Traumatic Atlanto-occipital Dislocation in Children Followed by Hydrocephalus – A Case Report and Literature Review

Deslocamento atlanto-occipital em crianças seguido por hidrocefalia – Relato de caso e revisão da literatura

Cleiton Formentin¹ Lucas de Souza Rodrigues dos Santos¹ Fernando Luis Maeda¹ Helder Tedeschi¹ Enrico Ghizoni¹ Andrei F. Joaquim¹

¹ Neurosurgery Division, Department of Neurology, Universidade de Campinas (UNICAMP), Campinas, SP, Brazil

Arq Bras Neurocir 2022;41(3):e262-e269.

Address for correspondence Cleiton Formentin, MD, Divisão de Neurocirurgia, Departamento de Neurologia, Universidade de Campinas, Rua Tessalia Vieira de Camargo, 126, Campinas, SP 13083-970, Brazil (e-mail: cleitonformentin@gmail.com).

 \bigcirc \bigcirc \bigcirc \bigcirc \bigcirc \bigcirc

Abstract

Traumatic atlanto-occipital dislocation (TAOD) are uncommon injuries associated with high immediate mortality rate and occurs more than twice in children than adults, due to biomechanical properties and immaturity of children's cervical spine. We report a pediatric patient with TAOD, who underwent occipitocervical stabilization and also developed a late hydrocephalus requiring a shunt procedure. A six-year-old boy was admitted to the emergency department after a car accident with refractory cervical pain. A cervical computed tomography (CT) scan showed an anterior C1-C2 level hematoma, and a dynamic CT scan demonstrated an increasing basion-dens interval on extension. Cervical magnetic resonance imaging (MRI) showed discontinuity of the tectorial membrane and diffused hyperintense signal on the left alar ligament. These findings were attributed to TAOD, and an occipitocervical fusion was performed. The pain and neurological status improved after surgery, but after 3 months he returned with persistent vomiting, headache, and a CT scan showing hydrocephalus. Then, a ventriculoperitoneal shunt was performed, improving the symptoms. One year after the injury, the patient remained asymptomatic, and a later radiography demonstrated satisfactory bone fusion. In conclusion, the decision-making process regarding treatment should consider several clinical and radiographic findings. Occipitocervical fusion is the treatment of choice, while hydrocephalus is not an unusual complication in children.

Resumo

Keywords

► trauma

► atlanto-occipital

dislocation

► pediatric spine

hydrocephalus

O deslocamento atlanto-occipital (DAO) é uma lesão incomum associada a uma alta taxa de mortalidade imediata que ocorre duas vezes mais em crianças do que em adultos, fato relacionado às propriedades biomecânicas e à imaturidade da coluna cervical pediátrica. Relatamos o caso de um paciente pediátrico com DAO traumático

received April 15, 2021 accepted June 8, 2021 DOI https://doi.org/ 10.1055/s-0042-1744430. ISSN 0103-5355. © 2022. Sociedade Brasileira de Neurocirurgia. All rights reserved. This is an open access article published by Thieme under the terms of the Creative Commons Attribution-NonDerivative-NonCommercial-License, permitting copying and reproduction so long as the original work is given appropriate credit. Contents may not be used for commercial purposes, or adapted, remixed, transformed or built upon. (https://creativecommons.org/ licenses/by-nc-nd/4.0/)

Thieme Revinter Publicações Ltda., Rua do Matoso 170, Rio de Janeiro, RJ, CEP 20270-135, Brazil

