

18th International Conference of the
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ROYAL MAXIM PALACE KEMPINSKI, CAIRO

**ORBIT AND
BEYOND**

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Orbital Meningioma

- Meningiomas
 - Arise from meningotheelial cells (arachnoid cap cells)
 - Comprising
 - 13%–20% of all brain tumors
 - 34–36% of all primary brain tumors.
 - Make up 4% of all orbital tumors
- Orbital meningiomas are divided into:
 - Primary Orbital Meningioma
 - Optic Nerve Sheath Meningioma
 - Sphenoid Wing Meningioma with Orbital Involvement

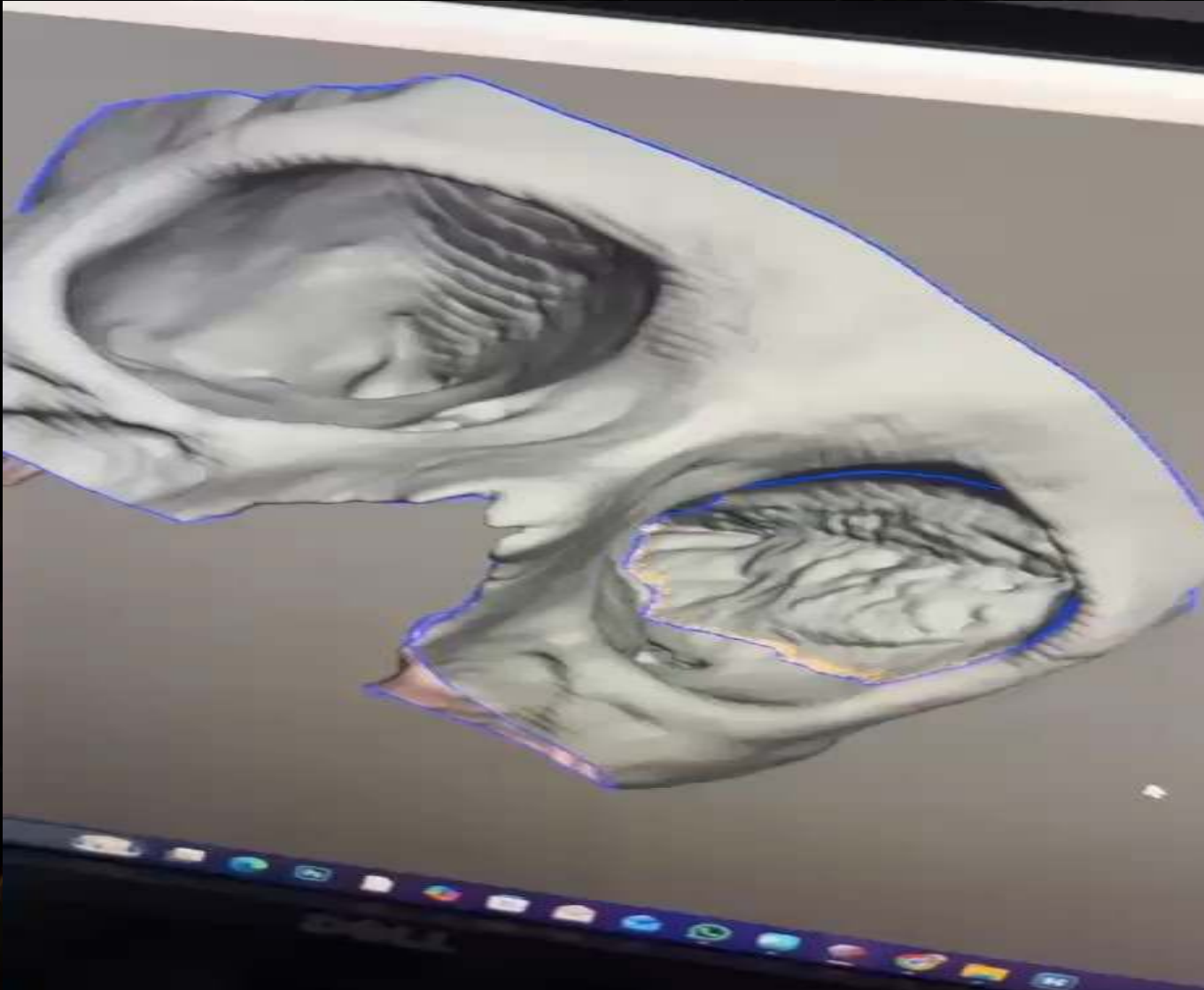
• Primary Orbital Meningioma

- Very rare tumours
- Intraorbital ectopic meningioma,
- Arising from ectopic arachnoid without any optic nerve sheath or meningeal connection.
- present around 30 years of age,
- Located Superomedially, medially, or inferomedially in the orbit.
- Trauma is thought to be responsible for some of these tumors,

• Optic Nerve Sheath Meningioma

- Arise from the optic nerve sheath arachnoid layer
- make up 2% of all orbital tumors and 1–2% of all meningiomas
- proptosis with classic triad of
 - vision loss,
 - optic atrophy, and
 - optociliary shunt vessels.
- 5% of cases are bilateral.
- CT scans show classic tram-track appearance.
- Managed conservatively or with Fractionated radiotherapy

Sphenoid Wing Meningioma with Orbital Involvement



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• **Sphenoid Wing Meningioma with Orbital Involvement**

- Spheno-orbital meningiomas (SOM), AKA
