The sellar region is one of the most common sites of intracranial disease. A wide range of lesions can affect the sella, including pituitary adenomas, Rathke cleft cysts, craniopharyngiomas, and, less commonly, metastases and lymphomas, among others. Pituitary adenomas, the most common sellar lesion, have an estimated prevalence as high as 20%, based on autopsy and radiological studies, and are often identified incidentally.

Although often asymptomatic, lesions of the sellar region can have devastating effects due to mass effect on surrounding structures, including the pituitary gland and the optic chiasm. Common presenting symptoms in patients with sellar disease include headache, visual disturbances, and hypopituitarism, which can progress to permanent disease or can resolve with appropriate treatment. Patients with sellar lesions requiring surgical intervention most commonly undergo transsphenoidal treatment, in which surgeons use either the operating microscope or endoscope.

Besides optimizing general preoperative cardiovascular conditioning, there are shared and unique modifiable variables of each subset of patients that can be leveraged to optimize surgical outcomes. The perioperative management of patients with sellar lesions is complex, requiring input from a multidisciplinary team of specialists for ongoing management of both endocrinological and neurological issues. Here, the authors reviewed the experience of a single multidisciplinary center over 10 years to identify key postoperative practices that ensure positive outcomes for patients with sellar lesions who undergo transsphenoidal surgery.
ment of patients with sellar lesions is complicated by the delicate nature of the affected structures and the complexity of the underlying disease. Shared variables include operating room ergonomics for the patient and surgeon, preservation of normal pituitary gland, proper reconstruction of the skull base, operative time, postoperative mobility, pain control, monitoring for new hypopituitarism, and prevention of hypotension. Unique modifiable variables are typically dictated by the preoperative diagnosis. The diagnoses that must require specific perioperative consideration are acromegaly, Cushing’s disease, and macro- or giant adenomas with compression of the optic apparatus, along with those patients requiring complex skull base reconstruction."},

"The most common adverse events after transsphenoidal surgery for sellar pathology include hypopituitarism, hypopituitarism, CSF leak, and epistaxis. Patients require careful monitoring for these problems, and institutional protocols in the perioperative period can help limit their occurrence. Many of these patients also require long-term follow-up for management of endocrinological and neurosurgical issues, but few studies have discussed the optimum strategy for postoperative management in this patient population."},

"We aimed to describe the experience of our institution in the pre-, intra-, and postoperative management of a broad range of patients with sellar pathology who undergo transsphenoidal surgery. Evolving during more than 1000 operations at our center, an effective and user-friendly routine has been developed. This routine has resulted in improved outcomes, fewer readmissions, and fewer complications. Patient compliance and satisfaction also continue to improve. An integral part is nursing and patient education and teamwork. The details of our practice are outlined, with commentary, and a review of the literature on select topics was performed.

Methods
We performed a retrospective review of all transsphenoidal operations carried out by the senior author at a single center from April 2008 through November 2018. We included only adult patients and recorded preoperative demographics and lesion characteristics. We also reviewed clinical practices at our center for the care of patients undergoing transsphenoidal surgery. These were divided into the pre-, intra-, and postoperative periods, including long-term follow-up. We also grouped important preoperative characteristics into shared and unique variables in order to underscore the importance of individualized patient management within a general, structured, perioperative institutional protocol. A literature review was performed for each topic to accompany the institutional perspective and results.

Results
Patient Demographics and Characteristics
From April 2008 through November 2018, 928 patients underwent 1023 operations for sellar pathology performed by the senior author (Table 1). Of these operations, 440 (43.0%) were in male patients, and the median age was 48 years (IQR 24.9–32.4). The most common lesion type was pituitary adenoma in 712 operations (70.0%), followed by Rathke cleft cyst in 122 operations (11.9%), craniopharyngioma in 37 operations (3.6%), arachnoid cyst in 17 operations (1.6%), and pituitary tumor apoplexy in 10 operations (1.0%). Other sellar pathologies were present in 125 operations (12.2%).

