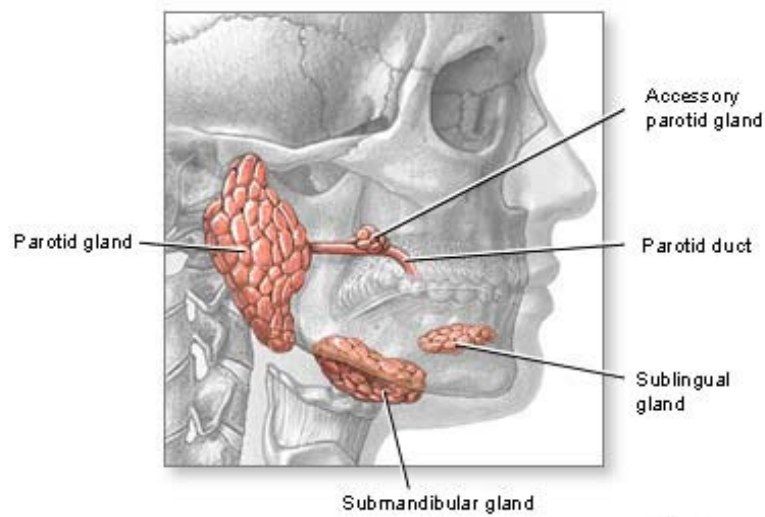
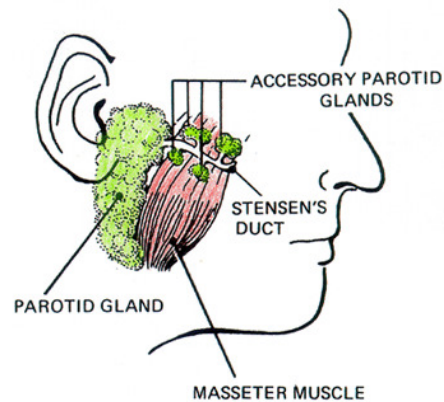


Salivary gland disorders



☑ Developmental anomalies:

- Aplasia
- Atresia
- Ectopic tissue



☑ Sialadenitis:

📖 Acute bacterial sialadenitis:

- Uncommon
- Parotid
- Xerostomia
- Seen in pts with:
 - ◆ Sjogren's syndrome
 - ◆ Tricyclic antidepressants



◆ Immunosuppression

◆ Obstructed glands

• **Microbiology:** mixed

• **Clinically:**

⤴ Rapid onset

⤴ Swelling

⤴ Pain

⤴ Fever

⤴ Malaise

⤴ Redness

⤴ LN

⤴ Pus

• **Hist:**

⤴ Intense AICI in acini, periductal tissue & within ducts

⤴ Ductal dilatation

📖 **Chronic bacterial sialadenitis:-**

• Ductal obstruction or ↓ secretion

• Submandibular G

• Unilateral

• **Clinically:**

⤴ Recurrent tender swelling

⤴ Inflamed orifice

⤴ Pus



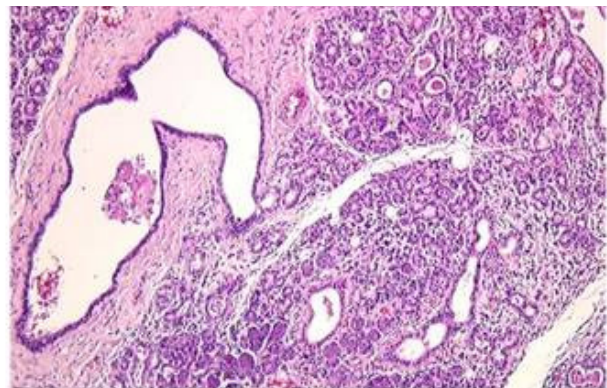
• **Hist:**

⤴ Destruction of acini

⤴ Duct dilatation & sq metaplasia

⤴ Scattered CICI

⤴ Interstitial fibrosis



📖 Recurrent Parotitis:

