Glioblastoma multiforme in conus medullaris with intracranial metastasis after postoperative adjuvant therapy

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Glioblastoma multiforme in conus medullaris with intracranial metastasis after postoperative adjuvant therapy

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Abstract
Spinal glioblastoma multiforme is not common among spinal cord tumors. According to our literature review, only 27 cases originating from the conus medullaris were reported. We herein reported a case of a 10-year-old child diagnosed with glioblastoma multiforme. The patient received adjuvant radiotherapy and standard temozolomide chemotherapy after total excision. Intracranial lesions were found 1 month after postoperative adjuvant therapy. We described the clinical characteristics and postoperative therapy of the patient, and reviewed all of the published cases of conus medullaris glioblastoma. Location, age, leptomeningeal spread, and secondary hydrocephalus may be predictive factors. Immunohistochemical factors such as p53 and Ki-67 are also important. Combined treatment of surgery and postoperative adjuvant therapy is commonly used, but is controversial.

Abbreviations: CgA = chromograin A, CNS = central nervous system, CSF = cerebrospinal fluid, EGFR = epidermal growth factor receptor, GBM = glioblastoma multiforme, GFAP = glial fibrillary acidic protein, MRI = magnetic resonance imaging, NF = neurofilament, NSE = neuron specific enolase, PNET = primitive neuroectodermal tumor, TMZ = temozolomide, WHO = World Health Organization.

Keywords: conus medullaris, immunohistochemistry, predictive factor, spinal glioblastoma multiforme, treatment

1. Introduction
In adults, spinal cord tumors mostly originate from extramedullary tumors (almost 80%).[1] whereas in children, the rate of primary spinal tumor is up to 35%. [2] Among these neoplastic spinal lesions in children, high-grade glioma is relatively rare (roughly 1%-3%). Spinal glioblastoma multiforme (GBM), defined as World Health Organization (WHO) IV in astrocytoma, is a highly malignant central nervous system (CNS) tumor that is clinically, histologically, and genetically heterogeneous.[3] A survey in 1989 revealed that spinal GBM accounted for 0.2% of all GBM and 1.4% of spinal glioma. [4] The total number of cases reported in the literature was less than 200.[1][5] According to the data in a single institution, spinal cord GBM accounted for only 1% of patients with intramedullary neoplasms.[5] Spinal GBM accounted for 7.5% of intramedullary glioma, and only 1.5% of all spinal tumors.[6] As we present a rare case of GBM located in conus, an effort was made to search for cases of conus GBM published ever (Table 1), to detect some similarities and to learn more about metastasis, pathology, and treatment.

2. Case presentation
A 10-year-old Chinese boy, without any past medical history, reported a 3-week history of weakness and a subsequent appearance of paresthesia of both lower limbs. He also complained of lower back pain with radiation to both legs, urinary disturbance, and weight loss. His examination of strength demonstrated a 3/5 at right leg and 4/5 at left leg. Sensory examination was significant for decreased sensation in both legs, whereas right leg was worse. His reflex was weakened at left patella and ankle, and was absent at right. Bilateral hyper-reflexia were presented of the lower limbs. There were no neurologic abnormalities of the upper limbs and cranial nerves.

