Os Odontoideum - “MR documented Craniocervical Ligamentous Injury in Early Childhood with delayed formation of Os Odontoideum on Sequential Imaging”. Case-based Review and Mini Database Analysis

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Abstract
The etiology of os odontoideum has been debated ever since its first description by Giacomini in 1886. There are proponents of congenital origin as well as reports of post-traumatic os odontoideum formation after early childhood craniovertebral junction (CVJ) trauma. We document CVJ ligamentous injury on MRI in an 18-month-old child with sequential imaging demonstrating the os odontoid formation. Our database of 260 surgically treated patients with os odontoideum was analyzed regarding etiology and associated abnormalities. The literature (1970-2022) is reviewed.

Introduction
Osodontoideum is radiographically described as an independent ossicle posterior to hypertrophic anterior atlas arch, with cortical margins, apart from a hypoplastic dens. Ever since its first description in 1886 by Giacomini, the origin has been debated. Numerous case reports and case series show both a possible congenital origin as well as secondary to early craniovertebral junction (CVJ) trauma. There is support for both causations. We document CVJ ligamentous injury on MRI in an 18-month-old child with sequential imaging demonstrating the os odontoid formation. The patient also had a Chiari I abnormality on the initial MRI. He became clinically symptomatic from craniovertebral junction instability and the Chiari abnormality needing both pathologies to be addressed at age 10.

Clinical Case
In 2006 this child presented at 18 months of age after having fallen out of a crib. He was referred to University of Iowa Hospitals & Clinics because of inability to move his trunk and poor hand movement. He did not move from side-to-side and had ecchymosis on the right forehead. A cervical collar was immediately placed, and the patient underwent magnetic resonance imaging. The cervico-medullary junction was enlarged and there was an 8-mm tonsillar herniation consistent with a Chiari I abnormality. There was an area of linear brightness on T2-W MRI lifting the anterior longitudinal ligament between C1 and C3. A small odontoid process was identified (Figure 1A, 1B). We felt that he had craniocervical ligamentous injury and cervico-medullary contusion with Chiari I abnormality. The clinical picture of neurological deficit, swollen cervico-medullary junction and MRI findings confirm the trauma to the craniovertebral area and ligamentous injury. Further evidence
is the presence of bony spicule off the posterior rim of the ossiculum penetrating the dura as found at transoral resection. He was maintained in a Philadelphia collar and started on Decadron with an immediate taper. He showed marked improvement neurologically and at 2 month follow up his neurological examination was normal.

Lateral cervical spine radiographs at 2 months showed an incomplete development of the anterior and posterior arches of the atlas with normal alignment. Magnetic resonance imaging showed soft tissue crowning of the odontoid process and cerebellar tonsillar ectopia. The prevertebral swelling had receded. The cruciate ligament and the alar ligaments were investigated as well as the tectorial membrane and found to be intact. Five months following injury his cervical spine radiographs showed no evidence of instability. An Aspen brace was maintained for a year following which a CT scan of the neck showed irregular ossification of the axis body, a hypoplastic dens on CT and a small odontoid process.

At age 9 he had lateral cervical spine radiographs made elsewhere following a school altercation that demonstrated the dystopic odontoideum. He was evaluated at age 10 for neck pain and occipital headaches, difficulty swallowing and bladder incontinence. He complained of numbness in his hands and tended to drop objects. Headaches were worse with Valsalva maneuver. Magnetic resonance imaging showed the Chiari I malformation with tonsils down to 20 mm below the plane of foramen magnum and a syrinx. A large os odontoideum was visible on CT. He had gross atlantoaxial instability on cervical spine dynamic radiographs. He underwent posterior fossa procedure for both the Chiari abnormality as well as dorsal occipitocervical fusion, which had to be revised. Despite this and a successful dorsal occipitocervical fusion, he had new findings of a Lhermitte’s phenomenon with neck movement a year later. This responded to an anterior transoral-transpalatopharyngeal resection of the odontoid and medullary decompression. He recovered. There was a bony spicule penetrating the dura from the posterior aspect of the os. This is only the second MRI documentation of early childhood CVJ trauma and sequential os odontoideum formation.

**Database Analysis**

We undertook a detailed analysis of 260 surgically treated cases of os odontoideum to document the age groups, first symptom presentation, radiographic findings and associated abnormalities to better confirm the etiology. A literature search (1970-2022) was made to correlate our findings.

