



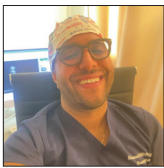
Case Report

Pediatric extra-axial glioblastoma with bone invasion leading to a subcutaneous mass: A case report

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ABSTRACT

Background: Pediatric glioblastoma multiforme (p-GBM) is an exceptionally rare and aggressive brain tumor, with even fewer reported cases with radiographic and intraoperative characteristics that mimic those of extra-axial lesions, often posing a diagnostic challenge. Despite advancements in imaging technologies, the diagnosis of GBM can still be intricate, relying primarily on histopathological confirmation.

Case Description: We present a unique case of a 15-year-old female who presented to our hospital with a new-onset focal-to-bilateral tonic-clonic seizure described as clonic movements of her left hemispheres; on clinical examination, a subcutaneous mass was evident in the right parietal region. Magnetic resonance imaging of the brain revealed a sizable extra-axial enhancing mass measuring 9 cm, located in the right parieto-occipital region with notable bone invasion. Moreover, the intraoperative findings revealed an extra-axial mass attached to the dura. Total *en bloc* resection was achieved. The histopathological analysis confirmed the diagnosis of glioblastoma multiforme. Subsequently, the patient underwent adjuvant radiotherapy in conjunction with temozolomide chemotherapy. Postoperatively, she exhibited clinical improvement and remained stable throughout the 6-month follow-up period.

Conclusion: We present the first case of extra-axial p-GBM in a young patient, which remarkably led to the destruction of the bone and finally resulted in a sizable parietal subcutaneous lesion in the absence of prior surgery or radiation.

Keywords: Bone invasion, Extra-axial, Pediatric glioblastoma, Subcutaneous mass

INTRODUCTION

Glioblastoma multiforme (GBM) is the most prevalent and aggressive primary malignant brain tumor in adults, accounting for approximately 45% of all primary brain tumors. However, it remains a rare entity in the pediatric population, constituting only 3–15% of primary central nervous system (CNS) tumors in children.^[8]

Typically, GBM is intra-axially located in the deep white matter of the supratentorial region, primarily within the frontal and temporal lobes.^[17] Extra-axial presentations are even rarer, with

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only a limited number of cases documented in the existing medical literature.^[1]

To the best of our knowledge, cases involving the destruction of the dura and calvaria caused by an extra-axial pediatric GBM (p-GBM) without any prior history of surgery or radiation have not been reported previously. This occurrence may raise considerations of extra-axial tumors, such as aggressive meningiomas, and highlights the exceptional nature of this case.

CASE DESCRIPTION

A 15-year-old female high school student presented with a 1-month history of persistent holocranial headaches and intermittent vomiting. She experienced a new-onset focal-to-bilateral tonic-clonic seizure characterized by clonic movements of her left hemicorps that secondarily progressed to a convulsion. On clinical examination, all findings were normal, except for the presence of a palpable subcutaneous mass in the right parietal region.

Initial laboratory tests yielded unremarkable results.

Brain computed tomography (CT) scan and magnetic resonance imaging (MRI) revealed a large, heterogeneous contrast-enhancing mass located in the right parieto-occipital region. This mass extended extra-axially, reaching up to the occipital horn medially and involving the cortical bone of the inner table laterally, displaying signs of bone invasion [Figure 1].

Based on these findings, a preoperative diagnosis of aggressive meningioma was initially considered.

Subsequently, a Mitre's flap and a right parieto-occipital craniotomy were performed. Intraoperatively, it was observed that the tumor had invaded and destroyed the surrounding bone [Figure 2]. The tumor exhibited vascularity with well-defined margins adherent to the inner layer of the dura. A total *en-bloc* resection was successfully achieved with duraplasty and cranioplasty.

Postoperative brain MRI confirmed complete tumor resection [Figure 3].

The histological analysis identified the tumor as a glioblastoma (World Health Organization – grade IV), characterized by small cells with marked angiogenesis and wide necrotic areas infiltrating both cerebral structures and meningeal sheaths [Figure 4]. Immunohistochemical analysis revealed a Ki-67 labeling index of 40–45% for all cells within the resected tumor area. The tumor displayed positive staining for ATRX, glial fibrillary acidic protein, and synaptophysin, while epithelial membrane antigen did not exhibit significant positivity. Tumor cells were negative for isocitrate dehydrogenase (IDH)1 and IDH2 genes. In addition, the tumor showed negative results for estimated glomerular filtration rate, c-Myc, and N-Myc amplification, with no MGMT promoter methylation detected.