| Characteristic                  | Total Operations (n = 1023) |
|--------------------------------|-----------------------------|
| **Demographics**               |                             |
| Male sex, n (%)                | 440 (43.0)                  |
| Age in yrs, median (IQR)       | 48 (35, 60)                 |
| BMI, median (IQR)              | 28.3 (24.9, 32.4)           |
| **Lesion type, n (%)**         |                             |
| Adenoma                        | 712 (70.0)                  |
| Nonfunctioning adenoma         | 309 (43.4)                  |
| GH                             | 125 (17.6)                  |
| ACTH                           | 120 (16.9)                  |
| PRL                            | 86 (12.1)                   |
| FSH/LH                         | 50 (7.0)                    |
| Adenoma, unspecified           | 16 (2.2)                    |
| TSH                            | 6 (0.8)                     |
| Rathke cleft cyst              | 122 (11.9)                  |
| Craniopharyngioma              | 37 (3.6)                    |
| Arachnoid cyst                 | 17 (1.6)                    |
| Apoplexy                       | 10 (1.0)                    |
| Other                          | 125 (12.2)                  |
| **Tumor characteristics**      |                             |
| Max diameter in cm, median (IQR) | 1.80 (1.1, 2.5)            |
| Tumor vol in cm², median (IQR)† | 2.28 (0.7, 5.5)            |
| **Hospital characteristics**   |                             |
| Length of stay in days, median (IQR) | 3.0 (2.0, 3.0)     |
| ICU admission, n (%)           | 128 (12.5)                  |

ACTH = adrenocorticotropin hormone; FSH = follicle-stimulating hormone; GH = growth hormone; LH = luteinizing hormone; PRL = prolactin; TSH = thyroid-stimulating hormone.
* Where applicable.
† Computed using ABC/2 method.

Preoperative Management
All patients undergoing transsphenoidal surgery at our multidisciplinary center undergo careful preoperative evaluation with both a neurosurgeon and a neuroendocrinologist. Physical examination focuses on the main presenting symptoms of sellar lesions, including visual fields and symptoms of hypo- and hyperpituitarism. Laboratory evaluation in all patients includes evaluation of all pituitary axes (Table 2), as well as serum sodium, in addition to standard preoperative laboratory examinations. Patients with evidence of low free thyroxine and morning cortisol are started on replacement therapy preoperatively. The biochemical diagnosis of acromegaly and Cushing's
Intraoperative Management

The intraoperative management of patients undergoing transsphenoidal surgery at our institution follows a standard protocol, according to a previously published surgical checklist. Principles of management include strategies to improve patient satisfaction, shorten the hospital stay, and reduce the incidence of postoperative complications. Specifically, patients undergoing transsphenoidal surgery at our center rarely require a Foley catheter or arterial line placement due to short operative times for routine cases (typically 3 hours), and thus intravenous fluids are used as sparingly as possible to avoid postoperative hyponatremia. Patients with acromegaly are at risk for airway complications due to glottal and laryngeal anatomical enlargement, so preoperative discussion with the anesthesia team is imperative. Patients are carefully and ergonomically positioned in order to optimize venous return from the head and neck (thorax is elevated 20°–30°), provide a direct working angle to the sellar pathology, and maintain surgeon orientation with the midline. Image guidance is used in all operations. Abdominal fat grafts are readily available, and careful diagnosis preoperatively is imperative. Depending on the size of the skull base defect, a vascularized nasoseptal flap can be harvested. One dose of antibiotic with primarily gram-positive coverage is given routinely within 60 minutes of incision and continued in the postoperative period only if nasal packing is placed, which occurs rarely. Patients with preoperative fasting cortisol levels below 10 μg/dl are given a steroid bolus preoperatively consisting of 50–100 mg hydrocortisone.

Postoperative Management

In the immediate postoperative period, all patients undergoing transsphenoidal surgery follow standard orders, regardless of underlying pathology (Table 3). Patients undergo assessments of serum sodium and urine-specific gravity every 6 hours and serum cortisol daily while in the hospital. Cortisol is best assessed in the fasting state in the early morning, as this is the typical physiological peak. In our patients with a diagnosis of Cushing’s disease, cortisol is checked every 6 hours, and replacement is only begun after cortisol levels reach subphysiological levels (typically cortisol < 5 μg/dl with symptoms of ad-
TABLE 3. Postoperative orders for patients who have undergone transsphenoidal surgery

| Order                                      | Rationale                                      |
|--------------------------------------------|------------------------------------------------|
| When tolerating clear liquids, discontinue IV fluids | Prevent SIADH/DI; allow for self-regulation     |
| Restrict oral fluid intake to 1 L daily    | Prevent SIADH                                 |
| Visual field checks 4 times daily          | Monitor for new visual field deficits, which may be indicative of postop hemorrhage or may require reop |
| Daily patient weights using the same scale & at the same time | Monitor fluid retention                         |
| Nose & abdominal wounds examined each shift | Monitor for bleeding, surgical site infection  |
| Afrin & saline spray start on postop day 2 | Improve patient satisfaction, wound healing    |