submetido à fixação occipitocervical, evoluindo com hidrocefalia e necessidade de um procedimento de derivação liquórica. Paciente do sexo masculino de 6 anos de idade admitido no pronto-socorro após um acidente automobilístico, apresentando dor cervical refratária. A tomografia computadorizada (TC) de coluna cervical demostrou um hematoma epidural ao nível de C1–C2, e a TC dinâmica evidenciou um intervalo basion-odontoide aumentado em extensão. A ressonância magnética (RM) da coluna cervical demonstrou descontinuidade da membrana tectorial e hiperintensidade difusa no ligamento alar esquerdo. Esses achados permitiram o diagnóstico de um DAO, sendo realizada uma fusão occipitocervical. A dor e o status neurológico melhoraram após a cirurgia, mas 3 meses após, o paciente evoluiu com vômitos persistentes, cefaleia e TC de crânio evidenciando hidrocefalia. Em seguida, foi realizada uma derivação ventriculoperitoneal, com melhora dos sintomas. Um ano após, o paciente permaneceu assintomático, e a radiografia demonstrou fusão óssea satisfatória. Em conclusão, o processo de tomada de decisão quanto ao tratamento deve levar em consideração diversos achados clínicos e radiográficos. A fixação occipitocervical é o tratamento de escolha, enquanto a hidrocefalia não é uma complicação incomum em crianças.

Palavras-chave

- deslocamento atlantooccipital
- coluna vertebral
- ► trauma
- hidrocefalia

Introduction

Traumatic atlanto-occipital dislocation (TAOD) is a rare injury that is associated with a high mortality rate, since it is frequently related to cardiorespiratory arrest at the trauma scene and severe neurological impairment due to injury of the cervicomedullary junction.^{1–3} Traumatic atlanto-occipital dislocation occurs more than twice in children than in adults. This is due to biomechanical properties and the immaturity of the pediatric cervical spine, such as smaller occipital condyles, and a horizontal atlanto-occipital joint, which is less resistant to shear forces.^{4,5} Some case series reported that a certain number of children who survive the initial trauma have a positive outcome despite presenting neurological deficits.^{6,7}

Because of a wide range of clinical presentations, from a neurologically intact patient to one with cardiac arrest at the emergency department, as well as the association with other traumatic brain injuries, the diagnosis of pediatric TAOD can be delayed or even missed entirely, risking irreversible damage to the upper cervical spinal cord and the brainstem.⁸ Determining the stability of the pediatric occipitocervical junction is not always straightforward, given that some patients may present subtle dislocation despite tremendous ligamentous injuries. As a consequence, treatment for less severe cases is not well established, especially in the setting of a normal computed tomography (CT) and only magnetic resonance imaging (MRI) findings.⁸

Additionally, occipitocervical stabilization in children remains a surgical challenge due to small bone dimensions, as well as the regional anatomical complexity, poor occipital bone purchase, significant thinness of the occipital bone, and the lack of instrumentation designed and sized for children.⁹ Furthermore, surgical fixation permanently limits upper cervical spine mobility and predisposes to long-term morbidity; thus, it should be reserved for truly unstable injuries. In the present study, we report a pediatric patient with TAOD who underwent occipitocervical stabilization and also developed a late hydrocephalus requiring a shunt procedure. The diagnosis of TAOD is discussed in detail.

Case Report

A six-year-old boy was admitted to the pediatric emergency department following a car accident, where he was on the backseat wearing a conventional seat belt, and the driver ran off the road and crashed. On admission, he was reporting abdominal pain and had undergone two episodes of emesis. He was otherwise stable, wearing a cervical collar, notifying neck pain. Neurological examination showed evident pyramidal signs, with mild left side weakness. An initial cranial CT scan revealed traumatic subarachnoid hemorrhage, and a cervical CT scan showed an anterior C1-C2 level hematoma (**Fig. 1**). We performed all the following radiological measurements to diagnose TAOD using a CT scan: Wholey densbasion interval (DBI),¹⁰ Powers' ratio,¹¹ Harris' basion-axis interval (BAI),¹² and Sun's interspinous ratio,¹³ but all results had normal values (>Fig. 1). Additionally, the abdominal CT scan showed a mesenteric rupture, treated non-operatively. He was then admitted to the intensive care unit with cervical immobilization. Despite the analgesic management, the cervical pain was refractory, leading us to perform a dynamic (flexion and extension) cervical CT and a MRI.

