Resection of Fibrous Dysplasia of the Sphenoid Bone and a Concomitant Calcified Pituitary Adenoma via an Endoscopic Endonasal Transsphenoidal Approach: Case Report

Benedict Tan, BS¹, Sanjay Yadla, MD², Peter G. Campbell, MD²
Marc R. Rosen, MD², James J. Evans, MD²
¹Jefferson Medical College, Thomas Jefferson University, Philadelphia, PA 19107 USA
²Department of Neurological Surgery, Thomas Jefferson University, Philadelphia, PA 19107 USA

Abstract

Background: Although pituitary adenomas have been described in association with polyostotic fibrous dysplasia in McCune-Albright Syndrome, no such relationship has been described with monostotic fibrous dysplasia. The authors describe a case of monostotic fibrous dysplasia of the sphenoid bone and concomitant pituitary adenoma in a 25-year-old male. To the authors' knowledge this is the first such case reported in the literature. Clinical presentation, pathology, and surgical approach are described in detail.

Case Description: A 25-year-old male initially presented with headaches, gynecomastia, and galactorrhea. Magnetic Resonance Imaging (MRI) revealed a sellar/suprasellar mass possibly consistent with a calcified pituitary adenoma or craniopharyngioma and a cranial base lesion obstructing the sphenoid sinus consistent with fibrous dysplasia. Both lesions were accessible via an endoscopic transnasal approach. The patient underwent resection of the affected sphenoid bone which allowed appropriate exposure for subsequent resection of the pituitary lesion without major complication. Pathology and immunohistochemical studies confirmed the diagnoses and postoperative imaging revealed gross total resection of the adenoma.

Conclusions: The authors describe a case of monostotic fibrous dysplasia of the sphenoid bone and pituitary adenoma. Resection of such dual lesions can be both safe and efficacious via an endoscopic endonasal transsphenoidal approach.

Key Words: Fibrous Dysplasia, Calcified Pituitary Adenoma, Pituitary Stone, Transsphenoidal

Abbreviations List: MRI: Magnetic Resonance Imaging, CT: Computerized Tomography, CAM: Cell Adhesion Molecule, EMA: Ethidium Moazeide, TSH: Thyroid Stimulating Hormone

Case Report

History and Physical

A 25-year-old African-American male presented with complaints of headaches and galactorrhea. There was no history of long bone fractures. On exam, the patient was noted to have intermittent esotropia of the left eye and gynecomastia. No café au lait spots or other skin lesions were noted. Humphrey visual field testing was normal. Endocrine evaluation revealed a low testosterone level and slightly elevated prolactin level.

Radiology

CT and MRI of the brain and skull were performed. A 1.4 cm calcified sellar/suprasellar mass was noted (Figure 1). This lesion extended superiorly to abut the left optic chiasm and optic nerve. Pituitary adenoma or craniopharyngioma were suspected based on these findings. Additionally, there was opacification of the left sphenoid sinus with hypointense material seen on both precontrast T1 and T2 weighted MRI. This material enhanced after contrast administration and imaging characteristics were consistent with fibrous dysplasia.

The patient was referred to our center for further evaluation and treatment. Based on the fact that the tumor could represent a symptomatic craniopharyngioma, resection was recommended. Although an underdeveloped or obstructed sphenoid sinus has historically been a relative contraindication to transphenoidal surgery, it was determined that an endoscopic transnasal approach would facilitate removal of the fibrous dysplasia involving the sphenoid sinus and also would be the best approach for resection of the sellar/suprasellar tumor.

Intraoperative course

Under general anesthesia, the patient’s nose was infiltrated with 5 ml of 2% Xylocaine with 1:100,000 epinephrine. Topicalizing cocaine pledgets were placed intranasally. A neuronavigation system with CT and MRI image fusion capability was used.

First, the middle turbinates were displaced laterally and outfractured to create an adequate operating pathway. The right sphenoid os was identified and opened. There was no identifiable sphenoid os on the left side as the sphenoid sinus in this region was completely replaced by abnormal bone. A posterior nasal septectomy was performed to allow binalar access to the posterior nasal cavity and sphenoid sinus. A Jansen-Middleton double action rongeur (C-Med Surgical, Inc.; Pompton Lakes, NJ) was used to resect the sphenoid rostrum. The bone...
was noted to be quite dense in areas and was grossly consistent with fibrous dysplasia. A Midas Rex drill (Medtronic, Inc.; Minneapolis, MN) with a 2.5mm matchstick cutting burr was used to perform drilling of the dense bone in the region of the sphenoid sinus, clivus, and laterally toward the left pterygoid. This provided endonasal access to the region of the sella.

Dense bone was encountered up to the cortical bone of the sella. Then, a 3 mm matchstick diamond drill was used to remove the cortical bone of the sella and expose the underlying dura. The dura of the sella was exposed from essentially the tuberculum sella down to the clivus and bilaterally toward the internal carotid arteries to allow adequate access to the suprasellar tumor.

