Tailored Surgical Approaches for Foramen Magnum Tumors

Abordagens personalizadas aos tumores do forame magno

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Abstract

**Objective** To describe our surgical techniques, analyze their safety and their postoperative outcomes for foramen magnum tumors (FMTs).

**Methods** From 1986 to 2014, 34 patients with FMTs underwent surgeries using either the lateral suboccipital approach, standard midline suboccipital craniotomy, or the far lateral approach, depending on the anatomic location of the lesions.

**Results** In the present series, there were 22 (64.7%) female and 12 (35.2%) male patients. The age of the patients ranged from 12 to 63 years old. We observed 1 operative mortality (2.9%). A total of 28 patients (82.3%) achieved a score of 4 or 5 in the Glasgow Outcome Scale (GOS). Gross total resection (GTR) was obtained in 22 (64.7%) patients. After the surgery, 9 (26%) patients developed lower cranial nerve dysfunction (LCNd) weakness. The follow-up varied from 1 to 24 years (mean: 13.2 years).

**Conclusion** The majority of tumors located in the FM can be safely and efficiently removed using either the lateral suboccipital approach, standard midline suboccipital craniotomy, or the far lateral approach, depending on the anatomic location of the lesions.

Keywords

- foramen magnum
- meningiomas
- tumors
- microsurgery
- retrocondylar approach
- suboccipital approach

Resumo

**Objetivo** Descrever as nossas técnicas cirúrgicas, analisar sua segurança, relatar e discutir os resultados para tumores localizados no foramen magnum (FM).

**Método** Este um estudo retrospectivo de 34 pacientes com tumores localizados no FM que foram submetidos a craniectomia suboccipital lateral, ou a craniectomia clássica suboccipital, ou ainda ao acesso extremo lateral, entre os anos de 1986 a 2014.

**Resultados** Este grupo de pacientes foi composto por 12 homens (35.2%) e 22 (64.7%) mulheres, cuja idade variou de 12 a 63 anos. A remoção total da lesão ocorreu em 22 (64.7%) indivíduos. A mortalidade cirúrgica foi observada em apenas 1 paciente (2.9%). Um total de 28 (82.3%) pacientes alcançaram 4 ou 5 pontos na Escala de Resultados de Glasgow. O acompanhamento médio foi de 13,2 anos.
Introduction

The foramen magnum (FM) comprises a bony channel formed anteriorly by the lower third of the clivus, the anterior arch of the atlas, and the odontoid process. The lateral limits are thejugular tubercle (JT), the occipital condyle (OC), and the lateral mass of the atlas. Lastly, the FM is limited posteriorly by the lower part of the occipital bone, the posterior arch of the atlas, and the two first intervertebral spaces. The FM encloses the brainstem, the vertebral arteries (VAs), the anterior and posterior spinal arteries, the IX, X, XI, XII cranial nerves and the roots of the C2. Despite the great development of microsurgery, the surgical management of tumors located in the FM continues to challenge neurosurgeons because of the complex anatomy of this region. These tumors grow in close contact with neural and vascular structures that cannot be sacrificed or retracted. It raises controversies related to the best surgical approaches to deal with these lesions.1–13

Method

Data Collection

A retrospective study was performed with 34 consecutive patients with foramen magnum tumors (FMTs) diagnosed and operated at the Department of Neurosurgery of the Public Servants Hospital, a tertiary referral center, and at Rede D’Or/São Luís hospital, a private unit at Rio de Janeiro, from 1986 to 2014. The files, operative notes, pre and postoperative imaging studies, pathological reports and, when available, intraoperative videos of the patients, were used for the analysis. A database was created, from which information pertinent to the present study was collected. The pathological review was performed based on the World Health Organization (WHO) guidelines. Postoperative imaging was performed within the first 72 hours after the surgery to document postoperative changes and the degree of tumor removal. Neuropsychological monitoring was used in the last 2 cases. The need for informed consent was waived due to the retrospective character of the study. The follow-up varied from 1 to 24 years. The first clinic visit was ~15 days after hospital discharge, and then at 2 and 6 months postoperatively; thereafter, the patients were reexamined as necessary. The Glasgow Outcome Scale (GOS) defined the outcome.