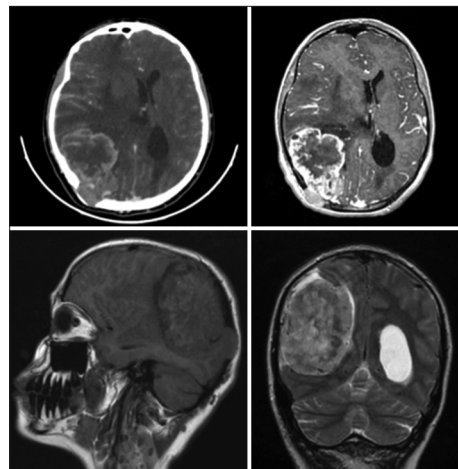


Figure 1: Preoperative brain computed tomography scan and brain magnetic resonance imaging show a large, heterogeneous contrast-enhancing mass located in the right parieto-occipital region. This mass extended extra-axially, reaching up to the occipital horn medially and involving the cortical bone of the inner table laterally.

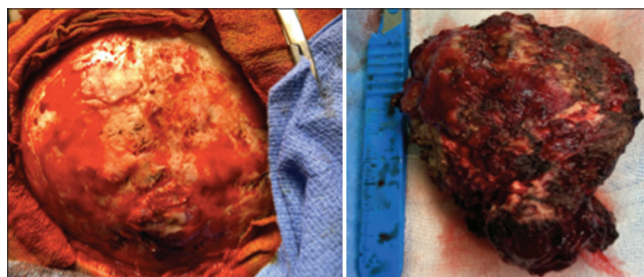


Figure 2: The bone invasion and the *en bloc* resected tumor.

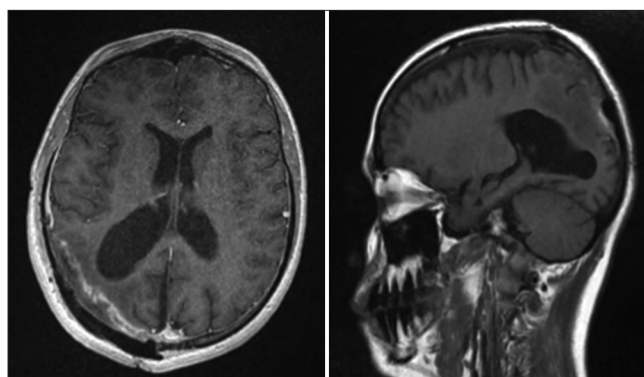


Figure 3: Postoperative brain magnetic resonance imaging showing complete total resection.

The postoperative period proceeded without complications, and the patient's recovery progressed smoothly. She was discharged on day 10 with no neurological deficits and experienced symptomatic improvement.

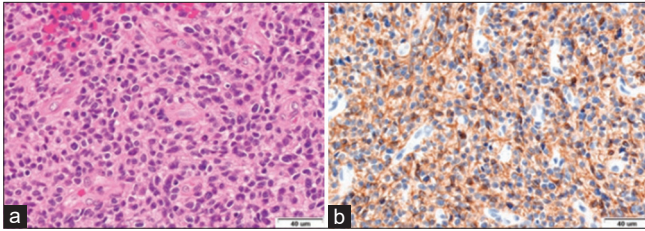


Figure 4: (a) Relative cellular uniformity is apparent, as well as frequent mitotic figures. Note how many cells display haloes (b) glial fibrillary acidic protein staining is strongly positive within tumor cells.

Following the surgical procedure, the patient underwent radiotherapy and received concurrent temozolomide treatment, followed by adjuvant temozolomide therapy. In the most recent examination conducted six months post-surgery, the patient was confirmed to be progressing favorably. She remained free from seizures and had successfully returned to school.

DISCUSSION

Glioblastoma multiforme (GBM) is the most prevalent and aggressive primary brain tumor in the adult population, accounting for approximately 45% of all primary brain tumors.^[6] In contrast, GBM is a rare occurrence in the pediatric population, constituting only 3–15% of primary CNS tumors.^[9]

Pediatric glioblastomas (p-GBMs) are typically diagnosed during the second decade of life, with rare cases even reported during fetal development. The peak incidence of p-GBMs is observed between the ages of 15 and 19, likely reflecting the cumulative impact of various genetic factors contributing to tumorigenesis.^[7]

From a pathophysiological standpoint, GBMs originate from astrocytes, the primary glial component of the CNS. Both macroscopic and microscopic characteristics of pediatric GBMs (p-GBMs) closely resemble those found in adults.^[7] Despite often appearing well-defined on imaging or during surgery, p-GBMs exhibit diffuse infiltrative properties.

GBMs are typically located in supratentorial cerebral lobes, such as the frontal and temporal lobes. Extracranial occurrences are rare; however, secondary dissemination within the brain or leptomeninges occurs in nearly 17% of patients.^[10]

Neuroimaging plays a pivotal role in diagnosing, managing, and prognosticating GBMs. CT and MRI are fundamental tools for radiological assessment of these tumors.