 - Orbitosphenoid meningiomas,
 - Meningiomas en plaque of the sphenoid wing
 - Sphenoid wing meningiomas with osseous involvement.
- Makes up 9% of all intracranial meningiomas
- Often associated with sphenoidal wing hyperostosis
- Meningiomas have two growth patterns:
 - **Meningioma en masse** which forms a space-occupying lesion
 - **Meningioma en plaque**, which is a flat spreading tumor.

Pathophysiology

- Ionized Radiation
- Genetic Abnormalities'
- Hormonal Factors
- Viruses
- Head Injury and Electromagnetic Fields

Epidemiology

- The average age at onset is 63 years
- The annual incidence of meningiomas can be as
 - low as 0.74 per 100,000 individuals younger than 34 years
 - high as 18.86 per 100,000 individuals older than 85 years.
- F:M 2.5:1
- NO racial privilege

Clinical Features

- Proptosis (86%). usually painless;
- Visual impairment (78%) . usually gradual but acute in 12%
- Ocular paresis (20%) with double vision .
- Papilledema and eventual optic atrophy .
- Headache is also a common manifestation
- Nausea if there is raised intracranial pressure.

Differential Diagnosis

- **Fibrous dysplasia**
- Paget disease
- Brown tumor
- Osteogenic sarcoma
- Eosinophilic granuloma
- Aneurysmal bone cyst
- **Osteoma**
- Hemangioma
- **Hemangiopericytoma**
- Epidermoid tumor and dermoid
- Multiple myeloma
- Plasmacytoma
- Giant cell tumor
- Metastatic cancer
- Exuberant arachnoid hyperplasia
- Fibrous histiocytoma
- **Juvenile ossifying fibroma**
- Solitary fibroustumor