Clinical Characteristics

Chronic headache and/or neck and arm pain were the symptoms observed in 19 individuals (55.8%). Gait disturbance was reported in 21 patients (61.7%). Pyramidal syndrome was observed in 19 patients (55.8%). Lower cranial nerve dysfunction (LCNd) was diagnosed in 11 individuals (32.3%). All of the patients underwent a computed tomography (CT) scan and/or a magnetic resonance imaging (MRI) exam. A total of 20 (58.8%) lesions showed enhancement after contrast injection. Two (5.8%) of them showed calcifications. Twenty (58.8%) of the tumors were inserted anteriorly or anterolaterally to the dentate ligament; 14 lesions (41.1%) were posterior or posterolateral. Twelve lesions (35.2%) crossed the midline. (→Table 1)

Surgical Technique

The same surgical technique was used following these general steps: anesthesia was induced with a carefully endotracheal intubation and standard anesthetic equipment was employed to detect and treat air embolism. A total of 21 (61.7%) patients were positioned on a semisitting position with the head slightly flexed and secured in the Mayfield

Table 1 Demographic features, histopathological types, and results of 34 patients with foramen magnum lesions

<table>
<thead>
<tr>
<th>Age of treatment: 12 to 66 years old (mean: 42.6 years old)</th>
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<tbody>
<tr>
<td>Gender: female = 22 (64.7%); male = 12 (35.2%)</td>
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<tr>
<td>Symptoms:</td>
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<tr>
<td>Headache: 20 (58.8%)</td>
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<tr>
<td>Gait disturbance: 22 (64.7%)</td>
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<td>Motor deficits: 20 (58.8%)</td>
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<tr>
<td>Lower cranial nerve dysfunction: 12 (35.2)</td>
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<td>Pathological entities:</td>
</tr>
<tr>
<td>Meningiomas: 16 (47.0%)</td>
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<tr>
<td>Brainstem astrocytomas: 8 (23.5%)</td>
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<td>Schwannomas: 4 (11.7%)</td>
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<tr>
<td>Epidermoids: 2 (5.8%)</td>
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<tr>
<td>Chordoma: 1 (2.9)</td>
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<td>Chondrosarcoma: 1 (2.9%)</td>
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<td>Ependymoma: 1 (2.9%)</td>
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<tr>
<td>Arachnoid cyst: 1 (2.9%)</td>
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<tr>
<td>Tumor size: 2.1 to 7.3 cm (mean: 3.6 cm).</td>
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<tr>
<td>Gross total resection in non-meningiomas: 12 (66.6%)</td>
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<tr>
<td>Gross total resection meningiomas: 12 (75.0%)</td>
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<tr>
<td>Cerebrospinal fluid fistula: 7 (20.5%)</td>
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<td>Lower cranial nerve deficits: 10 (29.4%)</td>
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<td>Recurrence: 9 (26.4%)</td>
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<td>Operative mortality: 1 (2.9%)</td>
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<td>Follow-up: 1 to 24 years (mean: 10.2 years).</td>
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<td>Glasgow outcome scale 4 and 5: 25 (73.5%)</td>
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head holder. In the other (38%) patients, the lateral decubitus or the park bench position was chosen. The procedure was initiated with the use of a 2.5 surgical loupe and coaxial lighting for soft tissue incision and bone work.

**Skin incision and muscle dissection:** For tumors located anterior to the dentate ligament, a 12-cm vertical linear incision was made 2 to 3 cm medially to the mastoid and centered ~2 cm above the mastoid tip. The incision was carried down through the galea and the periosteum over the suboccipital bone and the posterior border of the sternocleidomastoid and trapezius muscles, proceeding to the C2-C3 level.

**Muscular Dissection:** The spinous process of the second cervical vertebra is a palpatory guide to the position of the FM and enables the subperiosteal dissection of the suboccipital region to be carried along the C1 posterior arch. The paravertebral muscles are detached from their attachment to the occipital scama and are progressively sectioned with a scalpel. A self-retained retractor is progressively inserted into the wound, exposing the suboccipital triangle and maintaining the paravertebral muscle in the appropriate position. At this point, the C1cposterior arch is identified and dissected free with a periosteum elevator until the mastoid process is exposed. The vertebral artery (VA) is kept undisturbed in the sulcus arteriosus. The ipsilateral lateral half of the arch of the atlas is then resected with the Laksell rongeur, as previously described.\textsuperscript{14}

**Craniotomy:** The suboccipital craniotomy is performed unilaterally; it is extended to the FM and to the posterior edge of the occipital condyle, but the condyle is preserved in most of the cases. This access provides sufficient midline and lateral suboccipital exposure to the tumor. If more exposure is needed, C2 and C3 laminectomy can be included. Emisary veins opened during subperiosteally dissection should be coagulated bipolarly and waxed immediately.