On the dynamic CT scan, the vertebral bodies and facet joints remained aligned, except for the increasing distance between the basion and the odontoid process on extension (**~ Fig. 2A-B**). The cervical MRI demonstrated a transfixing rupture between the posterior arches of C1 and C2, an anterior arch subluxation with discontinuity of the anterior longitudinal ligament on the medium third of the dens and on the tectorial membrane. The diffuse hyperintense signal on the nuchal ligaments and on the left alar ligament was attributed to a distension/partial

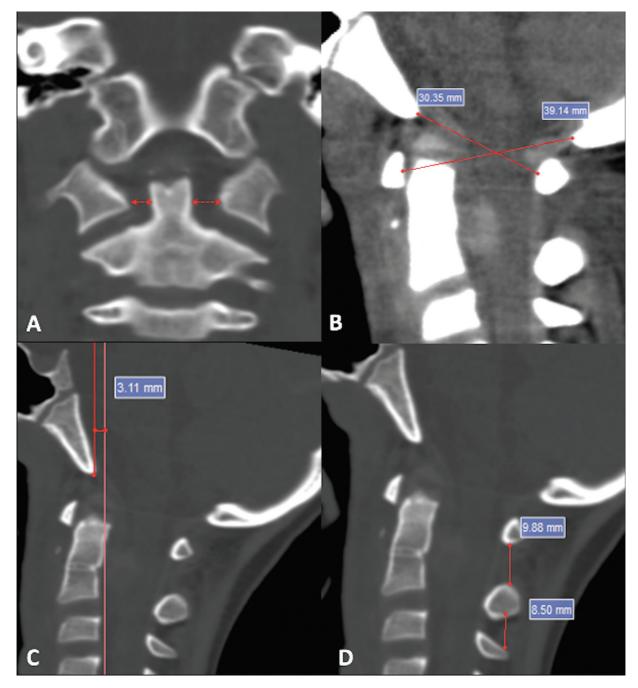


Fig. 1 Admission cervical computed tomography (CT) scan. (A) Slightly Left–right asymmetry between the atlas and vertebrae axis in the coronal plane. (B) Sagittal CT scans showing a normal Powers ratio measurement (0.79; normal \leq 1), but an epidural collection in the upper cervical canal. (C) Normal Harris' basion-axis interval (3.11). The normal distance is between 12 and 0 mm in children. (D) Normal Sun'si ratio (1.16). The interspinous ratio is indicative of atlanto-occipital dislocation (AOD) by a C1–C2/C2–C3 ratio of more than 2.5.

lesion (**- Fig. 3A-B**). The transverse ligament was unimpaired as on method (**- Fig. 3C**). Additionally, an epidural hematoma of 1.9 mL was attached to the left anterolateral spine canal. The entirety of the findings allowed us to diagnose the TAOD, despite the near normal CT findings. Based on the exams and clinical presentation, a multidisciplinary case discussion was held, and a decision was made to perform an occipitocervical fusion (OCF).

While under general anesthesia, the patient was prone positioned, the neck was kept neutral using a Mayfield head holder, and the shoulders were retracted caudally. We performed a posterior median incision from the inion to C3. A subperiosteal dissection exposed the squamous part of the occipital bone, the posterior tubercle of C1, and the spinous process of C2, and then the lamina, and the inferior articular process of C2 and the C2–C3 joint. The posterior arch of the C1 was exposed further with some bleeding from the vertebral plexus, controlled by hemostatic agents and bipolar cauterization. We found and drained the epidural hematoma, secondary to the traumatic avulsion of the right C1 nerve root, with a dural injury and a high debit cerebrospinal fluid (CSF) leakage, fixed with a fat graft and fibrin

