Transsphenoidal surgery in Cushing disease: The challenging microadenoma (Local experience)

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Abstract: Background: Cushing disease is uncommon challenging disease. The adenomas are usually small in size in most case making the disease diagnosis and management is sometimes difficult. In some cases, the tumor cannot be identified on imaging studies and in many cases the adenoma is eccentric in location adding more difficulties to the trans-sphenoid approach for excision of such tricky tumors. Object: to evaluate the challenge of diagnosis and trans-sphenoid surgery for ACTH pituitary microadenomas. Methods: 21 patients with ACTH secreting adenomas were involved in this retrospective study. All cases were operated on via trans-sphenoid approach during the period from January 2009 till October 2015. All of them were microadenomas. All patients had obvious Cushing manifestations. In all cases magnetic resonance (MR) imaging and CT paranasal sinuses were performed. In this study: cases with macroadenomas and recurrent cases after previous trans-sphenoid approach or gamma knife radiosurgery were excluded. Results: Among the 21 patients; 14 were female and 7 were male with age ranging from 21 to 44 years. The imaging study was positive for microadenoma in 18 patients while in 3 cases; no tumor was seen on the MRI imaging. All were operated via the trans-sphenoid approach (12 patients via the microscopic technique and other 9 patients via the endoscopic technique). The adenoma was successfully removed in 15 out of the 18 patients with evident adenoma on pre-operative MRI imaging while partial hypophysectomy were done in 5 cases without an evident adenoma on pre-operative MRI study or difficult adenoma identification. In one patient; the procedure was aborted because of sphenoid sinus abnormalities and the tumor was successfully removed via trans-cranial approach. Disease remission was achieved in 17 patients. In 4 patients; complete remission was not achieved and gamma knife radiosurgery was done after surgery. Disease progression occurred in two patients with initial remission and was treated gamma knife radiosurgery. 13 patients had an ACTH deficiency after surgery while other hormonal replacement therapy is needed in only 5 cases. CSF rhinorrhea occurred in 2 patients and was managed with transient lumbar drain without any subsequent morbidities. Transient diabetes insipidus occurred in 6
cases and was treated conservatively. Conclusions: The transsphenoidal approach ACTH secreting pituitary microadenomas is considered safe and effective for achieving disease remission despite the challenge of diagnosis and intra-operative tumor identification.

Key words: Adrenocorticotrophic hormone, Magnetic resonance, Computed tomography, Cerebrospinal fluid

Introduction

Surgery is considered the golden standard treatment for ACTH secreting adenomas causing Cushing disease. These benign adenomas are mostly microadenoma and easily achievable via the classic trans-sphenoid approach. However, there is still considerable challenges in the disease diagnosis management. In many cases (around 40%), the tumor cannot be identified on the pre-operative imaging studies (post-contrast magnetic resonance imaging) and intra-operative identification of the adenoma is dependent on the experience of the surgeon. Even if the inferior petrosal sinus sampling is feasible; its accuracy in identification the site of the adenoma is questionable. Another challenge is eccentric location of many adenomas is eccentric and despite its small size; it can be very close the carotid artery adding more difficulties to the trans-sphenoid approach for excision of such tricky tumors. The goal of surgery shouldn’t only to remove the tumor but also to preserve the pituitary function to prevent post-operative hypopituitarism necessitating long term hormonal replacement therapy. The aim of this study is to evaluate the challenge of diagnosis and potential difficulties and outcome of the trans-sphenoid approach for ACTH secreting pituitary adenomas.

Patients and methods

21 patients with ACTH secreting adenomas were involved in this retrospective study. All the patient had obvious clinical manifestations of Cushing disease. All these cases were operated up on in the neurosurgery department via the trans-sphenoidal approach during the period from January 2009 to October 2015.

