Outcome Assessment After Surgical Treatment of Tuberculum Sellae Meningiomas-A Preliminary Report

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ABSTRACT

AIM: To evaluate the endocrinological and ophthalmological results of the tuberculum sellae meningioma surgery.

MATERIAL and METHODS: A total of 18 patients diagnosed with tuberculum sellae meningioma received surgical treatment at Neurosurgery Clinic at Military Medical Institute in Warsaw from January 2010 to July 2012. This analyzed group of patients included 15 females and 3 males at a mean age of 50.5 years (ranging from 30 to 73; SD ± 13.4). In the pre-operative and post-operative periods, all patients underwent magnetic resonance imaging of the head according to a uniform protocol, eye examinations with assessment of visual acuity and field of vision, as well as endocrine tests to evaluate the hypothalamic-pituitary axis (including urine specific gravity and osmolality as well as blood cortisol, TSH, fT3, and fT4 levels). All patients underwent surgical treatment of the tumor via unilateral subfrontal craniotomy, achieving macroscopically complete tumor removal (Simpson grade II resection).

RESULTS: The use of the unilateral subfrontal approach helped improve vision in 88% of the treated patients. Endocrine tests revealed no hypothalamic-pituitary axis dysfunction. The most commonly diagnosed meningiomas of the area were meningothelial (77%) and transitional (12%) meningiomas.

CONCLUSION: Surgical treatment of tuberculum sellae meningiomas via the unilateral subfrontal approach is a safe technique with no significant complications. Visual improvement was observed in 88% of the patients who had received this treatment. There were no hormonal disturbances in patients operated via the subfrontal approach either prior to or after the procedure, which suggests that surgical treatment of tuberculum sellae meningiomas has no effect on pituitary endocrine function.

KEYWORDS: Tuberculum sellae meningioma, Frontal craniotomy, Pituitary hormones

INTRODUCTION

Tuberculum sellae meningiomas constitute 5–10% of intracranial meningiomas (12,25). Their location in the parasellar region, in close vicinity to hypothalamic structures and anterior vessels of the circle of Willis, results in the characteristic clinical signs. This location is also the reason for surgical treatment challenges. The most commonly used approach is the fronto-temporal, or pterional approach; with the fronto-orbital, fronto-orbitozygomatic, frontolateral (unilateral subfrontal), and bifrontal approaches being used less often (4,5,10,17,23,30,31).

We present the outcomes of tuberculum sellae meningioma surgery conducted by a single neurosurgeon, with a particular focus on visual disturbances and hypothalamic-pituitary function (21,22).

MATERIAL and METHODS

A total of 18 patients diagnosed with tuberculum sellae meningioma were treated surgically at the Neurosurgery Department between January 2010 and July 2012. The study group consisted of 15 females and 3 males at the mean age of 50.5 years (ranging from 30 to 73 years; SD ± 13.4). Table I presents demographic characteristics of the patients.
Clinical assessment

None of the evaluated patients were found to have hypothalamic endocrine dysfunction in the pre-operative period. All patients had visual disturbances in the form of reduced visual acuity and/or limited field of vision, including two patients (2 females) reporting only impaired visual acuity and two others (a female and a male) reporting only visual field defects. The history of these visual disturbances ranged from 1 to 36 months (mean: 14 months). Four patients additionally reported headaches in the pre-operative period.

Table II presents clinical data of the evaluated group.

Assessment of visual disturbances

A uniform protocol was followed to evaluate visual acuity and field of vision (the latter via Goldmann visual field exam) in all patients in the pre- and post-operative period.