260 patients underwent surgical management of a referral database of 520 cases (1978 to 2022). All surgically treated patients underwent neurodiagnostic imaging. A history of early childhood (less than 6 years of age) CVJ trauma was investigated including obtaining emergency departments initial radiographs from the national referral database.
and subsequent follow up. Associated radiographic and systemic abnormalities were noted. There were 156 of the 260 patients who had a previous history of early childhood trauma. We were able to obtain cervical spine documentation of a normal odontoid process at the initial injury in 54 of the 156 who subsequently developed an os odontoideum.

Discussion

The etiology of os odontoideum is debated. Proponents of the congenital os odontoideum claim that this may be the result of failure of fusion of the odontoid apex or the main part of the odontoid or failure of separation of C1 and caudal migration. However, there is always a hypoplastic dens below the os and the gap between the two is above the lateral C1-C2 articulation and hence the etiology of failure of fusion of C1 centrum to the axis body is difficult to accept. There are reports of congenital origin with the presence of ossicle in utero images in Down syndrome. It is also associated with syndromic abnormalities such as Morquio syndrome, Klippel-Feil syndrome, spondyloepiphyseal dysplasia, metaphyseal dysplasia, Down syndrome and other syndromes with ligamentous laxity. There have been reports of familial os odontoideum as well as in twins as seen in our database as well as in the literature. In our own database the ages range from 2 to 58 years. Associated abnormalities were 94/260 patients (Table 1). It appears that dystopic os odontoideum is associated with traumatic origin.

The true congenital os odontoideum has a very large ossicle and could represent a large ossiculum terminal persistens. None of the post-traumatic origin had associated bony abnormalities.

Early childhood trauma was documented in our database of surgically treated cases (156/260) with a documented normal odontoid process at the first initial encounter in the emergency facilities in 54/156 with a subsequent development of os odontoideum. It was Fielding et al.3 who initially brought up the possibility of acquired post-traumatic lesion caused by odontoid injury followed by what he felt was avascular necrosis and an osseous remodeling. Zygourakis et al. in 2011 illustrated MR findings in the initial trauma of a 2-year-old child who sustained ligamentous injury and subsequent formation of os odontoideum. It has been proposed that the initial trauma led to vascular compromise as with a type II odontoid fracture. The apical vascular arcade continues to supply the apex of the odontoid process which enlarges as also the anterior arch of the atlas with posterior arch becoming smaller. This would be a tell-tale sign of a traumatic origin since the original cervical spine radiographs showed a normal anterior and posterior arch of the atlas. The final stage is the osseous remodeling in the development of the os odontoideum. The initial injury was likely between the upper and the lower central synchondrosis of the odontoid. The associated abnormalities (94/260) in the database points also to a possible congenital origin. In our case, the increased descent of the cerebellar tonsils was likely attributed to decrease in posterior fossa volume that was corrected after resection of the os odontoideum and decompression of the medulla. Posterior craniocervical fusion has been proposed as treatment for the os odontoideum but it appears that this did not protect this child.

The variants from the literature with the population base is likely due to our referral pattern and that of syndromic and pediatric population.

Acknowledgment

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Table 1: Associated conditions with Os Odontoideum (260 patients)

<table>
<thead>
<tr>
<th>Pathology</th>
<th>#</th>
<th>Age at presentation</th>
<th>Orthotopic</th>
<th>Dystopic</th>
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<tr>
<td>Klippel-Feil syndrome*</td>
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<td>All ages</td>
<td>9</td>
<td>17</td>
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<tr>
<td>Chiari I malformation*</td>
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<td>All ages</td>
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<td>20</td>
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<td>Down syndrome</td>
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<td>4</td>
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<td>2-5 years</td>
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<td>3</td>
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<td>Bipartite atlas arches</td>
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<td>4-6 years</td>
<td>0</td>
<td>6</td>
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<tr>
<td>Spondyloepiphyseal dysplasia</td>
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<td>6-10 years</td>
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<td>Craniofacial dysostosis*</td>
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<td>Dandy-Walker malformation</td>
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<td>Charge syndrome</td>
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<td>0</td>
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</table>

TOTAL 94

*had combination or other abnormalities
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Conflict of Interest

No conflict of interest was reported by the author.

References