However, MRI findings associated with p-GBM lack specificity. Typically, these tumors appear as heterogeneous masses with ill-defined borders, exerting a mass effect on neighboring structures and displaying varying degrees of enhancement (complex, variable, or occasionally absent). Regarding signal characteristics, p-GBM may exhibit

iso- to hypointense T1 signals relative to gray matter and heterogeneously hyperintense T2 signals, often accompanied by surrounding edema, visible on fluid attenuation inversion recovery images. In cases with hemorrhage, distinct signal characteristics may emerge, including T1 hyperintensity, T2 hypointensity, low signal on T2*, and susceptibility-weighted imaging features.^[4] In addition, magnetic resonance spectroscopy often reveals a prominent choline peak with reduced N-acetyl aspartate in the tumor region, providing valuable metabolic information.^[3]

On the other hand, primary extra-axial involvement of glioblastoma (GBM) is exceedingly uncommon, with only ten/nine documented cases, including the current one. Instances of GBM with bone invasion are even scarcer. Sakata *et al.* reported a singular case of GBM in an elderly patient, where the disease naturally progressed to invade and destroy the skull bone.^[14] Zahir *et al.* reported a similar case, all without prior surgical intervention or radiation therapy.^[17]

To the best of our knowledge, this is the first reported case of pediatric GBM with bone invasion.

The bone erosion is related to a chronic mass effect.^[2] The mass exerted by the tumor displaces the cerebrospinal fluid (CSF), which normally acts as a cushion, dispersing brain pulsations across a broad area. When the CSF space is compressed, the brain can directly transmit these pulsations to the inner table of the skull. Over time, this localized increase in pressure may lead to the erosion of the cortical bone of the inner table.^[1] The infrequency of bone invasion may be explained by the relatively short median survival duration of GBM patients. Long-term survivors may face an increased risk of bone invasions, primarily due to the extended duration of their disease course.^[6]

To date, bone invasion in primary GBM has predominantly been observed in patients who had previously undergone surgical procedures, biopsies, or radiation therapy. These iatrogenic interventions facilitated a path for the tumor to extend beyond the confines of the brain into extracerebral structures.^[6]

Based on the updated WHO classification of brain tumors, GBM is now classified as per the IDH gene mutation status.

GBM can, therefore, be IDH wild type or IDH mutation positive. The latter group comprises *de novo* lesions that primarily affect elderly patients, whereas the former group represents secondary GBMs. However, research on pediatric high-grade gliomas, such as GBMs, has shown that IDH mutations are extremely rare, especially in younger children. In other words, p-GBM is virtually always IDH wild type, but, as Pollack *et al.* have shown, teenagers and younger adults may have a higher frequency of secondary GBM (IDH mutant).^[13]

The current gold standard for the treatment of adult GBMs involves maximal surgical resection followed by a combination of chemoradiotherapy and/or radiotherapy.

No well-defined standard of care has been established for pediatric glioblastomas, particularly concerning surgical management.^[12,14]

Based on the largest single-center experience with pediatric glioblastoma (p-GBM), achieving maximal tumor removal is a strong predictor in improving both progression-free survival and overall survival in p-GBM.^[11]

However, the selection of adjuvant treatment remains a subject of debate in pediatric glioblastomas due to concerns about the potential adverse effects of radiation therapy on the developing brain, as well as the variability in outcomes associated with different chemotherapy regimens.^[7]

Radiation therapy has emerged as the standard of treatment, especially for children aged three and older who are diagnosed with a new GBM. Nevertheless, younger children, due to their increased susceptibility to the adverse effects of radiation therapy, typically receive chemotherapy as a standalone treatment, often in combination with radiation-sparing approaches.^[12]

Recent advancements in oncology treatment have introduced novel therapeutic approaches, including molecular-targeted therapy and immunotherapy, which have become valuable additions to the arsenal for managing cancer patients.^[7] Immunotherapy encompasses a range of agents, including chimeric antigen receptor T-cell therapy, immune checkpoint inhibitors, virotherapy, cancer vaccines, and dendritic cell therapy.

In the context of pediatric brain tumors, despite numerous clinical trials, molecular-targeted therapy has, with a few exceptions, not yet demonstrated a substantial impact on the survival or quality of life for children with these tumors.^[11] However, the future holds significant promise, as ongoing research leads to the development and inclusion of more effective agents in clinical trials, offering renewed hope for improved outcomes in the treatment of pediatric brain tumors.

Irrespective of the treatment approach chosen, “pediatric glioblastoma” continues to present as a devastating disease, with median survival durations spanning from 13 to 73 months and a 5-year survival rate of <20%.^[5,8,16] Nevertheless, several studies have highlighted a relatively more favorable prognosis and the potential for long-term survival among pediatric patients when compared to adults.^[15,16]

CONCLUSION

We report a unique case of pediatric glioblastoma (GBM) located in an extradural location, presenting with an atypical

manifestation that led to the invasion and subsequent destruction of the skull bone, all without any prior surgical or radiation intervention.

Ethical approval

The Institutional Review Board approval is not required.

Declaration of patient consent

Patient’s consent not required as patient’s identity is not disclosed or compromised.

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Conflicts of interest

There are no conflicts of interest.

Use of artificial intelligence (AI)-assisted technology for manuscript preparation

The authors confirm that there was no use of artificial intelligence (AI)-assisted technology for assisting in the writing or editing of the manuscript and no images were manipulated using AI.

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