Symptomatic cerebral vasospasm following posterior fossa hemangioblastoma resection: illustrative case

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BACKGROUND Symptomatic cerebral vasospasm following posterior fossa intra-axial tumor resection is a rare phenomenon with only seven cases previously reported in the literature. The condition appears distinct to vasospasm following supratentorial tumor resection and extra-axial tumor resection of the posterior fossa. It shares, however, similarities with vasospasm following aneurysmal subarachnoid hemorrhage.

OBSERVATIONS The authors describe their experience with a 23-year-old female who developed delayed symptomatic vasospasm following resection of a left parapontine cerebellar hemangioblastoma. Tumor resection was complicated by rupture of a fragile arterialized vein, resulting in significant hemorrhage. The patient developed several episodes of focal and variably reversible neurological deficit. These clinical signs corresponded with angiographically confirmed vasospasm, which responded to standard therapies for vasospasm post aneurysmal subarachnoid hemorrhage.

LESSONS This case and literature review highlight that symptomatic vasospasm is a rare, potentially highly morbid complication of posterior fossa intra-axial tumor resection. This phenomenon may be related to significant intraoperative or postoperative hemorrhage. Postoperative radiological findings such as high risk modified Fisher scale hemorrhage could alert clinicians to this condition.

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KEYWORDS cerebral vasospasm; tumor; posterior cranial fossa; neurosurgery

Although a well-known and highly morbid complication of aneurysmal subarachnoid hemorrhage (aSAH), cerebral vasospasm has been reported as a rare complication of intracranial tumor resection, with 40 cases identified in a 2013 systematic review.1 Since this time a greater appreciation for this phenomenon has occurred, and further cases have been reported. Our literature review sourcing English articles from PubMed until December 3, 2021 has identified 19 cases2–17 of symptomatic vasospasm following posterior fossa tumor resection, as well as two case series of asymptomatic vasospasm documenting a total of 70 cases.16,18 Overall, a minority of these cases occur following resection of intra-axial posterior fossa tumors, with only seven cases reported.2–6 The characteristics of vasospasm observed in cases of intra-axial posterior fossa tumor resection differs from those seen in tumor resection at other sites. All cases were symptomatic, and vasospasm occurred bilaterally and involved vascular territories remote to the surgical resection. This remote and diffuse distribution of vasospasm is less commonly observed in vasospasm associated with extra-axial and supratentorial tumor resections.1 Moreover, factors such as hypothalamic dysfunction20 and intraoperative vessel manipulation,21,22 which are associated with vasospasm following supratentorial and extra-axial tumor resection, were not described in the intra-axial posterior fossa cases. The unique manifestations and rarity of vasospasm following intra-axial posterior fossa tumor resection lends to delays in recognition, which may be associated with worse outcomes.12 We describe our experience with an additional case, in which a patient
developed delayed and aggressive cerebral vasospasm following resection of a posterior fossa hemangioblastoma.

Illustrative Case

An independent 23-year-old female with unremarkable medical history presented with a 12-month history of progressive headaches. Magnetic resonance imaging (MRI) scan identified a large left parapontine cerebellar lesion with a contrast enhancing mural nodule and a smaller medially located cyst consistent with a hemangioblastoma (Fig. 1A). There were no other lesions identified on imaging her whole neuraxis. The tumor was approached via a left retrosigmoid craniotomy. The tumor was highly vascular with multiple draining arterialized veins; all peripheral arterial feeders were systematically obliterated, and the tumor was resected en bloc. Following tumor removal, a large fragile arterialized superior petrosal vein ruptured, producing significant arterial hemorrhage, with hemostasis ultimately achieved with packing and hemostatic agents. There were no further intraoperative complications.

An immediate postoperative noncontrast computed tomography (CT) scan was performed that demonstrated acute blood within her basal cistems, ventricular system, and a midline hematoma adjacent to the superior cerebellar vermis (Fig. 1B). Upon extubation she was oriented and obeying commands but demonstrated a left upper limb monoparesis and partial left sixth nerve palsy.

