Does General Comorbidity Impact the Postoperative Outcomes After Surgery for Large and Giant Petroclival Meningiomas?

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Abstract

Objective

We assessed the role of the general condition of the patient in addition to usual anatomical reasoning to improve the prediction of personalized surgical risk for patients harbouring a large and giant petroclival meningiomas.

Methods

Single-center, retrospective observational study including adult patients surgically treated for a large and giant petroclival meningioma between January 2002 and October 2019 in a French tertiary neurosurgical skull-base center by one Neurosurgeon. Inclusion criteria were: 1) histopathologically proven meningioma; 2) larger than 3cm in diameter; 2) located within the upper two-thirds of the clivus, the inferior petrosal sinus, or the petrous apex around the trigeminal incisura, medial to the trigeminal nerve. Clinical and radiological characteristics were gathered preoperatively including ASA score, the modified Frailty Index and the Charlson Comorbidity Index. Post-operative severe neurological and non-neurological complications were collected.

Results

A total of 102 patients harbouring a large and giant petroclival meningioma were included. The rate of postoperative death was 3.0% related to a congestive heart failure (n=1), a surgical site hematoma (n=1), and an ischemic stroke (n=1). A severe neurological impairment was found in 12.8% and a severe non-neurological morbidity was found in 4.0%. The overall rate of severe morbidity and mortality was 15.7% after large and giant petroclival meningioma surgery. The presence of brainstem peri-tumoral edema (adjusted OR, 4.83 [95% CI 1.84–7.52], p=0.028) was independently associated with a history of postoperative severe neurological morbidity. Male gender (adjusted OR, 7.42 [95% CI 1.05–49.77], p=0.044), major cardiovascular morbidity (adjusted OR, 9.5 [95% CI 1.05–86.72], p=0.045), and an ASA score ≥ 2 (adjusted OR, 11.09 [95% CI 1.46–92.98], p=0.038) were independently associated with a history of postoperative severe non-neurological morbidity. A modified Frailty index ≥ 1 (adjusted OR, 3.13 [95% CI 1.07–9.93], p=0.047), and a low neurosurgical experience (adjusted OR, 5.38 [95% CI 1.38–20.97], p=0.007) were independently associated with a history of postoperative overall morbidity and mortality.

Conclusions

This study suggests to add scores assessing the patient general condition in daily practice to improve the selection of patients eligible for surgery. Collaborative international multicenter studies will be necessary to confirm these results and allow their implementation in clinical routine.

Introduction
Petroclival meningiomas (PCMs) are rare and enduring tumour requiring one or more complex surgical resection[28]. Large and giant PCMs are defined as the main tumour diameter larger than 2.5cm and originate from the upper two-thirds of the clivus, the petrous apex around the trigeminal notch or the inferior petrosal sinus or, medial or around to the trigeminal incisura[38]. Surgical removal of a PCM has a high morbidity and mortality profile, related to: 1) the deep seated location, which requires complex skull base approaches[32]; 2) the induration and the adherences of meningiomas, which potentially make their intraoperative manipulation difficult; 3) the propensity to engulf nerves and blood vessels, to invade the cavernous sinus and to extend to cranial fossae foramina; and 4) the large tumour volume at time of surgery. In reported series, mortality ranged 0–10%, the incidence of cranial nerves deficits ranged 29–76%, and major complications ranged 8–45%[2, 5, 7, 11, 23, 26, 28, 31, 34, 40]. In order to predict the risk of the PCM surgery, Adachi et al. in 2009, developed a dedicated scoring system, i.e. the ABC Surgical Risk Score, based upon pure anatomical criteria[1]. However, ABC Surgical Risk Score score did not encompass the general condition of the patient and, especially, comorbidities. Other scores assessing patients’ comorbidities have been developed and are widely used, as the American Society of Anaesthesiologists (ASA) score[33], the modified Frailty Index[25], and the Charlson Comorbidity Index[9]. Unfortunately, neurosurgical and co-morbid predictors are rarely associated in the assessment of perioperative risk factors. To date, no study assessed the impact of scores assessing patient comorbidities for large and giant PCMs. However, the neurosurgeon intuitively assesses the surgical risk by taking into account anatomical data, as suggested by Adachi et al.[1], but also by taking into account the patient’s comorbidities. Indeed, it is common to maintain a close clinico-radiological monitoring for a large and giant PCM in an elderly patient with numerous comorbidities.