- Children
- Aetiology: ?
- **Clinically:**
 - ↗ Uni/bilateral recurrent painful swelling
 - ↗ ± Pus
 - ↗ **Most cases:** resolves spontaneously



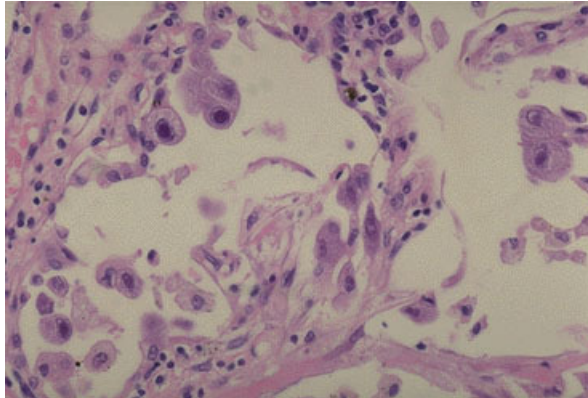
📖 Mumps:

- Common
- Children
- Highly infectious: airborne droplets
- Parotid
- **Clinically:**
 - ★ IP →
 - ↗ Fever & malaise
 - ↗ Rapid bilateral swelling
 - ↗ Pain
- Duration
- **Prognosis:**
 - Orchitis, Oophoritis, Encephalitis, Pancreatitis
- **Dx:** clinical, amylase, Abs to “s” & “v” Ags
- Immunity



📖 Cytomegalic inclusion disease:

- Asymptomatic
- large, doubly contoured “owl-eye” inclusion bodies



📖 Radiation induced sialadenitis:-

- Sensitive
- Serous acini
- ↑ dose → Irreversible fibrous replacement & sq metaplasia
- Complications

📖 Sialadenitis of minor SGs:-

- MEC
- Necrotic stomatitis
- Sarcoidosis
- Sjogren's syndrome

📖 Obstructive & traumatic lesions:

- **Causes:**

- **Sialolithiasis:**

- Submandibular 80%, parotid 6%, sublingual & minor SGs 2%

- Gland or duct

- **Clinically:**

- Adults, 2M: 1F

- Unilateral

- Can be multiple

- Most cause symptoms: pain & swelling, retrograde infection



- **Dx:**

- Palpation

- Rx: parotid 40%, SMG 20%



- Sialography



- **Pathogenesis:** Ca salts around a nidus of organic material

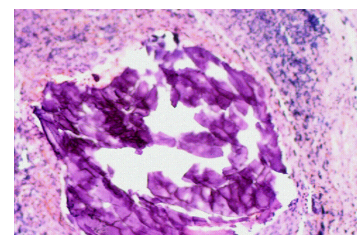
- **Hist:-**

- ★ **Stone:**

- **Grossly:** yellowish-white, round/oval, rough/smooth

- **Decalcification:** laminated w concentric rings w bacteria

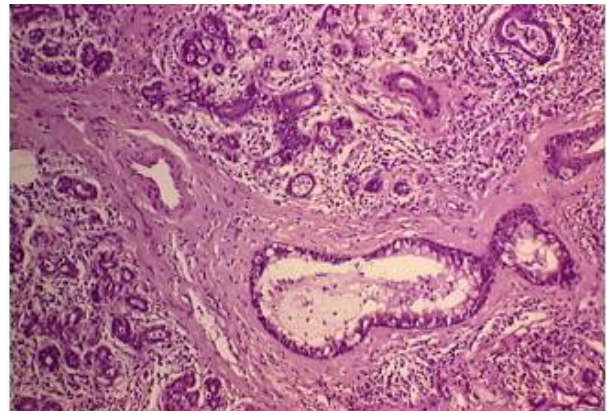
- Ca, PO₄, Bicarbonate



- ★ **Gland changes:**

- Duct dilatation w sq metaplasia

- Acinar atrophy & replacement fibrosis
- Periductal CICI & fibrosis
- CICI of lobules