On postoperative day 6 the patient had a decline in conscious state. Computed tomography angiogram (CTA) demonstrated vasospasm of the left middle cerebral artery (MCA) territory (Fig. 2A). The patient was readmitted to the intensive care unit (ICU) for hypertensive therapy with return of consciousness.

Over the coming days the patient experienced fluctuating neurological function, with several episodes of acute neurological deterioration attributed to angiographically confirmed vasospasm in bilateral anterior and posterior circulation territories (Fig. 2B and C). Nimodipine delivered via nasogastric tube, direct injection of intra-arterial verapamil with invasive angiography, and microcathether angioplasty were used with varying success to manage the episodes of vasospasm. Noradrenaline and saline infusion were used to maintain a mean arterial pressure of $>100$ mm Hg in the ICU. The patient ultimately demonstrated gradual clinical improvement and was transferred to a general ward on postoperative day 23. The patient was discharged to a rehabilitation facility on day 38 with modified Rankin scale (mRS) 4 graded disability. Outpatient follow-up at 3 months identified a stable level of disability.

Discussion

Cerebral vasospasm following tumor resection is a rare phenomenon. Nineteen cases of symptomatic cerebral artery vasospasm have been detected from tumor resection in the posterior fossa (Tables 1 and 2). Seven of these cases follow the resection of intra-axial tumors. There are several proposed mechanisms for this phenomenon. High arterial blood load in the subarachnoid space, hypothalamic dysfunction, and vessel injury or manipulation are suggested in the literature. Vascular encasement and manipulation are believed to be more relevant to extra-axial tumor resection, and hypothalamic insult is considered to have a greater role in vasospasm following suprasellar and middle cranial fossa resection. Vasospasm following posterior fossa intra-axial tumor resection appears to be related to high-volume arterial subarachnoid hemorrhage, drawing similarities with vasospasm following aSAH. In aSAH, the release of vasoactive arterial blood products into the cerebrospinal fluid is thought to irritate both local and distant vasculature. Complex interactions then promote vasospasm that largely manifests 3 to 14 days following aSAH. The presence of

![FIG. 1. A: Preoperative magnetic resonance axial section demonstrating tumor enhancement adjacent to a well-circumscribed void region. B: Immediate postoperative computed tomography axial section demonstrating mFS 4 hemorrhage.](image)

![FIG. 2. A: Day 6 computed tomography angiogram with focal area of proximal left middle cerebral artery stenosis (red arrow). B: Day 9 digital subtraction angiography showing minor vasospasm of the right carotid (red arrow). C: Day 7 angiographic reconstruction demonstrating mild bilateral intracranial vertebral and basilar artery vasospasm (red arrow).](image)
vasospasm in vascular territories remote to the site of tumor resection, the time delay from surgery, and the high perioperative blood load observed in some cases following intra-axial posterior fossa tumor resection echoes vasospasm post-aSAH. Recognizing this association may allow management strategies already utilized for aSAH to be applied in select patients post-tumor resection.

**Observations**

Vasospasm was found to be bilateral in all seven previously reported cases of vasospasm post-intra-axial tumor resection, and in the case we encountered. A comparatively smaller proportion of vasospasm post-extra-axial tumor resection occurred bilaterally, with five of 12 cases demonstrating this. To the best of our knowledge, neither direct vessel manipulation nor hypothalamic injury occurred in these eight cases of vasospasm post-intra-axial tumor resection. Rather, three of these total eight cases identified notable perioperative hemorrhage, each graded as equivalent to modified Fisher scale (mFS) 4 by two members of our team (Table 1).²,⁵ There is a 40% risk of clinical vasospasm in aSAH with mFS 4 hemorrhage,²⁷ and it is possible that the same degree of perioperative hemorrhage following posterior fossa tumor resection carries similar vasospasm risk. Of the three cases with mFS 4 equivalent hemorrhage, two had hemangioblastoma related intraoperative bleeding, and in the other case hemorrhage occurred on postoperative day 2. Hemorrhage preceded recognition of vasospasm between 6 and 12 days in these cases, mirroring the interval between aSAH and subsequent vasospasm frequently described.²⁶