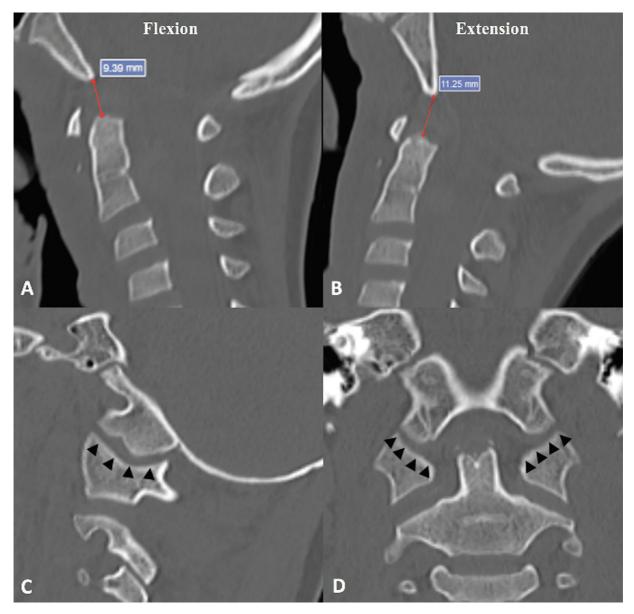


Fig. 2 (A and B) Dynamic computed tomography (CT) scan showing the Wholey's dens-basion interval. Note an increasing dens-basion interval on extension, in this case. (C and D) Conventional CT scan with placement of the measurement points (arrowheads) on the coronal (A) and sagittal (B) planes for calculation of the occipital condyle–C1 interval (CCI) in this case. A value of 4.05 mm was obtained (normal is < 4 mm).

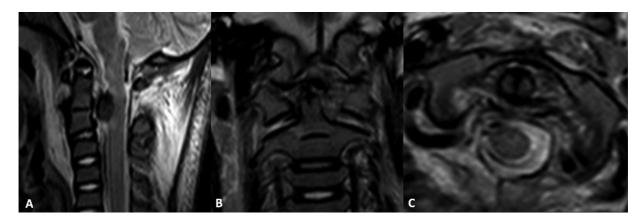


Fig. 3 Cervical magnetic resonance imaging (MRI). (A) Evident hypointense collection on anterior spine canal, causing spinal cord edema, compatible with subacute hemorrhage. There is diffuse hyperintensity on nuchal ligaments. (B) On coronal plane, a left-right asymmetry and also a left alar hyperintensity are evident. (C) The transverse ligament was preserved as on method.

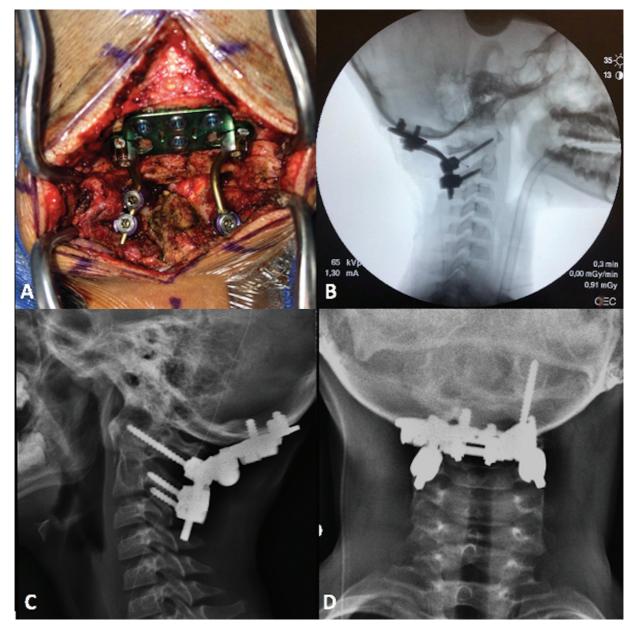


Fig. 4 (A) Intraoperative picture of the occipital plate and bilateral polyaxial pars screws at C2, and a unilateral left C1 lateral mass screw. (B) Lateral radioscopy confirms adequate positioning of the construction. (C and D) Follow-up X-rays with adequate screws positioning and fusion.