● Parotid papilla & duct stricture:

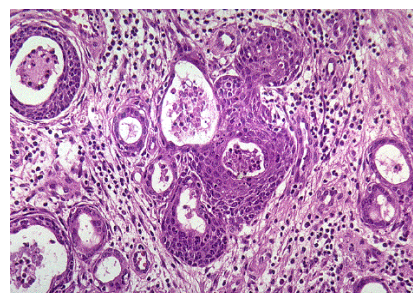
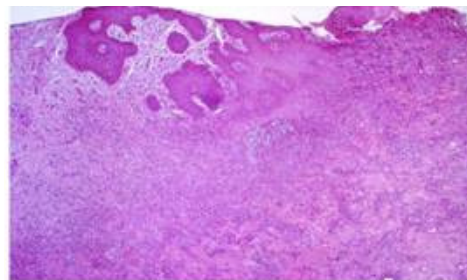
- Chronic trauma to parotid papilla
- Surgery
- Ulceration around a stone

○ Necrotizing sialometaplasia:

- ? aetiology
- Spontaneous necrosis
- Site: H-S palate
- **Clinically:**
 - Middle age
 - Deep ulcer ≈ 2cms
 - Painless
 - 2 months



- **Hist:**
 - Necrosis of SG lobules
 - Sq metaplasia of ducts & acini
 - ME
 - CICI
 - Pseudoepitheliomatous hyperplasia



○ Sialadenosis (Sialosis):-

- **Definition:**
- **Aetiology:** ?, defect in neurosecretory control
- Alcoholism, DM, pregnancy, Bulimia N., drugs, malnutrition, cirrhosis
- **Hist:** hypertrophy of acini & oedema of CT



📖 Sjogren's syndrome:-

- **Pathogenesis:**
 - Immune-mediated CI disease
 - Progressive lymphocytic infiltration & replacement of glandular parenchyma
 - Xerostomia & Xerophthalmia
- **Clinically:**
 - ≈ 1% of population
 - Middle age
 - Females 80%
 - **Two types:**
 - ❶ **Primary SS (Sicca S):** xerostomia & Xerophthalmia
 - ❷ **Secondary SS:** xerostomia & xerophthalmia + CT disease
 - ✧ RA (15%)
 - ✧ SLE (30%)
 - ✧ Systemic sclerosis
 - ✧ Primary biliary cirrhosis
 - ✧ Dermatomyositis
 - ✧ Mixed CT disease
 - **Eyes:** Keratoconjunctivitis sicca
 - ✧ Corneal keratotic lesions



✘ Failure of tear secretion: **Schirmer test**

✘ Conjunctivitis

➤ **Mouth:**

✘ Discomfort

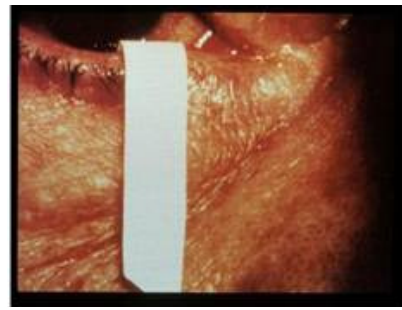
✘ Difficulties in eating, swallowing & speech

✘ Disturbed taste sensation

✘ Candidosis, sialadenitis, root caries

✘ Dry, red & shiny mucosa

✘ Red, atrophic, lobulated (cobble-stone) tongue



➤ **Others:**

✘ Severe tiredness & Arthralgia

✘ Xeroderma, nasal dryness, vaginal dryness

✘ Sinusitis, tracheitis, dysphagia, atrophic gastritis, pancreatitis, purpura

✘ Anaemia, leucopenia, thrombocytopenia

✘ SG enlargement



• **Hist:-**

✘ Lymphocytic infiltrate (T)