In the remaining five cases there was limited imaging-based evidence of intraoperative hemorrhage. The authors in one case stated that their postoperative imaging was potentially too late to detect subarachnoid blood,² and another described 100 mL of intraoperative bleeding related to tumor resection.⁴ In both instances, the authors recognized hemorrhage as a potential etiology of vasospasm, and vasospasm was diagnosed at postoperative day 15 and 13, respectively. In the remaining three cases, the authors denied the presence of significant perioperative hemorrhage,²⁶,⁷ instead proposing a potential paraneoplastic etiology in one,⁶ and the release of tumor cyst products in another.⁶

In the 12 cases of symptomatic vasospasm following extra-axial posterior fossa tumor resection (Table 2), there were five cases in which significant perioperative hemorrhage was noted.¹⁰,¹³,¹⁵,¹⁷ In three of these cases vasospasm occurred contralateral to the site of the tumor resection,¹⁰,¹³,¹⁷ possibly supporting the mechanism of vasoactive product release producing distant vasospasm. Furthermore, the interval between vasospasm and hemorrhage ranged between three and 14 days, echoing aSAH-related vasospasm. Notably, vessel manipulation occurred in one of these cases, and may be responsible for the early vasospasm seen on day 3.¹⁷ In the remaining eight cases in which the presence of perioperative hemorrhage was not explicit, only two gave evidence for a lack of hemorrhage. One stated that there was minimal intraoperative bleeding,³ and the other provided a postoperative MRI demonstrating minimal hemorrhage.¹⁰ Despite no specific mention of perioperative hemorrhage occurring in three individual cases reported within one case series,¹² the authors noted that intraoperative hemorrhage was marked in many of their cases. These authors also observed that preoperative embolization was a statistically significant risk factor for vasospasm and hypothesized that this may result from a selection bias and predisposition to intraoperative hemorrhage in cases in which preoperative embolization is utilized.¹²

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<table>
<thead>
<tr>
<th>Authors &amp; Year</th>
<th>Tumor Type</th>
<th>Sex</th>
<th>Age</th>
<th>mFS</th>
<th>Postoperative Day at Vasospasm Diagnosis*</th>
<th>Relative Morbidity at Last FU†</th>
<th>Intraoperative or Early Postoperative Bleeding as Evidenced by Postoperative CT Imaging</th>
</tr>
</thead>
<tbody>
<tr>
<td>Brau et al., 1986²</td>
<td>Posterior fossa epidermoid tumor</td>
<td>F</td>
<td>57 yrs</td>
<td>15</td>
<td>None evident</td>
<td>Disability</td>
<td></td>
</tr>
<tr>
<td>Dalvi et al., 2014³</td>
<td>Grade IV medulloblastoma</td>
<td>M</td>
<td>17 yrs</td>
<td>16</td>
<td>Insufficient information to comment</td>
<td>Recovery</td>
<td></td>
</tr>
<tr>
<td>Gocmen et al., 2021⁴</td>
<td>Vermis &amp; fourth ventricle medulloblastoma</td>
<td>F</td>
<td>9 yrs</td>
<td>13</td>
<td>Insufficient information to comment</td>
<td>Recovery</td>
<td></td>
</tr>
<tr>
<td>Jacob et al., 2011⁵</td>
<td>Grade II ependymoma</td>
<td>M</td>
<td>23 yrs</td>
<td>12</td>
<td>mFS 4</td>
<td>Recovery</td>
<td></td>
</tr>
<tr>
<td>Kinoshita et al., 2020⁶</td>
<td>Hemangioblastoma</td>
<td>F</td>
<td>62 yrs</td>
<td>1</td>
<td>None evident</td>
<td>Disability</td>
<td></td>
</tr>
<tr>
<td>Lee et al., 1998⁷</td>
<td>Primitive neuroectodermal tumor (PNET) with epithelioid &amp; neuronal differentiation</td>
<td>F</td>
<td>15 mos</td>
<td>17</td>
<td>Insufficient information to comment</td>
<td>Death</td>
<td></td>
</tr>
<tr>
<td>Prontera et al., 2014⁸</td>
<td>Hemangioblastoma (VHL)</td>
<td>F</td>
<td>35 yrs</td>
<td>12</td>
<td>mFS 4</td>
<td>Disability</td>
<td></td>
</tr>
<tr>
<td>Present case</td>
<td>Hemangioblastoma</td>
<td>F</td>
<td>23 yrs</td>
<td>6</td>
<td>mFS 4</td>
<td>Disability</td>
<td></td>
</tr>
</tbody>
</table>

