Pituitary Macroadenoma (Fungal Hyphae): A Case Report and Literature Review

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Abstract

The Aspergillosis is a well-recognized form of fungal infection, the involvement of the pituitary gland by aspergillosis is extremely rare. In which the main pathogen of the fungal seller is aspergillus classes. The pituitary infectious fungi consisted of different groups and dimorphic fungi. The leading pathogen of fungal seller abscesses is the type of aspergillus. The possible route of infection through sphenoid sinus with a thin seller roof. A pituitary tumor differentiates through its size. Pituitary tumors undesirably are known to grow in the sphenoid bone, cavernous sinus, mid-nasal ductus, and left cerebral hemisphere. In our case presentation patient presented with pituitary macroadenoma, which diagnosed through magnetic resonance imaging. Presented with nasal mass, and acromegaly features headache. Right eye vision loss. Through endoscopy, biopsy report shows the nasal fungal infection. Surgical treatment, the trans-sphenoidal procedure will be planned as recommended in literature reviews.

Keywords: Pituitary Macroadenoma, Nasal obstruction, Optic Chiasm, Fungal Hyphae.

INTRODUCTION

The etiologically of intracranial vascular contamination because of fungal infection is extremely uncommon. Furthermore, most cases occur in immunocompromised patients with hostile fungal disease, most commonly originating in the paranasal sinuses [1]. Since the first case pituitary aspergillus reported by Simmonds in 1914. The infecting fungi can be grouped into molds, yeasts, and dimorphic fungi.

The core pathogen of fungal seller infection is aspergillus classes. Aspergillus remain universal fungi produce in soil and organic materials and can start saprophytic progression inside the respiratory tract after inhalation [2].

The pathophysiologic pathway responsible for hypothalamic-pituitary dysfunction following acute meningitis is not fully unstated. In some patients, anti-pituitary and anti-hypothalamus antibodies are noticed. It is planned that acute infection aggravates an autoimmune process and may cause axonal injury with consequent neuroendocrine dysfunction (CNS infection) [3]. Initial identification and treatments are significant, and it is accordingly significant study the likelihood of seller aspergillosis in patients without primary risk factors. Grocott’s methenamine-silver stain (GMS) optimistic for septated hyphae growths in which the opportunity of an Aspergillus diagnosis and PCR (Polymerase Chain Reaction) can be used as a confirmatory test.

Discreetly adapted medical interventions with oral voriconazole seems to be effective after surgical excision. The Seller aspergillosis is an uncommon illness which is often originally misdiagnosed as an adenoma [4]. Pituitary aspergillosis has an extremely rare manifestation of invasive aspergillosis. A contrast MRI can often, but not always, help in suspecting this diagnosis. A trans-sphenoidal approach with debridement of the lesion is recommended. The optimum duration of therapy is unknown [5]. In this presenting case report, patient was admitted with sphenoid sinus infection, inflammatory signs of seller mass extension, optic chiasm involvement. Therefore, assumed that the pituitary fossa aspergillus contagion or infection was secondary to sphenoid sinus infection.
CASE PRESENTATION

A 36 years old female client was presented as an out-patient department, with complaint of headache, blurred vision, nasal obstruction, amenorrhea, acromegaly features, and high blood glucose levels >300mg/dl. Her complaint duration was 9 months. She had been seeking health care from district hospitals for nasal obstruction, and headache. She was literate (matriculation level). She had married since 7 years, even no kids or history of conceived pregnancy. Amenorrhea had been started 2 years back. She was a housewife. She had a history of the surgical procedure of appendicectomy 10 years back. In major medical problems history of diabetes Insipidus, Insulin dependent. Recent vital signs were recorded in normal ranges. No significant history of weight loss or gain. No history of food or drug allergy. She had no significant family history including (genetical, communicable and no-communicable diseases).

Physical examination has revealed typical acromegalic features in her face and extremities. Hard skin surface from palm of the hand soles. Right sided bulging eye. Cranial nerves intact.

After clinical evaluation, to rule out the cause of acromegaly features which develops in a short period of time. Her sleep rest pattern affected due to disease process and worried about changing of face features. After referral to the services, she started to be followed up regarding endocrinology (medical /U-II), neurosurgery, ophthalmology, and ENT consultations. On physical examination, acromegaly was evident. BSL= 300→350→270→250→450mg/dl. Weight = 53kg.