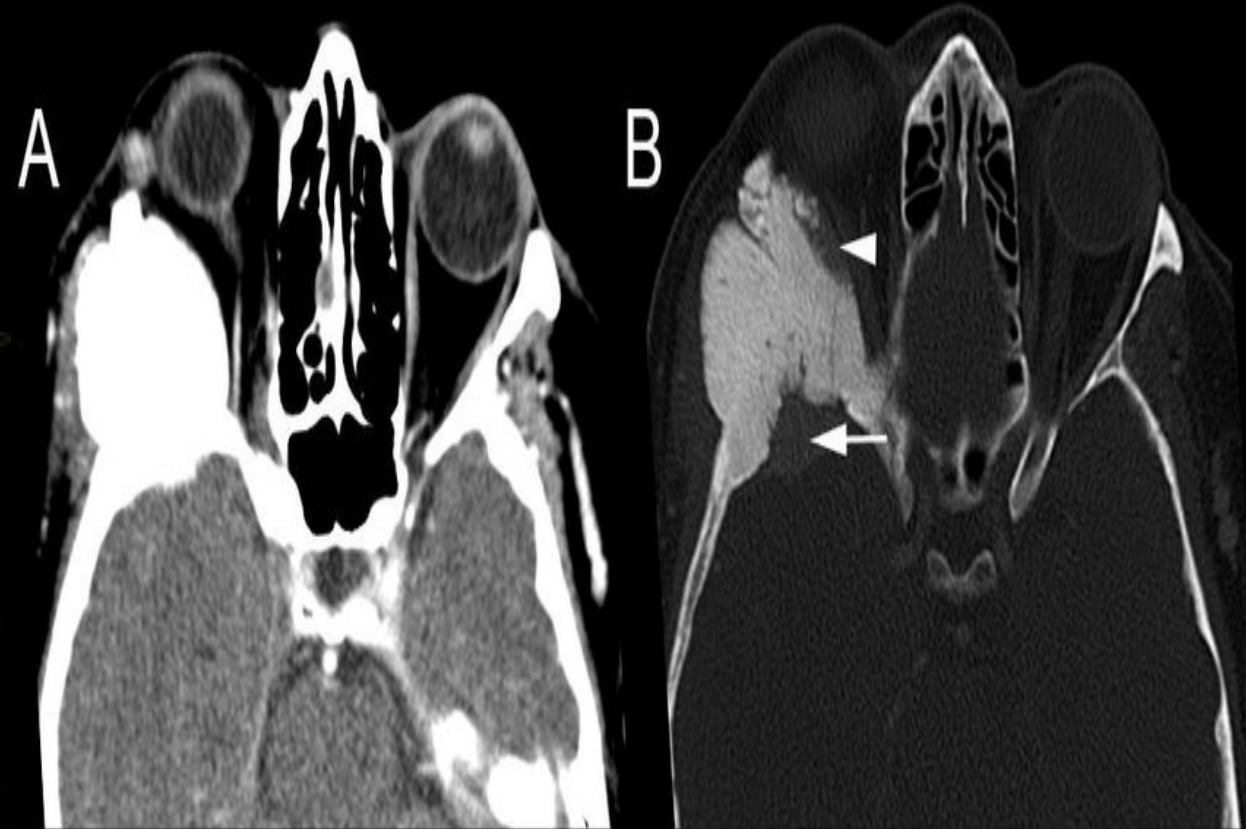
Diagnostic Evaluation

- Laboratory Studies
 - Calcium, bone-specific alkaline phosphatase, urine hydroxyproline)
- Imaging
 - CT ,MRI
- Other Tests
 - VEP, visual field, PET scan

Imaging

- CT Scan

- Bone windows shows a thick, hyperdense, intra-diploic lesion expanding the calvaria and destroying the cortical layers of the skull.
- The bony expansion with ground-glass appearance of sphenoid wing, with irregularity of the inner table associated with a dural reaction compared to smooth inner surface in fibrous dysplasia



Imaging



- MRI Scan
- MRI allows better delineation of the soft tissue component of the tumor and dural involvement, as well as delineation of intraorbital extension.
- A dural tail in sphenoid wing meningioma is typically identified but can be absent especially in the en plaque variant.

Management

- Surgery
 - Trans cranial
 - Trans-orbital
- Medical Treatment
 - Reserved for atypical and malignant meningiomas
 - Tamoxifen, an antioestrogen hormone
 - Mifepristone, (Mifeprex), is an antiprogesterone agent.
- Radiotherapy
 - Not indicated as primary treatment unless surgical resection is not possible,

Histology

- According to the WHO, three types of meningiomas exist based on malignant behaviour:
 - **Grade I (benign)**, with a recurrence rate of 7%–25%, does not meet atypical or malignant
 - **Grade II (atypical)**, with a recurrence rate of 29%–52%, includes atypical, increased mitotic activity (4–19 mitoses per 10 high-powered fields) or brain invasion
 - **Grade III (malignant)**, with a recurrence rate of 50%–94%, Anaplastic with 20 or more mitoses per 10 high-powered fields