CT = computed tomography; FU = follow-up.
* Defined as day of angiographic confirmation of vasospasm.
† Compared to status preceding the first diagnosed episode of vasospasm.
TABLE 2. Symptomatic vasospasm characteristics following extra-axial posterior fossa tumor resection

<table>
<thead>
<tr>
<th>Authors &amp; Year</th>
<th>Total No. of Cases</th>
<th>Tumor Location &amp; Type</th>
<th>Age (yrs), Sex</th>
<th>Postoperative Day at Vasospasm Diagnosis</th>
<th>Location of Vasospasm as Identified by Angiogram (unless otherwise specified)</th>
<th>Vascular Manipulation or Tumor Encasement of Vessels Found to Undergo Vasospasm</th>
<th>Intraoperative or Early Postoperative Bleeding as Evidenced by Postoperative CT Imaging or Author Report</th>
<th>Relative Morbidity at Last FU*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Afshari et al., 2014⁹</td>
<td>1</td>
<td>Lt hypoglossal schwannoma</td>
<td>9, F</td>
<td>8</td>
<td>Bilateral ICA, MCA, &amp; ACA.</td>
<td>Authors denied manipulation.</td>
<td>Intraoperative bleeding reported to be minimal.</td>
<td>Recovery</td>
</tr>
<tr>
<td>Almeida et al., 1985¹⁰</td>
<td>1</td>
<td>Rt acoustic neuroma</td>
<td>26, M</td>
<td>14</td>
<td>Lt ICA, ACA, &amp; MCA.</td>
<td>Authors denied manipulation.</td>
<td>Significant intraoperative blood loss. No further distinction provided by authors.</td>
<td>Disability</td>
</tr>
<tr>
<td>Aw et al., 2010¹¹</td>
<td>1</td>
<td>Lt CPA epidermoid cyst</td>
<td>43, M</td>
<td>14</td>
<td>Rt MCA, ICA, &amp; VA.</td>
<td>No comment made by authors.</td>
<td>Insufficient information to comment.</td>
<td>Disability</td>
</tr>
<tr>
<td>Bejani et al., 1999¹²</td>
<td>3</td>
<td>Lt foramen magnum meningioma</td>
<td>65, F</td>
<td>15</td>
<td>Rt VA &amp; BA.</td>
<td>VBA encased by tumor.</td>
<td>Insufficient information to comment.</td>
<td>Disability</td>
</tr>
<tr>
<td>Clivus chordoma</td>
<td>37, M</td>
<td>ICA, MCA &amp; ACA (laterality not specified)</td>
<td>11</td>
<td>Tumor compressing ICA, w/ vessel narrowing documented by authors.</td>
<td>Insufficient information to comment.</td>
<td>Insufficient information to comment.</td>
<td>Insufficient information to comment.</td>
<td></td>
</tr>
<tr>
<td>Clivus chordoma</td>
<td>57, M</td>
<td>Rt MCA (M1 segment)</td>
<td>1</td>
<td>Tumor encased ICA &amp; VBA bilaterally. ICA narrowing bilaterally.</td>
<td>Insufficient information to comment.</td>
<td>Insufficient information to comment.</td>
<td>Insufficient information to comment.</td>
<td></td>
</tr>
<tr>
<td>Cervoni et al., 1996¹³</td>
<td>1</td>
<td>Lt anterior, middle &amp; posterior fossae meningioma</td>
<td>51, M</td>
<td>4</td>
<td>Bilateral MCA vasospasm identified using TCD.</td>
<td>No comment made by authors.</td>
<td>Copious intraoperative blood loss from cavernous sinus. No further distinction provided by authors.</td>
<td>Disability</td>
</tr>
<tr>
<td>LeRoux et al., 1991¹⁴</td>
<td>1</td>
<td>Lt CPA acoustic neuroma</td>
<td>69, M</td>
<td>7</td>
<td>Bilateral VA, BA, &amp; Rt ICA.</td>
<td>No comment made by authors.</td>
<td>Insufficient information to comment.</td>
<td>Disability</td>
</tr>
<tr>
<td>Pan et al., 2021¹⁵</td>
<td>2</td>
<td>Rt cavernous sinus, middle &amp; posterior cranial fossae chordrosarcoma</td>
<td>63, M</td>
<td>10</td>
<td>BA.</td>
<td>No comment made by authors.</td>
<td>Postoperative CT indicated subdural &amp; subarachnoid hemorrhage, attributed to bleeding vein.</td>
<td>Disability</td>
</tr>
<tr>
<td>Lei et al., 2014¹⁶</td>
<td>1</td>
<td>Rt CPA anaplastic medulloblastoma</td>
<td>10, F</td>
<td>6</td>
<td>Lt ACA &amp; MCA.</td>
<td>No comment made by authors.</td>
<td>Insufficient information to comment.</td>
<td>Recovery</td>
</tr>
<tr>
<td>Qi et al., 2015¹⁷</td>
<td>1</td>
<td>Lt ventral medulla oblongata schwannoma</td>
<td>16, M</td>
<td>3</td>
<td>Bilateral siphon segment of ICA &amp; BA.</td>
<td>Authors suggested direct mechanical trauma may have been responsible for vasospasm.</td>
<td>300 mL intraoperative blood loss, 50 mL attributed to tumor resection.</td>
<td>Disability</td>
</tr>
</tbody>
</table>