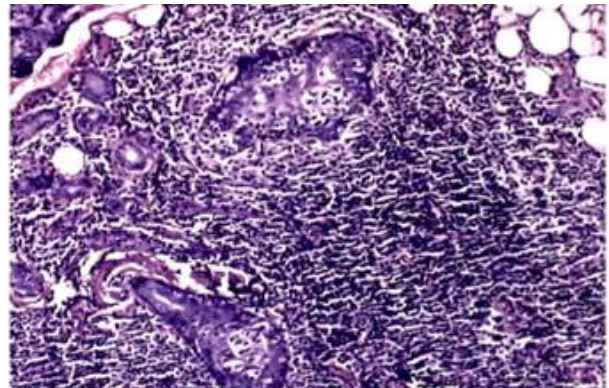
✘ Intralobular ducts

✘ Glandular tissue

✘ Acinar atrophy

✘ Epimyoepithelial islands

✘ Interlobular septa & capsule



• **Dx:**

✘ Mixed SFR

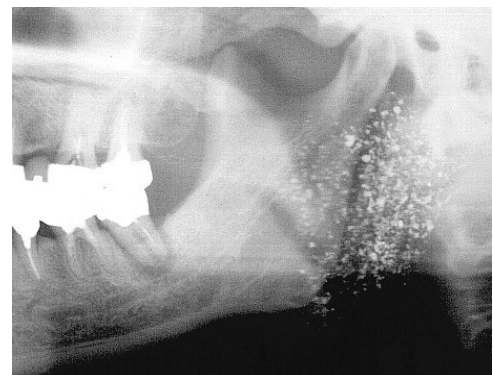
✘ Tear

✘ **SG biopsy**

✘ **Sialography:** sialectasis, “snowstorm”

