A Rare Case of Spontaneous Intrauterine Skull Fracture

Caso Raro de Fratura Espontânea Intrauterina do Crânio

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ABSTRACT
Skull fractures are rare in newborns and normally caused by maternal abdominal trauma or complicated deliveries. However, in rare cases, these fractures are found in neonates born after an uneventful pregnancy and delivery. We report a case of a primigravida who underwent cesarean delivery due to failure of descent and malpositioning of the fetal head. After birth, a right temporoparietal fracture and congenital muscular torticollis were diagnosed. The newborn’s blood tests showed hypocalcemia and relative hypoparathyroidism. Both mother and newborn presented low vitamin D levels. Serial imaging control showed gradual resolution of the lesions, with the newborn being discharged at the 10th day of life with vitamin D supplementation. This is an interesting case because the combination of three conditions - maternal and fetal hypovitaminosis D, congenital torticollis and malposition of the cephalic pole during labor – may have synergistically contributed to a spontaneous intrauterine skull fracture.

Keywords: Hypocalcemia; Infant, Newborn; Skull Fractures; Torticollis/congenital; Vitamin D Deficiency

INTRODUCTION
Skull fractures found in newborns can be linear, usually affecting the parietal bones, or depressed, the so called 'ping-pong fractures'. Linear fractures, unlike depressed fractures, are not generally accompanied by underlying brain lesions and do not require specific treatment.1

Cerebral computed tomography (CT) scan, with pediatric low-dose radiation protocol, is the gold standard for the diagnosis.2

Most common causes are maternal abdominal trauma during pregnancy and head injuries caused by instruments or obstetrical maneuvers in complicated deliveries.3 However, in rare cases, these fractures are found in neonates born after an uneventful pregnancy and delivery and are, in these instances, called ‘spontaneous’.4

It is likely that these 'spontaneous' fractures may result from continuous intrauterine mechanical forces upon the soft fetal skull caused by bony structures.5

Additionally, some vitamin deficits may result in a higher probability of fracture. Vitamin D plays a central role in calcium and phosphate homeostasis and bone metabolism. In the presence of vitamin D deficiency, the intestine is not capable of absorbing calcium and phosphate adequately, leading to secondary hyperparathyroidism.6

Severe deficits of vitamin D during pregnancy have been associated with compromise of skeletal fetal homeostasis, congenital rickets and fractures in the newborn, as well as neonatal hypocalcemic seizures, since the newborn levels of vitamin D are largely dependent on maternal vitamin D status.6 One of the signs of maternal vitamin D deficiency is craniolabes, a softening of skull bones, which makes the bone more prone to fractures.7 Moreover, maternal vitamin D insufficiency correlates with higher risk of pre-eclampsia, gestational diabetes, preterm birth or low birth weight.6

CASE REPORT
We report a case of a 31-year-old primigravida with no significant medical history. She took a multivitamins including 5 µg of vitamin D during pregnancy. The pregnancy was uneventful until 33 weeks, when she presented with threatened preterm labor and did a course of antenatal corticosteroids for fetal maturation. No trauma was reported during pregnancy.

She presented to our emergency department at 38 weeks and four days after premature rupture of membranes

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in labor. At admission, cervix was 2 cm dilated and 100% effaced with cephalic presentation. Cardiotocography was reassuring.

A cesarean was performed due to failure of descent and fetal malpresentation (no engagement and head hyperextension). No forceps or vacuum extractor were used.

Newborn Apgar score was 9/10 at one and five minutes, respectively. Weight was appropriate for gestational age. Physical examination showed right hemicranium tumefaction of soft consistency, with crackling in the right temporoparietal transition and left congenital muscular torticollis (CMT). Neurologic examination was adequate to the gestational age. Skull x-ray suggested a fracture in the right temporoparietal region (Fig. 1).

CT scan confirmed the diagnosis showing a solution of continuity in the right parietal convexity, with misalignment of bony tops, compatible with a linear fracture and also three focus of haemorrhage: a right temporoparietal haemorrhagic contusion measuring 9.3 x 6.7 x 4.3 mm, a smaller one laterally and linear traces of adjacent sulcal subarachnoid haemorrhage (Fig. 2).