Prognosis

- Following complete excision patients should be followed closely for recurrence rates can be as high as 27.7%

Transorbital and Endoscopic Approaches

- **Superior Eyelid Approach (SEA)**
 - Relief of proptosis and optic nerve compression are the primary goals of surgery
- **Inferolateral Transorbital Endoscopic Approach (ILTEA)**
 - Efficiently manage the infraorbital component of the lesions in cases of limited intracranial extension and without involvement of the optic canal
- **Combined Multiportal Approaches**
 - Multiportal transorbital and transnasal approaches have been proposed and described in order to overcome some limits of the pure transorbital or lateral approaches

Superior Eyelid Approach (SEA)

- **Incision:** Superior eyelid crease with lateral extension; skin-orbicularis flap raised
- **Orbital dissection:** Suborbicularis dissection supero-laterally to orbital rim;
- **Exposure landmarks:** (SOF), (IOF), optic canal identified.
- **Periorbita:** Must remain intact to prevent orbital fat protrusion.
- **Vessels:** Bridging vessels may be sacrificed; Hyrtl's (meningo-orbital) foramen vessel cauterized and cut if present.
- **Extended approaches:** Lateral orbital rim osteotomy, wider incision,
- **Bone drilling:** Greater sphenoidal wing (GSW) drilled laterally → medially to dura; temporalis muscle landmark.
- **Boundaries:**
 - Superomedial → SOF
 - Inferomedial → IOF
 - Lateral → temporalis muscle
- **Intracranial work:** Dura opened under neuronavigation;
- **Orbital tumors:** Periorbita opened if involved; preserve extraocular muscles and neurovascular structures to avoid diplopia.