Here, we suggested that integrating scores assessing patient comorbidities together with the usual anatomical reasoning could improve the prediction of personalized surgical risk for patients harboring a subtotal resection for a large and giant PCM.

Material And Methods

1. Data source

We performed a single-center, retrospective cohort study of adult patients (≥ 18 year-old) with a subtotal resection, which is our attitude, for large and giant PCM between January 2002 and October 2019 in a French tertiary neurosurgical skull-base center by the same senior neurosurgeon (PHR). Inclusion criteria were: 1) histopathologically proven meningioma; 2) larger diameter ≥ 3 cm; 2) location within the upper two-thirds of the clivus, the inferior petrosal sinus, or the petrous apex around the trigeminal incisura, medial to the trigeminal nerve. We used the Strengthening the Reporting of Observational Studies in Epidemiology (STROBE) Statement, as recommended[14].

2. Data collection

Data were obtained from the medical records using a protocol designed for the study.
Clinical characteristics were: sex, age, past medical history, the ASA physical status classification system score, the modified Frailty index, and the Charlson comorbidity index (Fig. 1). All patients underwent extensive pre- and post-operative neurological examinations with detailed medical history (including major diseases such as thromboembolism, respiratory, neoplastic, and other neurologic illness), the presence of a major cardiovascular morbidity (heart failure, rhythmic heart disease and coronary artery disease), functional independence using the Karnofsky Performance Status (KPS) score, and detailed cranial nerve (CN) testing. Swallowing and auditory function were assessed by an Otolaryngologist.

Meningioma imaging characteristics, systematically gathered from preoperative MRI, were: side, main insertion, secondary extension, largest diameter, volume, peri-tumoral edema, bone involvement, hydrocephalus, and mass effect on brainstem.

Surgical characteristics were: period of inclusion (the first half cases versus the second half cases) and surgical approach (anterior petrosectomy versus combined approaches).

Postoperative characteristics were: histopathological report, day 1 post-operative neurological examination, post-operative complications within the first month. Severe neurological impairment was defined as the presence of a postoperative: 1) surgical site hematoma requiring a new surgical procedure; 2) stroke; 3) refractory epileptic seizures; 4) long tracts deficits without stroke and/or 5) IX-X-XI palsy. Severe non-neurological morbidity was defined as postoperative: 1) congestive heart failure; 2) lung infection and; 3) pulmonary embolism.

3. Statistical analysis

To determine factors associated with a history of postoperative complications, univariate analyses were performed, computing unadjusted Odds Ratios (OR) and using the Chi square or Fisher’s exact tests for comparing categorical variables, and the unpaired t-test or Mann–Whitney rank-sum test for continuous variables, as appropriate. Variables associated at the p < 0.200 level in unadjusted analysis were then entered into backward stepwise logistic regression models.

We performed a backward stepwise selection of variables, in a subgroup of 73 patients according to available imaging data, removing the least significant variables one after the other, and defining the least significant variable as having the highest p-value in the model. The final model retained only the variables significant at the P < 0.05 level. Statistical analyses were performed using JMP software (version 14.3.0, SAS Institute Inc, Cary, USA).

4. Standard protocol approvals, registrations, and patient consents

Informed consent was obtained from all patients. This study received required authorizations (IRB#1: 2019/20) of the French College of Neurosurgery (IRB00011687).

Results
1. Clinical, imaging, histopathological, and treatment-related characteristics

