



# **MENINGEAL TUMOURS**

- **Primary**
- **Secondary**



Primary tumours: **MENINGIOMAS**

Frequency: 15-20% all primary intra-cranial tumours

Slowly growing

*Benign clinical course with possible recurrences*

Age: >50 ys.

Occurrence at **younger age, multiplicity** and **ponto-cerebellar** location

→ **Neurofibromatosis**

Sex: F (particularly spinal tumours)

Symptoms:     **Mass effect**  
                  **Focal defects**  
                  **Epilepsy**



## Histogenesis:

- Nests of arachnoidal cell within the Dura
- Pacchioni's granules
- Parasagittal location

## Gross:

- Size: 1-10 cm
- Colour: Whitish  
Reddish areas due to haemorrhage  
Yellow spots due to lipidization
- Consistency: Fibrous to hard, calcifications
- Growth pattern: expansile



## Localizations

Parasagittal

Olfactory douche

Sphenoid wing → *carpet-like*

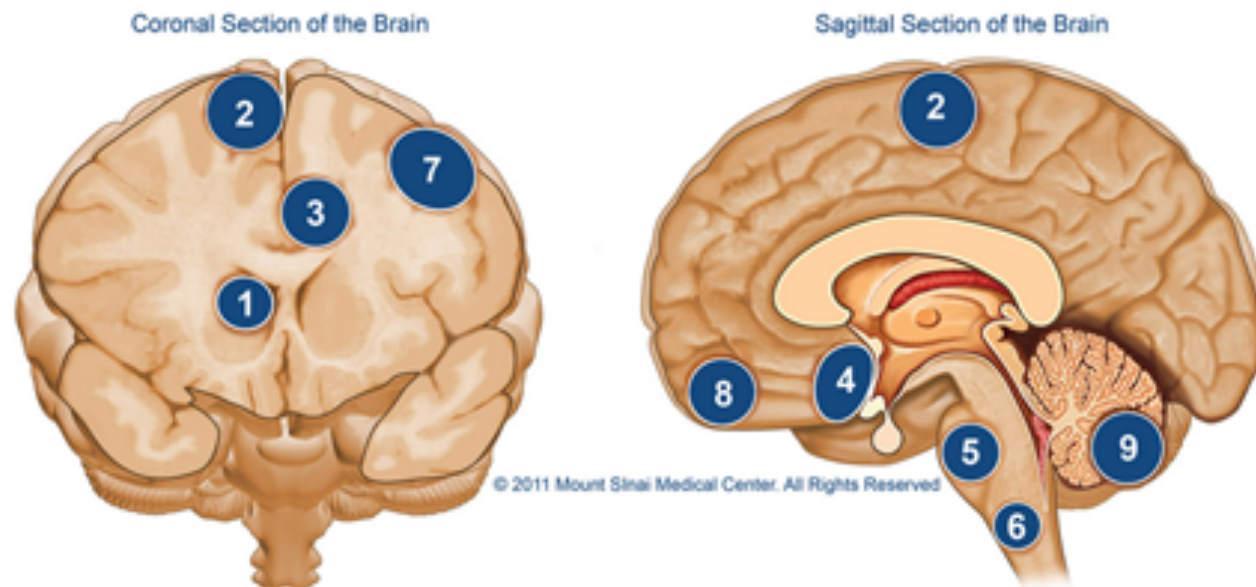
Tubercle of the sella turcica

Tentorium

Ponto-cerebellar


Intraventricular (from choroidal plexes)

## Common Location (Types) of Meningiomas



Meningiomas are often described by their location within the brain. The most common are:

1. Intraventricular
2. Parasagittal
3. Falcine
4. Suprasellar
5. Clivus
6. Foramen Magnum
7. Convexity
8. Olfactory Groove
9. Cerebellar



## Tumour expansion may cause:

- Compression of the underlying nervous tissue

***“en niche”***

- Compression of the inner cortical bone

**Hyperostotic** reaction and new bone deposition

**Erosione** and ***pseudoinfiltration*** (more rarely, invasion of adjacent soft tissues)

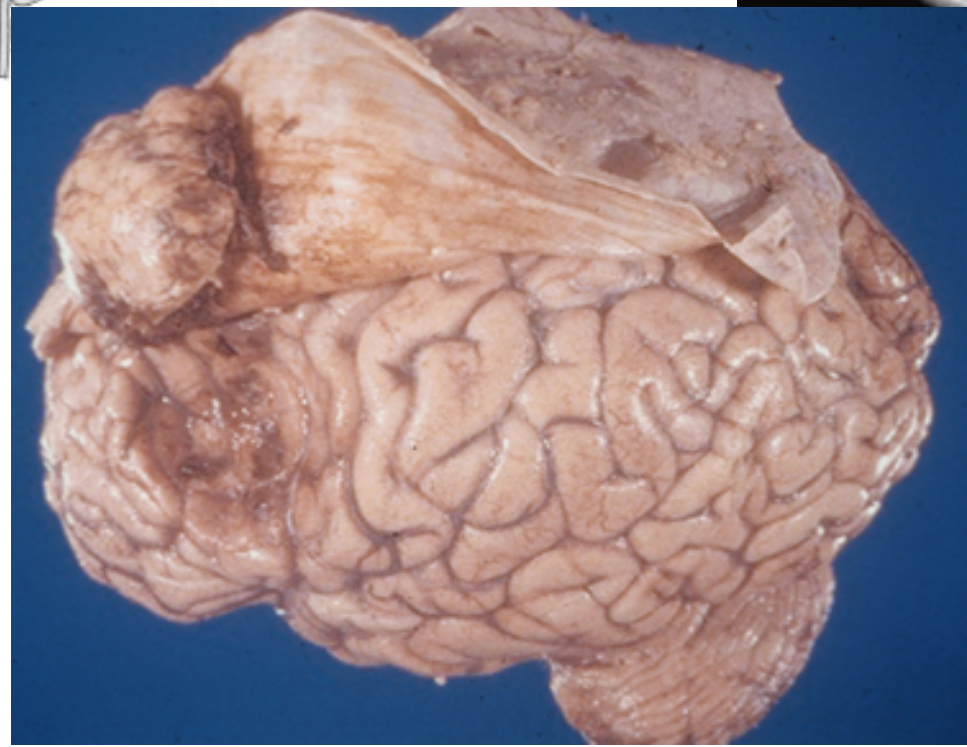
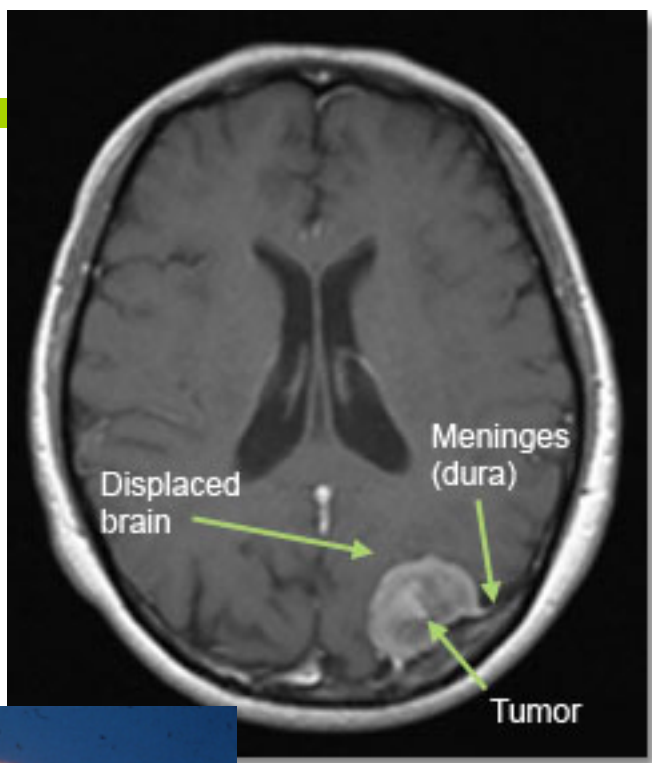
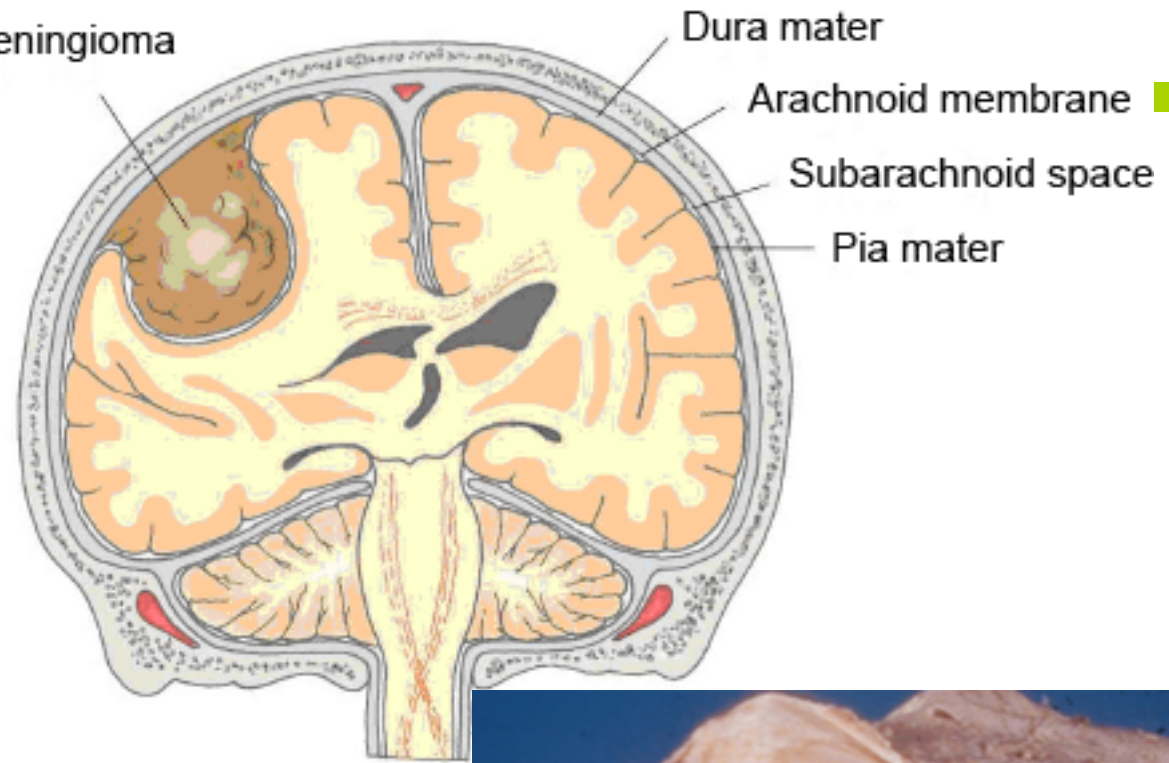
Falx