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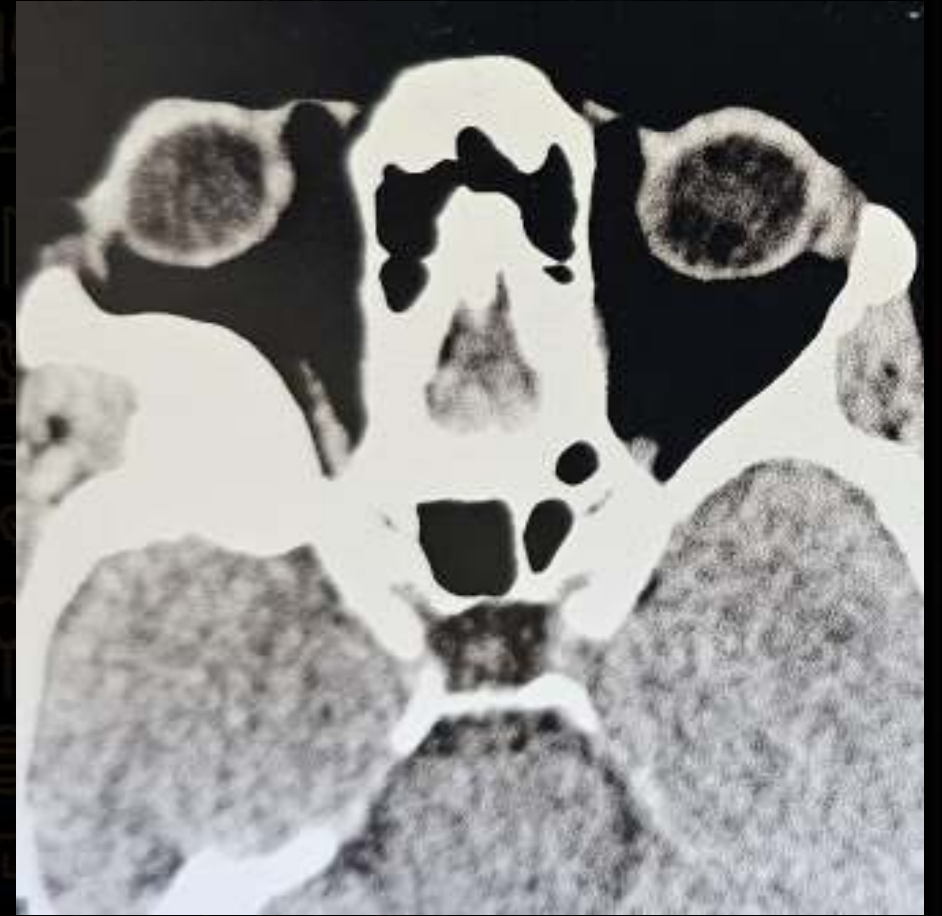
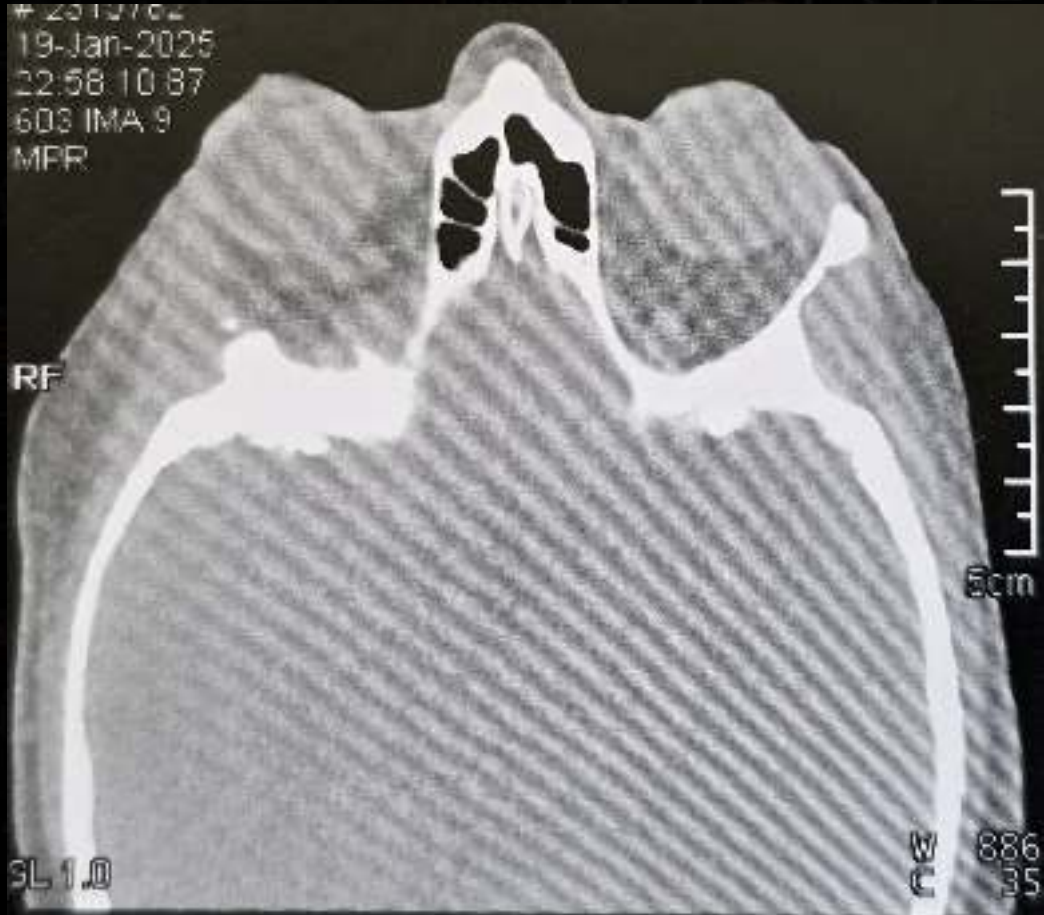


