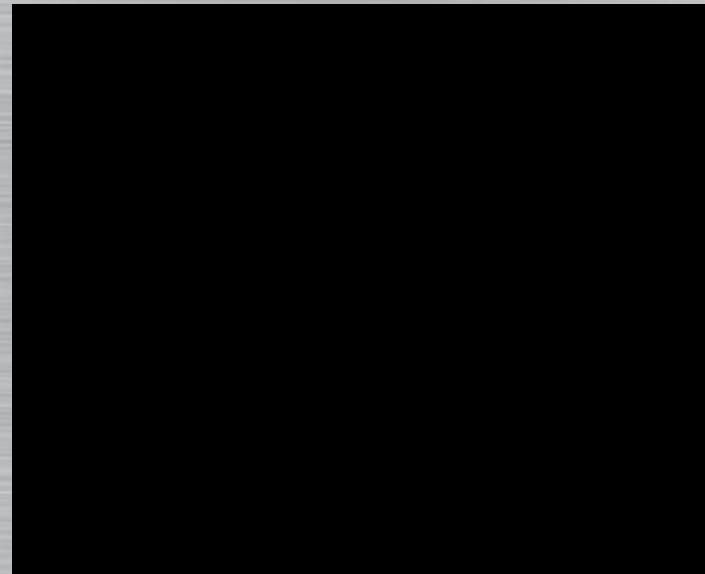
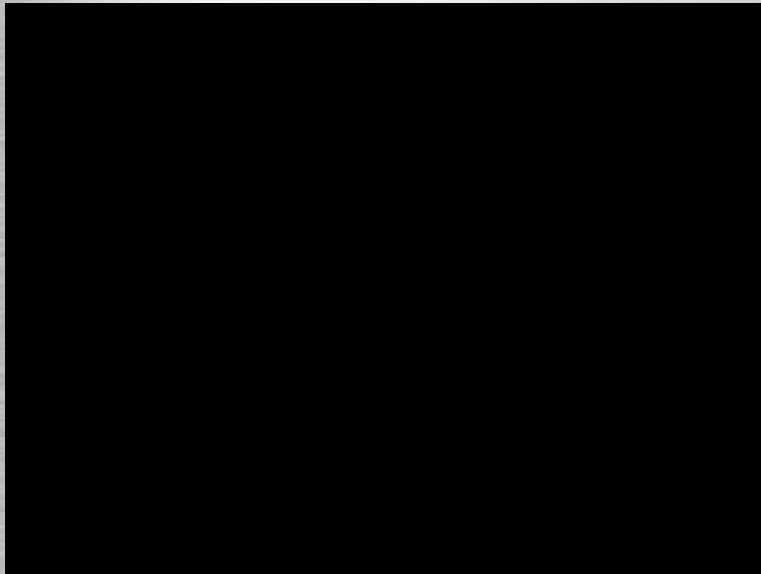