✘ Salivary **scintiscanning** [^{99}Tm] pertechnetate

✘ **Ab screening:** RF, ANA, SS-A, SS-B



☒ **Serology:** ↑ESR, hypergammaglobulinaemia

- **Aetiology:**

- ☒ Unknown

- ☒ Autoimmune: ?Ag

- ☒ CMV, EBV, HHV-6, Retrovirus

- **Complications:**

- ☒ B-cell malignant lymphoma

☒☒ **Salivary gland tumours:**

- Uncommon

- **80% in major Gs:** parotid ≈ 90%, SMG ≈ 10%, SLG ≈ 0.3%

- **20% in minor Gs:** palate ≈ 55%, UL ≈ 20%, others ≈ 25%

- **Malignancy:**

- ☒ **Parotid:** ≈ 15%

- ☒ **SMG:** ≈ 30%

- ☒ **SLG:** ≈ 86%

- ☒ **Minor SGs:** ≈ 50%

- **Classification:**

- ⇒ **Adenomas: slow growing, soft or rubbery, x ulcer, x N**

- Pleomorphic adenoma

- Warthin tumour

- Basal cell adenoma

- Oncocytoma

- Canalicular adenoma

- Ductal papillomas

- ⇒ **Carcinomas: ≈ fast-growing, ± hard, ± ulcerate, ± invasion, ± N**

- Mucoepidermoid Ca

- Adenoid cystic Ca

- Acinic cell Ca

- Ca in PA
- Polymorphous low-grade adenoCa
- Others

☒ **Pleomorphic A:**

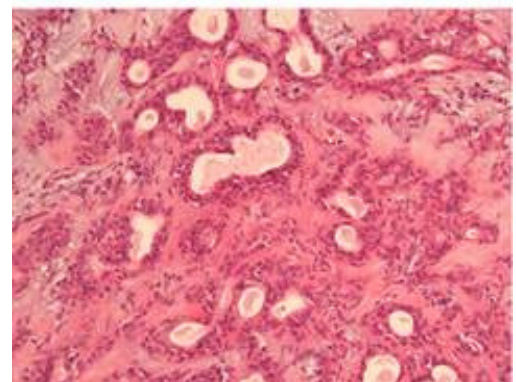
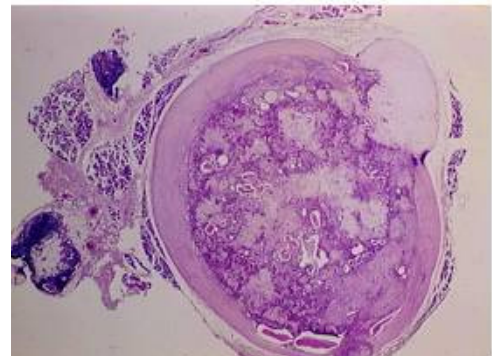
- Commonest
- 70% of PG, 50% of SMG, 7% in minor Gs (palate 55%)
- “Pleomorphic”, “mixed tumour”.
- Origin: epithelial &/or myoepithelial Cs
- **Clinically:**



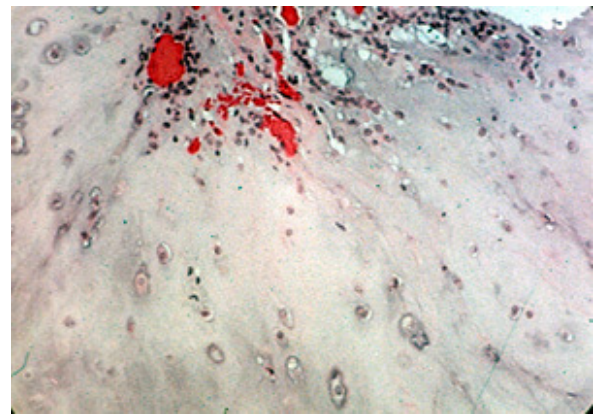
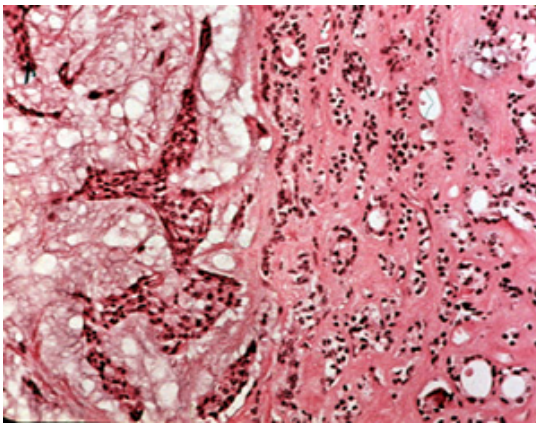
- ◎ 5th & 6th decades, F>M
- ◎ Slowly-growing, w-d, painless, rubbery swelling
- ◎ Superficial lobe: spherical mass overlying angle of M
- ◎ Deep lobe: lateral pharynx
- ◎ Lobulated or multinodular

- **Hist:**

- ÷ Extensive variations
- ÷ Fibrous capsule
- ÷ Epithelial & stromal components
- ÷ **Epithelial duct Cs:** line duct-like structures
- ÷ **Myoepithelial Cs:** spindle/stellate-shaped → sheets, clumps & strands
- ÷ Plasmacytoid Cs
- ÷ Sq metaplasia
- ÷ **Abundant stroma:**
 - Fibrous/hyalinization
 - Myxoid
 - Chondroid



- **Malignant transformation:** < 1%, Ca ex PA



☒ **Warthin tumour:**

- Hamartoma
- **Origin:** duct Cs in LNs → Adenolymphoma
- **Clinically:**
 - © Almost exclusively in parotid, 9% of parotid tumours
 - © M>F
 - © 5th & 6th decade
 - © Bilateral in 10% of cases
 - © Multiple in same gland



