Case Report

Increasing Head Circumference from Hydrocephalus, Not Only in Young Children: Case Report of a 7-Year-Old

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Increasing head circumference is a common presentation of hydrocephalus within the first 2 years of life but is extremely rare in older children or adults. Here, we present a rare case of rapidly increasing head circumference, as the sole symptom, as a new diagnosis of hydrocephalus in a 7-year-old boy, who then re-presented with an increasing head circumference at the time of a VPS malfunction a year later.

Case: A 7-year-old was referred to neurosurgery for papilledema and increased headaches. The mother reports a rapidly increasing head circumference necessitating changes in hat size at least twice in the last few months. The CT gave the diagnosis of a Dandy-Walker spectrum malformation and hydrocephalus. A VPS was therefore inserted, with a stabilisation of his head circumference for a few months. He presented again with a rapidly increasing head circumference over the course of a few months, from 57 cm up to 59.5 cm, in the setting of progressive headaches consistent with increased ICP. The patient subsequently underwent a shunt valve replacement to improve CSF diversion.

Methods: The patient was followed over 18 months from diagnosis to shunt revision, with MRI of the head. Volumetric measurements of the ventricles and head circumference are compared over the same time period.

Conclusion: We present a unique case of increasing head circumference in a 7-year-old boy with Dandy Walker spectrum malformation, who then had another rapid increase almost two years later with a shunt dysfunction. To our knowledge, no similar cases of that age were reported in the literature.
Within 2 months, his head circumference increased once again up to 59 cm, although his MRI’s demonstrated relatively stable appearance of the ventricles (Figure 1). As his head circumference continue to increase to 59.5 cm, and headaches increased, the patient underwent a VP shunt revision with a revision of his valve to a programmable valve (programmable valve set at 80 mm H$_2$O) in December of 2018. There was no shunt obstruction noted intraoperatively. The patient’s headaches soon resolved following his second operation. His head circumference decreased then stabilized at 59 cm (Figure 2), which was still the case 22 months after the shunt revision (October 2020). Clinically, his headaches were much better, and the papilledema was resolved.

**Figure 1**: A) Ventrilocemegaly at diagnosis; B) Post shunt insertion, pre-revision MR demonstrating ventriculomegaly (Dec 2018); C) Post shunt revision MR demonstrating smaller ventricles (July 2019).

**Figure 2**: Timeline of increase in head circumference and change in Evan’s ratio pre and post operatively.

**Discussion**

The development of the human cranium is a complex chronological process. To compensate for rapid brain growth, cranial sutures stay open in early childhood. The rate of suture fusion/closure varies according to the anatomical location. The metopic suture tends to fuse first between the ages of 3 months to 2 years, while the lambdoid, sagittal, and coronal suture may take up until the age of 40 before complete fusion [1]. However, head circumference growth typically slows down after 24 months of age but continues at a slow rate until adolescence [2].

A rapidly increasing head circumference is one of the first signs of hydrocephalus in infancy. This marker is used as an indication for CSF diversion, either with ventricular tapping through the anterior fontanelle, or with a definitive solution such as VP shunt insertion [3]. However, head circumference is rarely affected after the age of 3, and patients usually present with increasing headaches, nausea, vomiting, visual changes, and fluctuating level of consciousness with VP shunt malfunction. Rapidly increasing head circumference is almost never the first presenting symptom in preteens or young adolescents. Only one other case was found in the literature with a similar presentation. Ganainingham et al. reported suture diastasis in a 17-month-old boy, with no significant change in ventricular volume in a case of VP shunt malfunction [4]. From our center’s experience, one might even expect some rapid increased head circumference until 36 or 40 months, but we had never experienced it on a 7-year-old.

After CSF diversion, ventricular compliance can often decrease, and in certain cases the ventricles may fail to dilate despite raised intracranial pressure (ICP). This could be due to sub-ependymal gliosis or Laplace’s law, which states that the pressure needed to expand a small volume is greater than that required to expand a larger volume [5, 6]. This may explain our finding of our volumetric analysis (Figures 2 & 3), which demonstrated relatively stable ventricular volume/Evan’s ratio, despite the patient’s symptoms of increased ICP. However, another explanation is that when the skull can increase in volume, one can have a substantial volumetric increase by adding a few millimeters in the subarachnoid spaces instead of the ventricles especially in communicating hydrocephalus. With no other changes a head circumference increase from 57.5 to 59 cm would lead to an increase of more than 360cc of CSF in the subarachnoid spaces.

**Figure 3**: Volumetric analysis of three separate MRI studies both pre and post shunt revision.

**Conclusion**

We present a unique case of increasing head circumference in a 7-year-old boy with hydrocephalus and Dandy-Walker spectrum malformation. While sutures are typically open until adulthood, rapid head circumference increase due to hydrocephalus is typically not seen after the toddler’s years. To our knowledge, only one similar case was reported in the literature but in a much younger patient.

**Conflicts of Interest**

None.

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