Endoscopy of the Airway

Dave Albert



Airway Endoscopy

The Basics

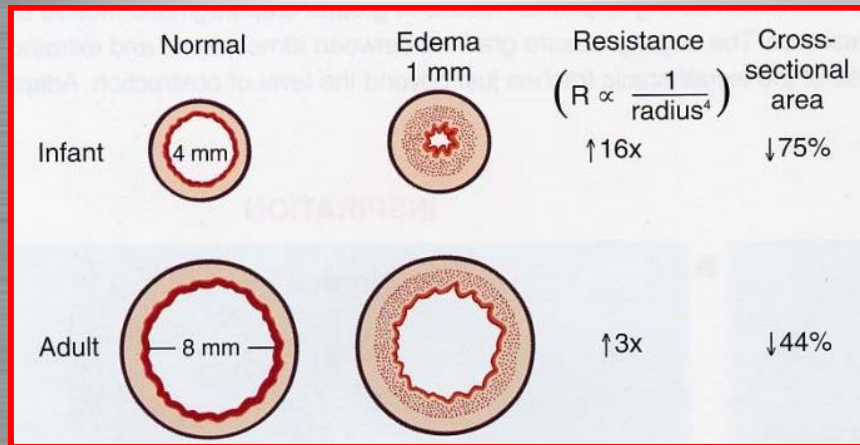
Neonatal

Progressive

Acute

Quiz

The laws of Poiseuille and Bernoulli

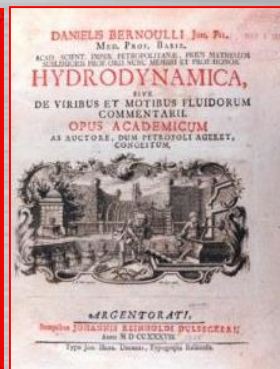


$$\Delta P = \frac{8\mu LQ}{\pi r^4}$$

French physician, physicist and mathematician

$$\frac{v^2}{2} + gy + \frac{P}{\rho} = \text{constant}$$

v = fluid velocity along the streamline
 g = gravitational constant
 y = elevation in g-field
 P = pressure along the streamline
 ρ = fluid density



Dutch Swiss mathematician
physician and physicist

Basics of Endoscopy

- Team

- Anaesthetist
- Nurse
- Surgeon

- Equipment

- Hopkins rod endoscopes
- Video/still digital recording
- Microdebrider rather than laser
- Balloon