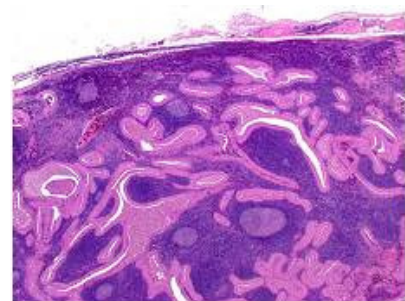
• **Hist:**

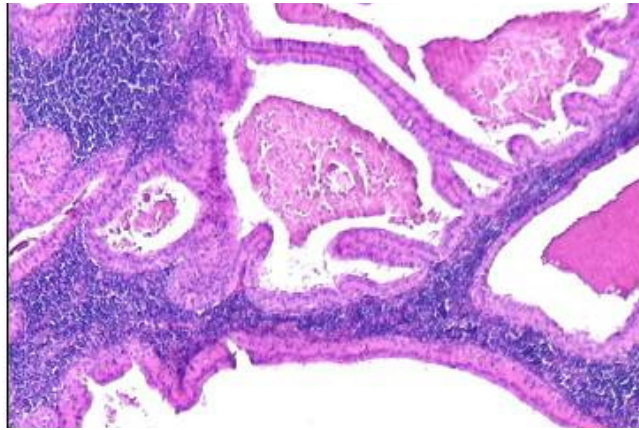
☒ **Grossly:** FC, irregular cystic spaces, mucoid material, papillary projections

☒ **Microscopically:**

Papillary fronds:

- Double layer of epith Cs
- Basal cuboidal Cs & superficial tall columnar Cs
- Marked, granular eosinophilic cytoplasm (Oncocytes)
- Supported by lymphoid tissue





☒ Basal Cell Adenoma:-

- 75% in parotid & 20% in UL

- 7th decade

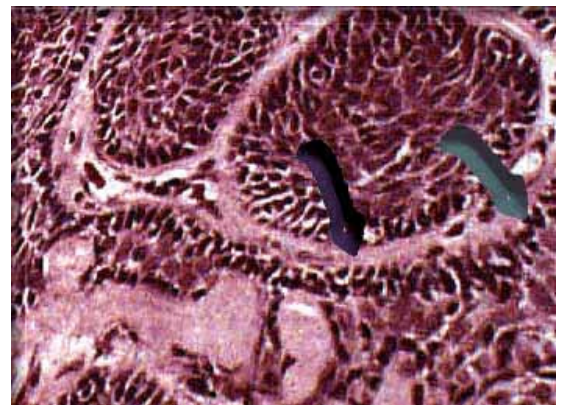
- **Clinically:** ≈ PSA

- **Hist:**

- Well-encapsulated

- Cytologically uniform basaloid Cs

- Different forms: solid, trabecular, tubular, membranous



☒ Oncocytoma:

- **Oncocyte:** abnormal C w prominent eosinophilic granular cytoplasm

- Rare

- Parotid

- F, 8th decade

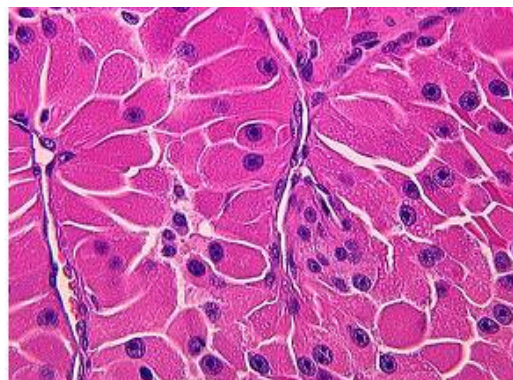
- **Clinically:** ≈ PSA

- **Hist:**

- Thin capsule

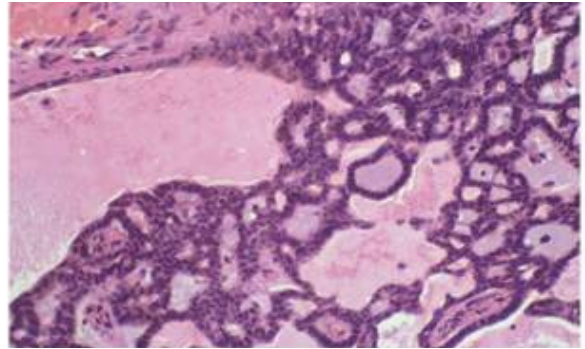
- Oncocytes arranged in solid cords or acinar pattern