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Meeting our limits

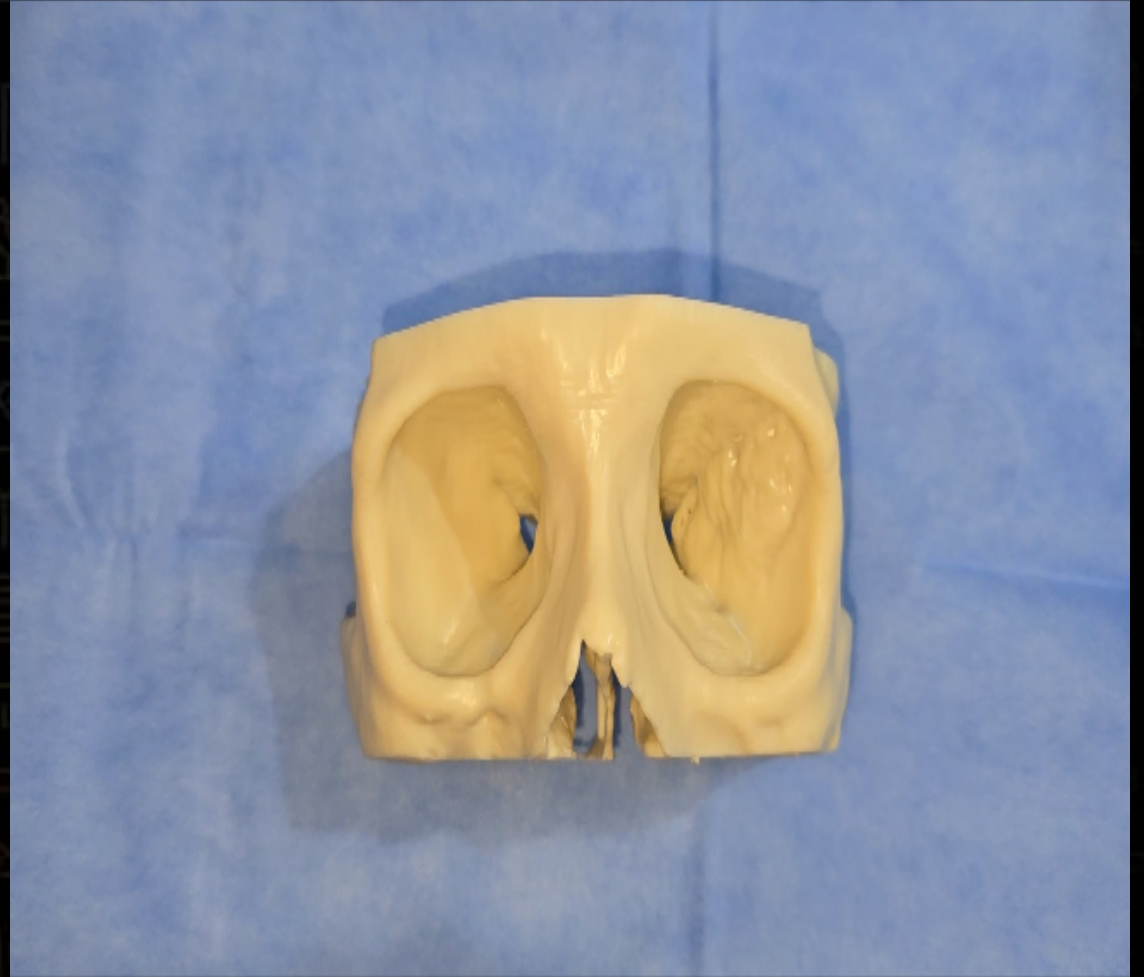
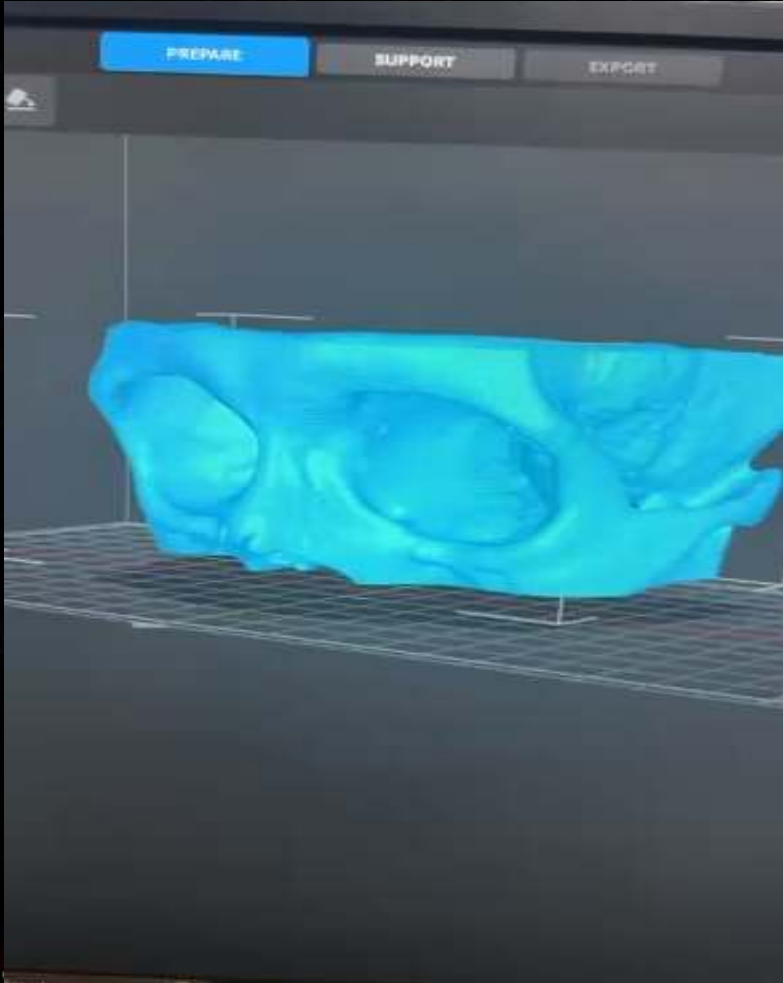


