BENIGN NECK DISEASES

BY

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CLASSIFICATION

- CONGENITAL: LYMPHANGIOMA, HEMANGIOMA, DERMOID CYST, BRANCHIAL CYST AND FISTULA, THYROGLOSSAL CYST, THYMIC CYST
- ACQUIRED : RANULA, LARYNGOCELE, PHARYNGO-ESOPHAGEAL DIVERTICULA
- INFECTED: BACTERIAL, VIRAL, TUBERCULAR LYMPHADENOPATHY & VARIOUS NECK SPACE INFECTION
- PARAPHARYNGEALTUMORS
- NEUROGENICTUMORS

- LYMPHATIC SYSTEM ARISES FROM 5 PRIMITIVE SACS
- o 2 JUGULAR SACS
- o 2 POSTERIOR SCIATIC SACS
- 1 RETROPERITONEAL SAC

- 3 TYPES
- i. SIMPLE LYMPHANGIOMA (40%)
- ii. CAVERNOUS LYMPHANGIOMA (35%)
- iii. CYSTIC HYGROMA (25%)



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- SIMPLE LYMPHANGIOMA
- o Thin walled capillary sized lymphatic channel
- Occur as pale fluctuant lesion where tissue planes are tight
- o Lips, tongue, cheek, FOM



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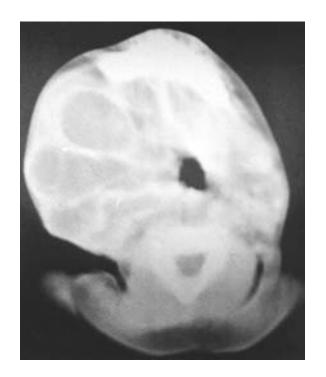
- CAVERNOUS LYMPHANGIOMA
- o Dilated lymphatic spaces often with fibrous adventitia
- o Mainly occur in tongue- BOT, lateral margin
- BOT lesions has to be differentiated from carcinoma, lingual thyroid, internal laryngocele
- CYSTIC HYGROMA
- o Earliest swelling one can see in neck
- Occur as cysts of various size
- o Commonly in neck where tissue planes are more lax
- o May spread into cheek, FOM, tongue, parotid, ear canal

- CLINICAL PRESENTATION
- No sex predominance
- o No side predominance
- o 35% oral, 25% cervical, 15% axillary
- o 60% by birth, 75% by first year, 90% by second year

- CLINICAL PRESENTATION
- o Usually painless neck swelling, more in posterior triangle
- Can encroach upper airway, pharynx, esophagus, to produce stridor or dysphagia
- o May compress brachial plexus to produce pain or hyperaesthesia
- o May have pain or sudden increase in size due to infection or hemorrhage

- ON EXAMINATION
- o Non-tender , ill defined, soft and spongy lesions containing clear fluid
- Translucent (cystic hygroma brilliantly translucent)
- o Fluctuation and fluid thrill present
- o Partially compressible
- o Impulse on cough present

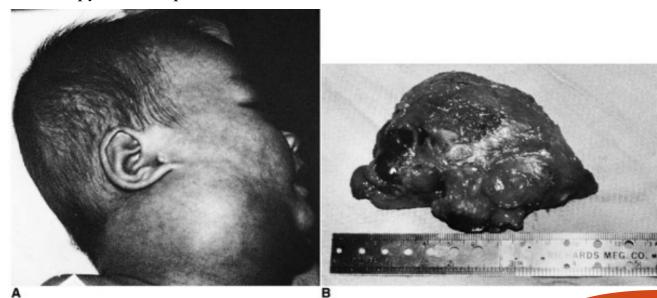
- DIAGNOSIS
- Clinical
- Cytology
- Radiology
- ❖ To know extent and size of swelling
- A chest radiology should invariably be done in case of neck lesions before surgery to rule out chest involvement



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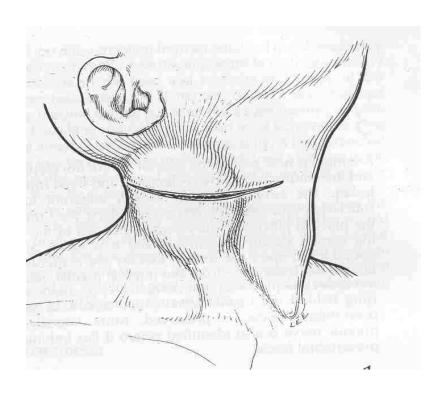
- TREATMENT
- Sclerosant injection- recurrence is more
- o Laser excision- for intraoral submucosal lesions
- o Cryoapplicatioin- for submucosal lesions
- o Surgical excision- definitive treatment
- o Radiotherapy- in un-operable recurrences