- Lacks a fibrous stroma



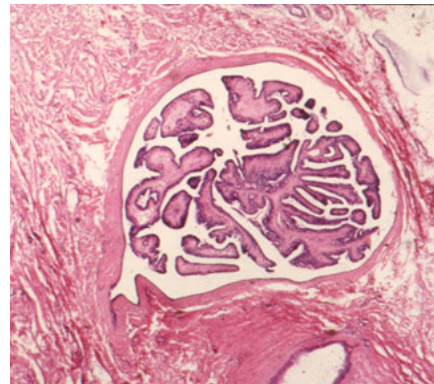
☒ **Canalicular adenoma:**

- 7th decade
- 75% in UL
- Rare in major SGs
- **Hist:**
 - Capsule
 - Single-layered cub/column ductal Cs
 - Arranged in elongated anastomosing cords
 - Myxomatous stroma & dilated capillaries
 - Degeneration of stroma



☒ **Ductal papillomas:**

- Rare
- **Types:**
 - ⇒ Sialadenoma papilliferum
 - ⇒ Inverted ductal papilloma

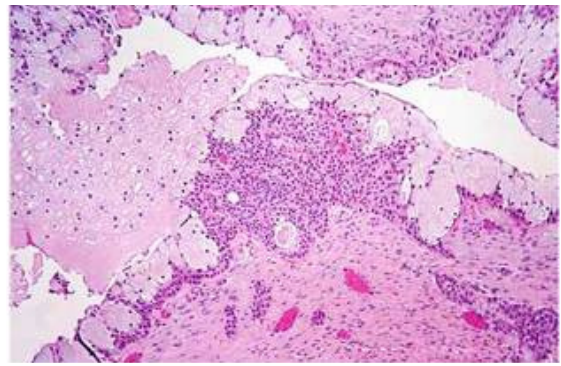


☒ **Mucoepidermoid Ca:**

- 4th & 5th decade, F
- 50% parotid, 20% palate
- 10-15% of minor SG tumours
- **Clinically:**
 - ≈ PSA or signs of malignancy
- **Hist:**

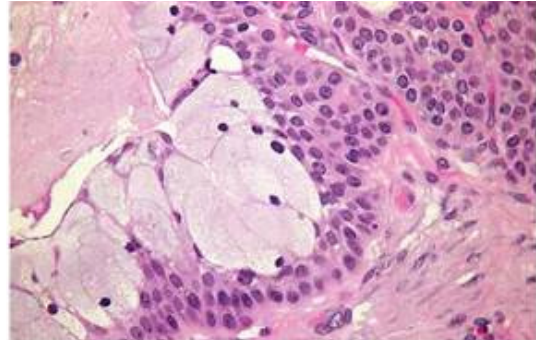


- No capsule
- Mucous, squamous & intermediate
- Nests, diffuse sheets & cystic spaces
- Invasion
- Low-grade or well-differentiated
- High-grade or poorly differentiated



- **Prognosis:**

- **Low-grade:** LRR < 10%, 95% 5-year SR
- **High-grade:** LRR 80%, 30% 5-year SR