Patients’ clinical characteristics are detailed in Table 1. A total of 102 patients (82 women, mean age 53.4±10.9) harboring a large and giant PCM were included. Ten patients (9.8%) underwent a previous surgical resection in another center and four of them (3.9%) had a postoperative radiotherapy. Two patients (2.0%) had a previous radiotherapy without surgery in another center. The median ASA score was 2 (range 1–3) with 45.1% of patients with a ASA score at 2 (mild systemic disease). The median modified Frailty Index and Charlson comorbidity index were 0 in 58.8% and 67.8% of patients with no significant disease, respectively.
| Parameters                                      | n  | %   |
|-----------------------------------------------|----|-----|
| **Clinical characteristics**                  |    |     |
| Sex                                           |    |     |
| Male                                          | 82 | 80.4|
| Female                                        | 20 | 19.6|
| Age, years (mean, SD) 53.4 ± 10.9              |    |     |
| < 55                                          | 49 | 48.0|
| ≥ 55                                          | 53 | 52.0|
| Karnofsky Performance Status                  |    |     |
| ≥ 70                                          | 3  | 2.9 |
| < 70                                          | 98 | 96.1|
| Cardiovascular comorbidities                  |    |     |
| Yes                                           | 96 | 94.1|
| No                                            | 6  | 5.9 |
| Previous surgical resection                   |    |     |
| Yes                                           | 92 | 90.2|
| No                                            | 10 | 9.8 |
| Previous radiotherapy                         |    |     |
| Yes                                           | 96 | 94.1|
| No                                            | 6  | 5.9 |
| ASA Score (mean, SD) 1.8 ± 0.7                 |    |     |
| 1                                             | 46 | 45.1|
| 2                                             | 20 | 19.6|
| 3                                             | 0  | 0   |
| ≥ 4                                           |    |     |
Patients’ neurological examination are detailed in Table 2. Cranial nerves deficits were the main symptoms at diagnosis, affecting 76.5% of patients: facial hypoesthesia in 42.2%, facial neuralgia in 21.6%, facial paralysis in 8.8%, and hearing loss in 31.4%. Only 4.9% of patients had an impairment of the IX-X-XI cranial nerves and no patient had a XII impairment at diagnosis. Diplopia was present at diagnosis in 21.6% of patients: Third cranial nerve impairment in 10.8%, fourth cranial nerve impairment in 4.9%, and sixth cranial nerve impairment 5.9%. Long tracts deficits were present in 30.4% and headaches were present in 19.6%. Epileptic seizures were present in 2.9% of patients at diagnosis.
### Table 2
Preoperative neurological examination of the study sample (n = 102)

| Parameters                        | n  | %   |
|-----------------------------------|----|-----|
| Cranial nerves deficits           | 78 | 76.5|
| II                                | 3  | 2.9 |
| III                               | 11 | 10.8|
| IV                                | 5  | 4.9 |
| V                                 | 49 | 48.0|
| Facial neuralgia                  | 22 | 21.6|
| Facial hypoesthesia               | 43 | 42.2|
| VI                                | 6  | 5.9 |
| VII                               | 9  | 8.8 |
| VIII                              | 35 | 34.3|
| Hearing loss                      | 32 | 31.4|
| Vestibular syndrome               | 11 | 10.8|
| IX-X-XI                           | 5  | 4.9 |
| XII                               | 0  | 0   |
| Long tracts deficits              | 31 | 30.4|
| Ataxia                            | 36 | 35.3|
| Proprioceptive ataxia             | 16 | 15.7|
| Cerebellar ataxia                 | 20 | 19.6|
| Hemiparesis                       | 12 | 11.8|
| Other neurological manifestations | 48 | 47.1|
| Headaches                         | 20 | 19.6|
| Epileptic seizure                 | 3  | 2.9 |
| Hydrocephalus                     | 28 | 27.5|

Imaging, histopathological and surgical characteristics are detailed in Table 3. Imaging characteristics were fully available for 73 patients. The median tumor volume was 21.8 cm$^3$ (range 7.5-164.2). A mass effect on brainstem was present in all cases. A peri-tumoral edema was present in 69.9% of cases.
Hydrocephalus was present in 38.4% of cases and hyperostosis was present in 23.3% of cases. Bone erosion was present in 4.1% of cases and a mixed erosion-condensation was present in 5.5% of cases.
Table 3
Main imaging, surgical and histopathological characteristics of the study sample (n = 102)

