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Rare regression of congenital brainstem high-grade glioma: case report and literature review

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Child's Nervous System

Aims and scope

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Abstract

Congenital infantile brainstem high-grade gliomas (HGGs) are extremely rare. Given the limited literature characterizing this disease, management of these tumors remains challenging. Brainstem HGGs are generally associated with extremely poor prognosis. Limited reports of spontaneous regression of radiologically diagnosed infantile brainstem tumors exist in published literature. In this case report, we document the first histologically proven congenital brainstem HGG with molecular characteristics that did not fall under any previously well-defined pediatric brain tumor classifications. The tumor underwent regression after biopsy, documented on neuroimaging up to 2 years of age. A review of the literature was also performed to identify previously reported infantile brainstem HGGs and the management for such tumors. Our case highlights the value of performing histopathological confirmation to guide management and the possible existence of a subcategory of a congenital brainstem HGG with better prognosis.



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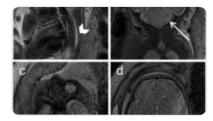
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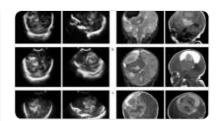
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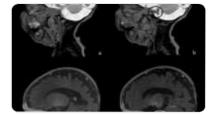
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Data availability

No datasets were generated or analysed during the current study.

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Contributions

All authors contributed to the study conception. Material preparation was prepared by Shi Hui Ong, Mervyn Jun Rui Lim, Miriam Santiago Kimpo, Balamurugan A Vellayappan, Ai Peng Tan and Vincent Diong Weng Nga. The first draft of the manuscript was written by Shi Hui Ong and all authors reviewed and revised the manuscript accordingly. All authors read and approved the final manuscript.

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Ethics declarations

Conflict of interest

The authors declare no competing interests.

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