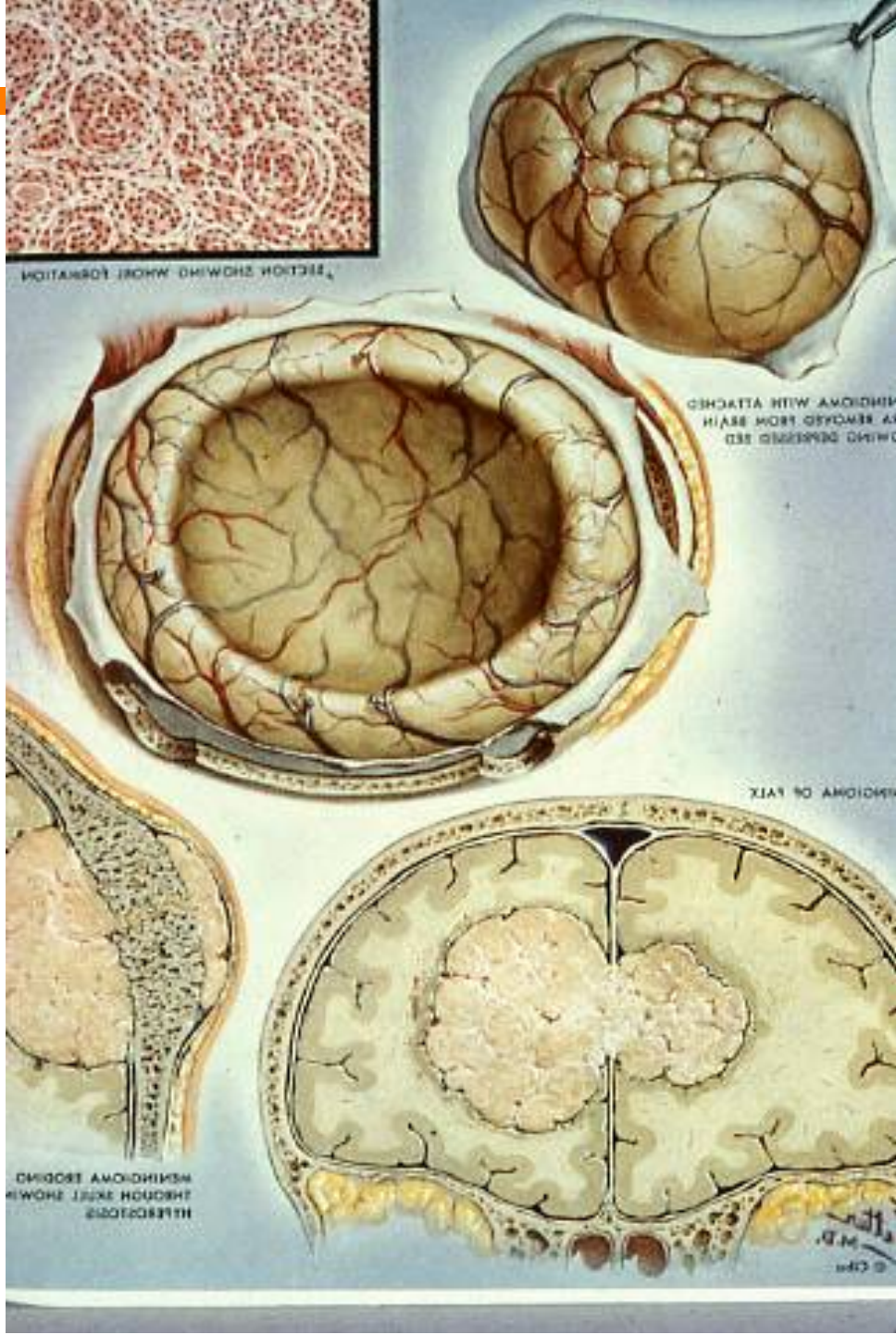
Tentorium

Dural sinuses

- ***En Plaque, carpet-like***

Meningioma

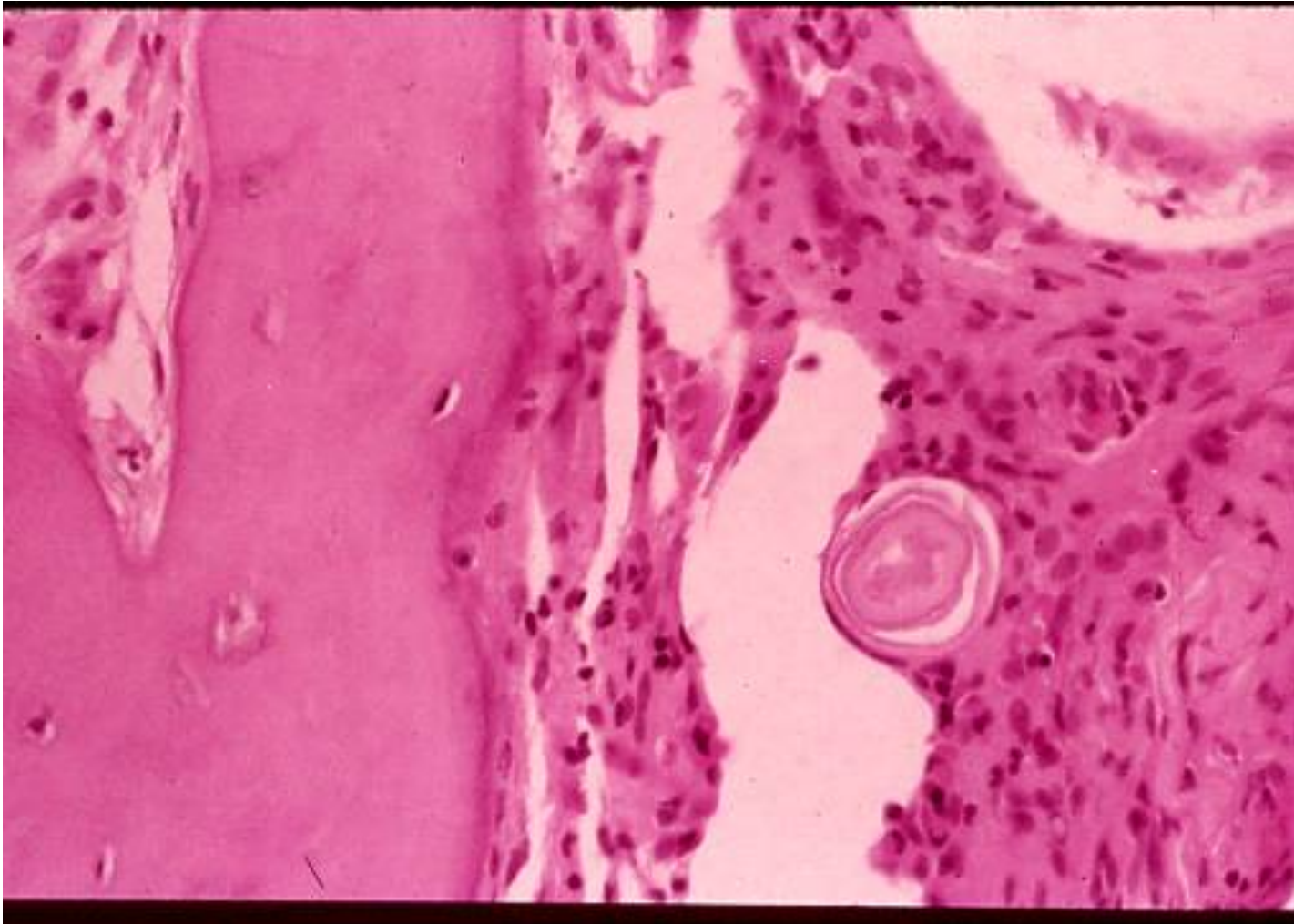


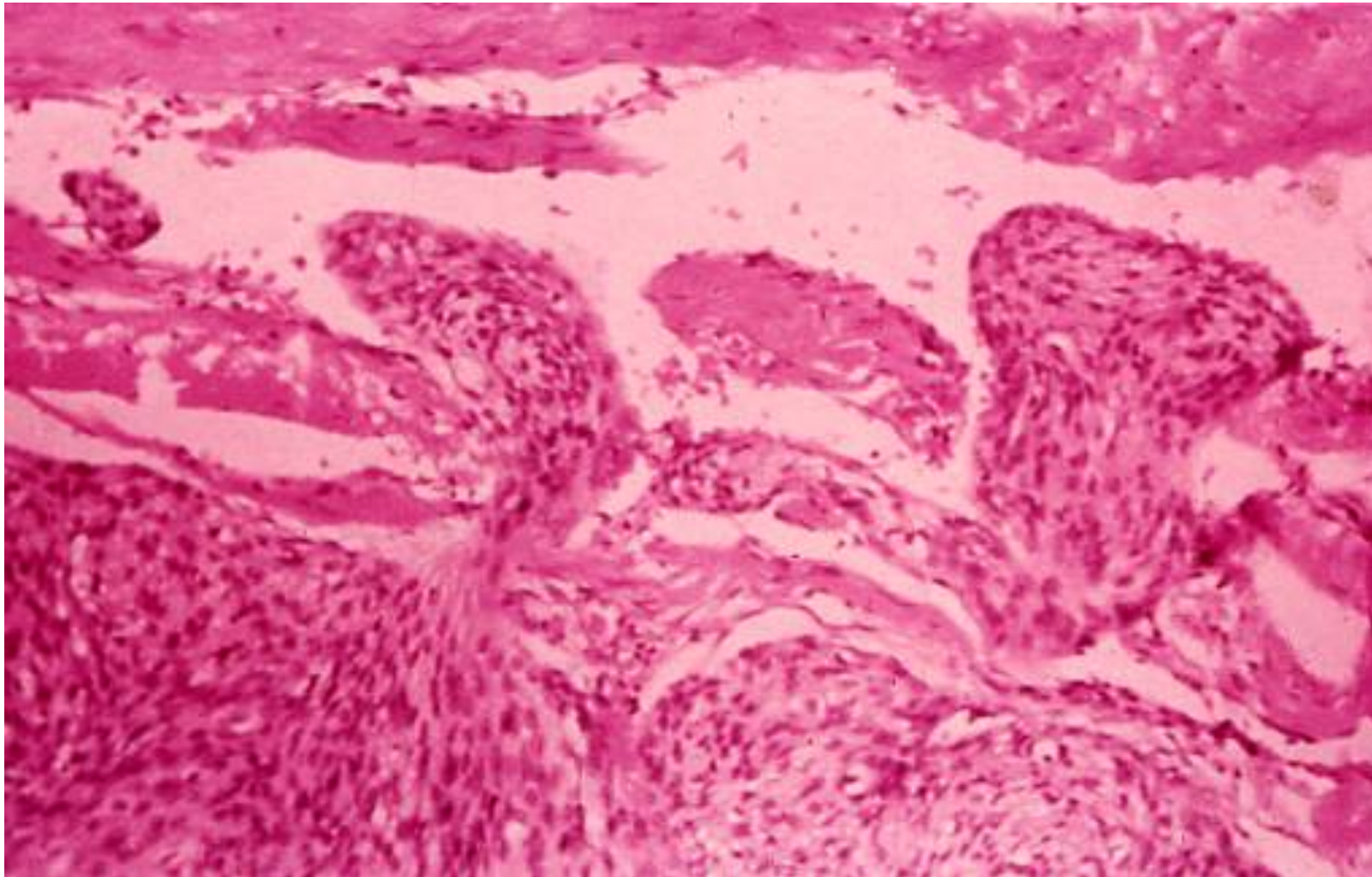




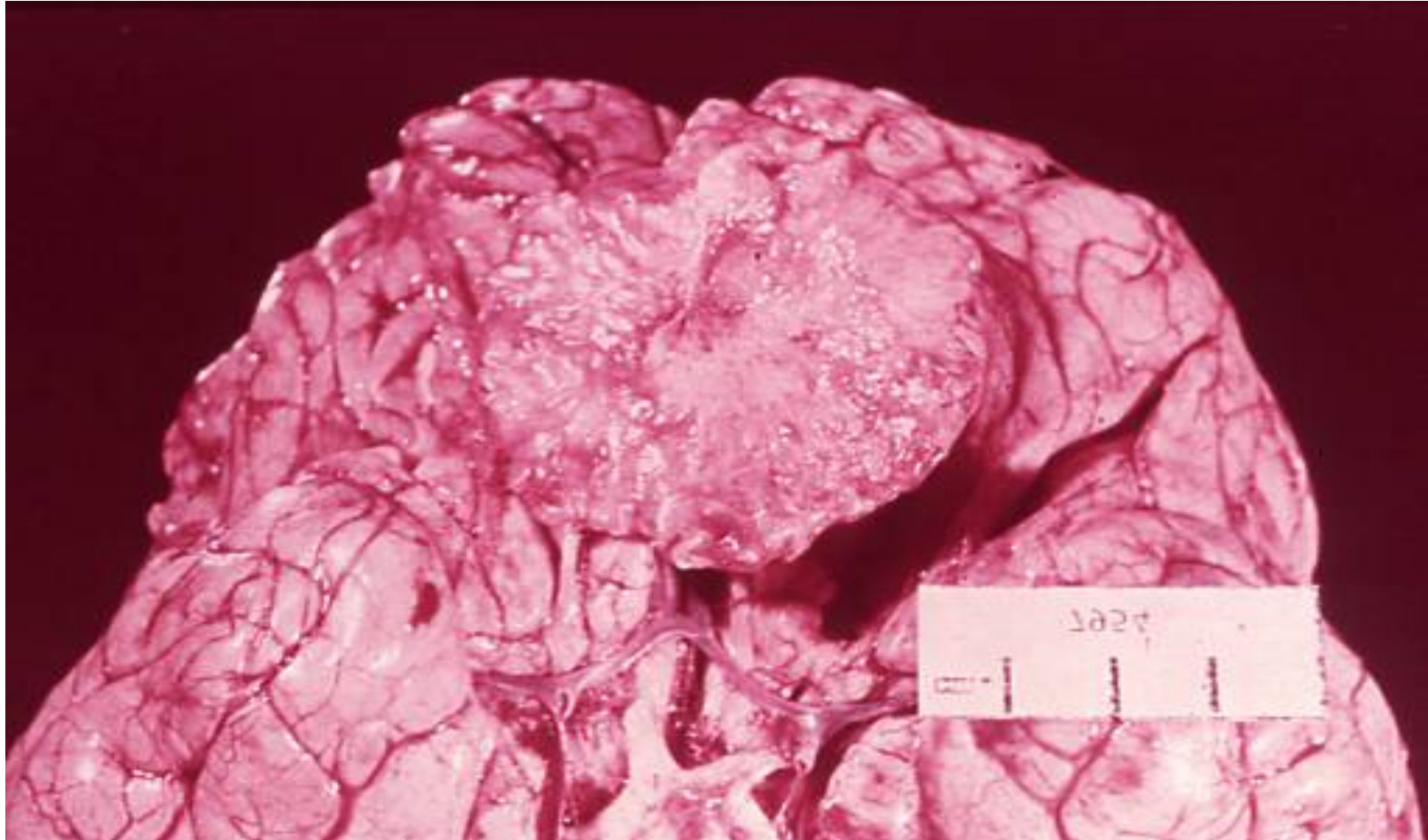


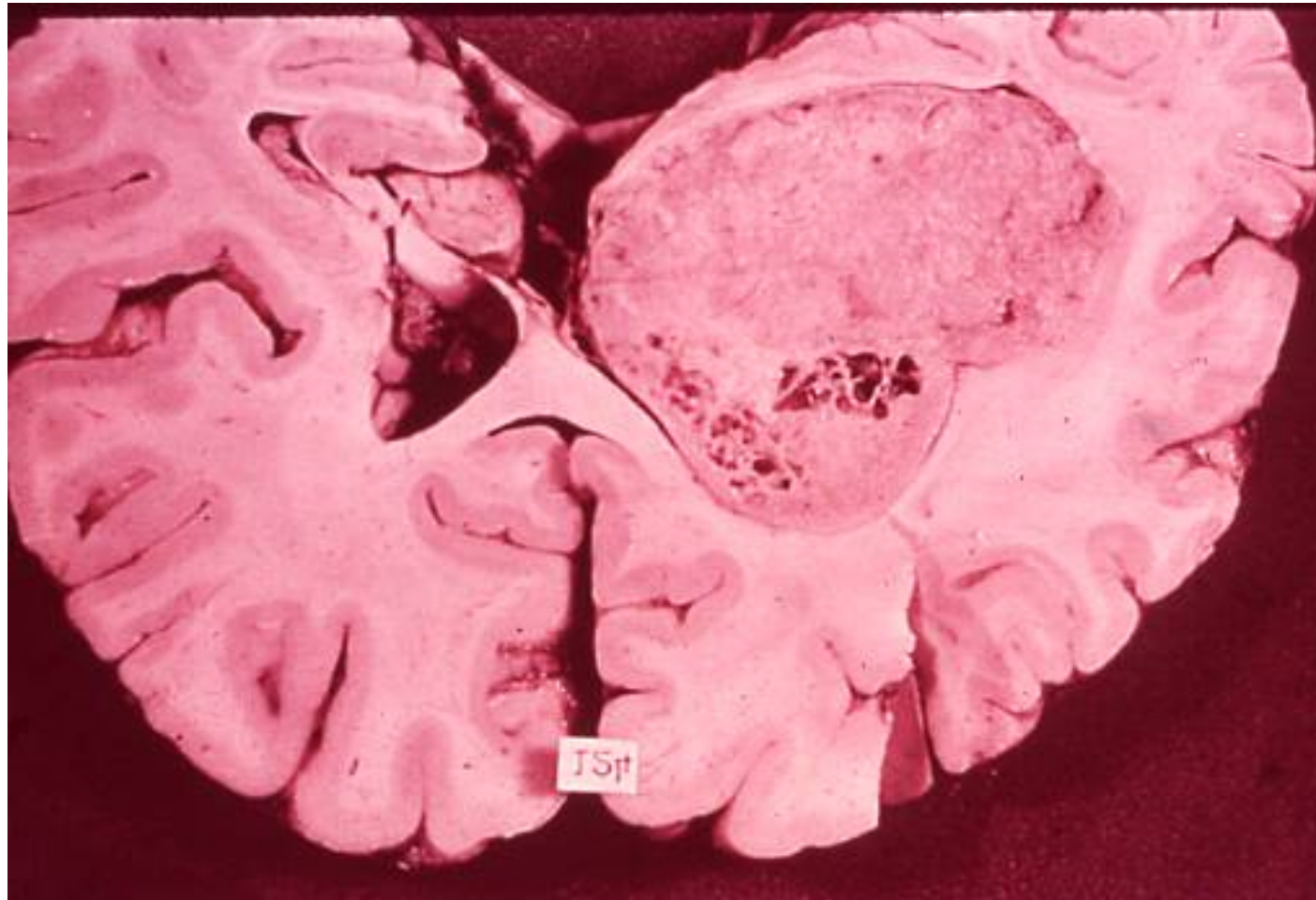


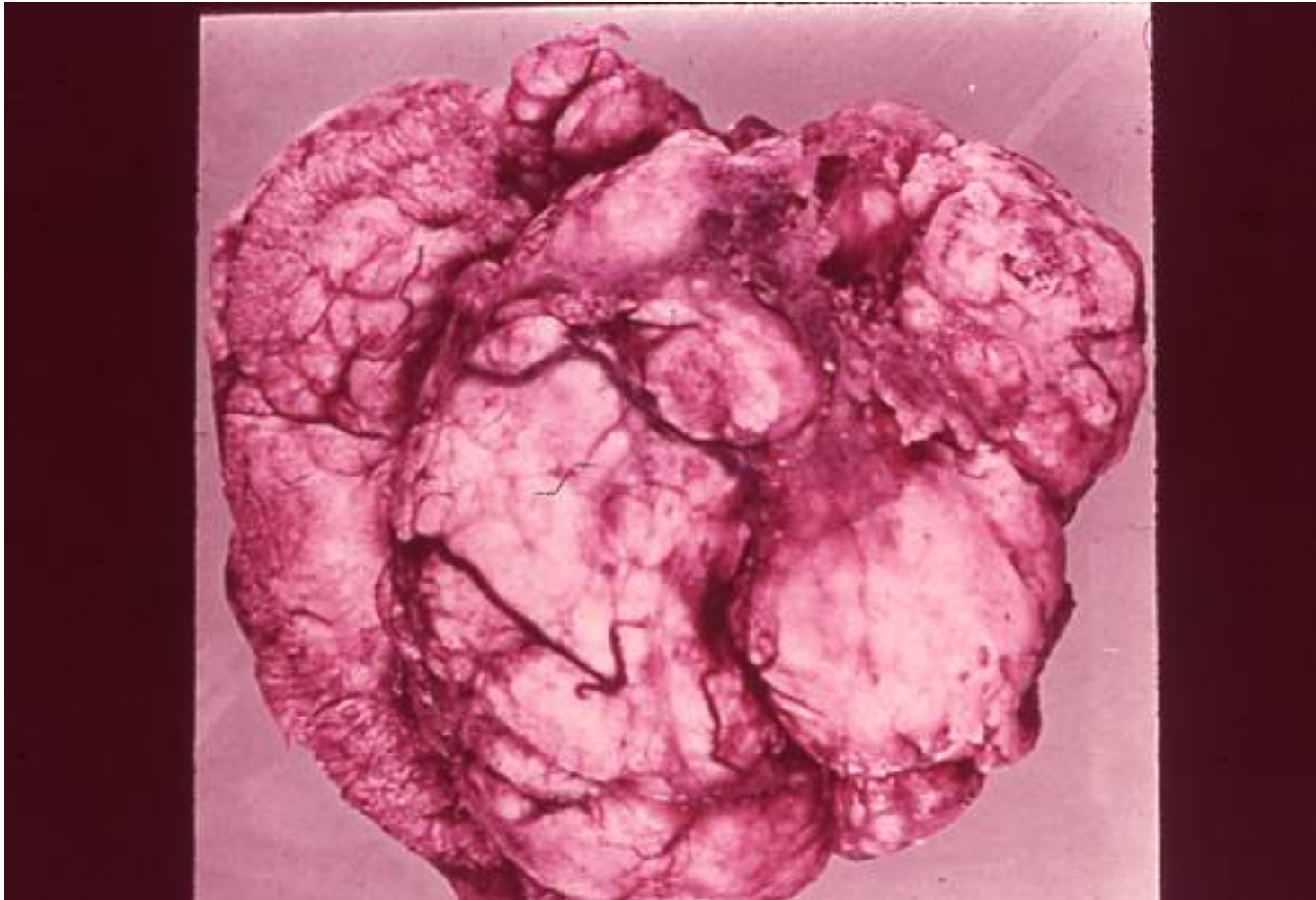


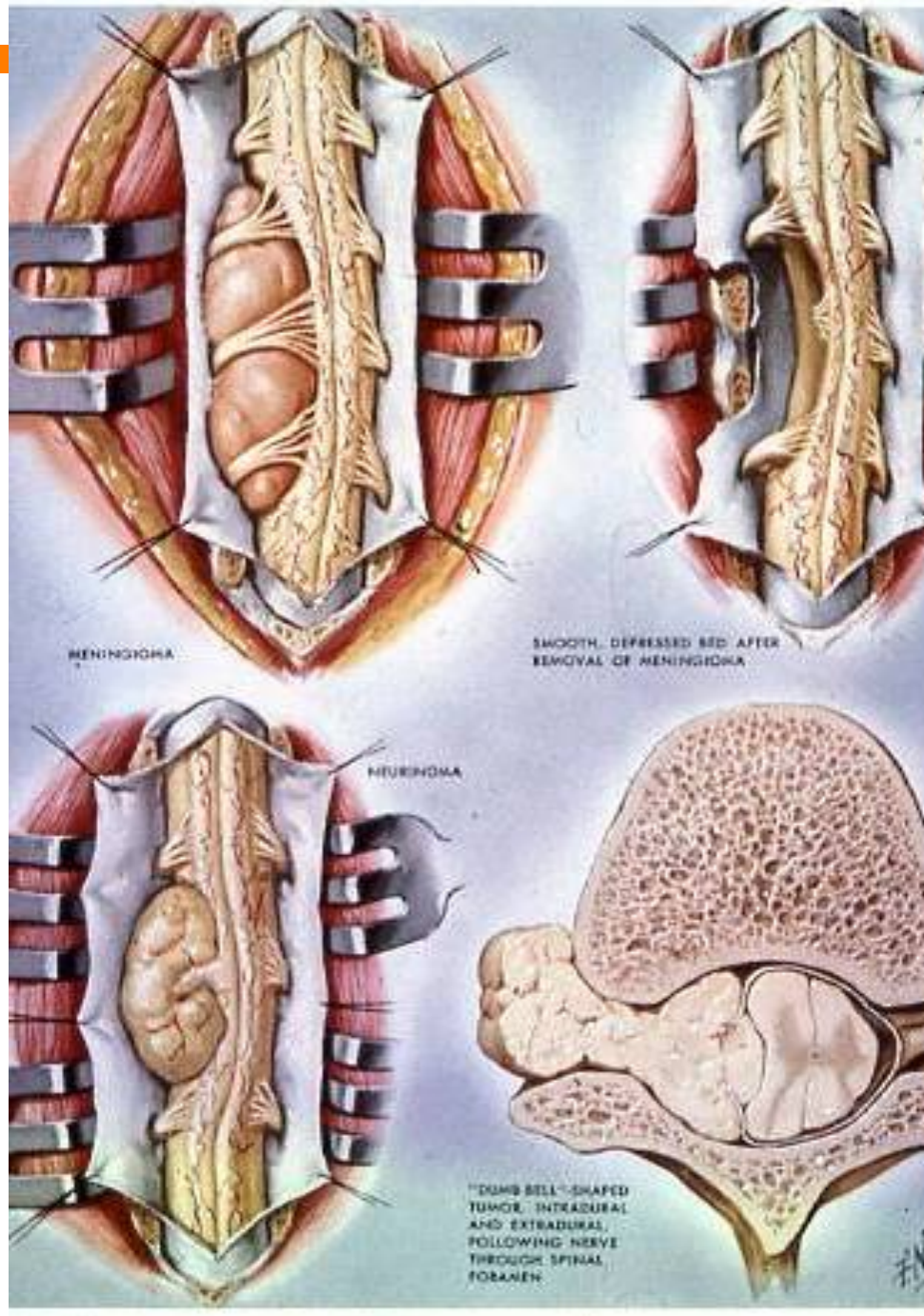


**Superior sagittal sinus**

















## Micro patterns (EMA+, CK-):

### Meningothelial (syncitial)

Lobules of meningotheial cells in a vorticooid pattern. Pseudonuclear inclusions, psammomatous bodies frequent

### Fibrous

Spindle cells in elongated fascicles, in perivascular locations; intranuclear inclusions rare, sclerosis and calcifications frequent

### Transitional

Intermediate between meningotheia and fibrous, with vorticooid pattern and spindle cells