IV = intravenous; SIADH = syndrome of inappropriate antidiuretic hormone.

renal insufficiency or nadir < 2 μg/dl at any point). The more quickly cortisol reaches a subphysiological level, the more likely the patient is to have achieved definitive surgical remission. Upon reaching nadir, hydrocortisone is typically begun at 40 mg in the morning and 20 mg in the evening and monitored closely postoperatively. If patients develop symptoms of hypocortisolism as steroids are begun or tapered, the daily doses are increased, typically not higher than 60 mg in the morning and 30 mg in the evening, and kept at this dose until a taper is tolerated by the patient. For patients with acromegaly, assessment of growth hormone level is an inpatient procedure, with insulin-like growth factor–1 measured at the 3-month follow-up visit. For patients with prolactinomas, serum prolactin is measured daily as an inpatient procedure until it reaches a nadir, and then is rechecked at the 6-week follow-up visit. Fluid intake and output are monitored closely during the postoperative period to assess for fluid retention or diabetes insipidus (DI). The use of desmopressin for treatment of DI should be reserved for patients with a sodium level above 145 mEq/L and inability to provide adequate fluid intake to achieve euvoolemia, or in cases where nocturia is interfering with sleep. Visual field checks are performed at least 3 times daily, both to identify postoperative adverse events, such as hemorrhage, and also to monitor for symptom improvement. Pain is managed without narcotics in the vast majority of cases, which prevents oversedation and promotes early mobility, which is highly encouraged. Patients with significant pain occasionally require intravenous ketorolac in addition to the standard regimen of acetaminophen and nonopioid analgesics. Postoperative NSAIDs (e.g., ibuprofen) are equally effective as opioids in most cases involving minimal trauma to the sinonasal structures. Patients with Cushing’s disease are at increased risk of thromboembolic events and thus we typically start low-dose (81 mg) aspirin administration for these patients on the 1st postoperative day. Aspirin, which has significant antiplatelet effects, has not resulted in a higher incidence of postoperative bleeding in our experience. Thus, NSAID use for pain starting on the 1st postoperative day is typically well tolerated. Because of the increased risk of postoperative bleeding in patients who receive NSAIDs, we use intravenous ketorolac rarely, and never in combination with aspirin. We are also careful to avoid NSAID use in patients who are prone to bleeding, either based on preoperative workup or on intraoperative assessment of bleeding propensity (even in the context of a normal coagulation and platelet profile). In patients without Cushing’s disease, venous thromboembolism prophylaxis is achieved with pneumatic boots and early ambulation, and therefore no subcutaneous heparin is routinely used. Only rarely will postoperative patients require admission to the neuroscience ICU, but having this level of care available is imperative should any patients experience significant intraoperative complications. In our practice, indications for admission to the neuroscience ICU include: 1) need for close postoperative monitoring of vision, 2) evidence of a particularly hemorrhagic lesion at risk for postoperative hematoma, and 3) patients with severe medical comorbidities. Admission to the ICU typically only occurs for patients with large lesions with suprasellar extension.