- Spontaneous respiration



Flexible Endoscopy

Good screening procedure

Good for dynamic conditions

May miss :-

Cricoarytenoid fixation

Cleft larynx

Subglottic pathology



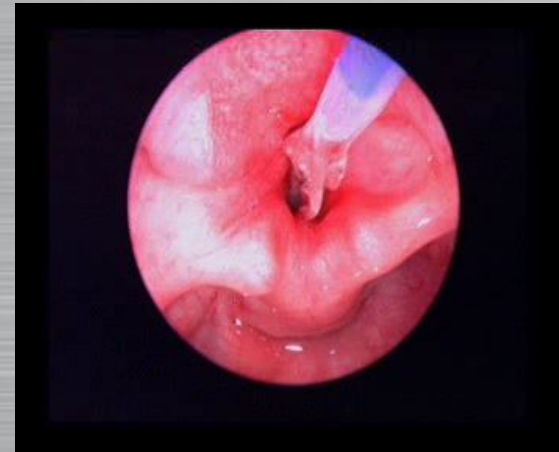
Rigid Endoscopy

Precairious airways -

–more control

Can probe arytenoids etc

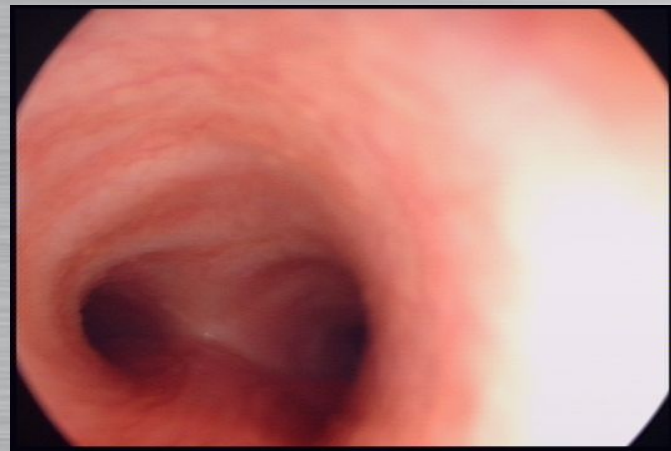
Can treat endoscopically



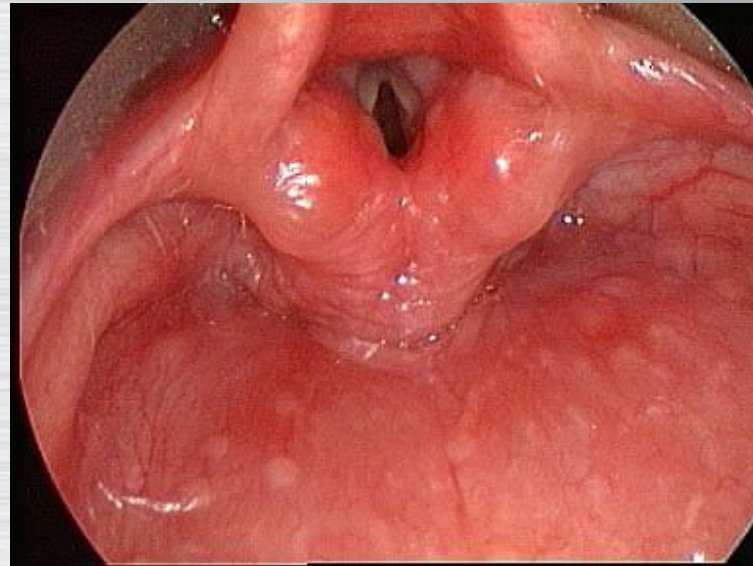
'FESS' Style Endoscopy



'Four Shot' Views



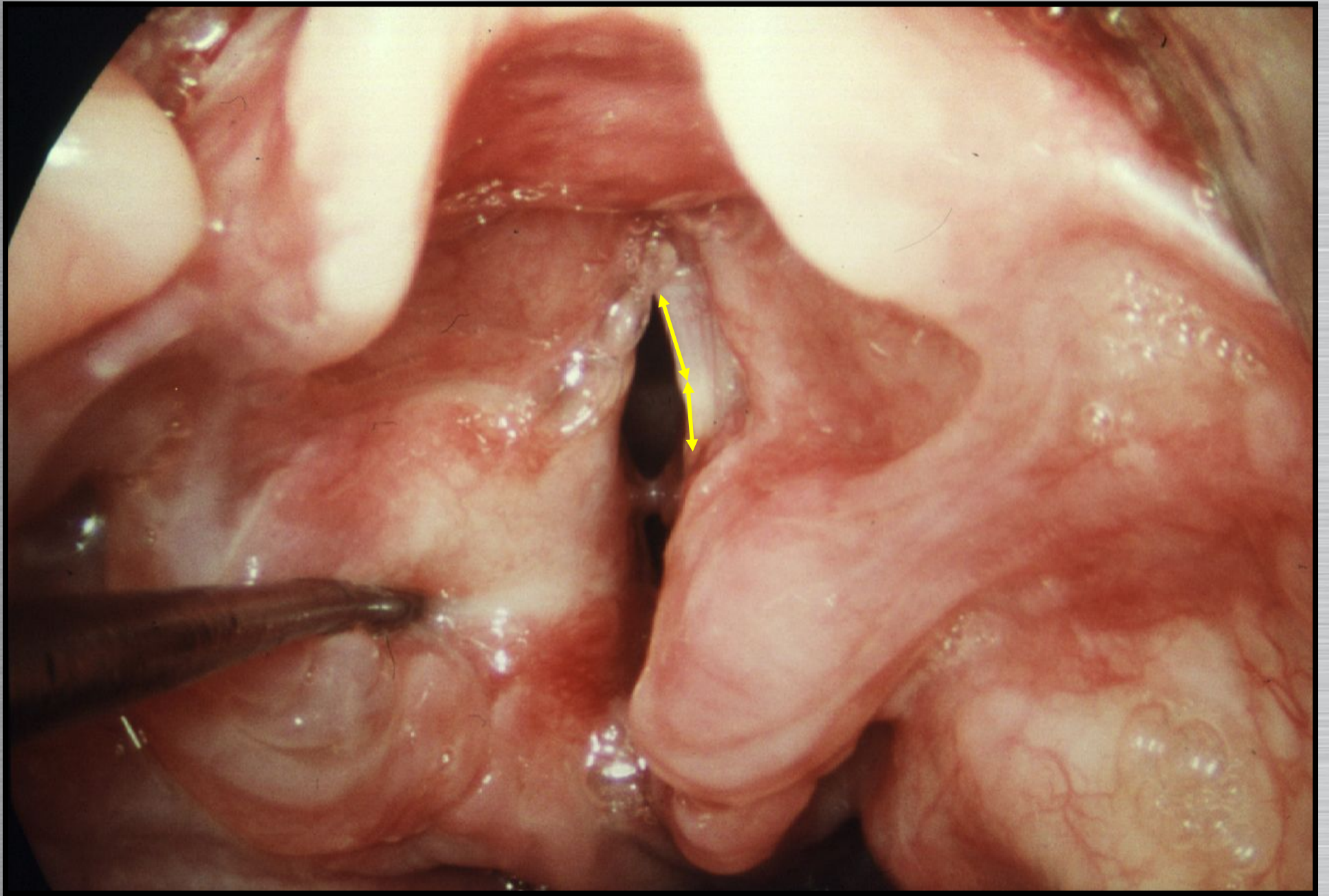
Dynamic View



Traditional Microlaryngobronchoscopy







NEONATAL

Laryngomalacia: History

Stridor

Not usually present at birth (first week)