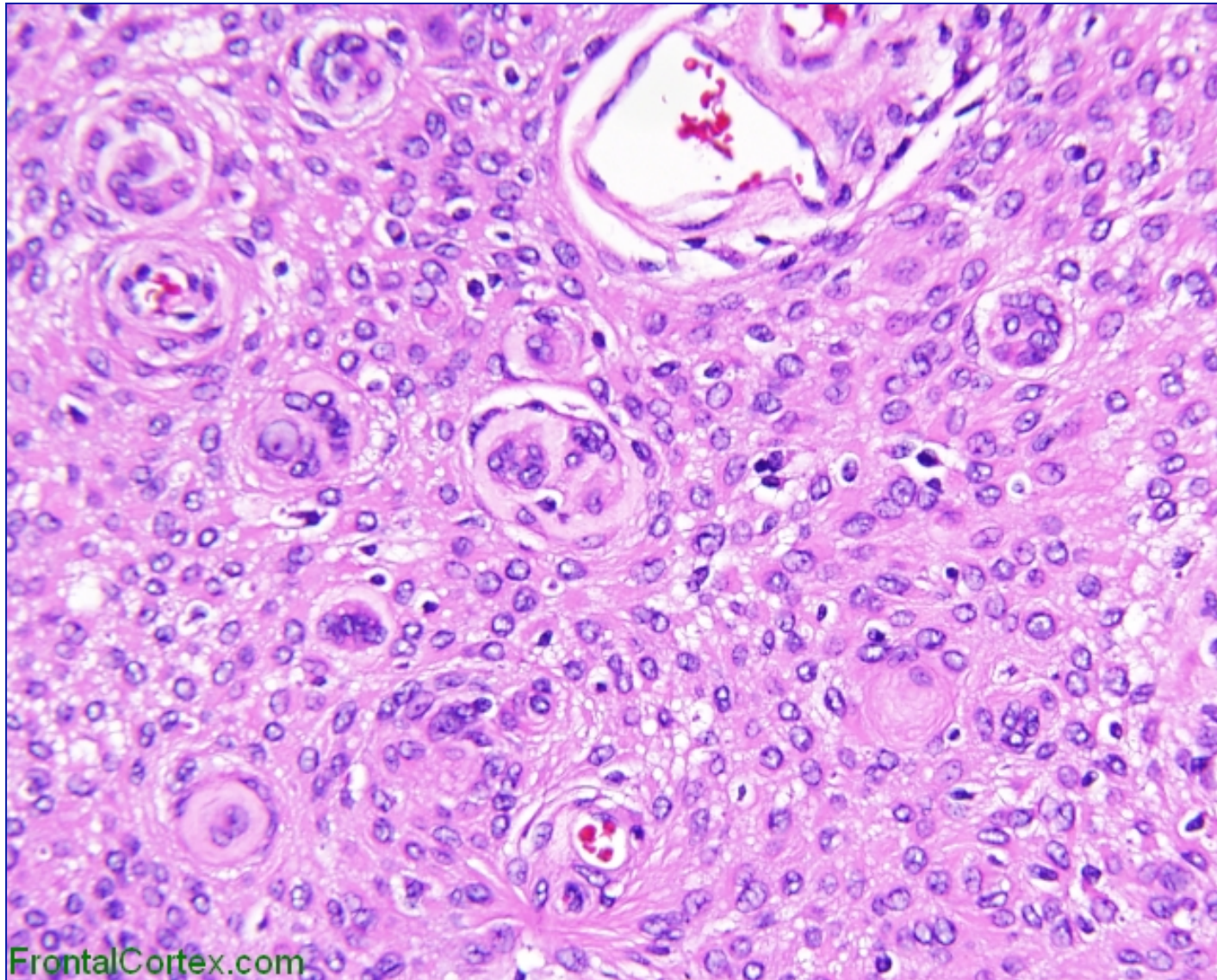
### Psammomatous

Olfactory douche, spinal; vorticooid pattern, psammoma bodies and calcifications

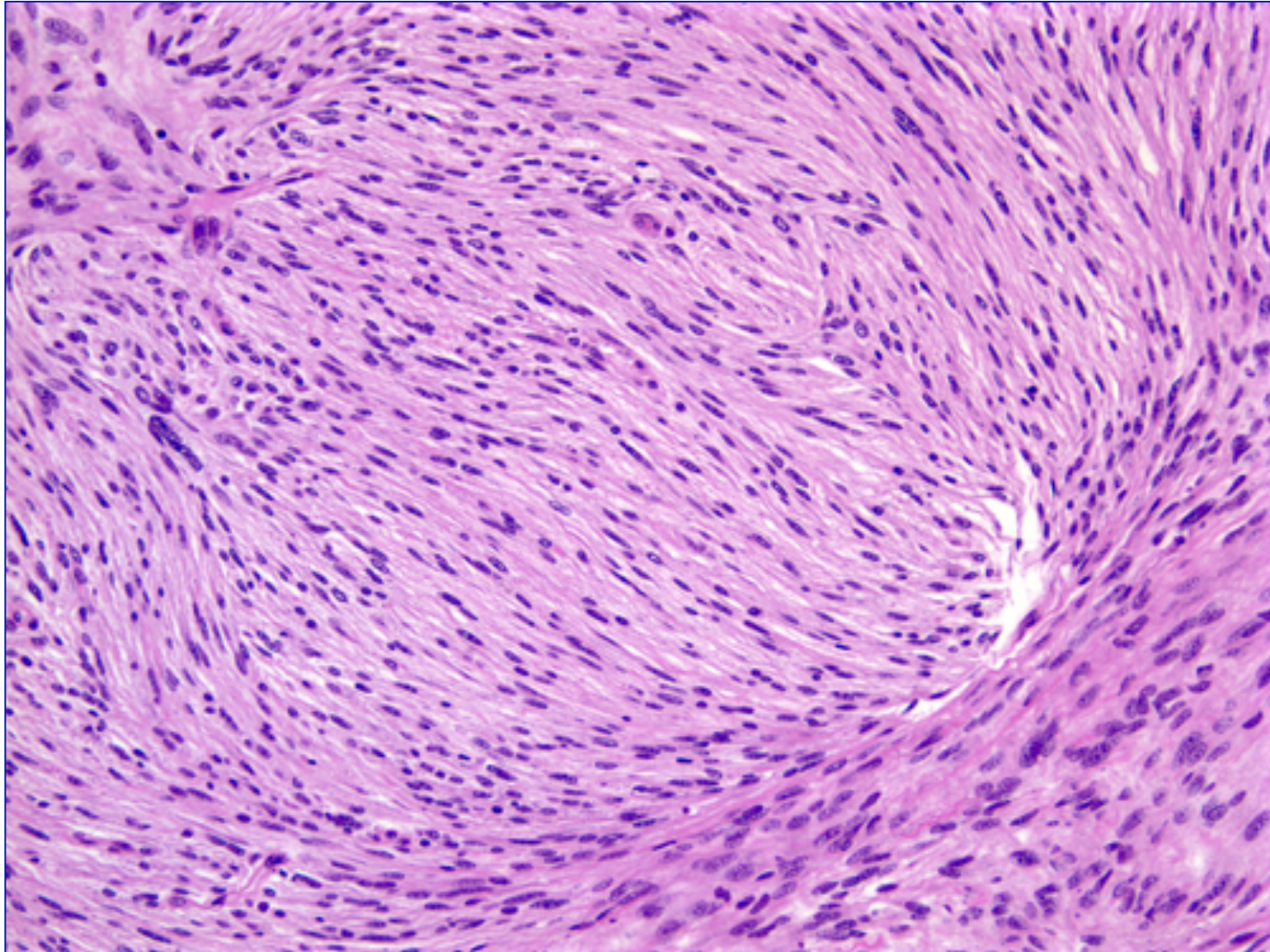
### Secretory

Intra-cytoplasmic lumina with eosinophilic (PAS+) bodies

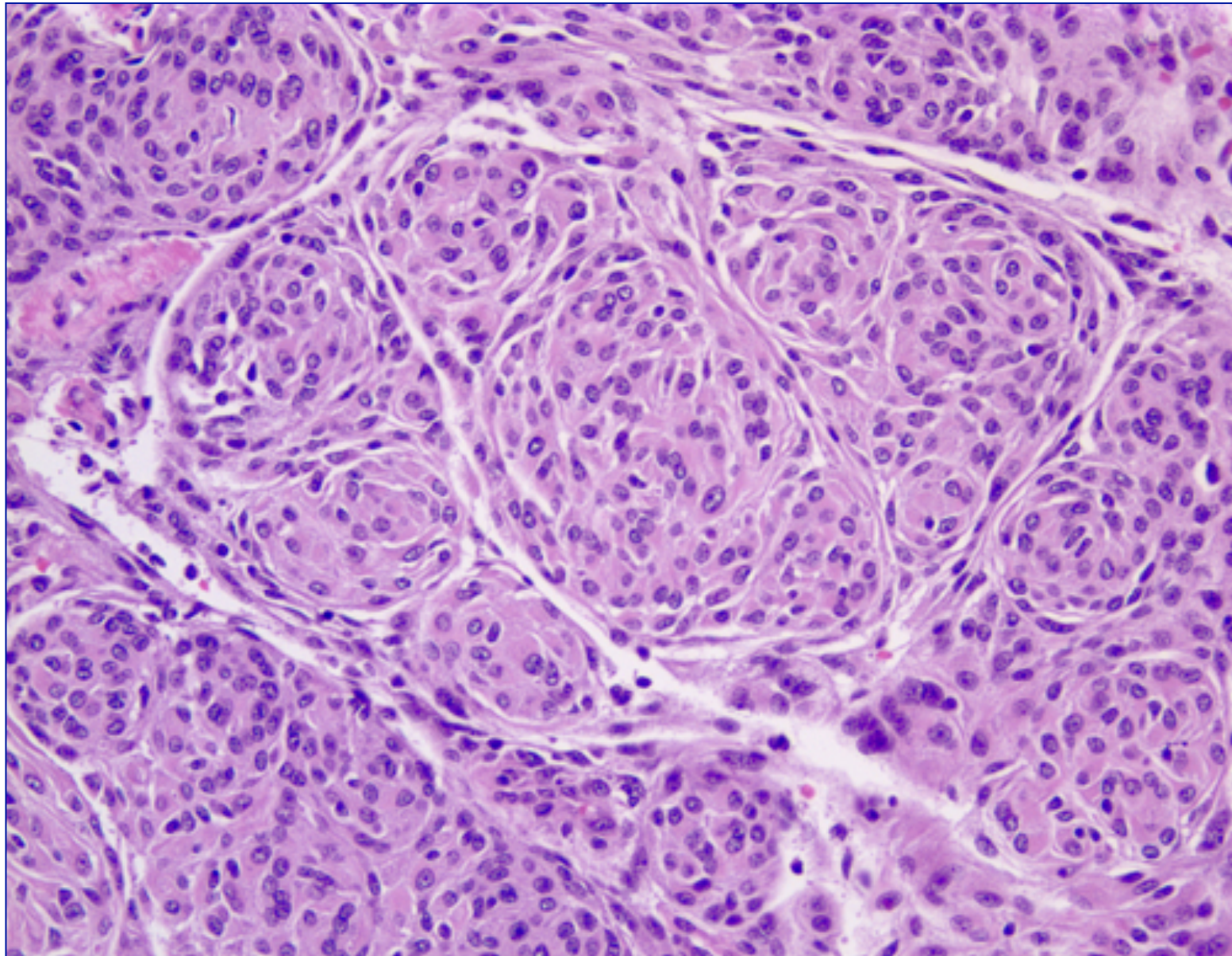
# Meningothelial



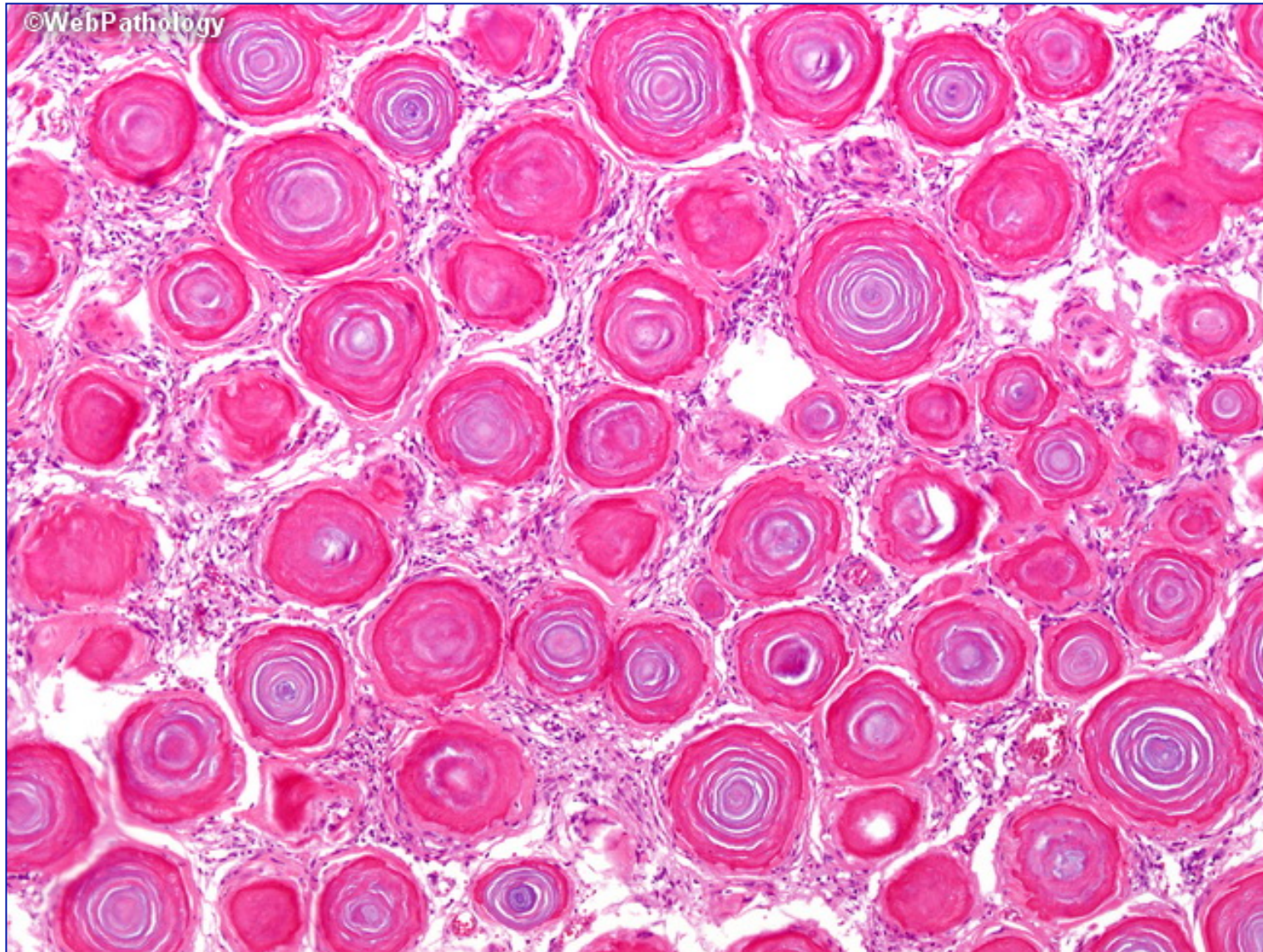