Neuroradiological assessment

All patients underwent magnetic resonance imaging (MRI) assessments pre-operatively and in the post-operative period (3-4 months after surgery). These assessments were

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Table I: Sex, Age, and Standard Deviation (SD) Values in the Operated Patients

<table>
<thead>
<tr>
<th>Sex</th>
<th>No. of patients</th>
<th>Mean age</th>
<th>Minimum</th>
<th>Maximum</th>
<th>SD±</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Age (years)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>F</td>
<td>15</td>
<td>48</td>
<td>30</td>
<td>73</td>
<td>14</td>
</tr>
<tr>
<td>M</td>
<td>3</td>
<td>54</td>
<td>46</td>
<td>65</td>
<td>10</td>
</tr>
</tbody>
</table>

* M: Male, F: Female.

Table II: Pre-Operative Signs and Symptoms in Patients Qualified for Surgery

<table>
<thead>
<tr>
<th>Pt No.</th>
<th>Pt initials</th>
<th>Age (years)</th>
<th>Sex</th>
<th>Headaches</th>
<th>Visual disturbances</th>
<th>Pituitary endocrine function</th>
<th>Tumor size in mm (MRI)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>History of visual disturbances (in months)</td>
<td>Diabetes insipidus</td>
<td>Corticotrophic function</td>
</tr>
<tr>
<td>1.</td>
<td>LS</td>
<td>45</td>
<td>F</td>
<td>0</td>
<td>1 1 1 12</td>
<td>0 0 0 0</td>
<td>20x20x26</td>
</tr>
<tr>
<td>2.</td>
<td>WJ</td>
<td>67</td>
<td>F</td>
<td>1</td>
<td>1 1 1 1</td>
<td>0 0 0 0</td>
<td>35x30x34</td>
</tr>
<tr>
<td>3.</td>
<td>IJ</td>
<td>73</td>
<td>F</td>
<td>1</td>
<td>1 1 18</td>
<td>0 0 0 0</td>
<td>23x24x36</td>
</tr>
<tr>
<td>4.</td>
<td>IB</td>
<td>37</td>
<td>F</td>
<td>1</td>
<td>1 1 4</td>
<td>0 0 0 0</td>
<td>20x20x20</td>
</tr>
<tr>
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<td>BT</td>
<td>65</td>
<td>F</td>
<td>0</td>
<td>1 1 24</td>
<td>0 0 0 0</td>
<td>34x30x20</td>
</tr>
<tr>
<td>6.</td>
<td>AD</td>
<td>56</td>
<td>F</td>
<td>0</td>
<td>1 0 36</td>
<td>0 0 0 0</td>
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</tr>
<tr>
<td>7.</td>
<td>JP</td>
<td>67</td>
<td>F</td>
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</tr>
<tr>
<td>8.</td>
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</tr>
<tr>
<td>9.</td>
<td>LT</td>
<td>46</td>
<td>M</td>
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<td>0 1 24</td>
<td>0 0 0 0</td>
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</tr>
<tr>
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<td>30</td>
<td>F</td>
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</tr>
<tr>
<td>11.</td>
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<td>40</td>
<td>F</td>
<td>0</td>
<td>0 1 24</td>
<td>0 0 0 0</td>
<td>18x19x14</td>
</tr>
<tr>
<td>12.</td>
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<td>52</td>
<td>M</td>
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</tr>
<tr>
<td>13.</td>
<td>IT</td>
<td>36</td>
<td>F</td>
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<td>1 1 1</td>
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<td>23x21x12</td>
</tr>
<tr>
<td>14.</td>
<td>JP</td>
<td>57</td>
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<td>15x23x20</td>
</tr>
<tr>
<td>15.</td>
<td>JK</td>
<td>40</td>
<td>F</td>
<td>0</td>
<td>1 1 36</td>
<td>0 0 0 0</td>
<td>25x25x25</td>
</tr>
<tr>
<td>16.</td>
<td>IO</td>
<td>65</td>
<td>M</td>
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<td>1 1 12</td>
<td>0 0 0 0</td>
<td>35x21x25</td>
</tr>
<tr>
<td>17.</td>
<td>MW</td>
<td>31</td>
<td>F</td>
<td>1</td>
<td>1 1 12</td>
<td>0 0 0 0</td>
<td>18x14x19</td>
</tr>
<tr>
<td>18.</td>
<td>US</td>
<td>47</td>
<td>F</td>
<td>0</td>
<td>1 1 12</td>
<td>0 0 0 0</td>
<td>27x24x24</td>
</tr>
</tbody>
</table>