glue. An occipital plate was fixed with four screws of $4.5 \times 6 \text{ mm}(2)$ and $4.5 \times 8 \text{ mm}(2)$, centered over the thickest portion of the occipital bone. Guided by fluoroscopy, bilateral polyaxial screws were positioned on the C2 pars $(3.5 \times 14 \text{ and } 3.5 \times 12 \text{ mm})$ and unilateral ones on the left C1 lateral mass $(3.5 \times 26 \text{ mm})$. The system was fixed with 2 adjusted bars, and positioned with an additional cross-link (**> Fig. 4A-B**). Bone graft was extracted from the iliac crest and placed between the occipital bone and C2. There was no significant intercurrence during surgery. A follow-up CT further confirmed the adequate screw positioning.

Thereafter, the patient was transferred to an intensive care unit with significant improvement of the pain, being able to sit on day 1, and having no further symptoms. After hospital discharge, he underwent the follow-up process at the outpatient unit, remaining pain-free. Three months after surgery, however, he presented repetitive nighttime vomiting and headache with no signs of fever. Following a CT scan, a communicating hydrocephalus was diagnosed (**~Fig. 5**). We collected a CSF sample and ruled out meningitis. Then, a frontal medium pressure ventriculoperitoneal shunt was performed, with radiological resolution of hydrocephalus and clinical improvement. One year after the injury, the patient remains asymptomatic, and later radiography showed satisfactory bone fusion (**~Fig. 4C-D**).

Discussion

Traumatic atlanto-occipital dislocations are uncommon injuries associated with high immediate mortality rates. However, the likelihood of a pediatric patient surviving this almost invariably lethal injury clearly improved with time, due to

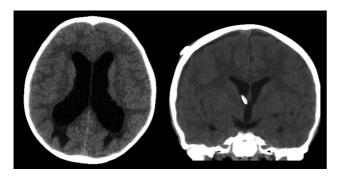


Fig. 5 Left: Head computed tomography (CT) scan showing moderate communicating hydrocephalus. Right: Postoperative CT scan with resolution of hydrocephalus.

advances in emergency resuscitation and prehospital care, as well as in the quality of radiological imaging.^{14,15} Children who survive initial injury may have a favorable outcome with early diagnosis and treatment, before irreversible damage occurs in the cervicomedullary junction.¹⁶

High-energy trauma is usually required to cause TAOD, typically in the form of sudden acceleration-deceleration forces on the head.¹⁷ The mechanism of injury most often reported is an automobile accident,¹⁸ although accident in which a pedestrian is struck by a motor vehicle is also a common cause in children. The high-energy mechanism of injury with TAOD frequently results in further additional injuries, especially traumatic brain injury, which may hamper the diagnosis of TAOD.¹⁹ Our patient had traumatic subarachnoid hemorrhage and blunt abdominal trauma in addition to TAOD.

Anatomical Background

The craniocervical junction is the most mobile part of the spine, and stability is provided mainly by the ligaments.³ The transverse ligament secures anteriorly the odontoid process against the anterior arch of the C1, while the alar ligament attaches the dens to the anterolateral part of the foramen magnum.²⁰ The tectorial membrane is the continuation of the posterior longitudinal ligament and connects the axis with the clivus.^{3,20}

Children younger than 10 years of age are particularly predisposed to TAOD because of the larger head-to-body ratio, smaller and flatter atlantooccipital joints and more flexible and weaker ligaments.²¹ Previous studies showed that rupture of the alar ligaments and the tectorial membrane are sufficient to result in TAOD, since the remaining ligaments that attach the upper cervical spine to the occiput are insufficient to maintain adequate stability, and these abnormalities were also noted in this case.^{22–24} There are usually no fractures associated with this injury, although in older children or adolescents, stronger ligaments can result in avulsion fractures at the ligamentous attachment of the skull base.

Imaging/Diagnosis

Although several radiographic methods to detect TAOD have been described, such as those proposed by Power, Harris, Wholey, and Sun, ^{10–13,25} none have been proven adequate as a single diagnostic criterion.¹⁸ Also, there have been reports of significant variances from the previously accepted normal values on plain radiographs compared with the CT scans for most of these methods.²⁶ High-quality MRI is valuable for detecting ligamentous injury and careful surgeon-supervised flexion-extension CT scan (or even a CT scan performed under cranial traction) can also be informative.