After a typical hospitalization of 2–3 days, patients with sellar pathology are discharged with consistent postdischarge instructions (Table 4). During the 1st week postoperatively, all patients are instructed to limit daily water intake to 1 L, and all patients are discharged with a standard 1-L hospital water pitcher. In patients with Cushing’s disease who enter adrenal insufficiency postoperatively, hyponatremia can occur due to release of antiuretic hormone as a vasopressor. Fluid restriction helps to limit the occurrence of hyponatremia in this setting, and careful steroid repletion, as above, precludes development of complications of adrenal insufficiency. During the first 5 weeks postoperatively, patients using obstructive sleep apnea, particularly common among patients with acromegaly, are instructed to refrain from using home CPAP (continuous positive airway pressure) machines until the nasal mucosa heals. Patients return to clinic 1 week postoperatively for evaluation, including repeat measurement of serum sodium and serum cortisol, and physical examination to assess for possible postoperative complications, including CSF leak and meningitis. Patients then return 6 weeks postoperatively for another repeat pituitary and endocrine assessment, often accompanied by an appointment with otolaryngology for a reevaluation of healing in the posterior nasal cavity. Patients then return for a 3-month follow-up visit, which includes definitive postoperative MRI. Among this cohort of patients, the overall readmission rate was 9.0% (92/1023). Prior to 2015, when mandatory postoperative fluid restriction was implemented, the rate was 10.4% (73/702) versus 5.9% after (19/321). Assuming there is no residual lesion and there have been no complications, typical follow-up in our practice entails annual follow-up visits with brain MRI for several years when indicated.
Patients who undergo transsphenoidal surgery for sellar disease require careful perioperative care due to the delicate nature of pituitary function and the risks of complications during the postoperative period. In particular, the convergence of numerous disciplines (neurosurgery, endocrinology, anesthesiology, neuroophthalmology, and otolaryngology) in the care of these patients necessitates effective communication about diagnostic evaluations and perioperative management. Our practice incorporates several evidence-based recommendations for the care of patients during the period of transsphenoidal surgery, but there is a lack of high-quality, randomized evidence because it is impossible to perform randomized controlled trials for each of the small variations in clinical practice that may affect patient outcomes. In the absence of this evidence, and in light of the implausibility of randomized studies clarifying the effectiveness of certain practices, we reviewed and presented here our own institution’s practices as a guide by which perioperative care of the patient with sellar pathology can be approached.

Given the complexity of lesions in the sella, multidisciplinary care, preferably at a tertiary or quaternary care center, is preferred.10,13,18 These centers allow for coordination of care from multiple specialists, with access to high-quality imaging and treatment technologies and a high level of neurocritical care, if necessary. Care for patients at our institution occurs through a multidisciplinary pituitary/neuroendocrine center, which incorporates care by neurosurgeons, neuroendocrinologists, neuroradiologists, otolaryngologists, neuroophthalmologists, and neuropathologists. When patients arrive for consultation, they are evaluated by both a neurosurgeon and a neuroendocrinologist, who each discuss with the patient the risks and benefits of surgery and other possible options for treatment. These relationships are carried through follow-up, with input from neuropathologists in the postoperative period and neuroradiologists at each radiological assessment. The benefit of such an arrangement is typically realized by improved accuracy of diagnosis in patients with acromegaly and Cushing’s disease, which requires the interpretation of numerous and often specialized diagnostic tests (oral glucose tolerance test, 24-hour urine free cortisol, late-night salivary cortisol, dexamethasone suppression test, inferior petrosal sinus sampling [IPSS]). For example, we perform IPSS only after the criteria for the diagnosis of Cushing’s disease are established according to current guidelines in order to distinguish a central pituitary source from an ectopic source.14,36 Interpretation of IPSS results can be challenging, and the inappropriate use of IPSS may result in false-positive results. Multidisciplinary follow-up is also essential for monitoring and diagnosing disease recurrence.

Overall, the care of patients who are undergoing transsphenoidal surgery should be directed according to the expected timeline for these patients (Table 5). Preoperative-ly, patients require careful evaluation to assess pituitary function and to plan operative strategies. In our practice, this includes multidisciplinary evaluation in the outpatient setting, with detailed laboratory and neuroradiological assessment. Often, additional consultations are needed to ensure appropriate management, and many of our patients are assessed preoperatively by otolaryngologists in the case of complex nasal and sinus anatomy. Intraoperative management is also carefully standardized according to a previously published surgical checklist.23 All operations incorporate intraoperative image guidance, which helps to ensure accurate and safe surgery.

During the immediate postoperative period, patients who have undergone transsphenoidal surgery are at risk of several complications, irrespective of their underlying sellar pathology. These include hyponatremia, hypopituitarism, and CSF leak, among others.11,31 Rates of hyponatremia reported in the literature vary widely, ranging from 3.6% to 19.8%.10 We recently reported an overall rate of readmission for hyponatremia after transsphenoidal surgery of 2.5%, with no readmissions in 203 consecutive operations after institution of mandatory fluid restriction of 1 L per day in all patients for 1 week postoperatively.6 As a result of this study, we now recommend mandatory 1 L per day fluid restriction for all posttranssphenoidal patients until the 1-week postoperative appointment, at which point the restriction is liberalized if serum sodium is normal.

