

Case Report

Isolated Supratentorial Intraventricular Recurrence of Medulloblastoma

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Medulloblastoma is a common pediatric tumor typically diagnosed before the age of fifteen. Initial therapy includes surgical resection and radiation of the entire neuro-axis. Recurrence is common and typically occurs within 2 years of initial diagnosis. Those fitting Collin's Law is considered tumor-free. We report a case of single supratentorial recurrence 13 years after initial diagnosis. Here we present a 22 year old male presenting 13 years after initial diagnosis with isolated septum pellucidum recurrence. He underwent complete resection of the tumor. Medulloblastoma is a common in the pediatric population. Late recurrence to the ventricular system is uncommon. Long term follow-up is recommended in these patients.

Key Words : Medulloblastoma · PNET · Recurrence · Hydrocephalus, Drop metastasis.

INTRODUCTION

Medulloblastoma (MB) is the most common malignant tumor in children, accounting for 15 to 25% of all childhood brain tumors¹⁹⁻²¹. Occurrence in the adult population is also well documented, but only accounts for 1% of adult tumors¹¹. Treatment includes surgical resection followed by radiotherapy of the entire neuro-axis and chemotherapy^{4,5,19,23}.

Recurrence of this tumor is well recognized and may require salvage therapy. Time to recurrence typically occurs within two years of initial diagnosis in the pediatric population. Tumors that follow Collin's Law (tumor free period of 9 months plus the age at diagnosis) are considered to be cured^{1,19,20}. However, late recurrence of MB has been documented in the pediatric population^{2,3,12,13,23}.

The locations of recurrences most commonly present as posterior fossa, spinal, supratentorial, or boney metastases²³. Supratentorial recurrence is reported to be more common in the sub-frontal region^{19,23}, while late recurrence of MB in the supratentorial intraventricular compartment is rare¹⁷.

CASE REPORT

History

A 22-year-old male with a history of a medulloblastoma at 9 years of age, removed surgically and treated with adjuvant cra-

niospinal irradiation and chemotherapy presented with 2.5 weeks of progressive gait ataxia and headache.

Examination

Patient was alert and oriented. The right pupil was sluggish. The left pupil was reactive and the left eye deviated uncontrollably on exam. The left upper extremity had 4/5 strength. CT scan showed an anterior septum pellucidum mass at the foramen of Monro with secondary obstructive hydrocephalus (Fig. 1).

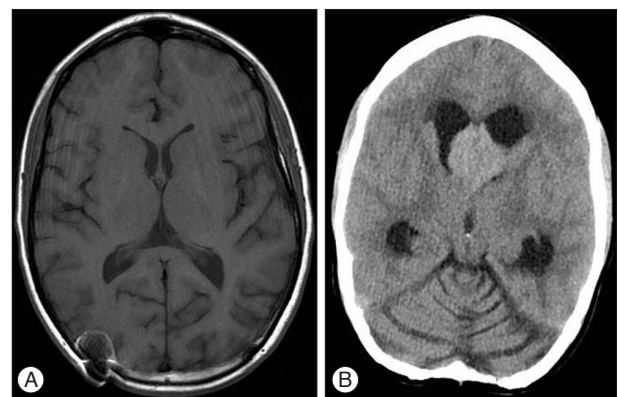


Fig. 1. CT brain without contrast (B) showing a septal lesion causing obstruction of the foramen Monro resulting in obstructive hydrocephalus. (A) MRI of the brain without contrast on 7/14/2009 last scan prior to recent presentation.

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Post-operative course

The patient underwent bilateral ventriculostomy and external ventricular drain placement with subsequent tumor resection (Fig. 2, 3). Pathology report was consistent with anaplastic/large-cell medulloblastoma (Fig. 4).

DISCUSSION

Medulloblastoma, a malignant tumor typically arising from the cerebellar vermis in young children and lobes in older children, represents 4–8% of all intracranial tumors, and is the most common malignant central nervous system tumor of childhood¹⁹⁻²¹, with approximately 80% occurring in patients under 15 years of age²¹. It accounts for 15–25% of all childhood brain tumors in comparison with only 1% of adult intracranial neoplasms, with a slight male predominance in both groups^{19-21,23,24}.