| Parameters                        | n   | %    |
|-----------------------------------|-----|------|
| **Imaging characteristics (n = 73)** |     |      |
| Tumor volume (cm³) (mean, SD) 27.7 ± 23.0 | 23  | 31.5 |
| ≥ 30                              | 50  | 68.5 |
| < 30                              |     |      |
| Peri-tumoral edema                | 51  | 69.9 |
| Yes                               | 22  | 30.1 |
| No                                |     |      |
| Bone involvement                  | 3   | 4.1  |
| Erosion                           | 17  | 23.3 |
| Hyperostosis                      | 4   | 5.5  |
| Mixed                             | 49  | 67.1 |
| No                                |     |      |
| Hydrocephalus                     | 28  | 38.4 |
| Yes                               | 45  | 61.6 |
| No                                |     |      |
| **Surgical characteristics (n = 102)** |     |      |
| Neurosurgeon experience (number of cases) | 50  | 49.0 |
| < 50                              | 52  | 51.0 |
| ≥ 50                              |     |      |
| Surgical approach                 | 75  | 73.5 |
| Transpetrosal approach            | 27  | 26.5 |
| Combined approach                 |     |      |
| **Histopathological characteristics (n = 102)** |     |      |

SD : Standard Deviation ; WHO : World Health Organization
The majority of patients were treated by an isolated transpetrosal approach (73.5%) and combined approaches were performed in 26.5% of cases. The histopathological analysis demonstrated a grade 1, grade 2, and grade 3 meningioma in 88.2%, 10.8%, and 1.0% of cases respectively.

2. Surgical outcomes

Adverse postoperative events data are detailed in Table 4. The rate of postoperative death was 3.0% in the whole series related to a congestive heart failure (n = 1), a surgical site hematoma (n = 1), and an ischemic stroke (n = 1). The overall rate of severe postoperative morbidity was 15.7%: a severe neurological impairment was present in 12.8% of cases, and a severe non-neurological morbidity was present in 4.0% of cases.

| Parameters                  | n  | %     |
|-----------------------------|----|-------|
| WHO grading                 | 90 | 88.2  |
| Grade 1                     | 11 | 10.8  |
| Grade 2                     | 1  | 1.0   |

Table 4
Adverse postoperative events of the study sample (n = 102)

| Parameters                                    | n  | %   |
|-----------------------------------------------|----|-----|
| **Postoperative events**                      |    |     |
| Severe neurological morbidity                 | 13 | 12.8|
| Ischemic Stroke                               | 2  | 2.0 |
| Cerebral Hematoma                             | 1  | 1.0 |
| Epileptic seizures                            | 1  | 1.0 |
| Long tracts deficits without stroke           | 8  | 7.8 |
| IX-X-XI palsy                                 | 1  | 1.0 |
| Severe non-neurological morbidity             | 4  | 4.0 |
| Congestive heart failure                      | 1  | 1.0 |
| Lung infection                                | 2  | 2.0 |
| Pulmonary embolism                            | 1  | 1.0 |
| Overall severe morbidity and mortality        | 16 | 15.7|
Table 5
Multivariate analyses (n = 73)

| Predictors                                      | Adjusted Hazard Ratio | 95% CI          | p-value |
|------------------------------------------------|-----------------------|-----------------|---------|
| **Severe neurological impairment**              |                       |                 |         |
| Peri-tumoral edema                              | 4.83                  | 1.84–7.52       | 0.028   |
| Yes                                            |                       |                 |         |
| No                                             |                       |                 |         |
| **Severe non-neurological morbidity**           |                       |                 |         |
| Sex                                            | 7.42                  | 1.05–49.77      | 0.044   |
| Male                                           |                       |                 |         |
| Female                                         |                       |                 |         |
| Major cardiovascular morbidity                  | 9.5                   | 1.05–86.72      | 0.045   |
| Yes                                            |                       |                 |         |
| No                                             |                       |                 |         |
| ASA Score                                       | 11.09                 | 1.46–92.98      | 0.038   |
| = 1                                            |                       |                 |         |
| ≥ 2                                            |                       |                 |         |
| **Overall morbidity and mortality**             |                       |                 |         |
| Modified Frailty Index                         | 3.13                  | 1.07–9.93       | 0.047   |
| = 0                                            |                       |                 |         |
| ≥ 1                                            |                       |                 |         |
| Pre-operative long tract deficits              | 1.43                  | 0.45–4.61       | 0.542   |
| Yes                                            |                       |                 |         |
| No                                             |                       |                 |         |
| Neurosurgeon experience (number of cases)      | 5.38                  | 1.38–20.97      | 0.007   |
| < 50                                           |                       |                 |         |
| ≥ 50                                           |                       |                 |         |

