A Case report of Nasal rhinosporidiosis presenting as an oropharyngeal mass.

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Abstract:

Rhinosporidiosis is a rare chronic granulomatous disease, characterised by polypoidal lesions of the mucous membrane. It commonly affects the mucous membrane of the nasopharynx, conjunctiva and palate. The disease is more prevalent in the Indian subcontinent, but remains quite rare. We hereby present a case description of a 19 year old male patient, native of Banaskantha district, diagnosed with polypoid nasal rhinosporidiosis. The patient presented with a history of nasal obstruction since 10 months and intermittent epistaxis for 2 months. Diagnosis was confirmed by histopathological examination and he was successfully operated with endoscopic resection of mass. This was a very unusual cause of nasal mass in our institute. Nasal rhinosporidiosis lesions commonly mimic other ordinary nasal polyps, angiofibroma and angiomatous polyp, it is therefore crucial for clinicians in our region to consider rhinosporidiosis as a differential diagnosis when assessing patients presenting with nasal swellings.

Key words: Nasal obstruction, Nasopharynx, Rhinosporidosis.

Introduction:

Rhinosporidiosis is an infectious, granulomatous, mucocutaneous disease caused by Rhinosporidium Seeberi, which affects mainly the naso-oropharynx and occasionally other mucosal surfaces.¹ It occurs worldwide and has been reported in the Americas, Europe, East Africa, and Asia, with the highest incidences registered in India and Sri Lanka.¹ Infection occurs after inoculation of spores found in stagnant water ponds, lakes, and rivers, and even in dust. In the nasal mucosa, the disease manifests as a reddish, friable, polypoidal, hyperplastic mass, which can be confused with malignant lesions, especially when ulcerated.²

The clinical diagnosis of rhinosporidiosis, affecting the nose and throat, is difficult and histopathological confirmation is essential. Radiological evaluation is often required to delineate the extent of the lesion, the status of underlying structures and maybe useful to rule out other common clinical differentials.

Case report:

A 19-year old male presented to our
hospital with long standing history of left-sided nasal obstruction since 10 months. He also gave history of occasional epistaxis, intermittent post-nasal drip, nasal discharge and foreign body sensation in oral cavity with occasional cough since 2 months. He reported no history of significant constitutional symptoms. The patient was a farmer. He reported no history of similar illness in the family.

On general examination, we observed that the young man had good nutritional status, was not pale or dyspnoeic and had poor general hygiene.

Intraoral examination revealed a pinkish red lobulated mass hanging from above, in the left lateral aspect of oropharynx, behind the faucial pillar, measuring about 3 x 2cm in size. The lesion was firm and bled on touch on palpation with the tongue depressor. On nasal examination there was a polypoidal mass filling the left nasal cavity, abutting the septum. The mass was erythematous, non-tender and bled easily on contact. The zero degree endoscope couldn’t be passed beyond the lesion. The right sided nasal cavity was normal. No enlargement of local-regional lymph nodes was observed. Other systemic examination was unremarkable.

The following differential diagnoses were considered: Nasopharyngeal angiofibroma, Vascular polyp, Nasopharyngeal carcinoma and Mycotic granuloma.

Work-up of the patient comprised of biochemical and haematological profiles as well as radiological work up. Biochemical and haematological parameters were essentially normal.

CECT PNS showed a soft tissue density lesion in the inferior part of left nasal cavity involving the inferior turbinate, inferior portion of nasal septum and floor of nasal cavity. The lesion causes bony erosion and rarefaction of anterior part of left inferior turbinate (Image 1). Posteriorly, the lesion showed extension into the nasopharynx and oropharynx. The lesion also showed involvement of inferior portion of left sided nasolacrimal duct with expansion of the bony nasolacrimal duct and enhancing soft tissue within it (Image 2). On post contrast study, the lesion showed intense heterogenous enhancement (Image 3).

Image 1: Coronal unenhanced bone window CT section showing soft tissue density lesion in inferior left nasal cavity. Bony erosion and rarefaction of anterior aspect of left inferior turbinate is identified (red arrow).
Image 2: Axial unenhanced bone window CT scan showing soft tissue density lesion in left nasal cavity extending posteriorly into nasopharynx on left side (yellow arrow). Expansion of inferior bony portion of left sided nasolacrimal duct with soft tissue within it (red arrow).

Image 3: Axial contrast enhanced soft window CT scan showing intense heterogeneously enhancing (red arrow) in left nasal cavity extending into nasopharynx on left side. Also note enhancing lesion in expanded left bony nasolacrimal duct.

Patient underwent endoscopic excision of mass coupled with electric cauterisation of the base. Anterior nasal packing was done which was removed after 2 days. Post-operative period was uneventful; the patient did remarkably well and was discharged with tablet dapsone for a 6 month period. Follow-up visit six months later revealed no signs of recurrence.