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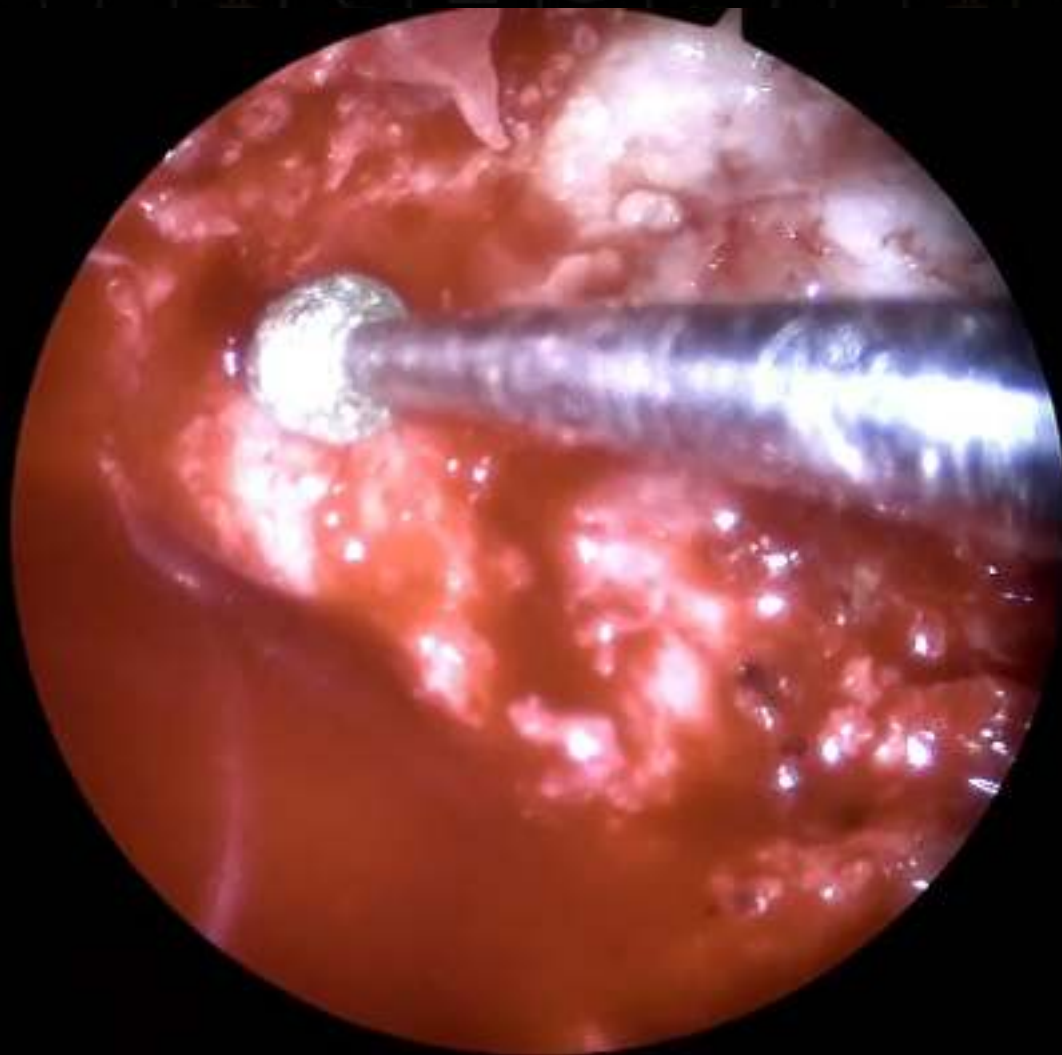
Poor man navigator