In 1979, Powers et al.¹¹ described the Powers-ratio as a tool for the evaluation of TAOD, diagnosed by a ratio of more than 1. However, this method is only sensitive for the diagnosis of anterior TAOD. A vertical dissociation injury could result in a normal value, like in this case, and consequently go undiagnosed.

Harris et al.¹² established a reliable and accepted method to diagnose TAOD on lateral cervical radiographs, the BAI. In all 50 children (100%) with no occipitocervical abnormalities, the BAI was less than 12 mm, which is considered as the upper limit of normal.¹² However, this test alone has been found to have a sensitivity of 50%.^{16–27}

The DBI was originally described by Wholey et al.,¹⁰ and the commonly accepted cutoff on plain radiograph is 12 mm. However, normal values on CT scans were significantly different from the accepted ranges of normal on plain radiographs.^{26,28} Considering the pediatric population, Bertozzi et al.²⁸ showed that the DBI was shorter than 10.5 mm in 97.5% of patients.

Pang et al. describe both the normal anatomy and radiographic findings suggestive of TAOD,^{15,16} but it has been discussed that these parameters cannot be applied to all age groups.⁸ They showed that the normal occiput–C1 joint in children has an extremely narrow joint gap (condyle–C1 interval or CCI).¹⁵ With a cutoff value of 4 mm, the CCI criterion had the highest diagnostic sensitivity and specificity for TAOD among all other radiographic criteria in their study.¹⁶ In this case, the combined CCI value (average of both the sagittal and coronal CCIs) was 4.05 and represents the only abnormal radiographic standard test, proving this test as an important tool for TAOD diagnosis in children with less evident dislocations (**~Fig. 2C-D**).

Treatment and Complications

Occipitocervical fusion in the pediatric population has been a challenging surgery even for experienced spine surgeons. Posterior spinal fusion is the pillar treatment for TAOD, and the use of screws has now become more common in young children, in whom other techniques, such as rib grafting with wiring followed by a halo vest, are used.^{29–34} In a systematic review in which 285 patients underwent OCF, Hwang et al.³⁵ found that both screw and wiring groups had very high fusion rates (99% and 95%, respectively, p = 0.08); however, wiring was associated with a higher complication rate.

Pediatric OCF carries serious risks, and the complication rates reported in the literature range from 7.5 to 26%.³⁶ Short-term complications include vertebral artery injury, blood loss, neurologic deterioration, dural tear, and CSF leak.³⁶ There are also long-term risks, including hardware-related complications, infection, pseudoarthrosis, and deformity.³⁶

In our case, the patient developed hydrocephalus, which is a complication has also been described in previous case reports of TAOD.^{18,37,38} In a series of 14 patients, the most common postoperative complication was hydrocephalus, and the authors hypothesize that it occurs as a result of posthemorrhagic scarring within the basal cisterns or outlets of the 4th ventricle.¹⁸ In our case, the CSF leak associated with the epidural hematoma may have played a role in the development of a CSF disturb, as well as the root injury.

Conclusion

In conclusion, TAOD is an uncommon and challenging subject in the pediatric population. Surgical stabilization is life-saving in cases of TAOD, while missing an unstable injury could have catastrophic consequences. The diagnosis criterion is not unique, generally requiring multimodal image, especially in less evident dislocations. Fixation of the occipitocervical junction with screws have a higher fusion rate and should be considered as the treatment of choice when feasible. Finally, hydrocephalus is not an unusual complication in children, and attention is necessary, with close clinical and radiological follow-up.

Conflict of Interests

The authors have no conflict of interests to declare.