Hypopituitarism after transsphenoidal surgery is more frequent in patients with large or invasive lesions and is common in the postoperative period for patients with functioning adenomas, who may experience substantial shifts in levels of pituitary hormones.13,18 Replacement of pituitary hormones is a crucial aspect of posttranssphenoidal surgery care, and consideration of replacement must begin even before the operation, by discussing with

| Order | Rationale |
|-------|-----------|
| Restrict oral fluid intake to 1 L daily until 1-wk postop appointment | Prevention of SIADH |
| Only drink when thirsty | Prevention of dysnatremia |
| Saline nasal spray at least twice per day | Improve patient satisfaction, wound healing |
| No bending or lifting >30 lbs for 4–6 wks | Prevention of CSF leak |
| Report any of the following symptoms: nausea, vomiting, fatigue, severe headache, stiff neck, fever | May indicate meningitis, hypocortisolemia, hyponatremia |
| Excessive thirst & frequent urination | May indicate dysnatremia |
| Clear nasal drainage with headache | May indicate CSF leak |

Discussion

Patients who undergo transsphenoidal surgery for sellar disease require careful perioperative care due to the delicate nature of pituitary function and the risks of complications during the postoperative period. In particular, the convergence of numerous disciplines (neurosurgery, endocrinology, anesthesiology, neuroophthalmology, and otolaryngology) in the care of these patients necessitates effective communication about diagnostic evaluations and perioperative management. Our practice incorporates several evidence-based recommendations for the care of patients during the period of transsphenoidal surgery, but there is a lack of high-quality, randomized evidence because it is impossible to perform randomized controlled trials for each of the small variations in clinical practice that may affect patient outcomes. In the absence of this evidence, and in light of the implausibility of randomized studies clarifying the effectiveness of certain practices, we reviewed and presented here our own institution’s practices as a guide by which perioperative care of the patient with sellar pathology can be approached.

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TABLE 5. Postoperative management of patients undergoing transsphenoidal surgery, including typical postoperative follow-up schedule, diagnostic evaluation, and monitoring of adverse events

| Preoperative | Inpatient | POD7 | POW6 | POM3 | POY1 |
|--------------|-----------|------|------|------|------|
| **Diagnostic** |            |      |      |      |      |
| Tests        | Physical exam; MRI brain/pituitary; CBC/BMP/PTI/INR; Serum ACTH, cortisol, PRL, GH, IGF-1, FSH, LH, testosterone/estrogen, TSH, FT4 | Serum sodium & cortisol; Visual field checks 4 times daily | Serum sodium; Serum cortisol; Physical exam | Full pituitary endocrine panel; Physical exam | MRI brain/pituitary; Pituitary hormones for functioning lesions; Physical exam | MRI brain/pituitary; Pituitary hormones for functioning lesions; Physical exam; Optic fields when applicable |
| Management   |            |      |      |      |      |
| Considerations | Consider additional consults as needed; Assess preop pituitary function | AMS, pain, visual field changes: consider hemorrhage or meningitis | Discuss preliminary pathology | Assess for return of pituitary function in patients w/ preop hypopituitarism | Assess response to surgery | Monitor for lesion recurrence; Continue annual MRI follow-ups, for at least 5 years of stability |

AMS = altered mental status; BMP = basic metabolic panel; CBC = complete blood count; FT4 = free T4; GTR = gross-total resection; INR = international normalized ratio; POD = postoperative day; POM = postoperative month; POW = postoperative week; POY = postoperative year; PT = prothrombin time.

patients the possibility of requiring lifelong replacement therapy (Table 6).31 Replacement of physiological cortisol can be achieved with either hydrocortisone (twice daily) or prednisone (once or twice daily), usually with lower doses in the evening to improve sleep. It is imperative to instruct patients to report symptoms of hypocortisolemia, including fatigue, weight loss, nausea, and syncope, and to increase intake during periods of illness. Thyroid replacement is achieved with daily levothyroxine dosing, and replacement of sex hormones can be achieved with testosterone in men (gel, patches, or depot injections are appropriate, depending on patient preference), versus oral contraceptive agents or conjugated estrogen tablets in women. We previously reported a rate of new-onset hormone deficiency after transsphenoidal surgery for pituitary macroadenoma of 6.3%, with patients with recurrent lesions and apoplexy at higher risk.24 Diabetes insipidus (DI), in this context often the result of abnormally low levels of antidiuretic hormone resulting in profuse, dilute urine, can be carefully treated with desmopressin if the serum sodium rises above 145 mEq/L. Although often transient, DI after transsphenoidal surgery can infrequently be a permanent issue. We recently reported an overall rate of DI after transsphenoidal surgery of 14.7% and a rate of permanent DI of 4.6%.7