Unadjusted of severe neurological, non-neurological and overall morbidity and mortality are detailed in Supplementary Tables 1–3. In a multivariable analysis, the presence of brainstem peri-tumoral edema (adjusted OR, 4.83 [95% CI 1.84–7.52], p = 0.028) was independently associated with postoperative severe neurological morbidity. Male sex (adjusted OR, 7.42 [95% CI 1.05–49.77], p = 0.044), major
cardiovascular morbidity (adjusted OR, 9.5 [95% CI 1.05–86.72], p = 0.045), and an ASA score ≥ 2 (adjusted OR, 11.09 [95% CI 1.46–92.98], p = 0.038) were independently associated with postoperative severe non-neurological morbidity. A modified Frailty index ≥ 1 (adjusted OR, 3.03 [95% CI 1.07–9.70], p = 0.040), and a low neurosurgical experience (adjusted OR, 5.17 [95% CI 1.32–20.18], p = 0.009) were independently associated with postoperative overall morbidity and mortality.

Discussion

1. Key results

In this large retrospective cohort study of 102 large and giant PCM surgically treated by the same senior neurosurgeon we identified: 1) peri-tumoral edema as an independent predictor of severe neurological impairment; 2) male sex, major cardiovascular morbidity, a high ASA score as independent predictors of severe non-neurological morbidity; and 3) a high modified Frailty index and a low experience in large and giant PCM surgery as independent predictors of overall morbidity and mortality.

2. Interpretation

In our study, the rate of postoperative death was 3.0% after large and giant PCM surgery with a severe neurological impairment in 12.8% and a severe non-neurological morbidity in 4.0%. In reported series, mortality ranged 0–10%, the incidence of cranial nerves deficits ranged 29–76%, and major complications ranged 8–45%[2, 5, 7, 11, 23, 26, 28, 31, 34, 40]. Thereby, our overall rate of severe morbidity and mortality was 15.7% after large and giant PCM surgery in accordance with the literature[2, 4, 5, 7, 11, 17, 19, 23, 26, 28, 29, 31, 34, 40].

Interestingly, Adachi et al. proposed a scoring system for predicting the extent of surgical resection and the neurological outcome for skull base meningiomas[1]. In accordance with their results, we suggested that peri-tumoral edema on preoperative MRI increased the risk of postoperative severe neurological impairment, as previously described[1, 4, 17, 38]. The presence of peri-tumoral edema is possibly related to the pial invasion of the brainstem[17], which may explain difficulties during tumour dissection resulting in vascular injuries and direct injury to the brainstem. However, Adachi et al. analyzed only the postoperative severe neurological impairment but not the non-neurological morbidity, which required the preoperative assessment of the patient general conditions.

Male sex was associated with a higher rate of postoperative non-neurological morbidity, which was consistent with a previous study published by Sekhar et al., where male sex was independently associated with early postoperative KPS deterioration[38]. We suggested that major cardiovascular morbidity increased the risk of postoperative non-neurological morbidity,

which remains a criterion found in all the scores assessing the patient general condition[9, 25, 33].

The ASA scores[33] is a widely used grading system for preoperative health of the surgical patients, which was correlated, for neurosurgical procedures, with non-neurological morbidity[16, 24, 30, 35–37],
postoperative length of stay[18] and survival[41]. The modified Frailty index[25] and the Charlson comorbidity index[9] were also studied in the neurosurgical literature with interesting results, especially concerning their highest accuracy for predicting postoperative complications compared to the ASA score[15, 22, 43]. Youngerman et al., published a large cohort of 9149 patients who underwent oncologic neurosurgery procedures with an estimation of the modified Frailty index[44]. They found that mortality, severe medical complications, prolonged length of stay, and unfavourable discharge increased incrementally with increasing levels of Frailty[44]. However, all these studies were not specifically dedicated to assess skull base and PCM surgery. Here we suggested, for the first time, the importance of assessing the patient general condition in order to improve the overall risk of postoperative complications. The ASA score and the modified Frailty index are significant predictors of post-operative severe non-neurological morbidity and overall severe morbidity and mortality, respectively, and remain convenient scores that can be achieved in daily clinical practice.