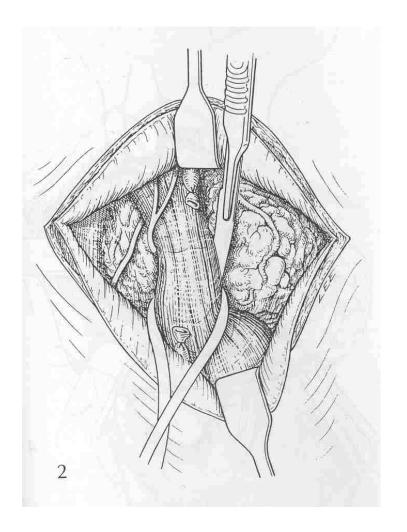


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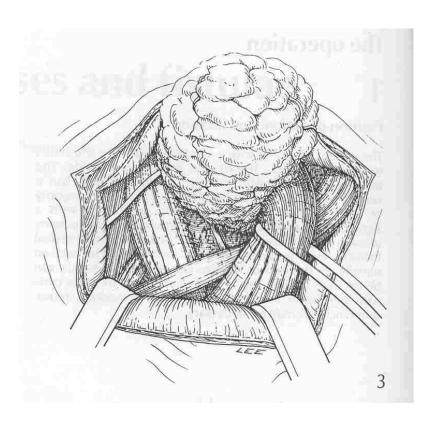
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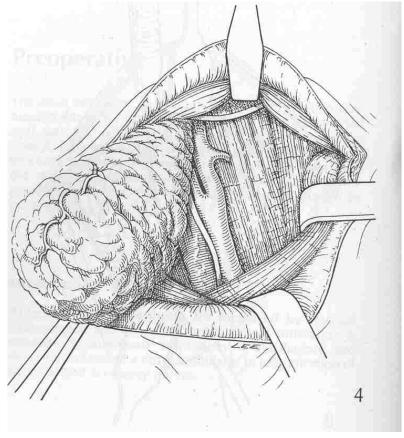
CYSTIC HYGROMA





CYSTIC HYGROMA





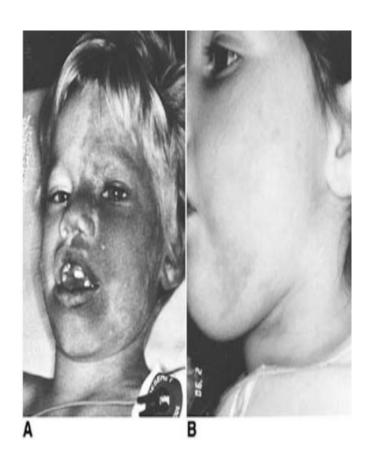
HEMANGIOMA

- Most common benign tumor of infancy
- 3 types : capillary, cavernous, arterial or plexiform
- 14-21% in head & neck
- Most often in trapezius or masseter
- 3 times more common in female



CAPILLARY HEMANGIOMA

- 4 types: salmon patch, port-wine stain, straw-berry angioma, vin-rose patch
- Present at birth or immediately after birth, all disappear with age expect port-wine stain
- These present as bright red or dark colored patch or swelling
- Typically compressible
- Treatment
 - Wait & watch upto 8 yrsTreatment if persists after 8 yrs:
 - Excision with skin grafting
 - Co2 laser application
 - Sclerosant injection
 - ✓ X-ray therapy
 - Steroid injection



CAVERNOUS HEMANGIOMA

- Present since birth
- Doesn't in volute, rather progress
- Always present as bluish localized swelling on face, cheek, ears, lips, mouth, tongue
- Classically compressible