Magnetic resonance imaging (MRI) revealed an ill-defined intramedullary mass filled the spinal canal between T11 and L1 (Fig. 1A and B) with inhomogeneous enhancement of the tumor area (Fig. 1C and D). Based on these findings, glioma was considered.
<table>
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<th>No.</th>
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<th>Radiotherapy and proposal</th>
<th>Chemotherapy</th>
<th>Metastasis</th>
<th>Survival time, mos</th>
<th>Other features</th>
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<td>1</td>
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<td>Biopsy</td>
<td>No</td>
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<td>Subarachnoid space</td>
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<td>Yes 7500r</td>
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<td>3</td>
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<td>12/F</td>
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<td>Andrews[10]</td>
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<td>Septal region</td>
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<td>No</td>
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<td>TR</td>
<td>Yes</td>
<td>No</td>
<td>—</td>
<td>17</td>
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<tr>
<td>8</td>
<td>Kawariish[12]</td>
<td>50/M</td>
<td>PR</td>
<td>Yes 40 Gy</td>
<td>No</td>
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<td>Shirato[13]</td>
<td>35/F</td>
<td>PR</td>
<td>Yes 65 Gy</td>
<td>Yes AGNMCR</td>
<td>Cerebellar hemisphere</td>
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<td>31/F</td>
<td>STR</td>
<td>Yes 45 Gy/68–L1 +9 Gy/tumor</td>
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<td>Suprasellar region</td>
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<td>Santi[20]</td>
<td>23/M</td>
<td>STR/PR</td>
<td>Yes 20–36 Gy</td>
<td>No</td>
<td>—</td>
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<td>58/M</td>
<td>Biopsy</td>
<td>Yes</td>
<td>No</td>
<td>—</td>
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<td>41/M</td>
<td>STR</td>
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<td>No</td>
<td>—</td>
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<td>14/F</td>
<td>STR</td>
<td>No</td>
<td>Yes BONJ</td>
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<td>Yes 36 Gy</td>
<td>Yes MTX</td>
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<td>Methkour[15]</td>
<td>20/M</td>
<td>STR × 2</td>
<td>Yes 65 Gy</td>
<td>Yes Unknown</td>
<td>Pontomedullary junction, Cerebellum, suprasellar cistern, Left lateral ventricle, Posterior fossa</td>
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<td>18</td>
<td>Stecco[16]</td>
<td>14/M</td>
<td>STR</td>
<td>No</td>
<td>No</td>
<td>Simultaneous cervical-medullary lesion</td>
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<td>20/M</td>
<td>STR × 2</td>
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<td>No</td>
<td>—</td>
<td>10</td>
<td></td>
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<tr>
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<td>Bonde[17]</td>
<td>16/M</td>
<td>STR</td>
<td>Yes 20–36 Gy</td>
<td>Yes</td>
<td>—</td>
<td>10</td>
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<td>Chol[18]</td>
<td>46/M</td>
<td>STR</td>
<td>Yes 48 Gy/79–L3</td>
<td>Yes</td>
<td>—</td>
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<td>22</td>
<td>Sun[22]</td>
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<td>STR</td>
<td>No</td>
<td>Yes TMZ</td>
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<td>36/F</td>
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<td>Yes 30 Gy/79–L3</td>
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<td>Gee[26]</td>
<td>9/F</td>
<td>STR</td>
<td>Yes 4500cGy/CM</td>
<td>Yes TMZ</td>
<td>T8–T8 vertebral bodies</td>
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<td>4/M</td>
<td>STR</td>
<td>Yes 4500cGy/CM</td>
<td>Yes TMZ</td>
<td>—</td>
<td>52+</td>
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<td>28</td>
<td>The present case</td>
<td>10/M</td>
<td>TR</td>
<td>Yes</td>
<td>Yes TMZ</td>
<td>Left apical lobe, right cerebellar vermis, corpus callosum, basal ganglia and lateral cerebral ventricle</td>
<td>14</td>
<td>CSF protein increased</td>
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</tbody>
</table>

ACNU = Nimustine, BONJ = Carmustine, CCNU = Lomustine, CM = Conus medullaris, CSF = Cerebrospinal fluid, CTX = Cyclophosphamide, DDP = Cisplatin, MTX = Methotrexate, PCB = Procarbazine, PR = partial resection, STR = Subtotal resection, TMZ = temozolomide, TR = total resection, VCR = Vincristine.
Open total excision was conducted under general anesthesia. The procedure presented a hypertonic dura. After opening the dura, the tumor was located in the conus medullaris, highly vascularized, and ill-defined.