**Opening the dura:** The surgical microscope is introduced in the operative field and the operation is performed with a magnification that varies from 10x to 16x until the end of the procedure. The dura is opened longitudinally, medial from the VA entry or in the midline in the cases of a standard suboccipital approach. The dural edges are tented up. The arachnoid space of the cisterna magna is opened to drain the cerebrospinal fluid (CSF). The tumor is exposed under the arachnoidal cleavage plane and is dissected free with a periosteum elevator until the mastoid process is exposed. The vertebral artery (VA) is kept undisturbed in the sulcus arteriosus. The ipsilateral lateral half of the arch of the atlas is then resected with the Laksell rongeur, as previously described.\textsuperscript{14}

**Debulking the tumor:** After bipolar coagulation with low current under saline irrigation, the tumor is partially devascularized; the capsule is incised with a scalpel, penetrated, and progressively debulked from within, with piecemeal tissue removal techniques. Rigorous homeostasis is maintained throughout the operation. Careful attention is paid to identify and respect the arachnoid plane at the tumor-brainstem interface, which facilitates complete tumor resection and minimizes small vessel and brainstem injury, as previously published.\textsuperscript{14,15}

**Dissecting the tumor:** The surgery proceeds within the space provided by the tumor growth. The meningoima is then dissected away from the LCNs and from the blood vessels by gentle meticulous microsurgical techniques, with microscissors and dissectors and multiple microscope angulations. As the debulking of the tumor proceeds, the brainstem relaxes progressively and provides additional working space for dissection around the capsule.

Then, the site of attachment of the tumor is identified, coagulated, and sectioned. A GTR is always attempted, but if the arachnoidal cleavage plane cannot be defined during surgery, or if dissection of the tumor from the VA, from its branches, from the brainstem or from the LCN may entail risk of damage, we leave a thin rim of tumor attached to these structures. In case of meningiomas, no attempts are made to resect the dura or to excise the bone involved.

In the cases of posterior or posterolateral lesions, as in brainstem gliomas, the skin incision is placed in the midline and the access to the suboccipital scama is made between the paravertebral muscles. The microsurgical removal of the lesions is made following the aforementioned technical rules.

The dura is closed either primarily or with a free pericranial graft. The closure of superficial planes consists of three layers of suture, with nylon stitches on the skin. If swallow deficits are noted in the postoperative period, an early tracheotomy is performed.\textsuperscript{15}

**Results**

In the present series, there were 22 (64.7%) female and 12 (35.2%) male patients, whose age ranged from 12 to 66 years old (mean: 42.6 years old). The follow-up varied from 1 to 24 years (mean: 10.2 years old). The pathological entities comprised: 16 meningiomas, 8 brainstem gliomas, 4 schwannomas, 2 epidermoid tumors, 1 chordoma, 1 chondrosarcoma, 1 ependymoma, and 1 arachnoid cyst. The diameter of the tumors ranged from 2.1 to 7.3 cm (mean: 3.6 cm).

**Mortality, Morbidity and Outcome**

There was 1 (2.9%) operative mortality (until 30 days postsurgery) that occurred in an adolescent with Neurofibromatosis type 1 (NF1). We completely removed 2 dumbbell C2 Schwannomas involving both sensory and motor rootlets with critical compression of the spinal-medullary junction. Several days after surgery, he was admitted with acute respiratory distress and died due to aspiration pneumonia. Three other patients (8.8%) died between the 60th and 180th postsurgical days resulting from aspiration pneumonia and its consequences. Immediate postoperative dysfunction or aggravation of previous lower cranial deficits was observed in the 10 patients with aggravation of previous lower cranial deficits, only 3 recovered in the follow-up. Three of these patients recovered partially from the LCN deficits during the follow-up period. One patient presented partial brachial plexus paralysis and in 7 (20.5%)
Discussion

Surgical Aspects

Based on the insertion in the dura, the FM meningiomas (FMMs) can be classified as anterior, if they were attached to the anterior rim of the FM and displaced the neuraxis in a posterior direction (► Fig. 1). Anterolateral meningiomas are those situated on the ventrolateral rim of the FM and that displace the neuraxis posterolaterally (► Fig. 2A and 2B). Tumors in both subgroups were located ventral to the dentate ligament. Lesions are classified as posterior if the insertion in the dura is posterior to the dentate ligament (► Fig. 3). Most of the FMMs arise anterolaterally, followed by those that arise posterolaterally.1,3,4