Postoperative CSF leak after transsphenoidal surgery can occasionally be treated conservatively with lumbar drainage, but it often requires reoperation for repair of the skull base defect.17,38 Previously identified predictors of CSF leak post–transsphenoidal surgery include body mass index, identification of intraoperative leak, and pre-

TABLE 6. Recommended hormone replacement strategies for patients requiring pituitary hormone replacement after transsphenoidal surgery

| Axis Deficiency | Replacement Medication | Initial Replacement Dose* | Notes |
|-----------------|------------------------|---------------------------|-------|
| Cortisol        | Hydrocortisone or prednisone | Hydrocortisone: 20 mg qAM, 10 mg qPM; prednisone: 5 mg qAM, 2.5 mg qPM | Cortisol best measured in fasting state, early morning |
| Thyroid         | Levothyroxine          | 75 μg qD                  |       |
| GH              | Somatropin             | Approximately 0.2 mg SC qD, depending on injection type | IGF-1 is more appropriate than GH for monitoring of GH function due to lower daily variability |
| Testosterone    | Testosterone replacement | Depends on replacement type (gel vs injection vs patch, etc.) |       |
| Estrogen        | OCPs or conjugated estrogens | Should be tailored based on replacement type & menopausal status |       |
| ADH             | Desmopressin           | 0.05 mg qHS, may increase to 0.1 mg qHS | Careful replacement & monitoring for transient vs permanent DI |

ADH = antidiuretic hormone; OCP = oral contraceptive pill; qAM = every morning; qD = every day; qHS = every day at bedtime; qPM = every evening; SC = subcutaneously.
* These represent initial doses for typical patients but should be tailored to patient characteristics, including weight and signs and symptoms of excess or deficiency.
operative hydrocephalus, with rates ranging from 2.6% to 11%. Our practice commonly incorporates abdominal fat grafting as a method of preventing CSF leak, and collaboration with colleagues in otolaryngology can help ensure adequate repair of the skull base defect using nasal septal flaps if needed. Lumbar drainage is rarely used due to the risk of infection with prolonged intrathecal catheter use, as well as the efficacy of bolstering the initial reconstruction in the operating room. Recent evidence suggests that use of perioperative lumbar drain is effective in reducing the rate of postoperative CSF leak in patients undergoing endoscopic endonasal surgery. Upon discharge, patients must be instructed to return to the clinic at any sign of continuous clear drainage from the nose.

Long-term follow-up in these patients is imperative to assess for recurrence of the initially resected lesion. Our practice recommends that all patients return to the clinic at 3 months and 1 year postoperatively, at a minimum, with most patients returning annually for follow-up subsequently. At each of these appointments, patients are reevaluated by both a neurosurgeon and a neuroendocrinologist, and the patients undergo MRI of the brain and pituitary. At the 3-month postoperative visit, this MRI scan serves to evaluate the effect of surgery and to assess for the extent of resection. At the ensuing annual visits, MRI scans help the clinician assess for lesion recurrence. Signs of recurrence of the initial lesion that would merit repeat imaging outside of this routine include worsening or returning symptoms, such as headache, visual field deficits, or endocrinological symptoms, which are often reported by the patient or detected on physical examination. In the case of functioning tumors, laboratory evaluation can occasionally identify recurrence prior to radiological evaluation. In some cases, patients who have undergone transsphenoidal surgery may require adjuvant treatment for residual or recurrent lesions. These treatments may include radiation therapy or medical therapy, as appropriate, but a complete review of these treatments is beyond the scope of this manuscript.

In this paper, we have reviewed the practices for perioperative care for patients who have undergone transsphenoidal surgery at our institution. This experience incorporates clinical data from over 1000 operations, and aims to serve as an overview for appropriate management of these patients, who often require complicated perioperative care. Several limitations are present in this study. First, this is a retrospective case series and is limited by this aspect of our data. To alleviate any bias, we presented here a complete record of consecutive operations by the senior author. It is also important to note that the outcomes reported at our institution may not necessarily be the result of the postoperative management strategies in place at our institution but rather may be confounded by the surgical experience of the senior author, or by other factors such as institution volume or experience. Minor variations in some clinical practices from center to center may not have substantial effects on clinical outcomes, and the practices presented here are meant to serve as a guide by which postoperative care after transsphenoidal operations can be approached. Given the diversity of lesions that can affect this region, perioperative care should of course be tailored to the specific patient according to medical history, lesion characteristics, functional status, and endocrine function.