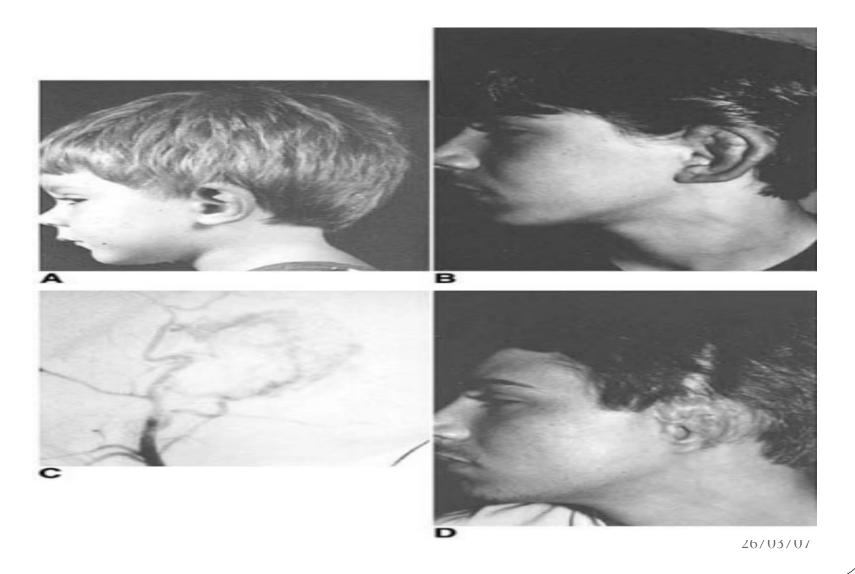


CAVERNOUS HEMANGIOMA

- TREATMENT
- Sclerosant injection
- Steroid injection
- Cryoapplication
- Laser excision
- Surgical excision

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AV Malformation



- Only 7% in head and neck
- 3 types
- Epidermoid cysts
- True dermoid cysts
- Teratoid cysts

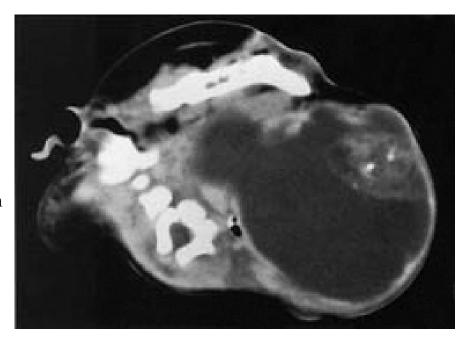
- Epidermoid cysts
- o Most common
- No adnexal structure
- Lined by squamous epithelium
- o Contain cheesy keratinous material

- True dermoid cysts
- Contain skin appendages like hair follicles, sebaceous & sweat glands
- Lined by squamous epithelium
- o Typically, seen in the midline of the neck, usually submental region
- They are attached to and move with the overlying skin and are painless unless infected.
- Management is by complete surgical excision.





- Teratoid cysts
- Rarest variety
- Contain elements from endoderm, ectoderm, mesoderm
- Lined by squamous or respiratory epithelium
- o A history of maternal polyhydramnios in up to 18% of neonates who have a cervical teratoma
- Management is by surgical excision;
 failure of complete excision may result
 in recurrence

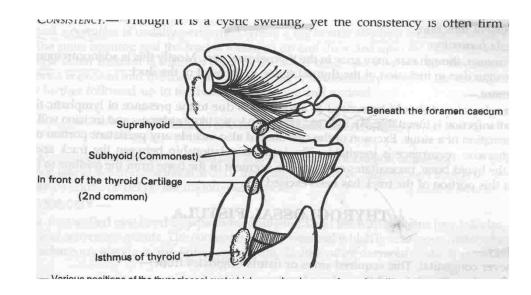


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- Commonest non-neoplastic neck swelling
- Most common midline neck cyst
- o Enlarges caudally as a bilobed diverticulum, which forms lobes of thyroid



- Clinical presentation
- o Male:female 1:1
- Mean age 5yrs
- o 90% midline
- o 10% lateral (95% lt & 5% rt)
- 2.1% intralingual24.1% suprahyoid60.9% thyrohyoid12.9% supra sternal

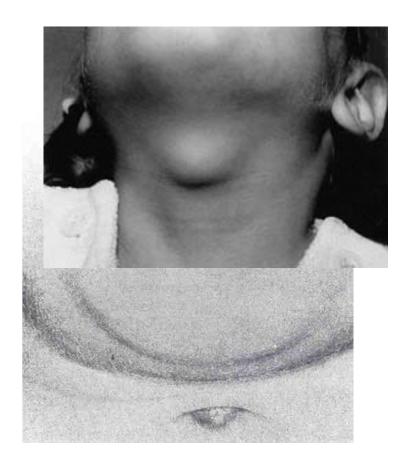


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- Clinical presentation
- Suprahyoid cysts may be a lingual cyst
- May be associated with lingual thyroid, when normal thyroid is often absent
- o Infrahyoid cysts often confused with thyroid adenomas
- A cyst may form on both sides of hyoid producing a dumb-bell cyst, the missing part often accounts for recurrence