* Headaches: 0= no; 1= yes; Visual acuity disturbances: 0= no; 1= yes; Limited field of vision: 0= no; 1= yes; Pituitary function: 0= normal; 1= abnormal. Pt: Patient, M: Male, F: Female, MRI: Magnetic resonance imaging.
conducted at Medical Radiology Laboratory with a Signa 1.5 T scanner (GE Medical Systems) with spin echo (SE) T1-weighted sequences (parameters: Repetition time (TR) =450–600 ms, Echo time (TE)=15–30 ms, three excitations, minimum matrix size of 256x256, field of view approximately 20 cm, slice thickness 2–3 mm in the coronal and sagittal planes, before and after intravenous administration of the paramagnetic contrast agent gadolinium DTPA (Gd-DTPA), as well as T2-weighted sequences in coronal sections

(Di Chro-Nelson formula);
The pre-operative assessments included:
1) Tumor size,
2) Possible concomitant cerebral edema,
3) Calcifications in the tumor,
4) Hyperostosis or degeneration of cranial bones near tumor attachment,
5) Tumor infiltrations towards the sella turcica,
6) Tumor position with respect to the circle of Willis.
The post-operative assessments included:
1) The extent of tumor resection,
2) Damage to the hypothalamic-pituitary system

Endocrine assessment
All patients qualified for operation underwent endocrine tests to evaluate the hypothalamic-pituitary axis function both before and after the procedure (1 day and 3–4 months following surgery). We focused primarily on: 1) finding any evidence of diabetes insipidus (24-hour urine collection, urine specific gravity and osmolality in mOsm/kgH2O), 2) measuring blood cortisol level to evaluate corticotropic function, 3) TSH, fT3, and fT4 to evaluate thyreotropic function.

Methods of qualification for surgery
After a clinical assessment and MRI evaluation, the patients were qualified to undergo surgery via the unilateral subfrontal approach

Anesthesiology
All patients were placed under general anesthesia according to standard neuroanesthesiologic practice with a focus on adequate cerebral perfusion and oxygenation. Therefore, total intravenous anesthesia (TIVA) was preferred for these procedures. As always, the patients were monitored (as dictated also by their general condition), which included electrocardiography, capnometry, pulse oximetry, as well as non-invasive, and direct blood pressure monitoring. Routine practice included the use of two intravenous lines.

Surgical procedure
All patients were operated on via a unilateral subfrontal approach. The medial part of the Sylvian fissure was routinely opened to drain the cerebrospinal fluid (CSF). After identification of the ipsilateral optic nerve (it was stretched by the tumor and appear in all cases), the internal carotid artery and precommunicating segment of the anterior cerebral artery (ACA), the blood supply to the tumor from the dura of the anterior fossa and tuberculum sellae was interrupted. Then, the tumor capsule was carefully microsurgically dissected from the optic nerves, chiasm, carotid arteries, and pituitary stalk after internal decompression of the tumor with cavitrul ultrasonic surgical aspirator and bipolar cautery. The tumor was also removed from the ipsilateral optic canal after its unroofing. Tumor resection was completed after identification and preservation of the postero-laterally displaced pituitary stalk. Finally, the basal dura (dural attachment) was removed and any enostosis on the limbus sphenoidale were drilled out. The tumor was totally removed in all cases (Simpson Grade I or II). The pieces of resected tumor were fixed in 10% formalin for the pathological examination.