ACA = anterior cerebral artery; BA = basilar artery; CT = computed tomography; ICA = internal carotid artery; VA = vertebral artery.

* Compared to status preceding the first diagnosed episode of vasospasm.
Despite a potential role of hemorrhage in symptomatic vasospasm, this is not clearly evident in two case series of asymptomatic vasospasm post-posterior fossa tumor resection (Table 3). Together, the two series reported 70 cases, and identified younger patient age, large tumor size, vessel encasement, and adhesion to the brainstem or trigeminal nerve as risk factors for vasospasm. In one series (n = 27), the role of hemorrhage was found to have poor correlation with subsequent risk of vasospasm. In the other study (n = 43), cases with postoperative imaging demonstrating hemorrhage were excluded from reporting.

The experience from our case and the available literature suggests that high-volume arterial blood within the subarachnoid space may contribute to symptomatic vasospasm post-posterior fossa tumor resection. In potential support of this are the observed similarities with vasospasm that high-volume arterial blood within the subarachnoid space may contribute to symptomatic vasospasm post-posterior fossa tumor resection. Methods used in the management of vasospasm post-aSAH may also be used to recognize and manage this condition.

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**References**


Disclosures
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Author Contributions
Conception and design: Hiwase, Ovenden, Wells. Acquisition of data: Hiwase, Kalyanasundaram, Bak, Laden, Wells. Analysis and interpretation of data: Hiwase, Ovenden, Wells. Review of submitted version of manuscript: Hiwase, Bak, Laden, Ovenden, Wells. Approved the final version of the manuscript on behalf of all authors: Hiwase. Administrative/technical/material support: Hiwase, Kalyanasundaram, Wells. Study supervision: Wells.

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