Presenting features of medulloblastoma are generally related to hydrocephalus (found radiographically in 97% of patients) and cerebellar dysfunction which include : headache, nausea/vomiting, truncal ataxia, and unsteady gait²³. Approximately 83% are found in the midline in children, while 49% of adult tumors are lateral²¹.

Differences in childhood and adult tumors can also be observed

in terms of histologic variant. Classical histologic presentation of medulloblastoma includes densely-packed primitive cells with hyperchromatic nuclei, scant cytoplasm, and nuclear molding^{1,20}. This classical tumor histology is more common in children^{8,21}. Homer-Wright rosettes may be seen¹⁰. Adults are more likely to have laterally-located tumors, which more often desmoplastic^{5,10,21}. Both tumor types include markers of neuronal lineage, such as class III beta tubulin and MAP-2 neurofilaments,

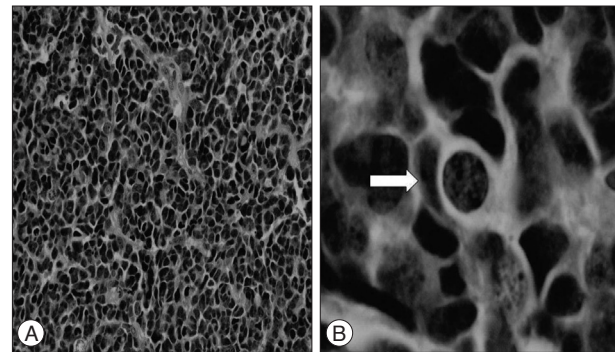


Fig. 4. A : Diffuse infiltrate of small to large pleomorphic cells with little cytoplasm (synaptophysin positive). B : High power view. Cells have hyperchromatic angular nuclei. Cell wrapping, a feature commonly found in anaplastic/large cell MB (white arrow).

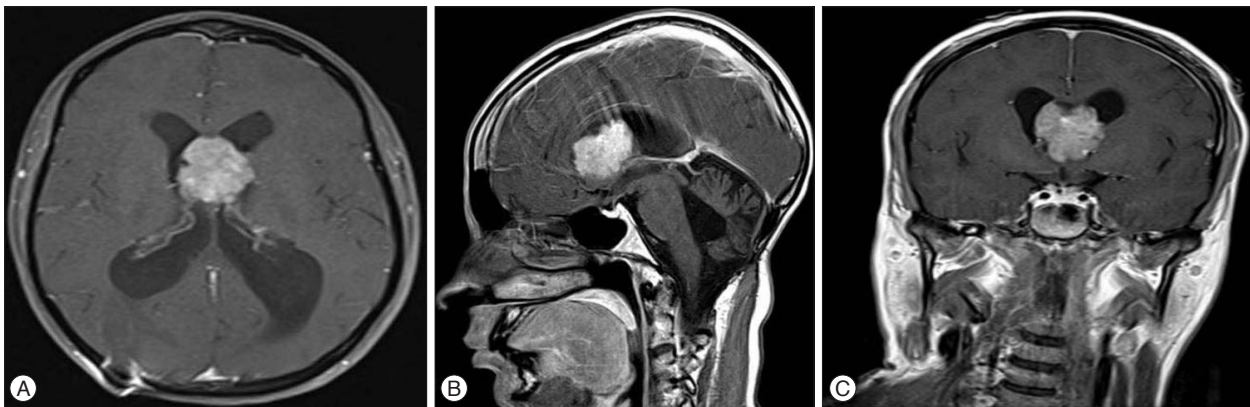


Fig. 2. MRI with contrast axial (A), sagittal (B), and coronal (C) showing a homogeneously enhancing septal mass obstructing the foramen Monro.