We evaluated all the medical data and imaging studies to assess all the potential difficulties in the disease diagnosis and the expected surgical challenge and this data was correlated with the operative finding and post-operative outcome particularly the rate of disease remission. Data analysis included patient age, gender, clinical manifestations, pre-operative and initial and late post-operative hormonal levels including serum cortisol and ACTH levels and pre-operative and post-operative imaging studies. In all cases, magnetic resonance (MR) imaging of the sellar region and CT paranasal sinuses were performed. In this study; cases with macroadenomas (greater than one CM in the size) and recurrent cases after previous trans-sphenoid approach or gamma knife radiosurgery were excluded. Outcome assessment was evaluated by improvement of the Cushing manifestation, hormonal assessment to ensure biochemical remission and post-operative contrast enhancing magnetic resonance imaging. Post-operative cortisol level evaluation was routinely done on
the first day after the surgery and then after one, three and six months thereafter. Then hormonal evaluation was done annually for assessment of any relapse. Remission was defined by either normalization of the serum cortisol level or those with low serum cortisol level (≤3 mg/dl) necessitating hormonal replacement therapy. Those patients with remaining elevated serum cortisol level were considered treatment failure. Post-operative imaging (MRI) was done initially after three months to ensure any residual tumor and then annually for five years to detect any recurrence. Post-operative morbidities (transient or permanent) were also analyzed.

Results

Among the 21 patients; 14 were females and 7 were males with age ranging from 21 to 44 years (mean 33.6). All patients had the clinical manifestation of Cushing disease (Table 1). Headache was manifest in eleven patients, 2 patients had ocular palsy concomitant with his Cushing manifestation (Figure 1), and in one patient; impotence was his early presentation for Cushing disease. In three patients; uncontrolled diabetes was the triggering for the diagnosis of the disease and in one patient refractory hypertension was the initial presentation. Depending on the pre-operative post-contrast magnetic resonance study; we have 2 categories of patients; those with positive microadenoma (18 patients) and those with negative finding (3 patients). All patients were operated up on via the trans-sphenoid approach (12 patients via the microscopic endonasal technique and other 9 patients via the endoscopic endonasal technique) (Table 2). The technique was exactly like what was widely described in the literature but we have faced with difficulties from mucosal changes rendering it more friable and bloody and difficulties from difficult control of the blood pressure. In one patient with conical shaped sinus; drilling of the bony sinus was performed under image guidance but bleeding was evident and the approach was aborted and the adenoma was removed three weeks later via the trans-cranial approach.

TABLE I

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<th>Patients characteristics</th>
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<td><strong>Patient sex</strong></td>
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<td><strong>Follow up period</strong></td>
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TABLE II

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<th>Imaging characteristic, Treatment plan and outcome of surgery</th>
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| **Pre-operative MRI of the sella** | Positive for adenoma: 18 patients  
Negative for adenoma: 3 patients |
| **Technique of surgery** | Microscopic trans-sphenoidal approach: 12 patients  
Endoscopic trans-sphenoidal approach: 9 patients |
| **Intra-operative finding** | Adenoma removal: 15 via the trans-sphenoidal approach  
One via additional trans-cranial approach after aborted Trans-sphenoidal approach  
Partial hypophysectomy: 5 patients |
| **Pathological confirmation of adenoma** | Positive: 18 patients  
Negative: 3 patients |
| **Post-operative treatment** | Gamma knife radiosurgery: 6 patients  
4 patients: treatment failure  
2 patients: disease progression after initial remission  
Medical treatment:  
Cortisol replacement therapy: 13 patients  
Other hormonal replacement therapy: 5 patients |
| **Surgery related morbidities** | CSF rhinorrhea: 3 patients (Transient)  
Transient diabetes insipidus: 6 patients |

TABLE III

**Disease remission (Biochemical markers)**

- **Disease remission:** 17 patients (80.9%)
  - Immediate after surgery: 8 patients
  - Within 6 months from surgery: 7 patients
  - Up to one year from surgery: 2 patients
  - Failure to achieve remission: 4 patients (19.1%)
  - Disease progression after initial remission: 2 patients (9.5%)