Conclusions

Patients with sellar lesions who undergo transsphenoidal surgery require complex, multidisciplinary postoperative care to monitor for common adverse events and improve outcomes, but there is a dearth of high-quality evidence guiding most perioperative practices. Here, we reviewed practices at our institution across more than 1000 transsphenoidal operations that may help ensure successful patient outcomes.

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References

1. Abou-Al-Shaar H, Zaidi HA, Cote DJ, Laws ER Jr: Bolstering the nasoseptal flap using sphenoid sinus fat packing: a technical case report. World Neurosurg 99:813.e1–813.e5, 2017
2. Almutairi RD, Muskens IS, Cote DJ, Dijkman MD, Kavouridis VK, Crocker E, et al: Gross total resection of pituitary adenomas after endoscopic vs. microscopic transsphenoidal surgery: a meta-analysis. Acta Neurochir (Wien) 160:1005–1021, 2018
3. Alzhrani G, Sivakumar W, Park MS, Taussky P, Couldwell WT: Delayed complications after transsphenoidal surgery for pituitary adenomas. World Neurosurg 109:233–241, 2018
4. Barbot M, Daidone V, Zilio M, Albiger N, Mazzai L, Sartori MT, et al: Perioperative thromboprophylaxis in Cushing’s disease: What we did and what we are doing? Pituitary 18:487–493, 2015
5. Brennan MD, Jackson IT, Keller EE, Laws ER Jr, Sather AH: Multidisciplinary management of acromegaly and its deformities. JAMA 253:682–683, 1985
6. Burke WT, Cote DJ, Iuliano SI, Zaidi HA, Laws ER: A practical method for prevention of readmission for symptomatic hyponatremia following transsphenoidal surgery. Pituitary 21:25–31, 2018
7. Burke WT, Cote DJ, Penn DL, Iuliano SL, McMullen K, Laws ER: Diabetes insipidus after endoscopic transsphenoidal surgery. Neurosurgery [in press], 2020
8. Burke WT, Penn DL, Castlen JP, Donoho DA, Repetti CS, Iuliano S, et al: Prolactinomas and nonfunctioning adenomas: preoperative diagnosis of tumor type using serum prolactin and tumor size. J Neurosurg [epub ahead of print June 14, 2019, DOI: 10.3171/2019.3.JNS19121]
9. Casanueva FF, Barkan AL, Buchfelder M, Klibanski A, Laws ER, Loefller JS, et al: Criteria for the definition of Pituitary Tumor Centers of Excellence (PTCOE): a Pituitary Society statement. Pituitary 20:489–498, 2017
10. Cote DJ, Alzarea A, Acosta MA, Hulou MM, Huang KT, Almutairi H, et al: Predictors and rates of delayed symptomatic hyponatremia after transsphenoidal surgery: a systematic review [corrected]. World Neurosurg 88:1–6, 2016
11. Cote DJ, Daseenbrock HH, Muskens IS, Broekman MLD, Zaidi HA, Dunn IF, et al: Readmission and other adverse events after transsphenoidal surgery: prevalence, timing, and predictive factors. J Am Coll Surg 224:971–979, 2017
12. Cote DJ, Wiemann R, Smith TR, Dunn IF, Al-Mefty O, Laws ER: The expanding spectrum of disease treated by the transnasal, transsphenoidal microscopic and endoscopic anterior skull base approach: a single-center experience 2008–2015. World Neurosurg 84:899–905, 2015
51. Zhang Y, Guo X, Pei L, Zhang Z, Tan G, Xing B: High levels of IGF-1 predict difficult intubation of patients with acromegaly. *Endocrine* 57:326–334, 2017
52. Ziu M, Dunn IF, Hess C, Fleseriu M, Bodach ME, Tumialan LM, et al: Congress of Neurological Surgeons systematic review and evidence-based guideline on posttreatment follow-up evaluation of patients with nonfunctioning pituitary adenomas. *Neurosurgery* 79:E541–E543, 2016
53. Zwagerman NT, Wang EW, Shin SS, Chang YF, Fernandez-Miranda JC, Snyderman CH, et al: Does lumbar drainage reduce postoperative cerebrospinal fluid leak after endoscopic endonasal skull base surgery? A prospective, randomized controlled trial. *J Neurosurg* 131:1172–1178, 2019

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