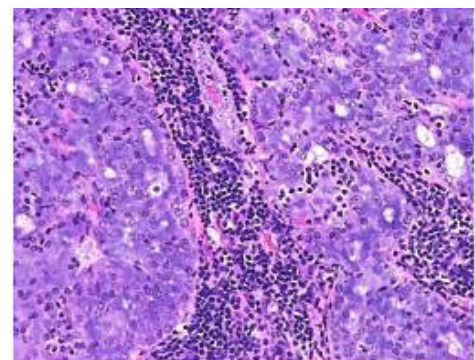


☒ **Acinic cell Ca:**

- Parotid
- **Clinically:**
 - Any age, F>M
 - ≈ PSA



- **Hist:**
 - Large Cs w granular basophilic cytoplasm often in acinar pattern
 - Solid, microcystic, papillary cystic, follicular
 - Very scanty stroma
 - W-d but lobulated
 - Some are poorly differentiated



- **Prognosis:**
 - Generally low-grade w slow growth potential
 - **Low-grade:** 80-100% 5y-SR
 - **Poorly differentiated:** 65% 5y-SR

⊗ Adenoid cystic Ca:

- **Clinically:**

- 7th decade
- ≈ 30% of minor SG tumours
- ≈ 3-6% of parotid G tumours
- May ≈ PSA
- Pain, ulceration, facial paralysis, paraesthesia



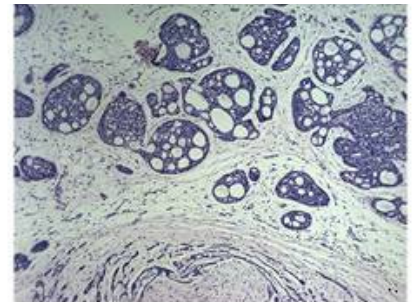
- **Hist:**

- Oval nests of cuboidal or polygonal Cs w hyperchromatic nuclei

- **Three main patterns:**

- **① Cribriform pattern:**

- Classic form
- Islands punctuated w multiple microcystic spaces
- “Swiss cheese” or “honeycomb” appearance
- Eosinophilic or basophilic substance



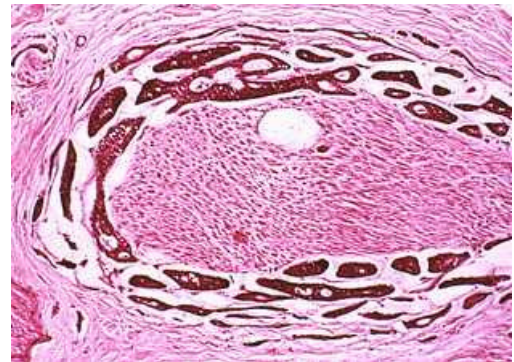
- **② Tubular pattern:**

- Small ductal elements lined by basaloid Cs

- **③ Basaloid pattern:**

- Solid nests of basal Cs

- Nerotropism
- Marrow spaces



- **Prognosis:**

- 75% 5-y SR, 40% 10-y SR, <20% 20-y SR
- Basaloid

⊗ Ca ex PSA:

- PSA (many ys & LR → pain, palsy, ulceration)

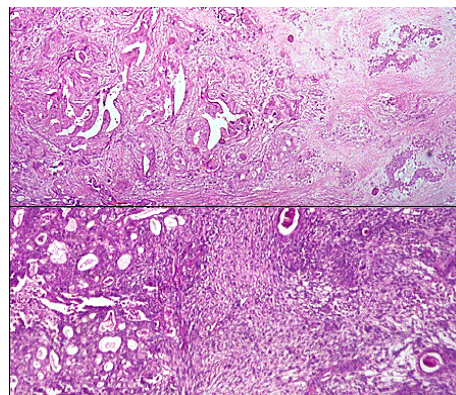
- Uncommon

- Parotid

- **Hist:**

- Ca features adjacent to PSA
- AdenoCa, undifferentiated Ca

- **Prognosis:**



☒ Polymorphous low-grade adenoCa:-

- Palate
- **Hist:**
 - Cytologically benign-looking pale-staining Cs
 - Tubular, lobular, papillary, papillary cystic, Cribriform
 - Invasion & spread



☒ Other Ca:

- AdenoCa (NOS)
- Basal Cell AdenoCa
- SCC
- Sebaceous Ca
- Undifferentiated Ca