Gross examination of the specimen depicted an intact elongated, polypoidal soft friable greyish mass measuring about 9 x 4 x 1cm (Image 4). Cut surface was whitish. Histopathological examination of the excised tissue showed granulation tissue containing plasma cells, lymphocytes and collection of histiocytes and neutrophils. Globular cysts containing spores (sporangia) in different stages of development were identified predominantly in the stroma of the polypoidal tissue. These findings were pathognomonic of rhinosporidiosis. Henceforth, a final diagnosis of rhinosporidiosis was established.
Case Report

Discussion:

Rhinosporidiosis has existed since ancient times. Initial account of this disease entity was made over a century ago in Latin America.\(^3\) Though it appears to occur universally, rhinosporidiosis remains largely endemic in Indian subcontinent. The mode of transmission to humans is not clearly understood, however most researchers believe that direct contact with spores through dust, soil or prolonged exposure to stagnant water are among the major risk factors for acquisition of the infection.\(^3\) Many patients may give similar history which points towards the diagnosis. We believe that in our case, the patient being a farmer by occupation, his frequent bathing in ponds and stagnant waters and overall lack of personal hygiene, were high risk factors contributing to contraction of such an infection.

Rhinosporidiosis is a non-contagious chronic granulomatous infection seen mainly in adult males aged between 20 and 40 years.\(^4\) Usually the patient presents with a history of gradual nasal obstruction, occasional epistaxis, nasal itching, sneezing, and at times post-nasal dripping.\(^1\) Clinically, nasal rhinosporidiosis is characterised by development of single pedunculated or sessile polyp, multiple sessile polyoidal lesions or a combination of both\(^3\). Our case had a single polyoidal lesion. The classical clinical finding in RS is the presence of

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Image 4: Excised polypoidal specimen (Soft friable greyish with irregular surface.)

Image 5 Multiple varying sized mature and immature sporangia

[H&E, 4X]

Image 6 Sporangia containing microspores

[H&E, 10X]
a reddish mass with greyish-whitish dots on the surface (representing sporangia); however this was not seen in our case. The lesion often bleeds on touch and has a soft friable consistency. Location of the lesion plays an important role in narrowing down the differential diagnosis. Contrary to ordinary polyps which often arise from the middle turbinate, rhinosporidiosis frequently involves mucosal lining of the naso-pharynx, anterior nares and inferior nasal cavity. Nasal polyps originating from these locations should always be treated with high index of suspicion. In the nasal cavity, the most common site involved was the inferior nasal cavity, comprising nasal floor, inferior turbinate, and inferior meatus. According to Banjara et al., the most common sites of involvement in rhinosporidiosis in order of frequency are nasal cavity, nasopharynx, lacrimal sac, and conjunctiva.

Radiological features, though nonspecific may provide a clue and may give rise to suspicion of rhinosporidiosis infection. The most common imaging appearance of nasal RS is that of a polypoidal lesion centred in the inferior nasal cavity, (as opposed to the nasopharynx and sphenopalatine foramen in JNAF) involving the nasal floor, inferior turbinate, inferior meatus and inferior part of septum, as was the case in our patient. Involvement of surrounding bones is common in rhinosporidiosis, as seen in our case. Bone involvement is seen as irregularity, rarefaction, partial or complete erosion of inferior turbinate, thinning of medial maxillary wall, and septal erosion. Maxillary sinus extension is uncommon and this feature may help in differentiating it from other nasal masses such as antro-choanal polyps and inverted papillomas. nasolacrimal duct involvement is common due to frequent presence of rhinosporidiosis in inferior meatus and is diagnosed when nasolacrimal duct is dilated with extension of soft tissue density into it, as was seen in our case.

Apart from history and clinical and radiological findings, histopathology is mandatory for definitive diagnosis of rhinosporidiosis. Definitive diagnosis of rhinosporidiosis depends upon identification of the pathogen in its diverse stages on biopsied or resected tissues. Histopathological sections show multiple sporangia in various stages of maturity, enclosed in a thin chitinous wall. The sporangia are 50-1000 µm in diameter, containing numerous endospores of diameter 5-10 µm. Overlying epithelium is usually hyperplastic, with or without areas of ulceration and loose fibrovascular stroma infiltrated with lymphocytes, macrophages, plasma cells and even polymorphonuclear leucocytes. Rupture of sporangia can cause giant cell reaction.

Surgical excision remains the mainstay of treatment for rhinosporidiosis lesions. Wide, complete and meticulous excision of the polyp followed by thorough electro-cautery of the lesion's base is recommended. It is hypothesized that cauterisation of the lesion's base may abate recurrence resulting from spillage of endospores on the adjacent mucosa.

Besides surgery, a variety of adjuvant medical therapies have been tried in the treatment of rhinosporidiosis. These include drugs like griseofluvin, amphoterecin B and dapsone (4, 4-diaminodiphenyl sulphone). Dapsone is a promising drug that is used as an adjunct to surgery as it arrests the maturation of sporangia and promotes fibrosis. However, by far there has been no tangible success with medical therapy.

Conclusion:
Nasal rhinosporidiosis seldom remains a diagnostic consideration in our region. However, with emerging reports of sporadic cases, it is consequently imperative for clinicians in our setting to consider rhinosporidiosis as a differential diagnosis when evaluating patients presenting with nasal growth.

References