Histopathologic examination, including immunohistochemical staining, was performed, and the diagnosis of GBM was made according to the WHO criteria (Fig. 2A and B). However, part of the tumor showed the structure of primitive neuroectodermal tumor (PNET). Immunohistochemistry showed positivity for glial fibrillary acidic protein (GFAP) (Fig. 2C), S100 (Fig. 2D), p53 (Fig. 2E), Ki-67 (proliferation index 60%; Fig. 2F), epidermal growth factor receptor (EGFR) (Fig. 2G), neuron-specific enolase (NSE) (Fig. 2H), CD34 (Fig. 2J), CD56 (Fig. 2J), and β-catenin (Fig. 2K), and negativity for chromogranin A (CgA), AE1/AE3, CD99, and neurofilament (NF). The mutation or amplification of EGFR is commonly observed in malignant gliomas, and these modifications are associated with increased cell proliferation and radiation resistance. NSE is a specific protein of neuron; high serum levels of NSE were noticed in the patients with malignant gliomas, and these modifications are associated with increased intracranial metastasis. New lesions were located in left apical lobe, right cerebellar vermis, corpus callosum, basal ganglia, and lateral cerebral ventricle (Fig. 3). We deemed them as the metastatic lesions from the tumor in conus medullaris. The patient and his parents refused further invasive treatments; thus the exact pathology of the intracranial lesions could not be confirmed. The patient died 14 months after his surgery.

3. Discussion

Primary spinal GBM located in conus medullaris is relatively rare; usually this kind of disease has a favor for the thoracic, cervical, or the conjunction area. In a literature review in 2011, the author found that conus medullaris is also a location where spinal GBM may generate, although it is not that usual. A study retrospect 128 cases from 1938 to 2015 performed that 42.2% of spinal GBM were located in thoracic spine and 29.7% were in cervical spine. Meanwhile, tumors located in the conus level only accounted for 14%, according to this study. A study retrospection 128 cases from 1938 to 2015 performed that 42.2% of spinal GBM were located in thoracic spine and 29.7% were in cervical spine. Meanwhile, tumors located in the conus level only accounted for 14%, according to this study. A study retrospect 128 cases from 1938 to 2015 performed that 42.2% of spinal GBM were located in thoracic spine and 29.7% were in cervical spine. Meanwhile, tumors located in the conus level only accounted for 14%, according to this study. Generally, the survival time of spinal GBM was limited to 12 to 24 months. Moreover, GBM of the conus medullaris ranged from 4 to 16 months. The mean follow-up time (most are equal to
survival time) in our study was 18.26 months, among which the
longest one was 67 months.\textsuperscript{[24]} Konar et al\textsuperscript{[23]} found that location
of the tumor had no influence to overall survival, and patients
with conus and thoracic tumors were less likely to die at
6 months. As to the influence of age, people have same opinions
that adults (over 18 years) were more likely to have longer
survival periods.\textsuperscript{[23,25]} But Wolff et al\textsuperscript{[26]} found that in children,
age less than 5 years may be a relatively positive prognostic
factor.

Among these predictive factors, Konar et al thought
leptomeningeal spread is the poor prognostic factor for survival.
The rate of CSF dissemination reaches up to 58\% in spinal GBM,
in comparison with 23\% to 27\% in cerebral GBM.\textsuperscript{[27]} In Konar
et al’s study, 60\% of the patients had CSF dissemination and
76\% had brain metastasis.\textsuperscript{[23]} We found that in those 28 cases of
conus GBM, 16 cases mentioned tumor metastasis, and among
these, 14 patients had brain metastasis. It is easily recognized that
brain metastasis occurred in most spinal GBM. Patchy nodular
lesions were discovered intracranially, whereas the specific
locations differed. According to our unpublished study, wall of
the ventricles could become a relatively welcome “home” of new
lesions, considering the anatomy. Strik et al\textsuperscript{[28]} noted that p53
may be a predictor of subsequent brain metastasis in spinal GBM.
But we lack more evidence about p53 accumulation for the
reason that few cases conducted immunohistochemical study. Of
note, regarding the pathology of the primary lesion in our case,
part of it showed the structure of PNET. In fact, primary PNET of the spine is unusual, with very few cases reported in the literature. Occurrence of primary spinal PNET in an intramedullary location is further uncommon. Because the patient and his parents refused further invasive treatments after the discovery of intracranial lesions, their exact pathology could not be confirmed.

Another important factor of overall survival period is secondary hydrocephalus. Six of the 28 patients in our study developed hydrocephalus. Higher concentration of protein in the CSF was thought to be related with hydrocephalus. Our patient also demonstrated to contain more protein in CSF, but he did not develop hydrocephalus. It is possible that the tumor had not blocked the subarachnoid completely. Most of the cases we found did not perform a lumbar puncture or did not display the results, but we can speculate that higher protein may be a sign of CSF dissemination.