# Fibrous



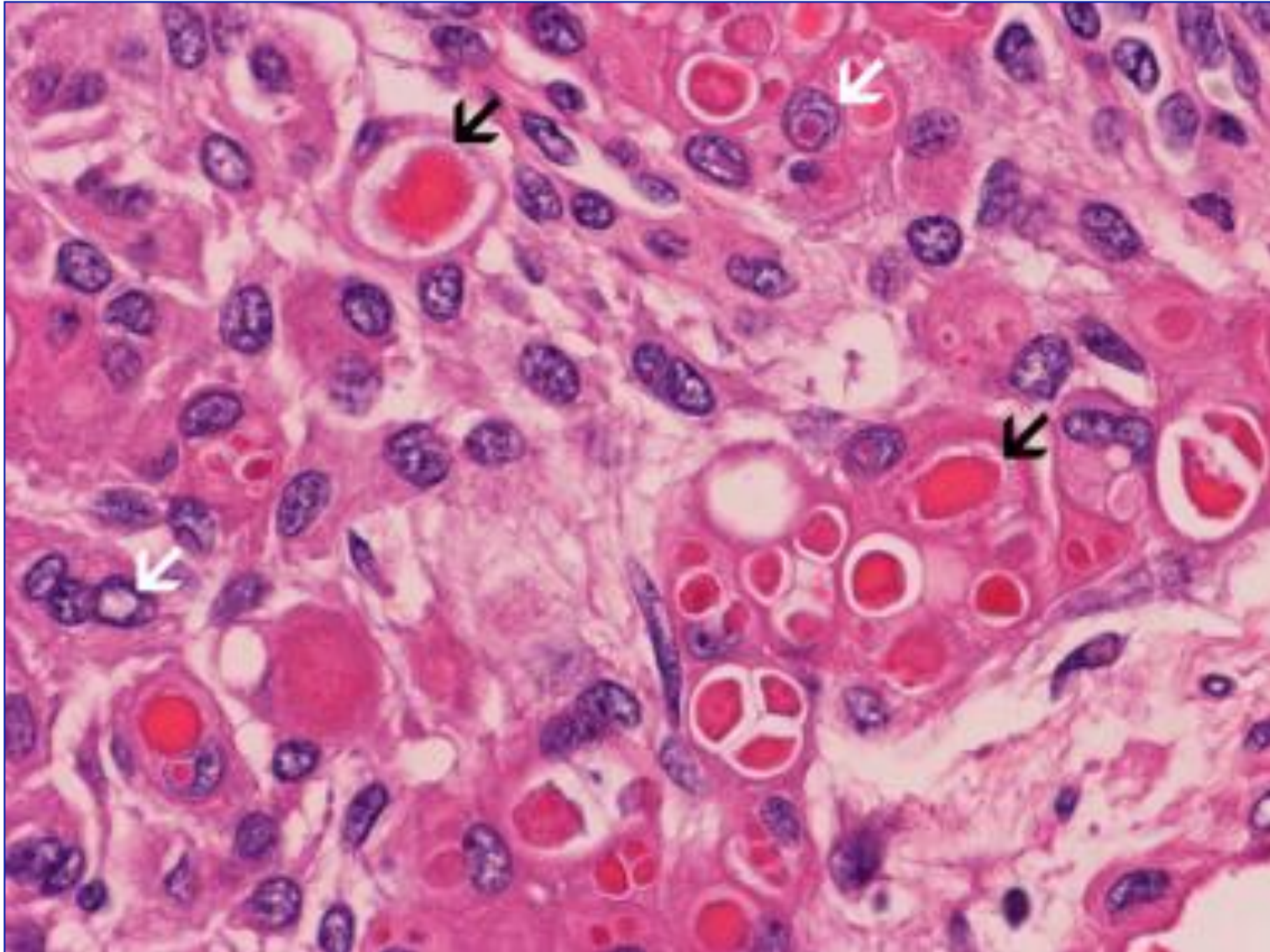
# Transitional



# Psammomatous



# Secretory





**Papillary**: younger age at presentation (childhood)  
aggressive course, recurrences, brain infiltration, metastases  
prominent papillary growth, perivascular rosettes