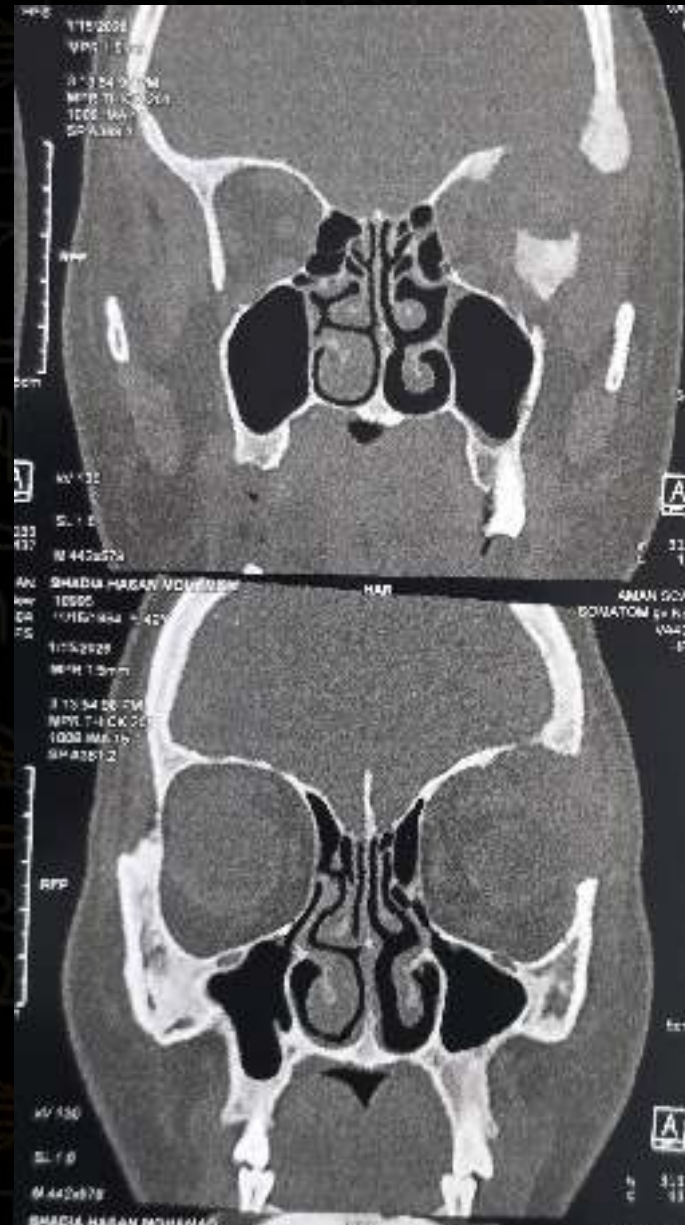


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Digging Deeper



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Thank you

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