Both mother and newborn presented hypocalcemia (8.6 mg/dL and 6.9 mg/dL) and vitamin D insufficiency (16 ng/mL and 11 ng/mL), respectively. Parathormone (PTH) levels in the newborn were within normal range, although inappropriately low (45.8 pg/mL) in the presence of hypocalcemia. The newborn also presented with hyperphosphatemia (10.4 mg/dL) and hypomagnesemia (1.45 mg/dL) (Table 1).

Serial brain CT scans (D3 and D8) were performed, with evidence of partial reabsorption of intra-axial haemorrhagic lesion and resolution of subarachnoid haemorrhage. The skeleton x-ray showed no evidence of additional fractures. The neurological examination was always normal.

The newborn was discharged at D10 with vitamin D and calcium supplementation and maintained multidisciplinary follow-up, with normal physical and psychomotor development. No sequels were identified.

DISCUSSION

The prevalence of spontaneous intrauterine skull fractures varies from 1 in 4000 to 1 in 10 000 deliveries. Establishing etiology can be challenging, especially when there is no evidence of abdominal trauma, no use of instruments or after a cesarean section delivery.

There are few similar cases in the literature, with the majority of studies describing depressed skull fractures. To our knowledge, this is the first case report of a linear fracture in a non-traumatic delivery of a newborn with docu-
Vitamin D deficiency screening is based on the measurement of serum concentrations of 25-hydroxivitamin D \([25(OH)D]\). According to the 2016 Global Consensus recommendations, levels comprised between 20-100 ng/mL are considered normal, with insufficiency in the range 12 - 20 ng/mL and deficiency when values are less than 12 ng/mL. There is no consensus on an optimal vitamin D level during pregnancy status and it remains unclear whether vitamin D supplementation should be routinely recommended in pregnancy or if it should be reserved for high-risk groups.

Vitamin D deficiency in neonates secondary to maternal deficiency may lead to hypocalcemia. PTH should increase in response to low serum calcium concentrations, but that was not found in our case, with PTH levels abnormally low for the level of hypocalcemia. This suggests a relative hypoparathyroidism, which may also have contributed to newborn hypocalcemia.

Birth trauma has been suggested as the main etiological factor for congenital muscular torticollis (CMT). However, CMT has also been described in cases where no trauma was reported. In our case, congenital muscular torticollis may have been caused by an abnormal intrauterine position during the third trimester which resulted in sternocleidomastoid muscle injury and deformity, as proposed by Stellwagen et al.

The fetal malpresentation, probably due to CMT, may have predisposed the fetus skull to collision with maternal body structures, especially during uterine contractions. The vitamin D deficit and hypocalcemia may have contributed to softening of the fetus skull, predisposing it to fracture.

This is a challenging case in which the combination of three conditions - maternal and fetal hypovitaminosis D, congenital torticollis and fetal malpresentation – may have synergistically contributed to a spontaneous intrauterine skull fracture.

**PROTECTION OF HUMANS AND ANIMALS**

The authors declare that the procedures were followed according to the regulations established by the Clinical Research and Ethics Committee and to the Helsinki Declaration of the World Medical Association.

**DATA CONFIDENTIALITY**

The authors declare having followed the protocols in use at their working center regarding patients’ data publication.

**PATIENT CONSENT**

Obtained.

**CONFLICTS OF INTEREST**

The authors declare that there are no conflicts of interest.

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### Table 1 – Blood laboratory findings of the newborn and his mother.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Newborn</th>
<th>Mother</th>
</tr>
</thead>
<tbody>
<tr>
<td>Calcium (mg/dL)</td>
<td>6.9 (7.6 - 10.4)</td>
<td>8.6 (8.9 - 10.0)</td>
</tr>
<tr>
<td>Vitamin D (pg/mL)</td>
<td>11 (16 - 65)</td>
<td>16 (30 - 100)</td>
</tr>
<tr>
<td>PTH (pg/mL)</td>
<td>45.8 (15 - 65)</td>
<td>---</td>
</tr>
<tr>
<td>Phosphorus (mg/dL)</td>
<td>10.4 (4.5 - 9.0)</td>
<td>---</td>
</tr>
<tr>
<td>Magnesium (mg/dL)</td>
<td>1.45 (1.6 - 2.3)</td>
<td>----</td>
</tr>
</tbody>
</table>

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**REFERENCES**