**Microcystic**: associated with severe oedema

**Clear cells**: glycogen-rich (PAS\*) (D.D: clear cell renal ca.)

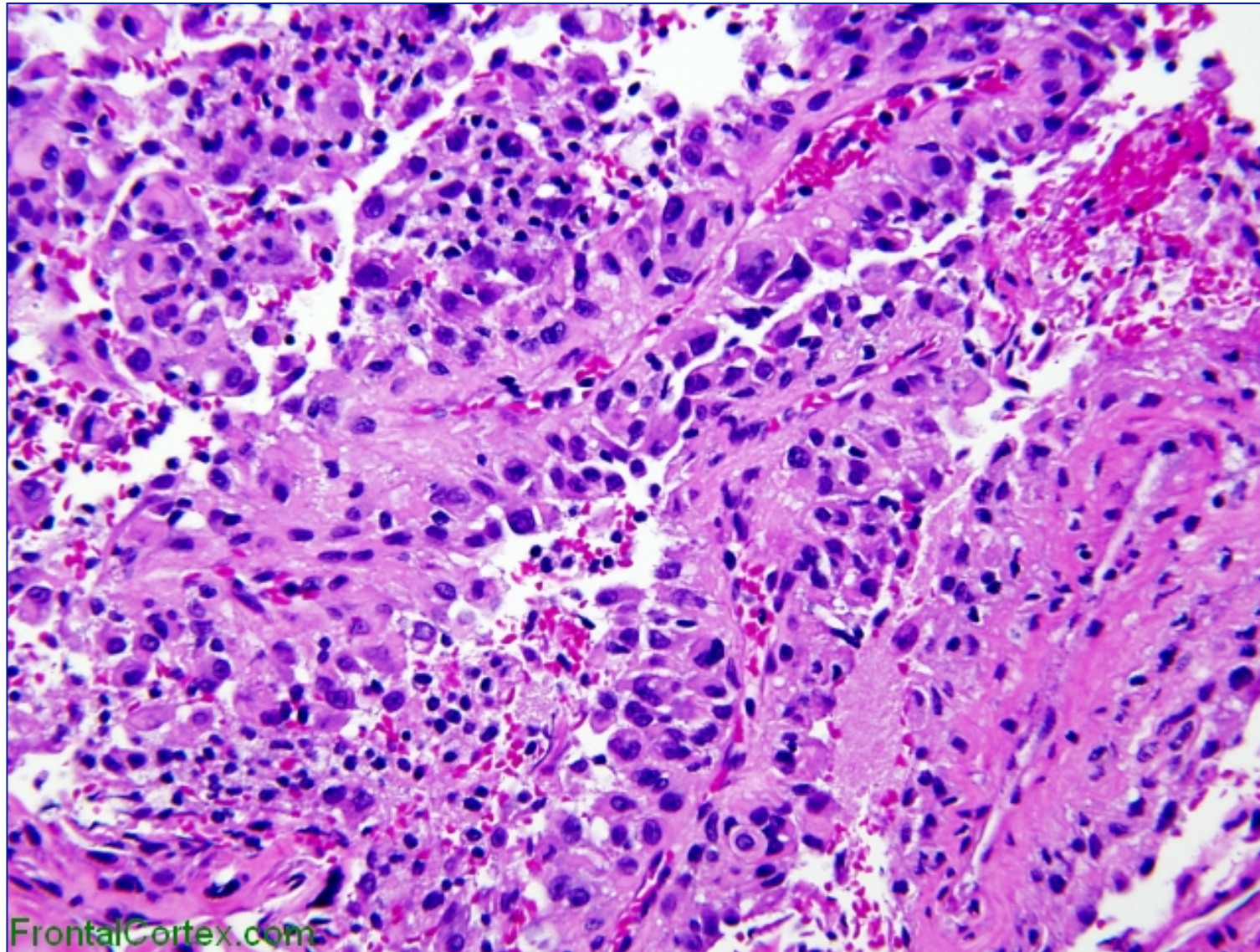
**Chordoid**: simulates chordoma

**Metaplastic**: bone, cartilage, myxoid tissue, xanthomatous cells

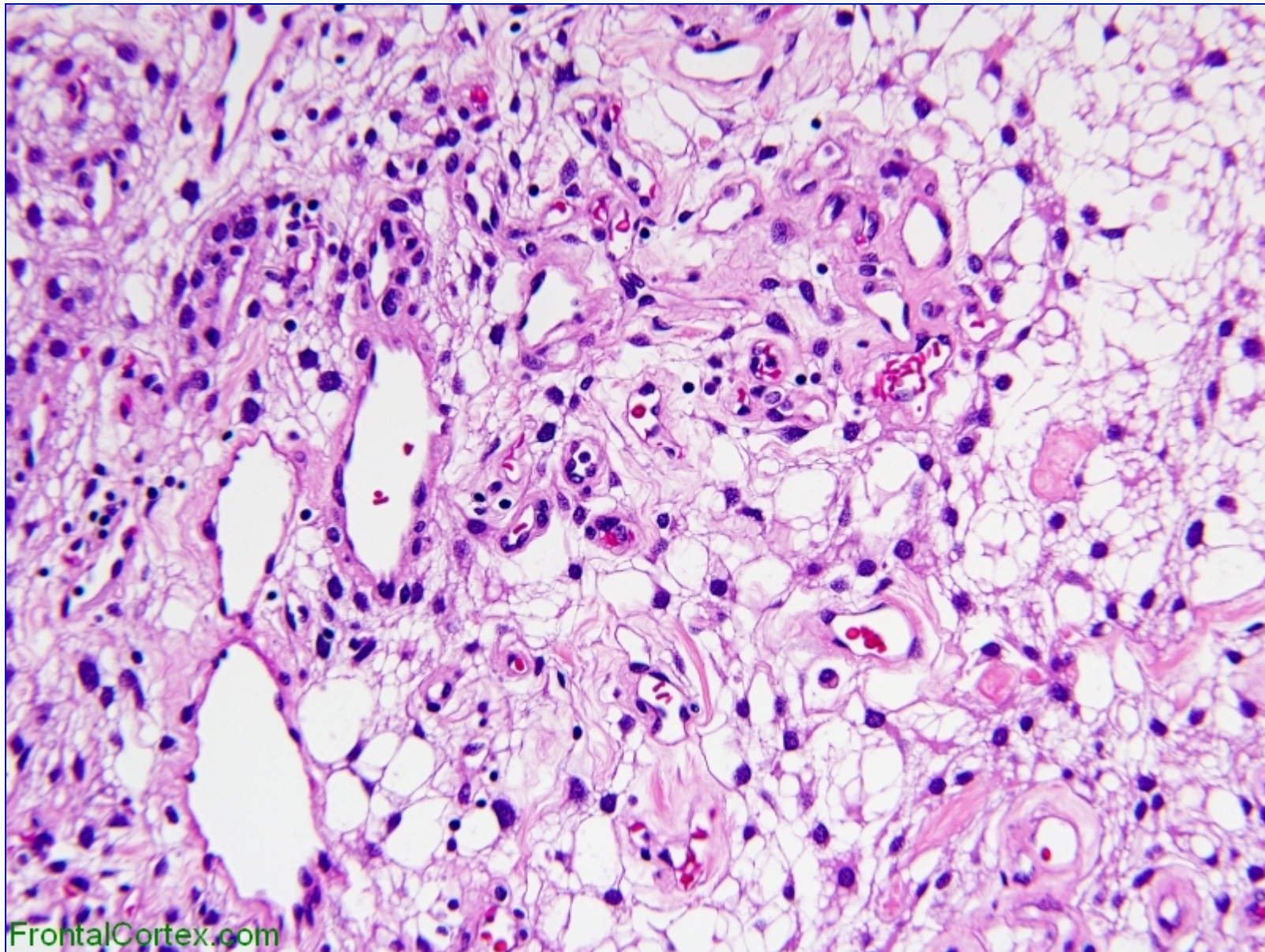
**Lympho-plasmacytic**: germinal centres, monoclonal gammopathy