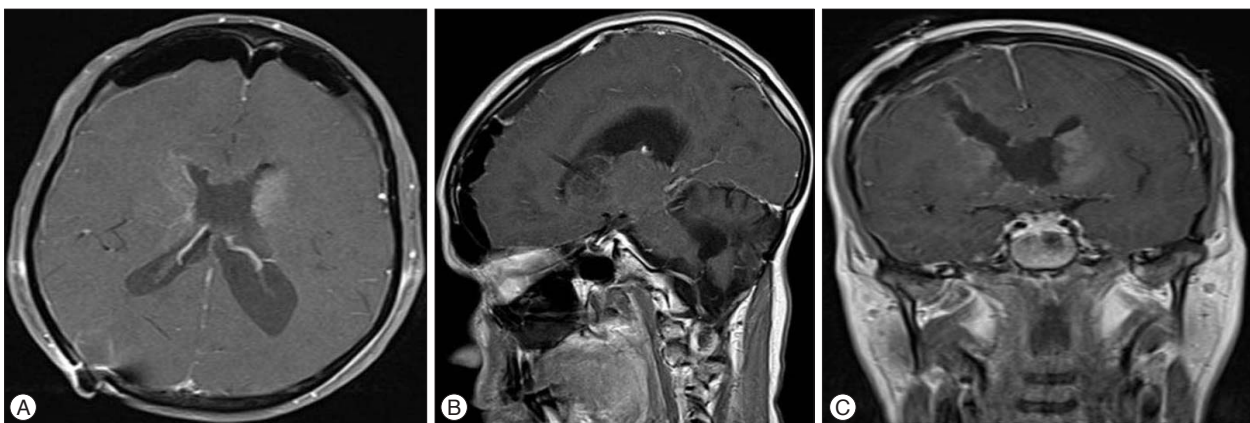


Fig. 3. MRI with contrast axial (A), sagittal (B), and coronal (C) showing right transcortical intraventricular approach for tumor resection. Gross tumor resection was accomplished.

whereas more adult tumors will express GFAP^{10,20}. The anaplastic/large cell variant is more frequently seen in older children and adults.

MB tends to seed the cerebrospinal fluid resulting in extensive leptomeningeal involvement, and treatment therefore requires surgical resection followed by neuro-axis radiotherapy and chemotherapy^{7,17,19,23}. Recurrence in the pediatric population differs from that seen in adults. Age at diagnosis has been shown to be a significant predictor of time until relapse²⁴. The majorities of pediatric recurrences are within two years of initial diagnosis, and are in the posterior fossa^{19,20,24}. Those with tumor-free period equal to the age at diagnosis plus nine months may be considered cured (Collins' Law)^{1,4,18-20}. While a good predictor of many childhood tumors, there are known exceptions to Collins law for MB^{12,6,9,13,18,22}. However, a recent review of 125 patients by Massimino et al.^{14,15}, gender, age at diagnosis, metastases, and therapeutic protocol were shown to have no prognostic impact.

In the event of recurrence, relapse at a single site and further out from initial time of diagnosis is considered a more favorable prognostic indicator¹⁵. Recurrences are most common in the posterior fossa, followed by spinal, supratentorial and bony metastases²³. Supratentorial dissemination is often found in the subfrontal area and may be due to overly generous radiation protection of the orbital roof to prevent irradiation of the cribriform plate, allowing for a nidus of recurrence¹⁶. Late recurrence of MB in the supratentorial ventricular compartment is uncommon. We present a case of late recurrent MB 13 years after initial diagnosis, again validating the need for long term follow-up in the pediatric population following diagnosis of MB.

CONCLUSION

Recurrence of MB is common and often occurs in the posterior fossa. Supratentorial dissemination of MB is more common in the subfrontal region. Supratentorial recurrence of MB not following Collins' Law is extremely uncommon, especially within the supratentorial ventricular system. We present a case of a 22 years old male who presented 13 years following his initial diagnosis of MB with symptoms of obstructive hydrocephalus from a septum pellucidum recurrence. Continued close follow-up is need in pediatric patients after resection and radiation for MB given their unexpected recurrence pattern.

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