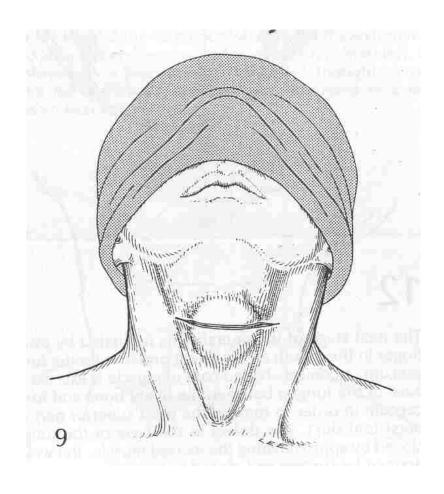
- On examination
- Well defined, smooth surface
- o Spherical, Non tender
- Cystic with fluctuation +
- Transillumination +
- Moves sideways, but not vertically
- Moves with deglutition and protrusion of tongue

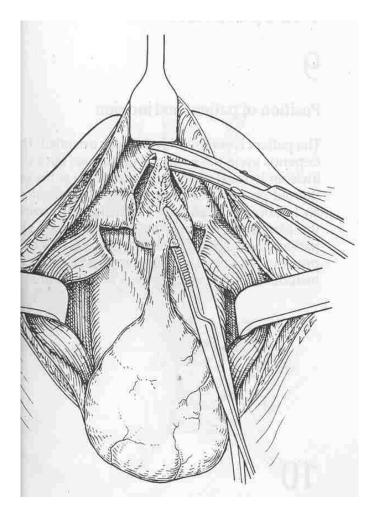


- Diagnosis
- Aspiration cytology
- o I^{131} scan in all suprahyoid to r\o lingual thyroid
- CT\MRI indicated when there is doubt of lingual thyroid
- o Treatment: sistrunk operation.
- Recurrence rate:
- o Simple excision − 85%
- o Sistrunk operation − 2-8%



SISTRUNK'S PROCEDURE

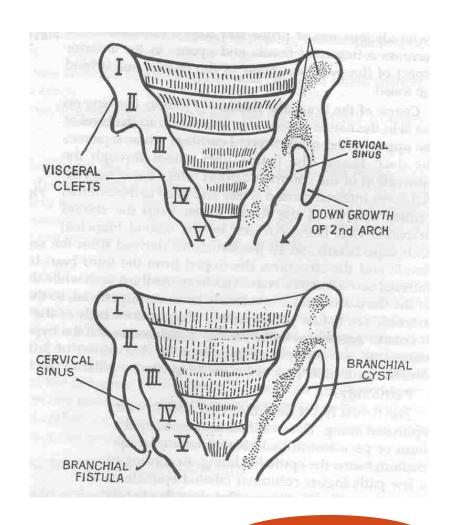




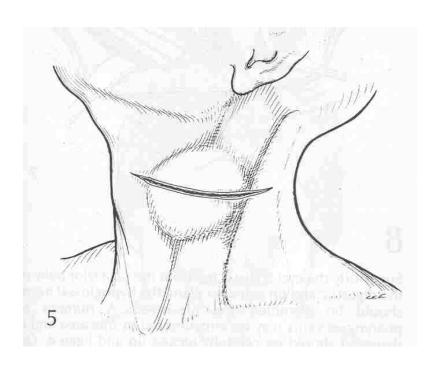
BRANCHIAL CYST, SINUS, FISTULA

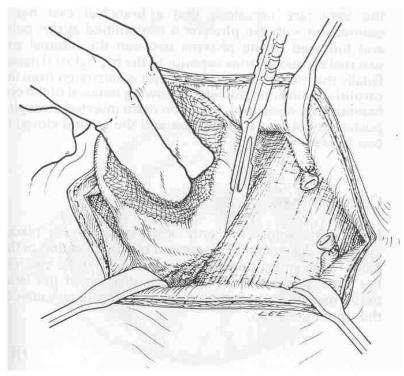
- Cysts present in lateral part of neck deep to SCM at junction of upper $1 \ 3^{rd}$ and lower $2 \ 3^{rd}$
- Sinus may have an external opening along line joining tragus and sternoclavicular joint or an internal opening depending on the origin
- Fistula has both external and internal openings

