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7	Palato-Antral Involvement of a Primary Extracranial Sinonasal
8	Meningioma
9	*Vasantha Dhara, ¹ K.P. Saamaja ²
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11	¹ Department of Dentistry, Government General Hospital, Kakinada, India; ² Department of
12	ENT, GSL Medical College, Rajahmundry, India
13	*Corresponding Author's e-mail: dg91dan@gmail.com
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15	Abstract
16	Primary extracranial sinonasal meningiomas are one of the rarest tumours involving the head
17	and neck region. Very few reports exist with additional involvement of the palato-antral
18	region. This paper describes a case of primary extracranial sinonasal meningioma with palatal
19	involvement and the management of the same. The patient was taken up for surgical excision
20	via intra-oral and endoscopic approaches. Postoperative healing was uneventful with no
21	recurrence noted over 2 years. A palatal obturator was used for rehabilitation. The diagnosis
22	of this pathology needs additional immunohistochemistry testing for confirmation and
23	treatment entails complete surgical excision which assures no recurrence or delayed
24	presentation of residual disease in follow-up.
25	Keywords: primary extra cranial meningioma; sinonasal; maxilla; palate.
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27	Introduction
28	Meningiomas are known to be common slow-growing benign tumours of the central nervous
29	system. Presentation of primary extracranial meningioma in the maxillo mandibular region is
30	very rare with an incidence of 2-0.9%. ¹⁻³ These are sporadic with suspected aetiology arising
31	from the meningiocytes of the central nervous system via cranial nerves or as embryonic
32	arachnoid cell rests. ⁴ Patients report complaints of nasal blockage, stuffiness, bleeding,
33	painless mass, facial swelling etc; which emphasises the importance of histological

34 confirmation of the diagnosis to separate from other sino nasal or maxillary tumours and proceed with management. The prognosis of this benign tumour is favourable with disease 35 36 recurring only if there is residual mass from the definitive treatment. ^{5,6} 37 38 Case report 39 A 45-year-old male reported with complaints of nasal blockage, frequent episodes of 40 epistaxis and nasal regurgitation for 8 months. The patient had no prior history of any allergies, sinusitis, rhinitis, trauma or headache. There was no history of any deleterious 41 42 habit. Clinical examination revealed a soft swelling over the right cheek region and a palatal 43 fistula with surrounding friable growth. (Figure 1A, B, C) Endoscopic evaluation showed a 44 reddish mass in the right nasal cavity. After routine blood investigations, biopsy and imaging were done to ascertain the nature of the mass. Contrast-enhanced computed tomography of 45 the head and neck region was suggestive of enhancing soft tissue attenuating mass of 46 33x34mm, involving the right naso-antral region with palatal erosion. No cranial pathology 47 48 was noted. (Figure 2A, B). Biopsy from the tissue showed islands of spindle cells and 49 fascicles of round cells with round nuclei showing a stippled chromatin pattern. (Figure 2C). 50 Hyaline stroma with areas of necrosis was also seen in a few sections. Bony trabeculae were 51 noted to be invaded by these cellular areas. Immunohistochemistry was positive for Vimentin, epithelial membrane antigen (EMA) and S-100 confirming the diagnosis of 52 53 atypical sinonasal meningioma. 54 55 The patient was taken up for definitive treatment under general anaesthesia. A combination of 56 intra-oral and endoscopic techniques was used for the excision. The vestibular approach 57 exposed the tumour which was seen to be involving the anterior, antero lateral wall, the floor of the maxillary sinus, the nasal floor and the palate. A complete excision of the tumour was 58 59 done. (Figure 3A, B, C). A titanium mesh was placed over the defect to provide support 60 followed by closure. A surgical obturator was used for initial rehabilitation and was planned 61 for a definitive prosthesis after healing. 62 The patient was followed up at regular intervals and was disease-free for 2 years. (Figure 4). 63 **Discussion** 64 65 Extracranial meningiomas are known to affect sites such as paranasal sinuses, orbit, scalp, 66 ear, and temporal bone. Primary extracranial sino nasal meningiomas are rare, ie. without

evidence of any intra-cranial pathology. Their origin is postulated to be as a direct extension