Intraoperative assessment
The extent of tumor resection was assessed with the Simpson grading scale:
I- Macroscopically complete tumor resection, with excision of its dural attachment and any adjacent abnormal bone;
II- Macroscopically complete tumor resection with coagulation of its dural attachment,
III- Macroscopically complete removal of the tumor without resection (or coagulation) of its dural attachment,
IV- Partial removal of the tumor,
V- Tumor biopsy

Histopathological assessment
The resected tumor fragments were preserved for a histopathological examination. The examination was conducted at the Pathomorphology Laboratory. Histopathology examinations were routinely performed according to the WHO classification, with immunohistochemical evaluation of meningioma diagnostic markers i.e.: epithelial membrane antigen (EMA), CK (cytokeratin), S-100, glial fibrillary acidic protein (GFAP), carcinoembryonic antigen (CEA), CD43, Vimentin. The material to be assessed under a light microscope was fixed in a 4% solution of formaldehyde in water with phosphate buffer (fixing agent pH 7.2).

Statistical analysis
The statistical analysis was conducted with Statistica 10.0. The data collected for the group were measured directly, constituting either objective or subjective aspects of the assessment. In two cases, the assessed variable values were altered:
1. Tumor volume was measured as the volume of an ellipsoid based on the tumor size graphed onto an X,Y,Z coordinate system, and calculated with the formula, where a,b,c represented half-axes of the ellipsoid;
2. Outcome was determined based on the sum of variables: Visual Acuity Outcome, Visual Field Outcome, Pituitary Function Outcome.
Considering the fact that the Shapiro and Wilk's W statistic was not statistically significant, there were no grounds to reject the hypothesis stating that distribution of the Age variable is not consistent with normal distribution. This conclusion was also confirmed by a normal distribution graph for this variable (Figure 3).

**Neuroradiological assessment results**

Tumor size was analyzed in all patients and stratified by sex. Tumors were divided into three groups depending on their size:

a) From 0 to 2 cm – in 3 patients (2 females and 1 male)

b) From 2 to 4 cm – in 15 patients (13 females and 2 male)

c) Over 4 cm – in 0 patients

The volume of the tumor was also evaluated in the entire study group, stratified by sex (Figure 4 and Table IV). Three patients had tumor volume in the 0–6,000-mm$^3$ range; 4 patients – in the 6,001-mm$^3$–12,000-mm$^3$ range; and in 11 patients tumor size exceeded >12,000-mm$^3$.

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**Figure 1:** Magnetic resonance images of the tuberculum sellae meningioma prior to surgical treatment (1-300, 2-300, 3-300).

**Figure 2:** Histogram of the Age variable.

**Figure 3:** Normal distribution of the Age variable.
Tumors found in female patients tended to be larger. However, the differences in tumor size (volume) between the female vs. male groups were not statistically significant.

**Results of assessing visual disturbances**

All patients were diagnosed with visual disturbances before their surgery, with two patients (2 females) reporting exclusively visual acuity abnormalities and two (1 female and 1 male) reporting exclusively field of vision abnormalities. The most common visual field defect was bitemporal hemianopsia, diagnosed in 10 patients. Four patients were diagnosed with mono-ocular (right) temporal hemianopsia and two with mono-ocular (left) temporal hemianopsia. Vision improved in all patients in the post-operative period versus their pre-operative status. There were no exacerbations in visual disturbances in any of the patients.

Although an analysis of statistical variables prior to surgical intervention revealed no statistically significant differences, it

![Figure 4: Tumor volume stratified by sex.](image)