**Remission in relation to pathological confirmation:**

- Positive adenoma (15 patients): 14 patients achieved remission
- Mixed adenoma and hyperplasia (3 patients): two achieved remission
- Negative specimen (3 patients): one achieved remission

**Post-operative cortisol level:**

- Hypocortisolemia: 13 patients
- Eucortisolemia: 4 patients
- Persistent hypercortisolemia: 4 patients
The adenoma was successfully removed in 15 out of the 18 patients with evident adenoma on pre-operative post-contrast magnetic resonance study (Figure 2) while partial hypophysectomy were done in 5 cases without an evident adenoma on pre-operative MRI study or difficult adenoma identification. Pathological confirmation for the adenoma was achieved in 15 patients, mixed adenoma and hyperplasia in 3 patients while specimen was negative in three cases. Follow up period ranged from 9 months up to 78 months. Disease remission (Table 3) was achieved in 17 patients (80.9%). In 4 patients (19.1%); complete remission was not achieved and gamma knife radiosurgery was done after surgery. Disease progression occurred in two patients (9.5%) after initial remission and was treated with gamma knife radiosurgery.

No approach related mortalities occurred. 13 patients had post-operative cortisol deficiency and was maintained on replacement therapy. Other hormonal replacement therapy was needed in only 5 cases. CSF rhinorrhea occurred in 2 patients and was managed with transient lumbar drain without any subsequent morbidities. Also, the case that was operated via the trans-cranial approach after failed trans-sphenoid surgery developed CSF rhinorrhea after 2nd surgery and was managed with transient lumbar drain. Transient diabetes insipidus occurred in 6 cases and was treated conservatively.

Discussion

Cushing disease is uncommon challenging disease. It occurs due to ACTH secreting adenomas which in majority of cases are microadenomas but many reports showed that macroadenomas can cause up to 13% of Cushing disease patients.⁶⁻⁸⁻¹¹⁻¹³⁻¹⁴ However, we decided to concentrate in this reports on challenge and outcome of surgery for de novo cases of ACTH secreting adenomas rather than macroadenomas or previously treated cases to stress on the challenge of treatment and outcome of such small lesion not to involve other potential challenge rather than the adenoma itself.¹⁻³⁻⁶⁻¹¹⁻¹²

The female predominance and age distribution in our study matches with the finding of other reports.¹⁻²⁻⁵⁻⁷⁻⁹ The challenge of this disease starts with disease diagnosis. Several factors are important in the endeavor to obtain satisfactory results. Preoperative detection of the ACTH secreting adenoma is crucial for the surgical outcome, achievement of cure and prevention of surgically induced hypopituitarism.⁸⁻¹⁰⁻¹⁵⁻¹⁶⁻¹⁷ Our 1st problem was with three cases (14.3%) with all manifestation of Cushing disease with exclusion with all potential differential diagnosis with negative finding of the post-contrast magnetic resonance of the sella. Our incidence of negative imaging is lower than what reported in other studies which may reach up to 40% of Cushing microadenomas which mostly due to lack of referral of MRI negative patients to our neurosurgical service.⁶⁻⁷⁻¹¹⁻¹³⁻¹⁷ Although inferior petrosal sampling played a role in identification the site of adenoma however we don’t have such facility and in many reports, its accuracy is up to 60% in identification the site of adenoma.³⁻⁴⁻⁹⁻¹⁰⁻¹¹ In such cases our decision was to explore the pituitary gland for the adenoma or do partial hypophysectomy.
The trans-sphenoid technique is exactly like what have been reported in different series and there was no difference between the microscopic versus the endoscopic approach for addressing such tiny adenomas. However, in our cases there was a potential challenge in the approach specific to this problem in the form of friable bloody nasal mucosa adding some difficulties to the nasal step of the approach. Another problem was the difficult control of the blood pressure during the procedure that increased the bleeding during the nasal phase of the approach. In one case; the sphenoid sinus was conical shaped type and under C arm guidance we tried to drill the bony sinus to reach the sellar floor but due to excessive bleeding and lack of navigation system under our facilities; we decided to abort the procedure and to come back from above via trans-cranial approach and the adenoma was removed. In five patients, no adenoma could be identified during pituitary gland exploration and partial hypophysectomy was performed. Many reports addressed the issue of negative exploration of pituitary gland for adenoma and advocated hemi-hypophysectomy of pituitary gland particularly on the side of positive inferior petrosal sampling with reported cases of disease remission even with absence of any pathological confirmation.