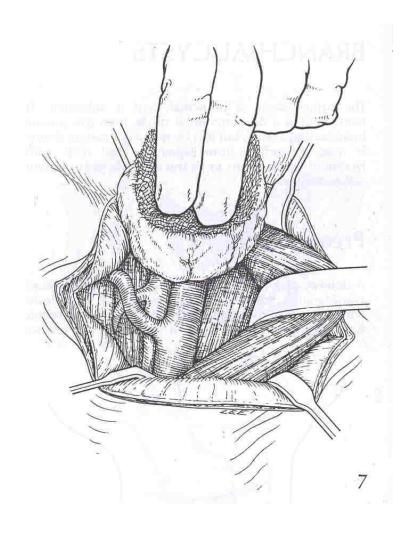
- Development
- A 2week embryo has 6 branchial arches with 5 branchial pouches and 5 branchial clefts
- o 5th and 6th arches are vestigeal
- o 2nd arch grows over 3rd and 4th arch to meet 5th arch element, thus forming cervical sinus of his

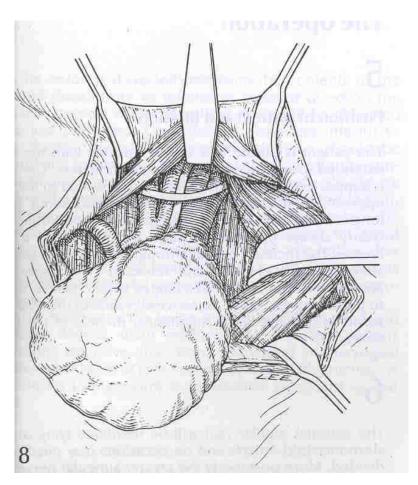


- Theories of origin
 - 1. Branchial apparatus theory
 - 2. Cervical sinus theory
 - 3. Thymopharyngeal theory
 - 4. Inclusion theory
- Pathology
 - o Lined by stratified squamous or non-cilliated columnar epithelium
 - o 80% have lymphoid tissue in the wall
- Clinical presentation
 - o Peak age 3rd decade
 - o $2 \setminus 3^{rd}$ left side and $1 \setminus 3^{rd}$ right side; 2% bilateral









• Congenital: most commonly from 2nd arch

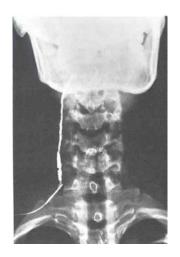
• ACQUIRED:

Infection---- burst

Inadvertent incision

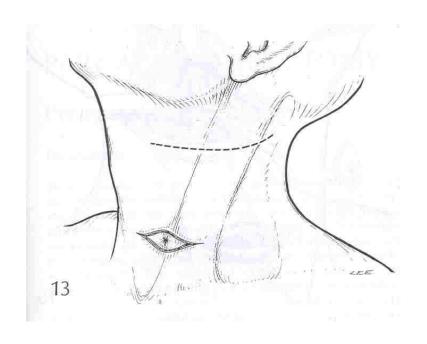
BRANCHIAL SINUS\FISTULA 2nd Arch Fistula

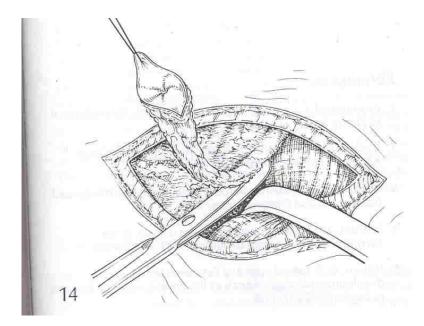
- 2nd arch fistula
- o Tract travells between the derivatives of 2nd arch (superficially) and 3rd arch (deep)
- Inferior-middle 2/3 junction of SCM, deep to platysma, lateral to IX, X, XII, between the internal and external carotid and terminate in the tonsillar fossa
- Surgical treatment may include tonsillectomy