**Table III: Results of Surgery**

<table>
<thead>
<tr>
<th>Pt No.</th>
<th>Pt Initials</th>
<th>Age (years)</th>
<th>Sex</th>
<th>Extent of the operation (Simpson grade)</th>
<th>Post-operative vision</th>
<th>Pituitary endocrine function</th>
<th>Histological diagnosis</th>
<th>Post-operative follow-up in months</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>LS</td>
<td>45</td>
<td>F</td>
<td>2</td>
<td>1</td>
<td>1</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>2</td>
<td>WJ</td>
<td>67</td>
<td>F</td>
<td>2</td>
<td>1</td>
<td>1</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>3</td>
<td>IJ</td>
<td>73</td>
<td>F</td>
<td>2</td>
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<td>0</td>
<td>0</td>
</tr>
<tr>
<td>4</td>
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</tr>
<tr>
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<td>0</td>
</tr>
<tr>
<td>6</td>
<td>AD</td>
<td>56</td>
<td>F</td>
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<td>0</td>
<td>0</td>
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<td>0</td>
<td>0</td>
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<td>1</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>9</td>
<td>LT</td>
<td>46</td>
<td>M</td>
<td>2</td>
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<td>1</td>
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<td>0</td>
</tr>
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<td>0</td>
<td>0</td>
</tr>
<tr>
<td>11</td>
<td>KL</td>
<td>40</td>
<td>F</td>
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<td>0</td>
<td>0</td>
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<tr>
<td>12</td>
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<td>52</td>
<td>M</td>
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<td>1</td>
<td>1</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
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<td>IT</td>
<td>36</td>
<td>F</td>
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<td>0</td>
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<tr>
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<td>0</td>
<td>0</td>
</tr>
<tr>
<td>15</td>
<td>JK</td>
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<td>F</td>
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<td>0</td>
<td>0</td>
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<tr>
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</tr>
<tr>
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<td>0</td>
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<td>47</td>
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<td>2</td>
<td>1</td>
<td>1</td>
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<td>0</td>
</tr>
</tbody>
</table>

**The extent of the operation (Simpson grade):** 1– macroscopically complete tumor resection, with excision of its dural attachment and any adjacent abnormal bone; 2– macroscopically complete tumor resection with coagulation of its dural attachment; 3– Macroscopically complete tumor removal without resection of its dural attachment, 4– Partial tumor removal, 5– Tumor biopsy. **Post-operative visual acuity:** -1–worsening; 0–no change; 1–improvement; **Post-operative field of vision (FOV):** -1–worsening; 0–no change; 1–improvement; **Post-operative pituitary function:** diabetes insipidus -1–worsening, 0–no improvement; 1–improvement; corticotropic function -1–worsening, 0–no change, 1–improvement; thyreotropic function -1–worsening, 0–no change, 1–improvement; **Histopathological diagnoses:** MM – meningothelial meningioma; PM – psammomatous meningioma; TM – transitional meningioma. **M:** Male, **F:** Female, **Pt:** Patient.
Similar analyses were conducted for post-operative variables. The results are presented in Table VI.

Analysis results indicated the existence of a slight correlation between tumor volume (size) and visual acuity results. This indicates that removal of an objectively larger tumor may be associated with greater expectations as to improvement in visual acuity. A similar, though weaker still, correlation was observed between the value of the Outcome variable and tumor volume. However, given the fact that these correlations were not statistically significant, they should not be subject to generalization but rather should be treated as an indicator in analyzing larger study groups.

Endocrine assessment results

In the pre-operative period, no patients presented symptoms of diabetes insipidus, or any abnormalities in corticotropic or thyreotropic pituitary function as manifestations of tuberculum sellae meningioma. A similar lack of pituitary abnormalities was observed post-operatively, with no patients having developed diabetes insipidus or cortico-/thyreotropic pituitary dysfunction.

Histological assessment results

Histopathological examinations of the resected tumors revealed only 3 meningioma subtypes: a) meningothelial, b) psammomatous, and c) transitional meningioma (Table VII). The most commonly diagnosed (in 12 females and 2 males [77%]) subtype of meningioma was meningothelial meningioma. Psammomatous meningioma was diagnosed in 2 females (11%) and transitional meningioma in 1 female and 1 male (11%). Post-operative follow-up revealed no cases of tumor recurrence, thus we were unable to compare these histological findings to those in recurrent tumors.