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68 of an intra-cranial meningioma via resorption of bone or arachnoid cell rests or as metastasis. ⁷ Literature shows a female predilection due to the presence of oestrogen, progesterone and 69 androgen receptors on these tumours. ⁸ In this case, the patient was a middle-aged male. 70 71 Common presenting symptoms are nasal stuffiness, epistaxis, sinusitis, anosmia, headache, 72 proptosis, periorbital oedema, etc; similar to any nasal benign or malignant mass. A 73 differential diagnosis in such sinonasal mass should include extracranial meningioma, which 74 can be confirmed by a biopsy. The World Health Organization (WHO) classified on a 75 histological basis as Grade I benign – consisting of the following types -fibroblastic, transitional, psammomatous, and angiomatous; which account for 90 % of meningiomas. 76 77 Grade II - consists of Atypical meningiomas which show hypercellularity, mitosis and 78 necrosis, with occasional brain invasion. Grade III - malignant meningiomas which resemble 79 melanomas, carcinomas, or high-grade sarcomas with definitive cerebral invasion and rapid recurrence. ⁹ Strong reactivity for anti-EMA and vimentin are consistent with the diagnosis 80 of meningioma, thus highlighting the importance of immunohistochemistry testing in the 81 diagnosis. ¹⁰ Our patient was diagnosed with atypical meningioma. 82 83 84 For studying the extent of the lesion, magnetic resonance imaging or contrast-enhanced 85 computed tomography is generally undertaken. Our patient showed enhancement in the nasoantral region, with additional erosion of the palate and the maxillary wall. There was no 86 87 breach into the orbital floor or any other cranial pathology which was noted. Involvement of 88 maxillo mandibular complex as separate or extension of sino nasal meningiomas has been reported, but are few. ^{2, 4,11-14} 89

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The definitive treatment for extracranial meningiomas is complete surgical excision with acceptable cure rates of 80-85 % for 5 years. ¹⁴ Any recurrence is rare and is attributed to incomplete removal in the primary surgery. Most sinonasal meningiomas are excised by an endoscopic approach. However, due to the sites of the antrum and palate being additionally involved in our case, an intra-oral approach was used to remove the bulk of the tumour via vestibular approach exposing the antral walls, pyriform rims, nasal floor and septum after initial endoscopic evaluation and excision. Closure was done after placing a titanium mesh for soft tissue support. Considering the patient's personal and economic preference, a surgical obturator was placed initially and later a definitive obturator was given, thus maintaining oral function and preventing nasal regurgitation.

101 A regular follow-up is advised especially in grade II meningiomas, with additional attention to any cerebral symptoms and subsequent associated imaging. ¹⁵ Healing was satisfactory in 102 103 our patient and was disease-free till the follow-up period of 2 years. The patient was able to 104 function well without nasal regurgitation during the consumption of food and had no nasality 105 in voice due to the palatal seal of the obturator. The 5-year disease-free survival rate ranges from 66.9% to 82.1%, and the 10-year disease-free survival rate ranges from 54.6% to 78.6%. 106 107 The recurrence rate after complete resection varies from 7% to 84% according to follow-up. 16

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Conclusion

Primary extracranial meningiomas involving sinonasal, maxillary and mandibular sites are rare and need a thorough diagnosis to identify the grade of tumour. Knowledge of this pathology is necessary for all maxillofacial and otolaryngologists as its presenting features are not uncommon. Adequate surgical excision by endoscopic or intra-oral or a combination of both gives satisfactory results and has good prognosis.

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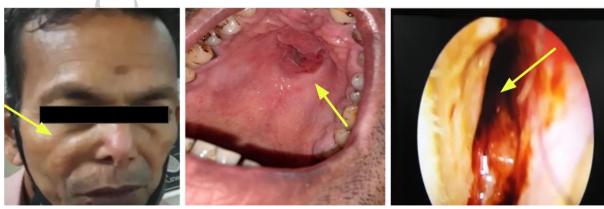


Figure 1: A: figure showing mild cheek swelling. B: growth in palate with fistula. C: endoscopic view of mass in right nasal cavity

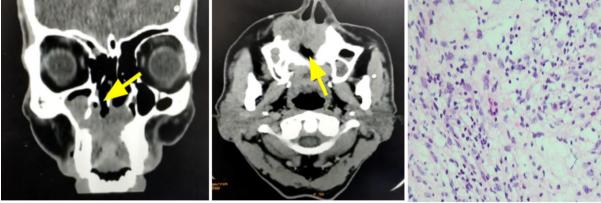


Figure 2: A: coronal view of lesion involving naso- antral region and palatal bone erosion. B: axial view of lesion. C: figure showing islands of spindle cells, round cells with prominent nucleus and chromatin pattern

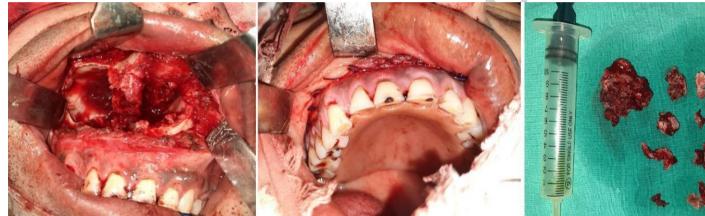


Figure 3: A: figure showing excision of mass from the antrum, nasal floor and cavity. B: figure showing closure of incision and obturator in place. C: figure showing excised specimen



Figure 4: follow up image showing no residual disease