In parallel, we suggested that a low neurosurgical experience was associated with a higher risk of overall postoperative morbidity and mortality. The major role of the surgical experience could be related to the assessment of the risk of neurological and non-neurological complications from an experience-based multimodal analysis. The reduction of postoperative severe neurological impairment could be associated with higher skills of the neurosurgeon and the reduction of postoperative non-neurological morbidity is possibly related to the better selection of patients eligible for surgery.

In addition to the PCM’s poor natural history[8, 10, 13, 42] -as relentlessly progressive, with frequent fatal outcomes- and with the natural increased comorbidities score linked to the aging, one may argue that the surgical indication may be proposed early in the patient PCM history. Moreover, the advancement of surgical techniques, the outcomes of PCMs are no longer as pessimistic as has been reported previously[3, 6, 20, 21, 26]. The mortality rate has decreased greatly to 3%; however, surgical morbidity remains high[3, 12, 20, 21, 27]. For selection of the ideal treatment modality in each case, understanding of the natural history and determination of preoperative predictors -such as comorbidity score- are mandatory[39]. A better understanding of the natural history of PCM in the elderly population may help to better wedged the surgical prognosis and the patient information, as comorbidities score appeared to be linked with poor outcomes.

Our results suggest incorporating general condition data into clinical and radiological data in order to improve the accuracy of predicting surgical risk for large and giant PCM.

3. Generalizability

This study captured the real-world practice of large and giant PCM surgery across a tertiary surgical skull base center. The present study controlled for patient-related and methodological biases by selecting a homogeneous population of large and giant petroclival meningioma in adults who all underwent surgical procedure by the same senior neurosurgeon. It represents a large single-center surgical experience.
The present study: 1) could help identify pre-operatively patients with a significant risk of postoperative neurological and/or non-neurological morbidity; 2) highlights the need for a new integrated scoring system to assess pre-operatively the risk of the surgical procedure.

4. Limitations

These findings should be interpreted with caution, given the retrospective and monocentric design, the small numbers of patients, the rarity of the events analyzed, and the lack of an external validation set that limited the generalizability of the results. Since all patients in this study were adults, harbored a large and giant PCM surgically treated by the same senior neurosurgeon, and operated on using the same surgical approach, we thus cannot extend the results to other surgical techniques, to other location, and to patients with a recurrent large and giant PCM. In an attempt to identify predictors available before the surgery, we decided not to integrate the extent of surgical resection. In addition, for large and giant PCM, our attitude is to perform, when possible, a subtotal resection followed by Gamma-Knife radiosurgery, as previously described[6]. Finally, the median modified Frailty Index and Charlson comorbidity index was 0, which suggests that patients who underwent surgery were already preselected by the neurosurgeon or by the referral physician.

Large and giant PCMs remain challenging surgical lesions requiring complex skull base approaches. The risk of postoperative severe neurological and non-neurological morbidity could be assessed pre-operatively following clinical examination, brain MRI analysis and by scores assessing patients' comorbidities. In this study, we propose to add scores assessing the patient general condition in daily practice to improve the selection of patients eligible for surgery. This holistic approach for surgical risk could optimize the management of patients as well as selecting those who can benefit from a surgical resection. Collaborative international multicenter studies are required to confirm these results and allow their implementation in clinical routine.

Declarations

Funding: This research did not receive any specific grant from funding agencies in the public, commercial, or not-for-profit sectors.

Conflicts of interest/Competing interests: None.

Availability of data and material: Anonymized data will be shared by request from any qualified investigator upon reasonable request.

Code availability: not applicable

Ethics approval: This study received required authorizations (IRB#1: 2019/20) of the French College of Neurosurgery (IRB00011687).

Consent to participate: Informed consent was obtained from all patients.
**Consent for publication:** Not applicable

**Authors’ contributions:**

Alexandre Roux: Conception and design of the study, acquisition and analysis of data, Drafting a significant portion of the manuscript and figures.

Lucas Troude: Conception and design of the study, acquisition and analysis of data, Drafting a significant portion of the manuscript.

Guillaume Baucher: acquisition and analysis of data, Drafting a significant portion of the manuscript.

Florian Bernard: acquisition and analysis of data, Drafting a significant portion of the manuscript.

Johan Pallud: Conception and design of the study, Drafting a significant portion of the manuscript.

Pierre-Hugues Roche: Conception and design of the study, acquisition and analysis of data, Drafting a significant portion of the manuscript or figures.

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