Surgery of FMTs, located anteriorly or anterolaterally to the brainstem constitutes a formidable challenge to neurosurgeons. A controversial discussion has risen concerning the utility of systematic occipital condyle drilling for approaching anterolateral FMMs. Sen et al8 and a group of surgeons stated that, in anteriorly or anterolaterally located FMMs, the extreme lateral or far lateral approach associated with VA medial transposition with a total or partial condylectomy improves the angle of visualization of the area ventral to the lower brainstem, facilitating the dissection of the interface between the neuroaxis and the tumor.4,7,8,16–18 On the other hand, George et al12 Bassiouni et al13 and others19–21 concluded that, as the tumors progressively displace the medulla and the spinal cord posteriorly, they create a space and, through this space, the tumor can be safely and completely resected via posterolateral suboccipital craniotomy, without condylar drilling. Most of the cases of incomplete tumor resections are not the result of inadequate exposure, but instead of invasion of the pia mater of the brainstem or of the involvement of the VA or of the cranial nerves by the tumor. In these patients, a subtotal tumor resection is recommended, which leaves a thin rim of the tumor.4–7,19–21 Wanebo et al22 concluded that, in patients with a small foramen magnum, with a short distance between the anterior rim of the foramen magnum and the brainstem and relatively large occipital condyles, the transcondylar approaches would be helpful. On the contrary, Spektor et al23 reported that total resection of the condyle provided very little additional exposure to the anterior FM and did not compensate for the significant level of possible additional morbidity. Silveira et al24 concluded that the extensions of bone removal should be adapted to the topography of the lesion: the retrocondylar approach for the lateral area of the FM; the partial transcondylar for the anterolateral portion; and the complete transcondylar for the anterior part of the FM.25,26 A recent review found 657 cases from 29 different neurosurgical centers revealed that in 6 they routinely performed condylar resection, in 4 they tailored the drilling of the condyle and 7 others centers they never resect it.1

We order the lateral sub-occipital retrocondylar approach without resection of the occipital condyle for 16 patients (47.0%), (► Fig. 1 and 2) the standard posterior midline access for 14 (41.1%) individuals (► Fig. 1, 4 and ► Fig. 5). For one patient with a chondrosarcoma, another with a chordoma, and for two others with huge bilateral epidermoid tumors
with encasement of the basilar artery, we elected the far lateral approach with partial or total condylar resection (11.7%). The retrocondylar approach provided satisfactory exposure in the majority of the patients, because most of these anterior or anterolateral tumors grow predominantly to one side, providing a corridor of exposure without need of condylar resection. These lesions belong to the group of tumors in which, paradoxically, it is easier to excise a large tumor than a small one, because the larger tumors provide more space anteriorly and, thus, lessen the need for a more lateral exposure. Extensive drilling of the occipital condyle, of the lateral mass of the atlas and of the jugular tubercle can lead injury to the hypoglossal nerve, to the VA, and promote spinal instability.\textsuperscript{3,6,7,13,14,20,21,24–26} Since our objective has
always been to keep the quality of life of the patients a priority, a subtotal removal might represent a very acceptable goal in fibrous or calcified tumors encasing the VA and perforating vessels or adhering to cranial nerves. In these patients, a subtotal tumor resection is recommended, which leaves a thin rim of the tumor. Most of the cases of incomplete tumor resections are not the result of inadequate exposure, but instead of invasion of the pia mater of the brainstem or of the involvement of the VA or of cranial nerves by the tumor. During the removal of the tumor, bipolar coagulation is kept to a minimal, used under continuous cold saline irrigation, and the aspiration is seated in soft suction mode. Total removal could be dangerous for some patients and difficult to achieve because these tumors are critically located, and there may be adherence or even encasement of vital structures by the tumor. If the surgeon persists with any attempt to remove every last residual lesion to achieve the cure of the patient, it could result in unwarranted lesions or injuries in the perforating branches. The main cause of postoperative morbidity is injury to the vertebral or basilar artery branches. Injury to even a single perforating vessel can result in brainstem infarction. On diagnosis, many lesions are already large or giant. Several authors reported increasing tumor size to portend higher rates of incomplete resection, higher rates of surgical morbidity, and an increased risk of recurrent disease. The decision on the extent of surgical resection can be difficult, particularly in older patients or in patients with minor symptoms. On the other hand, experience clearly emphasizes that, when tumors are incompletely removed, they tend
to regrow, and the next surgeon operating on the patient will be confronted with severe adhesions of blood vessels and brain tissue.

However, even when GTR is performed, tumor recurrence can occur. Radiosurgery might be considered as an alternative therapy for residues or recurrences, or even for patients deemed poor candidates for resection.26

The role of the endoscopic endonasal approach in the management of ventral posterior fossa meningiomas is still restricted due to limited surgical indications and experience. It can be used for the rare cases of meningiomas with most parts of its dural base at the midline clival region. Most of these patients were operated on by two senior neurosurgeons that gave to this sample a homogenous surgical technique. The limitation of the present study is that, compared with other published series, the present series is relatively small.

We can conclude that, in most of the cases of anterior or anterolateral FMTs, the lateral suboccipital approach without drilling the occipital condyle is less invasive, but it is sufficient for tumor removal. This approach is safe and effective. However, tailoring the surgery, including drilling of the occipital condyle, based on the size and shape of the FM, and on the local of the dural origin of the meningioma, also seems a reasonable option. In posterior or posterolateral located lesions, the standard posterior midline is sufficient.

Conflict of Interests
The authors have no conflict of interests to declare.

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