Quiet at rest/asleep

↑ with feeding and crying

musical quality

cry : normal

cyanotic episodes: unusual

growth/weight : affected if severe

Laryngomalacia: Examination

- Inspiratory stridor ↑ with crying
- Recession
- Check for
 - cutaneous haemangiomata
 - signs of syndromes:
 - small jaw, wide spaced eyes, low set ears etc



Laryngomalacia: Endoscopy

to confirm diagnosis of laryngomalacia
to exclude co-existent airway pathology

Fibre-optic in office – screening

? Now minimal “standard of care”

MLB under GA for full assessment

When to perform rigid endoscopy in suspected laryngomalacia

- Intubation history
- Traumatic birth
- Stridor from day 1
- Cyanotic episodes
- Aspiration
- **Failure to thrive**
- Biphasic stridor
- Severe recession
- Other congenital abnormalities
- Abnormal neurology

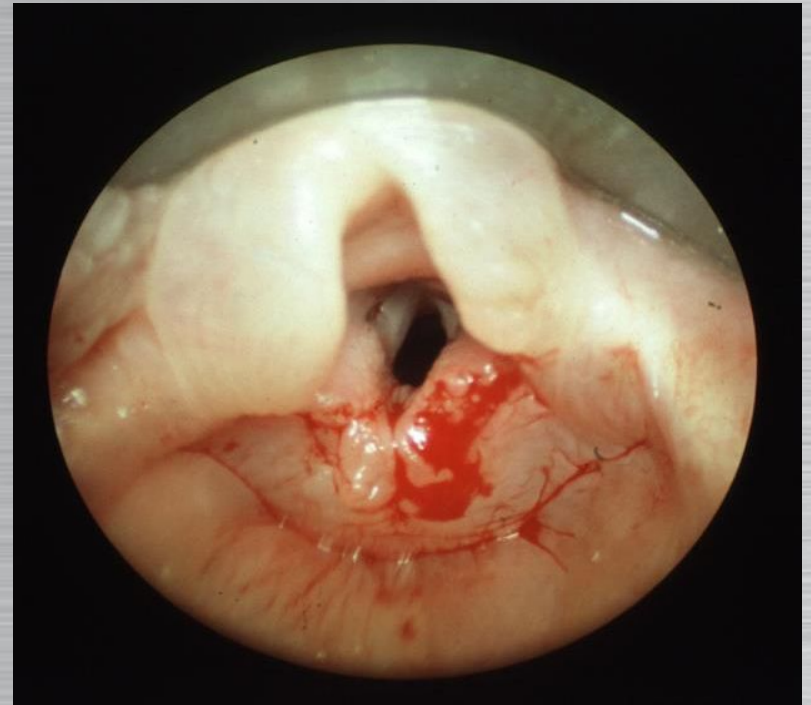
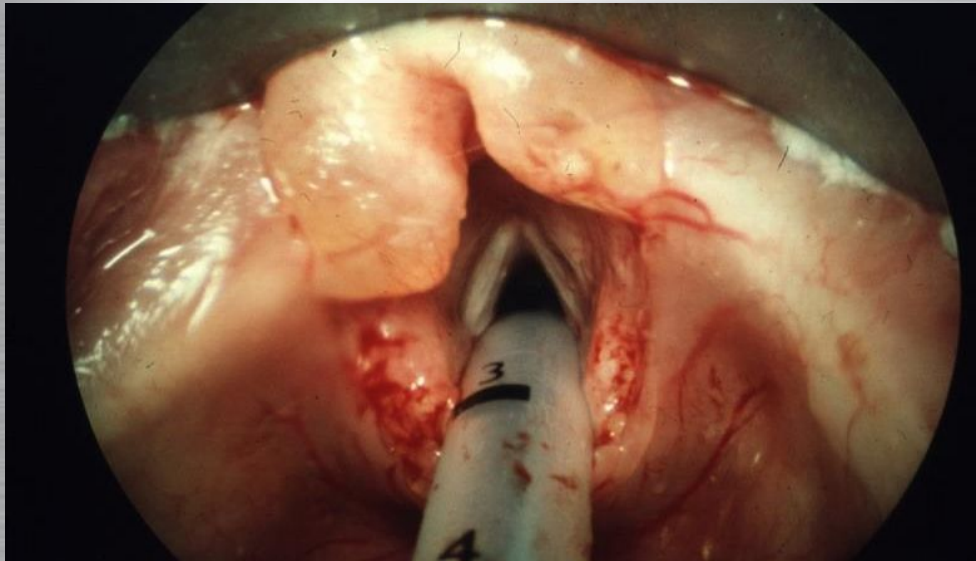
Posterior Laryngomalacia