<table>
<thead>
<tr>
<th>Labs test</th>
<th>Normal values</th>
<th>Labs Values</th>
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</thead>
<tbody>
<tr>
<td>Glucose (FASTING)</td>
<td>90-109</td>
<td>84mg/dl</td>
</tr>
<tr>
<td>Growth Hormone (GH)</td>
<td>2.0-5.0 ng/ml</td>
<td>107 ng/ml</td>
</tr>
<tr>
<td>Prolactin (PRL)</td>
<td>5.18 - 26.53</td>
<td>15.95ng/ml</td>
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<tr>
<td>TSH</td>
<td>0.388IU/ml</td>
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<tr>
<td>T4 (serum)</td>
<td>5.5-11.0</td>
<td>4.7</td>
</tr>
<tr>
<td>ACTH</td>
<td>Normal</td>
<td></td>
</tr>
<tr>
<td>Cortisole</td>
<td>Normal</td>
<td></td>
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</tbody>
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All routine blood labs values normal in range. Anti-nuclear factors: Negative. Magnetic resonance imaging (MRI) done (February 12, 2019) protocol plain- T1W1-12W1 (axial, sagittal, and coronal) depicted a pituitary gland is enlarge measuring (21 x 22mm), appearing is on T1W1, hypo on T2W1. MRI suggested of bulky pituitary gland possibly of macroadenoma should be considered. MRI imaging re-reporting from Shaukat Khanum Memorial Cancer hospital, showing large pituitary mass suggest pituitary macroadenoma. Computerized Tomography of Para Nasal Sinuses shows Sino-nasal Polyposis. Endoscopic histopathology report showing the result of extensive infarcted tissue with non-septate branching fungal hyphae.

Mechanism of infectious, significantly the sphenoid sinusitis can be due to infection of bacterial or fungal, and the risk of hostile fungal sinuses, especially in patients who are immunocompromised and existing with acute complications of sinusitis. Visual symptoms, including vision loss due to hyper pituitary adenoma [6].

Medically treated with, Tab Metformin 500mg/TDS, Tab Itraconazole 100mg/BD. Tab Thyroxin 25mg/OD, Insulin “R” according to sliding scale, Hivit-spray 1 puff/both nostrils/TDS, Tab Deltacortil 2+2, Tab Vorif 200mg, Cap Sporanox 100mg/BD. Treatment was ongoing and planned for Trans-sphenoidal surgical procedure.

DISCUSSION

Pituitary aspergillus is very extremely uncommon condition which occurs in almost exclusively in immunocompromised patients [7]. Pituitary aspergillus infection shows a variability of appearances signs and symptoms when entering the sphenoid sinus and nasal cavity [8]. Endocrine disorders not involving the gonads, but strongly influence the reproductive functions. The surgical procedure is suitable treatment for those patients which they are presented with visual field deficits because it suitable for immediate decompression of the optic chiasm [9]. The combine radiation treatment may be more beneficial for patients to treat from further visual injury. Furthermore, the effective treatment of fungal abscess is endoscopic trans-sphenoidal surgery [10].

The optic chiasm is a key anatomical arrangement of ejection along the visual pathway, situated at the intersection between the optic nerves and regions. A wide range of diseases can affect the optic chiasm and its adjacent delicate region at the base of the brain [11]. It has been endeavoring to highlight the importance of sound clinical evaluation and multidisciplinary cooperation (neurosurgery, otorhinolaryngology, ophthalmology, and endocrinology) in the diagnosis and (neurosurgery, otorhinolaryngology, anesthesiology, endocrinology, and radiation oncology) in the management of a rare clinical entity through this case report [12].

Table: Laboratory Results

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Uncertainty one is faced with an uncommon seller lesion, including hypo-intensity on T2WI, a peripheral contrast enhanced circumstance, sphenoid sinusitis and the possibility of aspergillus infection [6]. Early diagnosis and early inception of treatment in acromegalic patients can prevent the progression of cardiovascular illness and diminish the risk of unexpected death [13].

CONCLUSION
Endocrine disorders lead to disfunction of the reproductive functions. As a described in the literature reviews pituitary aspergillus is very rare condition which occurs in immunocompromised patients. Antifungal medication Voriconazole preferred after surgery, and surgical treatment suggested for those patients which diagnosed with visual field deficits the reason is to, it allows for immediate decompression of the optic chiasm. Aspergillus seller abscess essential considered in the differential diagnosis of a seller mass, especially in the immune deficiency patients.

Even though, the accurate diagnosis of Aspergillus seller abscess can only be resolute by histopathological investigations, MRI and CT remain the best methods for pre-operative diagnosis. In this case presentation, treatment is on continue in the light of trans-sphenoidal surgical procedure.

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REFERENCES