The dura was then cauterized in a cruciate fashion using concentric suction bipolar electrocautery. A retractable endoscopic scalpel was then used to open the dural plane of the tumor. Zero degree and 30 degree endoscopes facilitated direct visualization of the tumor and surrounding structures for extracapsular dissection of the tumor. Finally, the tumor capsule was dissected away from the overlying dura of the sella and the dura in the region of the dorsum sella with careful preservation of these structures. A gross, total resection of the tumor was achieved. Valsalva maneuvers were performed to confirm the absence of any continued venous bleeding or cerebrospinal fluid leakage. Closure was performed using a sheet of DuraGen (Integra Life Sciences Corp.; Plainsboro, NJ) dural substitute placed in an inlay fashion within the dura followed by a dural substitute onlay graft. A synthetic polyethylene glycol based dural sealant was used to complete the repair. Next, the middle turbinates were medialized and sheets of sterile absorbable gelatin film were placed lateral to the middle turbinates in order to keep the middle meatus from becoming obstructed. Also, sheets of gelatin film were placed medial to the inferior turbinates to prevent synechiae formation and nasal obstruction. A fat graft or lumbar drain was necessary.

Postoperative course

Postoperative MRI imaging of the brain revealed complete resection of the pituitary tumor. There was no longer mass effect upon the optic chiasm. The patient did experience transient diabetes insipidus which resolved and did not require chronic treatment. The patient had complete resolution of his headache, resolution of his oculomotor abnormalities, and did not develop any new endocrinopathy. He was discharged to home on postoperative day three after an extra day in the hospital due to transient diabetes insipidus.

Pathology

Histologic sections of the pituitary mass showed small nests of cells situated in areas of dense mineralization (Figure 2). Numerous microcalcifications were noted. The small nests of tumor cells showed positive immunohistochemical staining for CAM 5.2, synaptophysin, chromogranin, and prolactin. They were negative for EMA and TSH, consistent with a calcified pituitary adenoma.

The sphenoid bone specimen contained irregular trabeculae of bone in a dense fibrous matrix. The bone spicules demonstrated a paucity of osteocytes and only scant osteoplastic activity was present, consistent with fibrous dysplasia (Figure 3).

Discussion

Fibrous dysplasia can occur in one (monostotic) or multiple (polyostotic) bones. It has also been reported in association with other diseases, as in McCune-Albright syndrome. Trabeculae of woven bone are contained in fluid-filled cysts embedded in a collagenous fibrous matrix, which contributes to the generalized hazy appearance of the bone on x-rays. The presence of fibrous dysplasia
of the sphenoid bone in this patient presented a challenge to accessing the sella through an endonasal endoscopic approach.

Given the patient’s preoperative complaints of headache, it was felt that the fibrous dysplasia could be responsible for these symptoms and therefore removing the fibrous dysplasia en route to the pituitary and suprasellar region could be beneficial. Given the imaging findings, and that the patient had galactorrhea, mild prolactinemia, and low testosterone level, it was felt that the sellar tumor was mostly likely a pituitary adenoma. Given the imaging findings, and that the patient had galactorrhea, mild prolactinemia, and low testosterone level, it was felt that the sellar tumor was mostly likely a pituitary adenoma. Based on the fact that the tumor was partially calcified and suprasellar, a craniopharyngioma was also possible.

Endoscopic resection of sellar and suprasellar tumors such as pituitary adenomas can result in a decreased length of stay, quicker recovery and fewer complications and was therefore the preferred approach. Neuronavigation was essential in this case to allow for identification of critical landmarks, including the carotid arteries and the sella, which was obscured by the fibrous dysplasia.

Fibrous dysplasia is a benign tumor and therefore when it is asymptomatic it can usually be observed. Surgical intervention is indicated for symptomatic lesions causing intractable bone pain, severely reduced mobility, cosmetic deformity, or local pressure on vital structures. In this case it was felt that fibrous dysplasia may have been responsible for the patient’s headaches. Furthermore, removal of the fibrous dysplasia was necessary in this case to gain access to the pituitary tumor.

Removing fibrous dysplasia in this region runs the risk of injury to the skull base, carotid arteries, and optic nerves and chiasm. Multiple approaches to fibrous dysplasia of the sphenoid bone have been described, including endonasal, transfrontal and subcranial. Eisenberg was the first to describe the endoscopic approach for the diagnosis of fibrous dysplasia of the ethmoid sinus. The current report is the first in the literature to describe an endoscopic approach to address fibrous dysplasia of the sphenoid bone.

Endoscopic endonasal removal of pituitary adenomas has been well described and is becoming the approach of choice at many institutions. However, endoscopic endonasal removal of fibrous dysplasia has not been widely described. With the advent of neuronavigation and ability to use CT, MRI and CT-MRI fusion, the approach to the sellar and suprasellar region even in the face of fibrous dysplasia or non-pneumatized sphenoid is feasible. We believe that osseous or fibro-osseous lesions involving the sphenoid are not necessarily a contraindication to endoscopic transsphenoidal surgery. In this case, the endonasal endoscopic approach allowed management of the fibrous dysplasia obstructing the sphenoid sinus facilitating access and gross total resection of the calcified pituitary adenoma.

References

Corresponding Author: Sanjay Yadla, MD
Department of Neurological Surgery
909 Walnut St, 3rd Floor
Philadelphia, PA 19107
Phone: 215-955-7000
Fax: Fax: 215-503-9170
Email: sanjay.yadla@jeffersonhospital.org