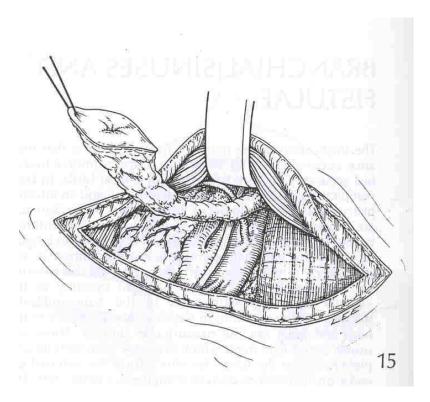


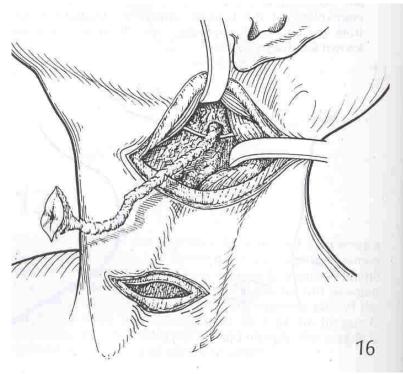


- 1ST ARCH FISTULA
- INTERNAL OPENNING IN EAC AT BONY-CARTILAGINOUS JUNCTION
- PRESENT AS DISCHARGING EAR
- 3RD ARCH FISTULA
- o INTERNAL OPENNING AT LATERAL WALL OF PFS
- 4TH ARCH FISTULA
- INTERNAL OPENNING AT APEX OF PFS
- ALMOST ALWAYS LEFT SIDED
- PRESENT AS SUPPURATIVE THYROIDITIS









RANULA

- Cystic mass in the FOM\tongue\neck
- Arise due to obstruction of ducts of minor salivary glands or sublingual gland
- Caused by trauma or ductal anomaly
- Newer studies quote that increased intraoral pressure may be a cause

RANULA

- On examination
- o Bluish cystic swelling in the FOM or submental region
- Nontender
- o Spherical in shape
- o Smooth surface with ill defined margins
- o Fluctuation +
- o Brilliantly translucent
- Mucosa easily mobile over swelling





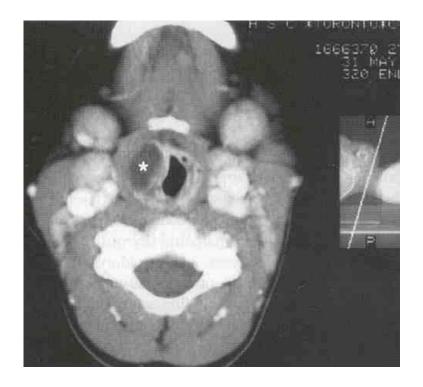


RANULA

- Treatment
- Marsupialisation
- o Excision of the lesion alone
- o Excision of cyst along with sublingual gland
- o Simple ranula excised by transoral approach including sublingual gland of origin
- Plunging ranula and recurrent cases are to be operated by transcervical approach

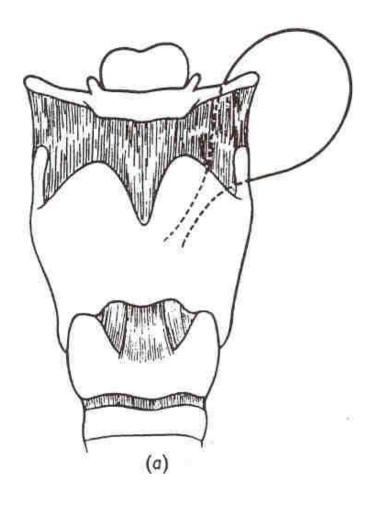
- Arises from enlargement of saccule of ventricle
- 3 types
- o External (30%)
- o Internal (20%)
- o Combined (50%)

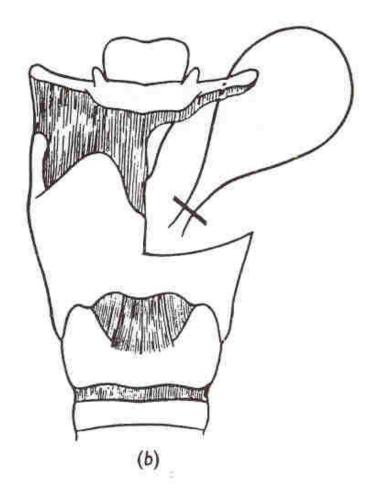
- Investigation
- Plain radiograph : stn (ap view)
- ✓ Air filled sac
- ✓ Upper boarder of thyroid cartilage is boarder line between a laryngocele and an enlarged sac
- ✓ Mucocele : air fluid level (d\d laryngeal cyst)
- ✓ Infected sac (d\d gas gangrene)
- o CT scan: to know the extent and confirm diagnosis
- o $D\L$ scopy : especially in u\l cases to r\o malignancy



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- Asymptomatic : no treatment
- Internal : endoscopic marsupialisation if recurrence excision
- External: transcervical excision
- Pyocele: antibiotics --- excision





• Thank you