**DISCUSSION**

The first case of tuberculum sellae meningioma was reported by J. Steward, based on a postmortem examination in 1899. In 1916, H. Cushing conducted the first complete removal of this tumor. In 1938, Cushing and Eisenhardt presented a series of 28 tuberculum sellae meningiomas and proposed their classification based on tumor size. They also introduced the term “suprasellar chiasmal syndrome” describing the clinical course of this tumor, which allowed for its differentiation from other tumors of the region (pituitary adenomas, craniopharyngiomas and gliomas). Since then, a number of tuberculum sellae meningioma case series have been described in the literature reporting the results of

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### Table IV: Tumor Volume in mm³ in the Total Study Group, and Stratified by Sex

<table>
<thead>
<tr>
<th>Sex</th>
<th>valid N</th>
<th>Mean</th>
<th>Minimum</th>
<th>Maximum</th>
<th>SD</th>
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<td>1072.33</td>
<td>18,692.48</td>
<td>4708.25</td>
</tr>
<tr>
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<td>6310.94</td>
<td>816.81</td>
<td>9621.13</td>
<td>4791.26</td>
</tr>
<tr>
<td>Total</td>
<td>18</td>
<td>6915.14</td>
<td>816.81</td>
<td>18692.48</td>
<td>4586.25</td>
</tr>
</tbody>
</table>

M: Male, F: Female.
surgical treatment. We presented an assessment of surgical treatment outcomes in 18 patients (i.e. 15 females [87%] and 3 males [15%] aged between 30 and 73 years (mean age: 50.5; standard deviation: 13.4; including 54.3 years in men and 49.7 in women) diagnosed with tuberculum sellae meningioma. According to Cushing and Eisenhardt, the mean age of patients diagnosed with central nervous system meningioma ranges from 42.9 years in females and 52 years in males (7). Fahlbusch and Schott (9) reported the mean age in affected females to be 54.3 years and that in affected males 54.9 years (7,9,28). In the present study, a surgical approach was selected for each patient and all patients who qualified for surgery underwent unilateral subfrontal craniotomy, as this approach was considered safe. The extent of tumor removal in all patients was classified as Simpson grade II. We did not observe any deaths or significant post-operative complications (such as cerebrospinal fluid leakage, infections, or intracranial hematomas) in the group operated via the unilateral subfrontal approach. According to the available literature reports, the mortality rate in tuberculum sellae meningiomas ranges from 0% to 8.6%. Interestingly, Nakamura et al. reported a correlation between mortality and the surgical approach used, with the following findings: frontal and pterional approach – 0%, bifrontal approach – 9.5% (8,29). No deaths were observed in our study.

Tumor volume was also analyzed. The mean tumor volume was 6915.14 mm$^3$ (7035.98 mm$^3$ in females and 6310.94 mm$^3$ in males). The study group showed a correlation between tumor size and visual (acuity) abnormalities (3,6,14,15). It was Rosenstein et al. who confirmed more favorable surgical treatment outcomes in the case of tumors below 3 cm in diameter, with greater odds for visual improvement. Andrew and Wilson noticed that in the case of tumors measuring over 6 cm in diameter, the chance of visual improvement is significantly lower (18).

