenital anomaly.

Keywords: Cervical Atlas, Magnetic Resonance Imaging, X-Ray Computed Tomography

ÖZ


Anahtar kelimeler: Servikal Atlas, Manyetik Rezonans Görülteme, Bilgisayarlı Tomografi.

Introduction

Congenital clefts and aplasias of the C1 vertebrae are rare. These anomalies develop in intrauterine life as a result of developmental failure of chondrogenesis which may variate from partial clefts to total agenesis of the atlas rings [1-3]. In patients with a history of trauma, these anomalies can mimic an injury of the craniovertebral junction (CVJ) [1,4].

We present a case with traumatic craniofacial injury which has a rare association of congenital anterior and posterior arch anomaly of C1 and coexist with an unilateral anterior atlantoaxial joint subluxation that was initially thought to be post-traumatic.

Case Report

A 56-year-old female, who had accidentaly fell down was admitted with head trauma. An urgent cranial computed tomography (CT) scan showed minor right frontotemporal-zygomatic superficial haemorrhage, and more caudal images of cranial CT depicted an abnormal atlas. At the time of admission, physical examination revealed no pathological findings with a full range of cervical movement and a normal neurological status. All laboratory investigations showed normal values. Due to the better demonstration of CVJ, a CT scan of the cervical spine was then performed. Total aplasia of posterior arch and midline cleft of the anterior arch of atlas were detected. There was partially anterior subluxation of right lateral mass of atlas over axis (Fig. 1,2). The pre-vertebral soft tissue at the upper cervical spine was not thickened (measuring about 4.9 mm at the C2 level on the mid-sagittal multiplanar reformat image). The distance between dens and atlas was measured 3 mm as normal. The rest of the cervical spine...
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Discussion

Congenital defects of the arches of C1 vertebra are rare anomalies and usually discovered incidentally. To have a detailed knowledge about embryological development is crucial to understand congenital atlas arch defects. A normal C1 vertebra can be divided into three parts including the anterior arch, the lateral masses, and the posterior arch. Three primary ossifications are responsible in formation of its parts including: an anterior ossification center which forms the anterior tubercle and two lateral centers which form the lateral masses and the posterior arch. In the seventh week of intrauterine life, ossification centers at the lateral masses extend posteromedially to form the posterior arch. A secondary center of ossification center which is the center for the posterior tubercle, may be present in about 2% of cases. Failure in any step of this complex process leads to abnormalities [1,3,4].

Curarrino et al. classified the posterior arch anomalies into five types depending on the absence of the posterior arch, the presence or absence of the posterior tubercle, and its clinical findings [1] (Table I). Based on the literature,
roughly 4% of patients present with congenital posterior arch defects, with the predominant (97%) type being Type A [1,4,5]. Type B through Type E congenital defects have been reported to occur in 0.69% of the population [1]. Type E defects are very rare. One study found two patients (0.18%) [3], while two studies did not find any patients with Type E defects and had patient populations of 1069 and 1153 respectively [4,5]. There have only been sporadic case reports of patients with Type E posterior arch anomalies. Anterior arch anomalies are much more uncommon than the posterior arch anomalies, and the reported incidence of the anterior arch anomalies is 0.1% [1,4,6,7]. The association of anterior and posterior arch defects, called a bipartite atlas, has been described in some studies [5,7]. Our case, may be classified as Type E and clinical subgroup 1, according to Currarino’s classification of posterior arch and coexistent with anterior arch midline cleft anomaly [1].

\[\text{Table I: Classification of the posterior arch anomalies} \text{[1]}\]

| Types                                  | 1) Asymptomatic incidental findings  
|                                       | 2) Neck pain or stiffness after trauma to the head or neck  
|                                       | 3) Chronic symptoms referable to the neck  
|                                       | 4) Various chronic neurological problems  
|                                       | 5) Acute neurological symptoms after minor cervical trauma  

Craniovertebral junction anomalies are commonly asymptomatic and found incidentally. When symptomatic, the symptoms typically begin insidiously and arise late, progress slowly, remain constant, and rarely relapse. The signs and symptoms of CVJ anomalies may be related to cervical spinal cord, the brain stem, cerebellum, cervical nerve roots, lower cranial nerves, or the vascular supply to these structures [1,3-5,8]. Our patient had no complaints or symptoms for detected C1 anomaly and, as in most cases reported in the literature, the anomaly was discovered incidentally by radiologic imaging after a trauma.

Posterior arch anomalies may be accompanied by other abnormalities, including clefts of the anterior arch, anterior atlantoaxial subluxation, rotatory atlantoaxial subluxation and cervical myelopathy are reported in the literature [1]. In our case, posterior arch anomaly of atlas exists together with both anterior arch cleft and anteriorly subluxed right atlantoaxial joint, and additionally with bilateral cervical ribs.

Imaging remains the first option for the evaluation of the patients and congenital anomalies of the CVJ are common incidental findings on cervical plain radiography. These anomalies should be kept in mind to avoid misinterpretation as fractures, luxation or osteolysis. Cervical lateral plain radiography with flexion and extension view may help to detect cases with instability where there is increased risk of spinal cord injury. CT is able to depict the nonossified and ossified portions of the arches of atlas (Fig. 1). It is especially useful for evaluating the integrity of the atlas rings, differentiating injury from a developmental cleft and demonstrating a small defect in the arch [1]. Additionally, CT with 3D reconstruction might display the topography of the upper cervical spine better, especially to clinicians (Fig. 2). Although, MRI is inadequate for diagnosing because of the lack of magnetic resonance signal from cortical bones, but osseous and/or periosteal soft tissue edema is easily visible and an important finding as indicating trauma or inflammation, thus making MRI a valuable tool for differentiating acute traumatic spinal lesions from other congenital and acquired abnormalities [9]. Furthermore, MRI study has advantages in evaluating the spinal cord, adjacent neural
structures and paraspinal soft tissue to exclude or confirm extrasosseous causes of clinical symptoms [1,2,5,9]. Our patient was diagnosed coincidentally in the setting of an acute trauma, while anomaly was easily detected by CT and MRI showing no signal alteration neither in bone nor in the joints as well as no soft tissue edema at the region of CVJ and no cord compression and/or signs of myelopathy (Fig. 3). This indicated that the findings, especially subluxed right atlantoaxial joint, was not acutely caused by recent trauma. It is thought to be nontraumatic chronic subluxation due to the existing congenital anomaly.

Surgery is the treatment of choice in symptomatic patients [10]. CVJ anomalies in patients with history of trauma, the possibility of spinal cord injury should always be considered. Neurological symptoms may become symptomatic after minor cervical trauma in patients with this anomaly [8]. For the current patient, since there was no symptoms due to this congenial anomaly, and there was no clinical and radiological sign of atlantoaxial or occipitocervical instability, the decision was made not to undergo for any non-surgical and surgical management. However, in such patients with this anomaly, it is recommended to avoid contact sports.

In this presented case, it was possible to evaluate the absence of associated soft-tissue and bone marrow edema by using MRI, which allowed non-invasive differentiation between the cases of acute injury and congenital anomaly. When upper cervical spine anomalies are found in a patient with acute trauma, the patient should be evaluated in detail with CT and MRI for avoiding misinterpretation as fractures, traumatic luxation, or instability. Advanced radiological studies could help making accurate diagnosis and deciding on therapeutic methods.

References