References

- Adams VI. Neck injuries: I. Occipitoatlantal dislocation-a pathologic study of twelve traffic fatalities. J Forensic Sci 1992;37(02): 556-564
- 2 Ahuja A, Glasauer FE, Alker GJ Jr, Klein DM. Radiology in survivors of traumatic atlanto-occipital dislocation. Surg Neurol 1994;41 (02):112–118
- 3 Beez T, Brown J. Traumatic atlanto-occipital dislocation in children-a case-based update on clinical characteristics, management and outcome. Childs Nerv Syst 2017;33(01):27–33
- 4 Alker GJ Jr, Oh YS, Leslie EV. High cervical spine and craniocervical junction injuries in fatal traffic accidents: a radiological study. Orthop Clin North Am 1978;9(04):1003–1010
- 5 Menezes AH, Traynelis VC. Anatomy and biomechanics of normal craniovertebral junction (a) and biomechanics of stabilization (b). Childs Nerv Syst 2008;24(10):1091–1100
- 6 Pang D, Sun PP. Pediatric vertebral column and spinal cord injuries. In: Youmans JR (ed): Neurological Surgery, 5th ed. Philadelphia: WB Saunders; 2004:3315–3357
- 7 Pang D, Wilberger JE Jr. Traumatic atlanto-occipital dislocation with survival: case report and review. Neurosurgery 1980;7(05): 503-508
- 8 Hale AT, Say I, Shah S, Dewan MC, Anderson RCE, Tomycz LD. Traumatic Occipitocervical Distraction Injuries in Children: A Systematic Review. Pediatr Neurosurg 2019;54(02):75–84
- 9 Hwang SW, Gressot LV, Chern JJ, Relyea K, Jea A. Complications of occipital screw placement for occipitocervical fusion in children. J Neurosurg Pediatr 2012;9(06):586–593
- 10 Wholey MH, Bruwer AJ, Baker HL Jr. The lateral roentgenogram of the neck; with comments on the atlanto-odontoid-basion relationship. Radiology 1958;71(03):350–356
- 11 Powers B, Miller MD, Kramer RS, Martinez S, Gehweiler JA Jr. Traumatic anterior atlanto-occipital dislocation. Neurosurgery 1979;4(01):12–17