The issue of Cushing disease remission both clinically and by biological confirmation has been addressed in many reports with overall remission ranging from 59 to 98% in different clinical series. Variable factors predict the potential remission rate; the size of adenoma, whether its de novo or recurrent, presence of cavernous sinus invasion and presence or absence of pathological confirmation. Remission rate is higher in newly diagnosed microadenomas without cavernous sinus invasion and slightly higher in pathologically proven adenoma rather than those with inconclusive pathological smear. We have achieved remission in 80.9% of cases but all of them is microadenomas. In one case; the adenoma was located laterally in close contact with internal carotid artery and despite pathological confirmation of being adenoma; remission wasn’t achieved and was sent for gamma knife (Figure 1). Among our three cases with negative pathological confirmation; only one case (33.3%) showed disease remission and other two cases were sent for gamma knife.

Figure 1 - MRI images of case with Cushing disease with uncontrolled diabetes presented with ptosis due to 3rd nerve palsy
In another case that post-operative remission wasn’t achieved; the pathological smear revealed mixed adenoma and hyperplasia and the patient was also referred for gamma knife treatment.

Recurrence of disease is not uncommon and it can happen over many years after initial remission hence long term follow up is necessary for those patients. Overall recurrence rate ranged from 5 % to 60 % in many case series and it can occur up to 10 years from remission.\textsuperscript{9,10,11,12,13,14} Recurrence is more expected in cases of macroadenomas, cavernous sinus invasion and with negative pathological confirmations or when the hyperplasia rather than adenoma is the cause of the disease.\textsuperscript{2,3,4,5,6,15} In our cases; we have two cases (9.5%) of disease progression after initial remission. The pathology in one of them was mixed adenoma and hyperplasia and was advised to have gamma knife treatment. The 2nd case showed recurrent adenoma on post-operative magnetic resonance study but the patient preferred to do gamma knife radiosurgery rather than the advised repeated trans-sphenoid approach. Complication after trans-sphenoid surgery for Cushing disease is relatively high compared to other pituitary adenomas and it may reach up to 53% in some reports.\textsuperscript{4,9,11,15} Also, mortality rate has been reported up to 8.4%. Complications could be minor and transient or it may be permeant or even life threatening. Reported morbidities
included Epistaxis, pituitary hypofunction, CSF leakage, pneuomocephalus, meningitis, visual impairment, ocular motility dysfunction, and occasional deep venous thrombosis and pulmonary embolism. Some reports claimed that morbidity was less with endoscopic procedures rather than with microscopic technique but in our study; there was no much difference. Our reported morbidities included transient CSF rhinorrhea (2 patients), transient diabetes insipidus (6 cases) and hypopituitarism (5 patients). Another case that was operated via the trans-cranial approach after failed trans-sphenoid surgery developed CSF rhinorrhea after 2nd surgery and was managed with transient lumbar drain. 13 patients had post-operative cortisol deficiency and was maintained on replacement therapy.

Conclusions

The transsphenoidal approach for ACTH secreting pituitary microadenomas is considered safe and effective for achieving disease remission despite the challenge of diagnosis and intra-operative tumor identification. Both microscopic and endoscopic techniques are of equal safety and efficacy. Both pre-operative tumor identification on magnetic resonance imaging of the sella and positive pathological confirmation of removed tissue play a role in achievement of remission. Disease progression may occur after initial remission hence long term follow up of those patients is necessary.

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References

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