We also analyzed visual disturbances – defects in visual field and visual acuity abnormalities prior to and after the operation. In the pre-operative period, visual acuity disturbances were observed in 16 patients (88%), and field of view defects were also observed in 16 patients (88%). We would like to emphasize that each patient had at least one type of visual disturbance. A study by Fahlbusch and Schott reported that 96% patients developed visual impairment as the first symptom of their disease. In our group, the pre-operative visual disturbances improved in all patients following the operation. No patient experienced further visual impairment – whether field of view or acuity. In 2 patients (12%, 1 male and 1 female) visual acuity did not change, and 16 patients (88%) experienced visual acuity improvement. In terms of visual field defects, no change was observed in two patients (12%; 2 females) and improved field of view was observed in 16 patients (88%). Similar findings were reported by Mathiesen and Kihlstrom in a group of 29 patients (16). In that study, 91% of patients experienced postoperative visual improvement. Somewhat less favorable results were described by Otanii et al. who reported visual field improvement in only 78% of 32 operated patients (2,11,20). Conversely, Fahlbusch and Schott reported no visual improvement in 36% of patients in their group of 47, and postoperative exacerbation in the existing visual acuity and field of view impairment in 20% of patients. Nakamura et al. reported an improvement in vision in 65% of 72 operated patients, and particularly emphasized that the surgical procedures conducted via lateral supraorbital access yielded vision improvement in 77.8% of patients. Bifrontal craniotomy yielded poorer results at 46.2% – still, this approach is used only in large tumors. However, Romani et al., who had used the lateral supraorbital approach, confirmed visual improvement in 52% out of 52 patients, with no improvement observed in 31% patients, and postoperative visual deterioration – in 2% (1,13,24,26,27). The rate of visual improvement in the presented literature ranges from 19% to 91% of operated patients, and the rate of visual deterioration ranges from 3% to 39%. These findings indicate that surgical treatment via frontal craniotomy is associated with a significant visual improvement versus the preoperative status. We also evaluated the duration of visual disturbances. Our
data was collected by means of a questionnaire completed by patients on a voluntary basis. Thus, these assessments were very subjective, as patients frequently failed to assign much significance to their early symptoms of visual impairment, treating them as part of the natural course of life. In reality, it was the emergence of subsequent clinical manifestations of the tumor that allowed for estimating the duration of visual disturbance history. The history of visual disturbances ranged from 1 to 36 months, with a mean of 14 months.

Our study included also an evaluation of endocrine pituitary function based on possible manifestations of diabetes insipidus, as well as corticotropic and thyreotropic function prior to, and after, surgery. Romani et al. reported endocrine disturbances in 10% out of their 42 patients (26). Recent literature in the field has seen an increased incidence of reported endocrine disturbances ranging from 2% to 43%. Our study group demonstrated no hormonal abnormalities either prior to or after the surgery. Apart from our study, a study by Fahrbusch and Schott including 43 patients who qualified to undergo hormonal tests (from a group of 47) was the only one to show no evidence of diabetes insipidus in this type of tumor either before or after surgery. We would like to stress that some patients were receiving long-term hormone replacement therapy prior to surgery and were euthyreotic until the time of the procedure, having been endocrinologically prepared for surgery (9).

We also analyzed histopathological findings, as they play a significant role in scheduling post-operative follow-up visits and possible referrals for radiotherapy. Fahrbusch and Schott (9) demonstrated in a group of 47 patients that 79% were cases of meningothelial meningioma and 11% represented transitional meningioma. Our study also showed a greater incidence of meningothelial (14 patients [77%]; 12 females and 2 males) and transitional meningioma (in 2 patients [12%]; 1 female and 1 male). These two tumor types constituted 89% of all meningiomas in the evaluated patients.

Our findings suggest certain correlations between the pre- and post-operative groups; however, these correlations must not be generalized. This may change when findings from a larger study group become available. Tumor location, with its proximity to the pituitary, suggests that, in a larger study group, surgical interventions in this region may have a discernible effect on pituitary function. The seeming lack of effects on the pituitary endocrine function may be amended in the future, as more pituitary hormones are considered and possible referrals for radiotherapy. Fahlbusch and Schott including 43 patients who qualified to undergo hormonal tests (from a group of 47) was the only one to show no evidence of diabetes insipidus in this type of tumor either before or after surgery. We would like to stress that some patients were receiving long-term hormone replacement therapy prior to surgery and were euthyreotic until the time of the procedure, having been endocrinologically prepared for surgery (9).

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## CONCLUSIONS

1. Surgical treatment of tuberculum sellae meningiomas via unilateral frontal craniotomy is a safe technique with no significant complications.

2. Visual improvement was reported in 88% of the operated patients.

3. No pre- or post-operative endocrine disturbances were observed in patients operated via the supraorbital approach, which suggests that surgical treatment of tuberculum sellae meningiomas has no effect on pituitary endocrine function.

## REFERENCES