For the reason that spinal GBM was not common in population, studies describing the histomorphological and molecular genetic alterations are not as many as cerebral GBM. Govindan et al[29] had the opinion that histopathological characteristics of spinal GBM are comparable with cerebral glioblastoma. In their 6-case study, GFAP was 100% positive, whereas p53 immunoreactivity was 83.3% (5 of 6).[29] Another case series study concluded that GFAP and p53 immunoreactivity was seen in all cases.[30] Studies also showed that p53 expression is seen in majority of glioblastoma. The pathology of our case was definitely positive for GFAP and p53, matching with results of those research. Eleven patients had immunohistochemical results in the list, and 8 of them expressed GFAP, but only 4 mentioned reactivity of p53. Even though the expression of p53 was not as expected, we are of the opinion that it may be a result of lack of data. Proliferative marker Ki-67 index/MIB-1 labeling should be a complement of pathologic features of GBM. It has been reported that Ki-67 index ranged from 12% to 34%.[30] The Ki-67 index ranged from 10% to 30% in our case series, with the values being comparable with those of other studies.[31] However, the boy in our case showed a very high Ki-67 index of 60%. It may be a hint why the patient had brain metastasis after radiochemotherapy, even though the prognostic value of Ki-67 was debatable.[32] Treatment of spinal GBM nowadays usually combines surgery with radiotherapy; most of the time chemotherapy is also considered. As for radiotherapy, although researchers did not find significant linkage between radiation and prognosis,[33] more people believe radiation can increase survival time in malignant spinal tumor.[34] But the optimal dosage is uncertain, although it is almost always used. It is reported that radiotherapy can prolong survival in some cases.[34] Shirato et al[35] recommended that radiation can be given in 2.5-Gy fractions 4 times weekly to total doses of 40 to 50 Gy over 4 to 5 weeks. Our patient received a total dose of 45.9 Gy for conus medullaris and an additional

![Figure 3. Follow-up magnetic resonance imaging (MRI) after 10 months from operation demonstrates intracranial metastasis. (A, B) T1-weighted and T2-weighted axial MR images show new lesions located in lateral cerebral ventricle and enlargement of ventricle. (C) T1-weighted sagittal MR image with contrast reveals heterogeneous enhancement. (D, E) Coronal and sagittal sections display several metastases. (F) Apparent diffusion coefficient (ADC) mapping shows heterogeneous signal.](image-url)
radiation to the whole brain. However, the tumor still metastasized to intracranial spaces. If patient tolerates the treatment well, Minehan et al. found that higher doses (59.4 Gy) could perform better in symptom improvement. Shirato et al. also reported a long survival case for 38 months with a total radiation dose of 65 Gy, but in consideration of adverse effects on growing and fertility, especially in teenager, we have to control the total dose of radiation. Whether chemotherapy is required or not remains controversial, but a retrospective series of 8 cases proved that both TMZ and bevaxizumab were useful in improving survival. As TMZ was recognized to be effective in intracranial GBM, it is generally used as adjuvant therapy to surgery and radiation in spinal GBM. For patients with spinal GBM, it is recommended that TMZ be used concomitantly during and after radiotherapy, but at different dosage. In a study consisting of 6 patients, TMZ was found to prolong survival time of primary spinal GBM, and at different dosage. In a study consisting of 6 patients, TMZ was found to prolong survival time of primary spinal GBM, but a retrospective series of 8 cases proved that both TMZ and bevacizumab were useful in improving survival.

In the development of molecular therapy, tumors can be investigated more and more thoroughly. In a study by Sharma et al., it was shown that children may have different molecular signatures from adults. So we can have the confidence that target therapies may become a powerful measure in treating spinal GBM.

4. Conclusions

Spinal GBM located in conus medullaris is rare. Several factors may be related to intracranial metastasis and prognosis. Immunohistochemistry currently plays a crucial role in diagnosis of CNS tumors. Adjuvant treatment composed of radiotherapy and chemotherapy are still under exploration.

Acknowledgments

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References