- 12 Harris JH Jr, Carson GC, Wagner LK. Radiologic diagnosis of traumatic occipitovertebral dissociation: 1. Normal occipitovertebral relationships on lateral radiographs of supine subjects. AJR Am J Roentgenol 1994;162(04):881–886
- 13 Sun PP, Poffenbarger GJ, Durham S, Zimmerman RA. Spectrum of occipitoatlantoaxial injury in young children. J Neurosurg 2000; 93(1, Suppl)28–39
- 14 Traynelis VC, Marano GD, Dunker RO, Kaufman HH. Traumatic atlanto-occipital dislocation. Case report. J Neurosurg 1986;65 (06):863–870
- 15 Pang D, Nemzek WR, Zovickian J. Atlanto-occipital dislocation: part 1-normal occipital condyle-C1 interval in 89 children. Neurosurgery 2007;61(03):514–521, discussion 521
- 16 Pang D, Nemzek WR, Zovickian J. Atlanto-occipital dislocationpart 2: The clinical use of (occipital) condyle-C1 interval, comparison with other diagnostic methods, and the manifestation, management, and outcome of atlanto-occipital dislocation in children. Neurosurgery 2007;61(05):995–1015, discussion 1015
- 17 Shamoun JM, Riddick L, Powell RW. Atlanto-occipital subluxation/dislocation: a "survivable" injury in children. Am Surg 1999;65(04):317–320
- 18 Astur N, Klimo P Jr, Sawyer JR, Kelly DM, Muhlbauer MS, Warner WC Jr. Traumatic atlanto-occipital dislocation in children: evaluation, treatment, and outcomes. J Bone Joint Surg Am 2013;95 (24):e194, 1–8)
- Labbe JL, Leclair O, Duparc B. Traumatic atlanto-occipital dislocation with survival in children. J Pediatr Orthop B 2001;10(04): 319–327
- 20 Tubbs RS, Hallock JD, Radcliff V, et al. Ligaments of the craniocervical junction. J Neurosurg Spine 2011;14(06):697–709
- 21 Klimo P Jr, Ware ML, Gupta N, Brockmeyer D. Cervical spine trauma in the pediatric patient. Neurosurg Clin N Am 2007;18 (04):599–620
- 22 Driscoll DR. Anatomical and biomechanical characteristics of upper cervical ligamentous structures: a review. J Manipulative Physiol Ther 1987;10(03):107–110
- 23 Werne S. Studies in spontaneous atlas dislocation. Acta Orthop Scand Suppl 1957;23:1–150
- 24 Harris MB, Duval MJ, Davis JA Jr, Bernini PM. Anatomical and roentgenographic features of atlantooccipital instability. J Spinal Disord 1993;6(01):5–10
- 25 Thiebaut F, Wackenheim A, Vrousos C. [Injuries and malformations of the cervico-occipital joint. Study of a basilar line]. Atlas Radiol Clin Presse Med 1961;141:1–4
- 26 Rojas CA, Bertozzi JC, Martinez CR, Whitlow J. Reassessment of the craniocervical junction: normal values on CT. AJNR Am J Neuroradiol 2007;28(09):1819–1823
- 27 Lee C, Woodring JH, Goldstein SJ, Daniel TL, Young AB, Tibbs PA. Evaluation of traumatic atlantooccipital dislocations. AJNR Am J Neuroradiol 1987;8(01):19–26
- 28 Bertozzi JC, Rojas CA, Martinez CR. Evaluation of the pediatric craniocervical junction on MDCT. AJR Am J Roentgenol 2009;192 (01):26–31
- 29 Bekelis K, Duhaime AC, Missios S, Belden C, Simmons N. Placement of occipital condyle screws for occipitocervical fixation in a pediatric patient with occipitocervical instability after decompression for Chiari malformation. J Neurosurg Pediatr 2010;6(02): 171–176
- 30 Chamoun RB, Relyea KM, Johnson KK, et al. Use of axial and subaxial translaminar screw fixation in the management of upper cervical spinal instability in a series of 7 children. Neurosurgery 2009;64(04):734–739, discussion 739
- 31 Haque A, Price AV, Sklar FH, Swift DM, Weprin BE, Sacco DJ. Screw fixation of the upper cervical spine in the pediatric population. Clinical article. J Neurosurg Pediatr 2009;3(06):529–533
- 32 Leonard JR, Wright NM. Pediatric atlantoaxial fixation with bilateral, crossing C-2 translaminar screws. Technical note. J Neurosurg 2006;104(1, Suppl)59–63

- 33 Coyne TJ, Fehlings MG, Wallace MC, Bernstein M, Tator CH. C1-C2 posterior cervical fusion: long-term evaluation of results and efficacy. Neurosurgery 1995;37(04):688–692, discussion 692–693
- 34 Smith MD, Phillips WA, Hensinger RN. Complications of fusion to the upper cervical spine. Spine 1991;16(07):702–705
- 35 Hwang SW, Gressot LV, Rangel-Castilla L, et al. Outcomes of instrumented fusion in the pediatric cervical spine. J Neurosurg Spine 2012;17(05):397–409
- 36 Martinez-Del-Campo E, Turner JD, Rangel-Castilla L, Soriano-Baron H, Kalb S, Theodore N. Pediatric occipitocervical fixation:

radiographic criteria, surgical technique, and clinical outcomes based on experience of a single surgeon. J Neurosurg Pediatr 2016;18(04):452-462

- 37 Vera M, Navarro R, Esteban E, Costa JM. Association of atlantooccipital dislocation and retroclival haematoma in a child. Childs Nerv Syst 2007;23(08):913–916
- 38 Naso WB, Cure J, Cuddy BG. Retropharyngeal pseudomeningocele after atlanto-occipital dislocation: report of two cases. Neurosurgery 1997;40(06):1288–1290, discussion 1290– 1291