Isolated congenital bilateral occipital perforations

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CASE REPORT

A seven-month-old male Asian baby presented with headache and a large right parietal scalp hematoma after falling from a bed onto a wooden floor. The infant was previously well, born via non-traumatic vaginal delivery following uneventful pregnancy. Normal developmental and growth milestones were met. On examination, Glasgow Coma Scale was 15 with no neurological deficits. In accordance with local guidelines, the infant was investigated via low-dose computed tomography (CT) scan of the head. This showed a right-sided linear non-depressed skull fracture with normal intracranial appearances, seen in a volume-rendered 3D-CT scan (Figure 1). This was managed conservatively via overnight admission to a pediatric unit for observation, and discharged home at 24 hours with post-head injury advice for his parents. Isolated bilateral symmetrical occipital perforations were noted as an incidental finding. The foramina could be palpated gently without discomfort, and were not visibly pulsatile. The volume rendered 3D-CT scan shows full thickness bilateral 6 mm circular foramina with smooth contours in the occipital bone (closed arrows), each 25 mm from the midline (Figure 1). At final follow-up, the infant was well, all scalp hematoma had resolved, however, perforations could still be palpated. No further investigations or treatment were performed.

DISCUSSION

Cranial foramina are holes within the skull, formed during development, permitting passage of blood vessels and nerves. Animal studies suggest they are formed...
during cranial development via a process independent of apoptosis, reliant upon local nerve/blood vessel-derived restriction of intramembranous ossification [1]. Historically, midline or slightly mesial symmetrical apertures of the occipital bone related to occipital emissary veins are described [2]. This is inconsistent with our findings. Occipital emissary veins connect the transverse sinus to the occipital vein, draining into the vertebral venous plexus. Larger studies of occipital emissary foramina have not shown bilateral symmetrical foramina, with single foramen consistently closer to the foramen magnum than the occipital protuberance in all specimens, not the case here [3]. Unlike occipital foramina, parietal foramina are well described, resulting from delayed or deficient intramembranous ossification of the parietal bone. Parietal perforations occur as an isolated autosomal dominant trait or as part of a syndrome, and this normally is of no clinical significance [4, 5]. Comparative osteoarchaeological studies describe occipital perforations occurring commonly in cattle but rarely in other species. These perforations are variously attributed as congenital, secondary to tumors, parasites, trauma or infection [6].

CONCLUSION

In view of radiological and clinical findings, we believe these isolated symmetrical occipital perforations are a rare congenital abnormality of the occipital bone, possibly resulting from delayed or deficient intramembranous ossification of the occipital bone. With no apparent precedent, we believe they are of no clinical significance and will resolve as the child grows. This could interest trainees as an anatomical variant they may see whilst reviewing pediatric imaging. To the best of our knowledge, this is the first described case of bilateral symmetrical occipital perforations, with this distribution in the cranium.

Keywords: Congenital, Occipital, Perforations

REFERENCE