**Oncocytic**

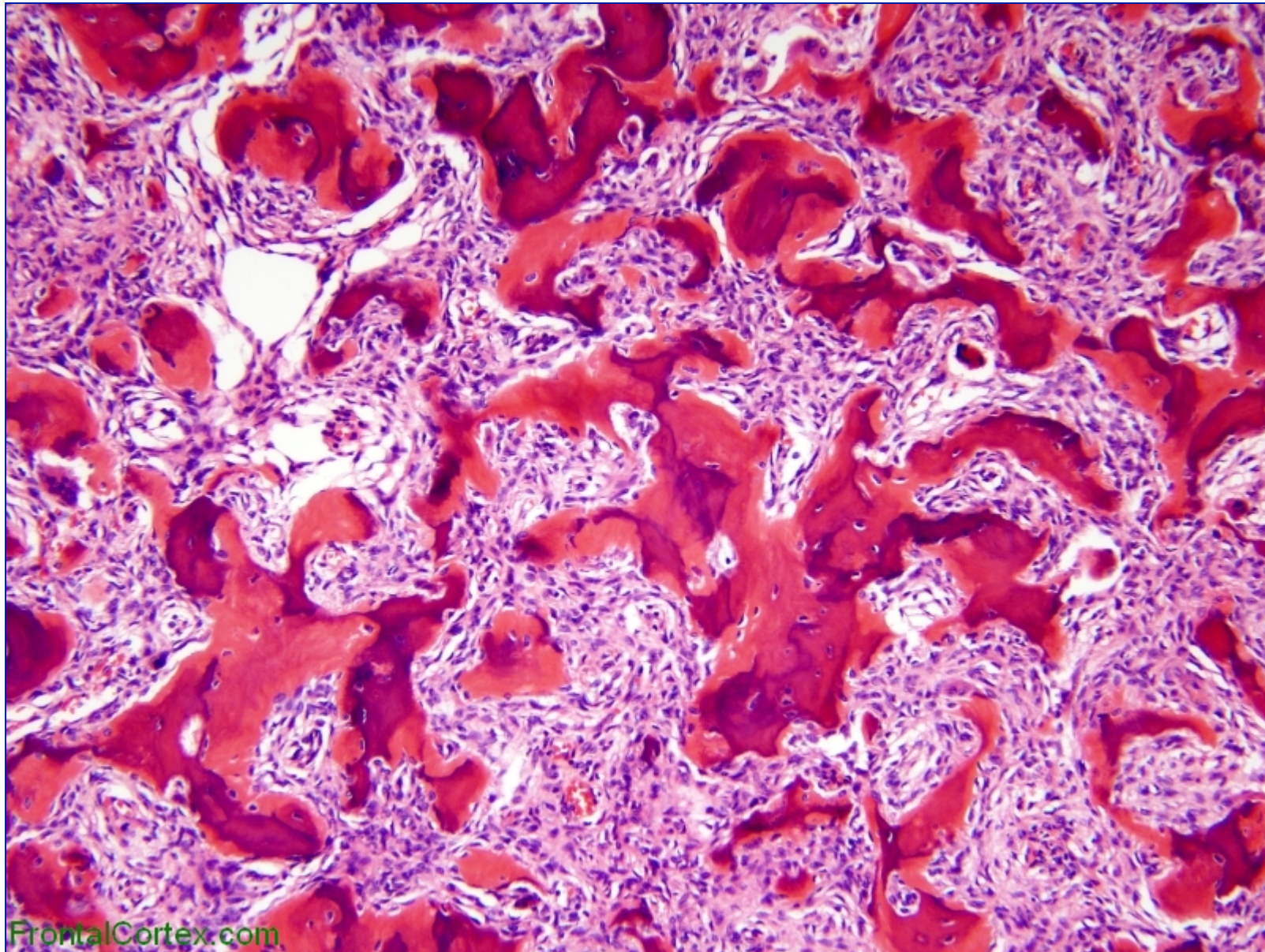
# Papillary



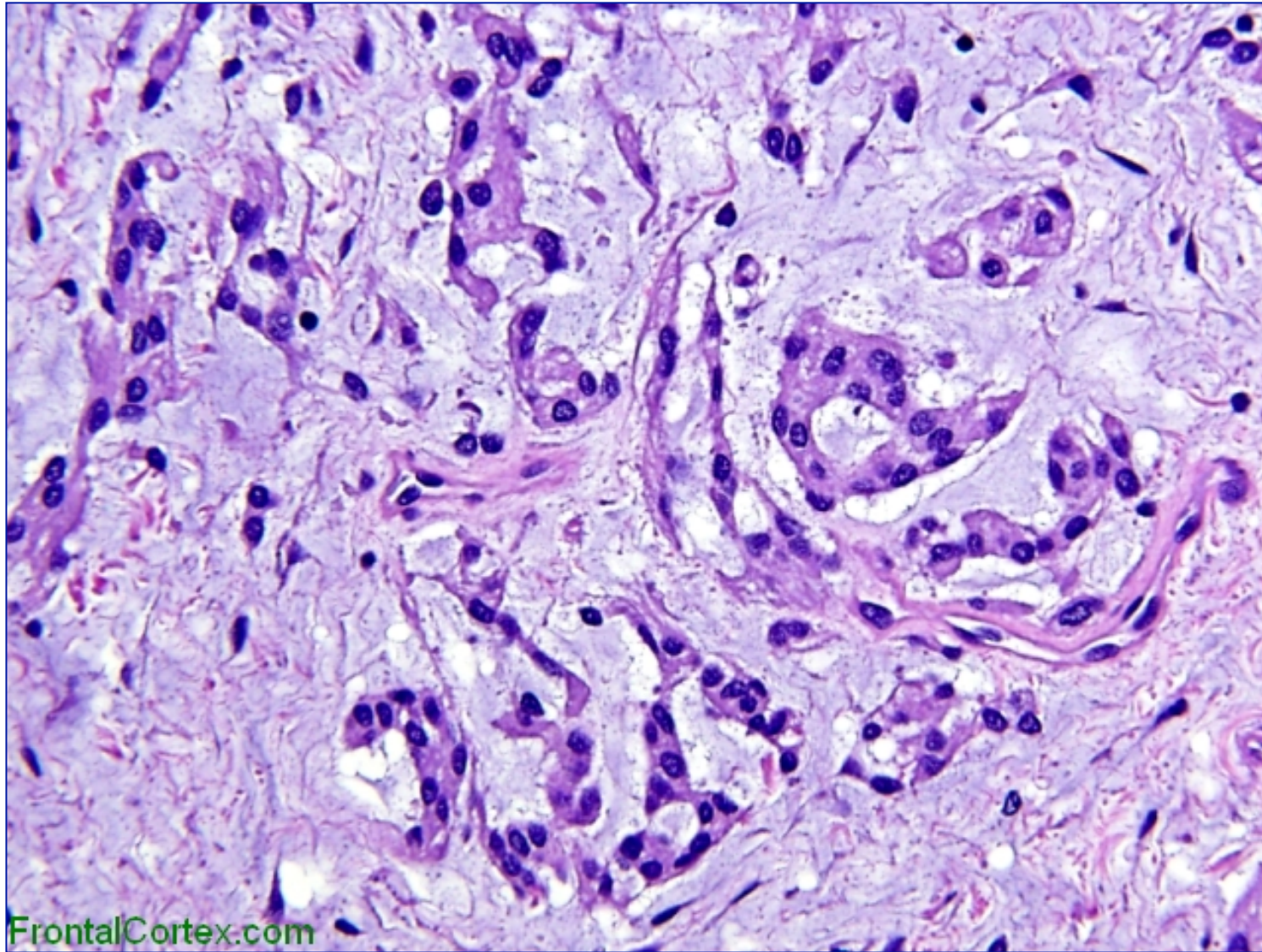
# Microcystic



# Metaplastic



# Chordoid





## Prognostic categories (WHO Grade):

**Classic (grade 1):** may occasionally recur

**Atypical (grade 2):** Focally hypercellular

Mitoses 5/10 HPF (40x)

Mild nuclear atypia and nucleoli

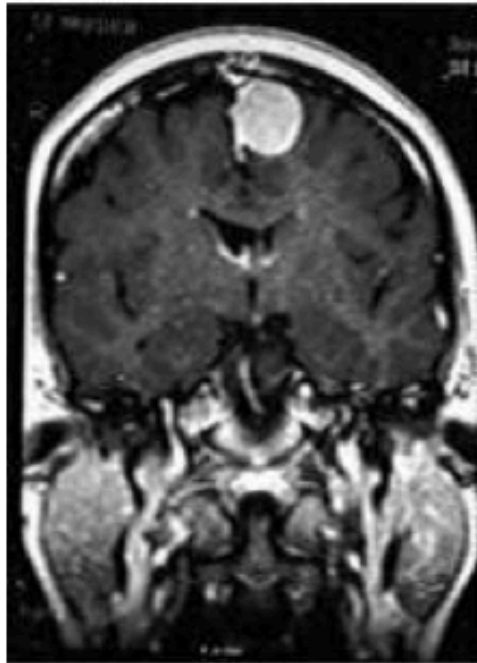
Small necrotic foci

Brain infiltration

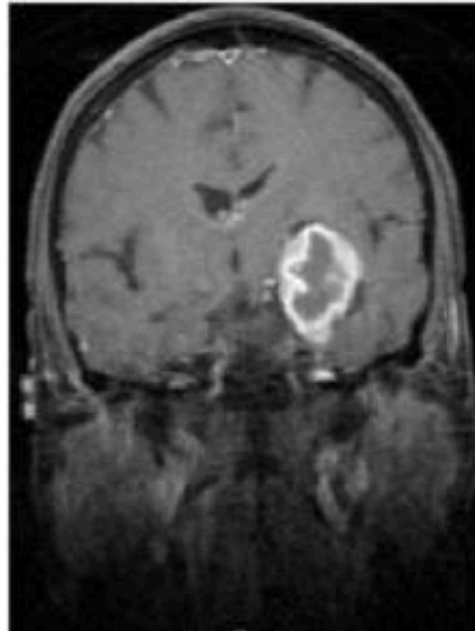
May recur, metastasize and  
progress to anaplastic

## How meningiomas are graded:

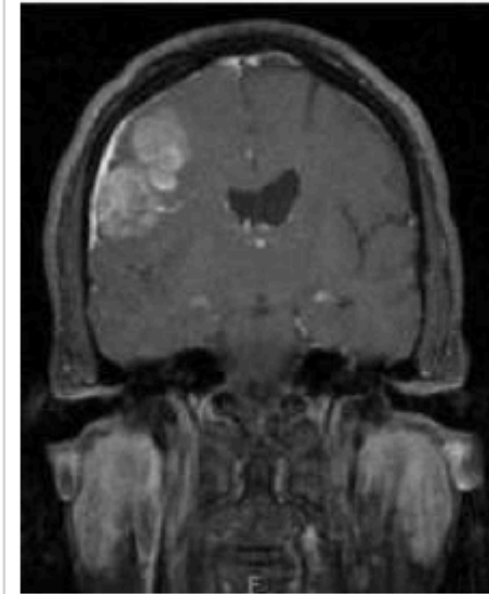
The grade (I to III) of a meningioma is based on the tumor cells' appearance under a microscope. Grade I is the most common and benign, and grade III is the most aggressive and is considered malignant.



*example of a grade I meningioma*



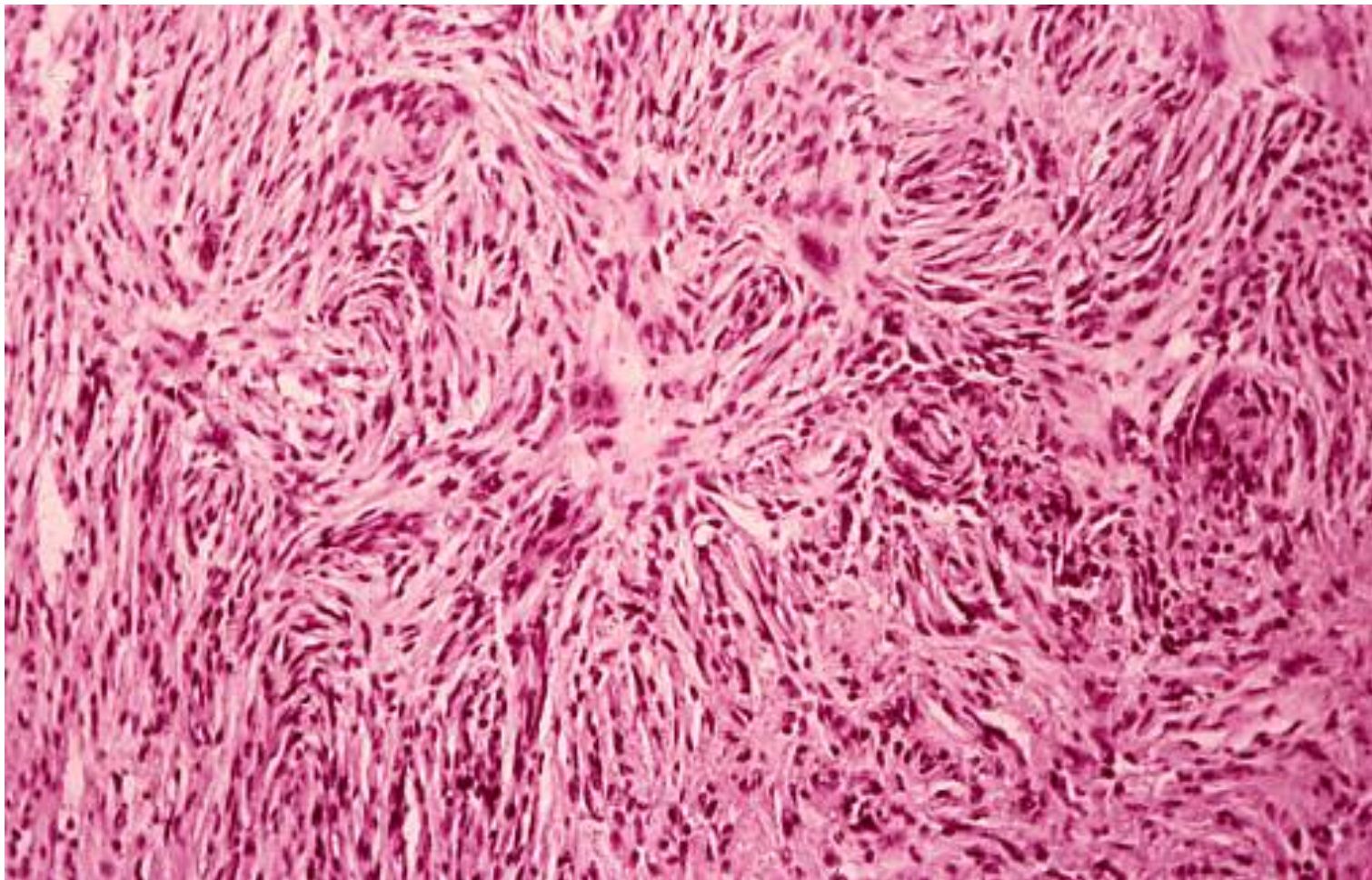
*example of a grade II meningioma*



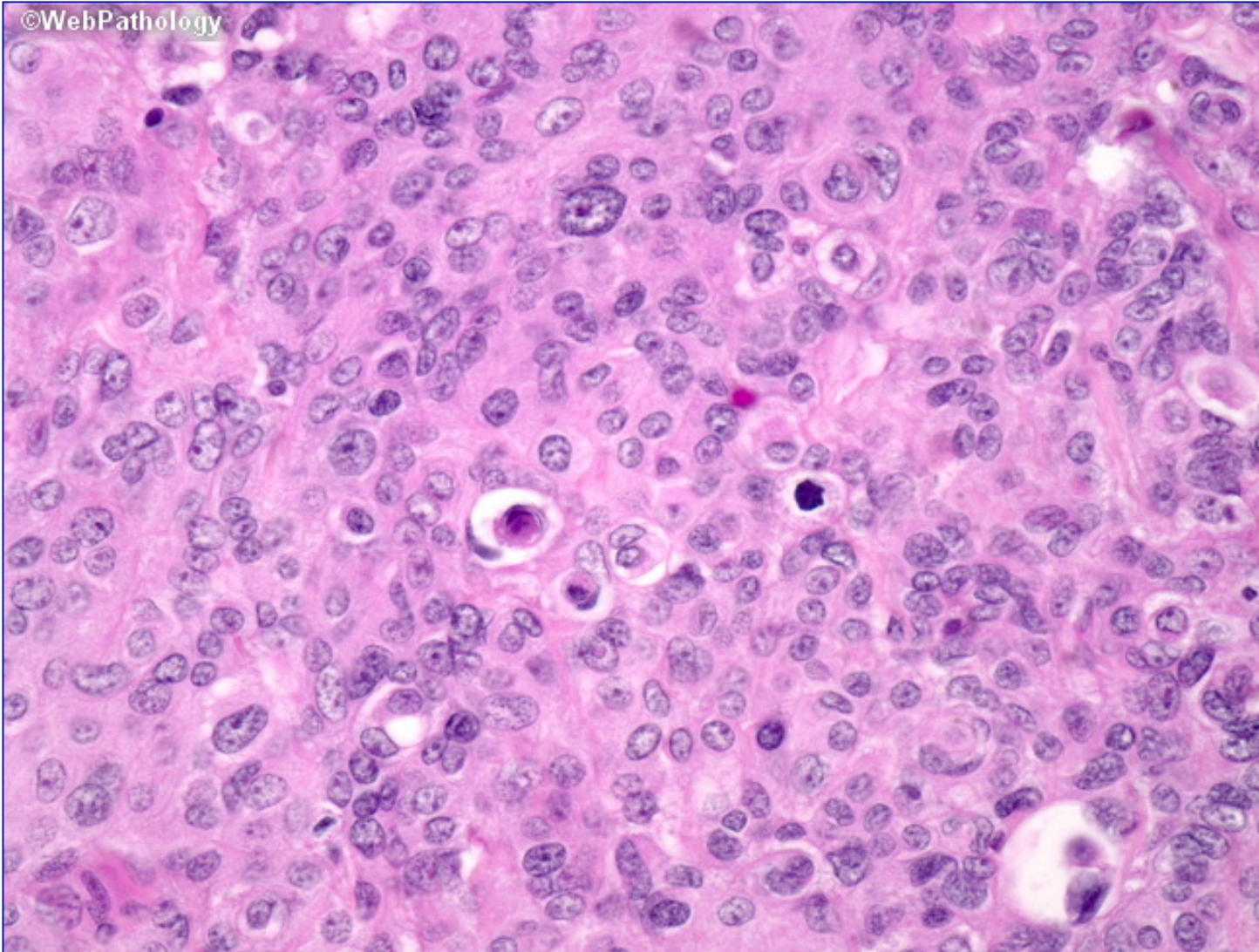
*example of a grade III meningioma*

There are three main grades (classifications) of meningiomas:

- **Grade I – Benign meningioma:** This non-cancerous type of brain tumor grows slowly and has distinct borders. Approximately 78-81% of meningiomas are benign (non-cancerous).
- **Grade II – Atypical meningioma:** Approximately 15-20% of meningiomas are atypical, meaning that the tumor cells do not appear typical or normal. Atypical meningiomas are neither malignant (cancerous) nor benign, but may become malignant. Grade II atypical meningiomas also tend to recur and grow faster.
- **Grade III – Malignant or anaplastic meningioma:** Malignant or anaplastic meningioma is an aggressive type of brain tumor that tends to invade the parts of the brain nearest to the tumor. Approximately 1-4% of meningiomas are grade III (cancerous).









## Prognostic categories:

**Anaplastic (grade 3):** Frankly hypercellular

Mitoses  $\geq 5/10$  HPF (40x)

Severe nuclear atypia and nucleoli

Extensive necrosis with palisades

Brain infiltration

Aggressive course with frequent metastases (meningiosarcoma)