- Prolapse of arytenoid into laryngeal lumen
- Trim or snip if:
 - Failure to thrive
 - Severe airway difficulties
 - eg desaturation



Aryepiglottoplasty: Surgery

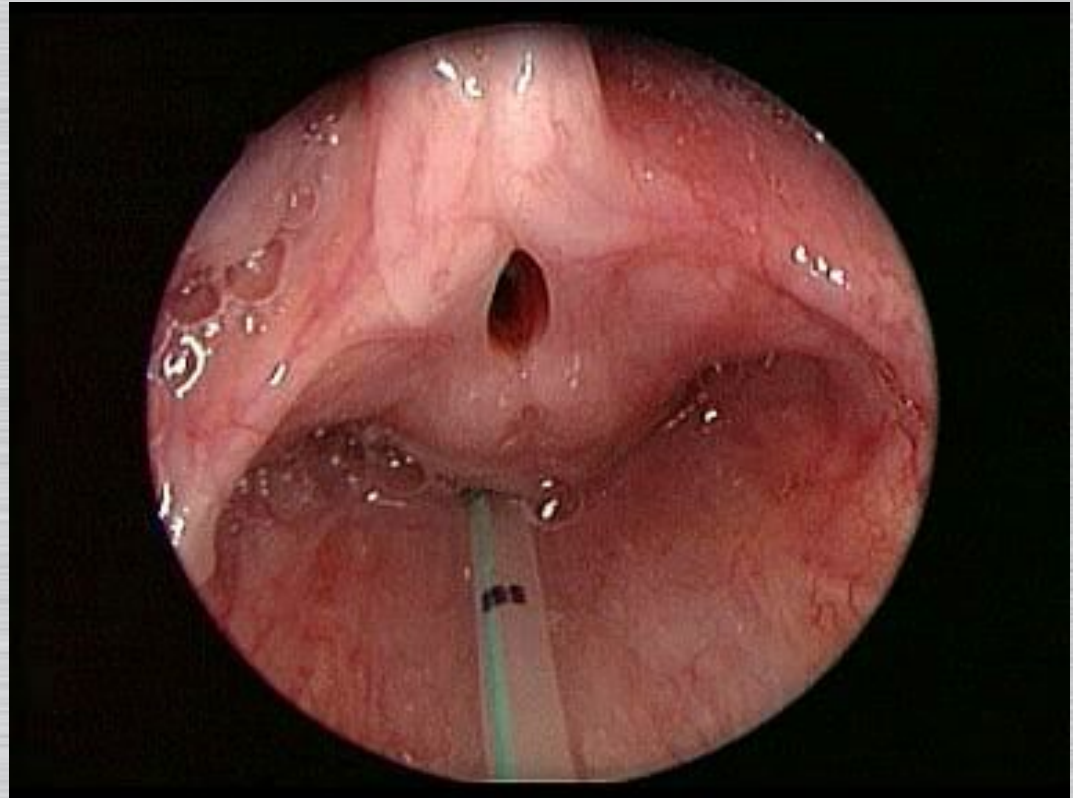
Mucosal excision



Aryepiglottoplasty

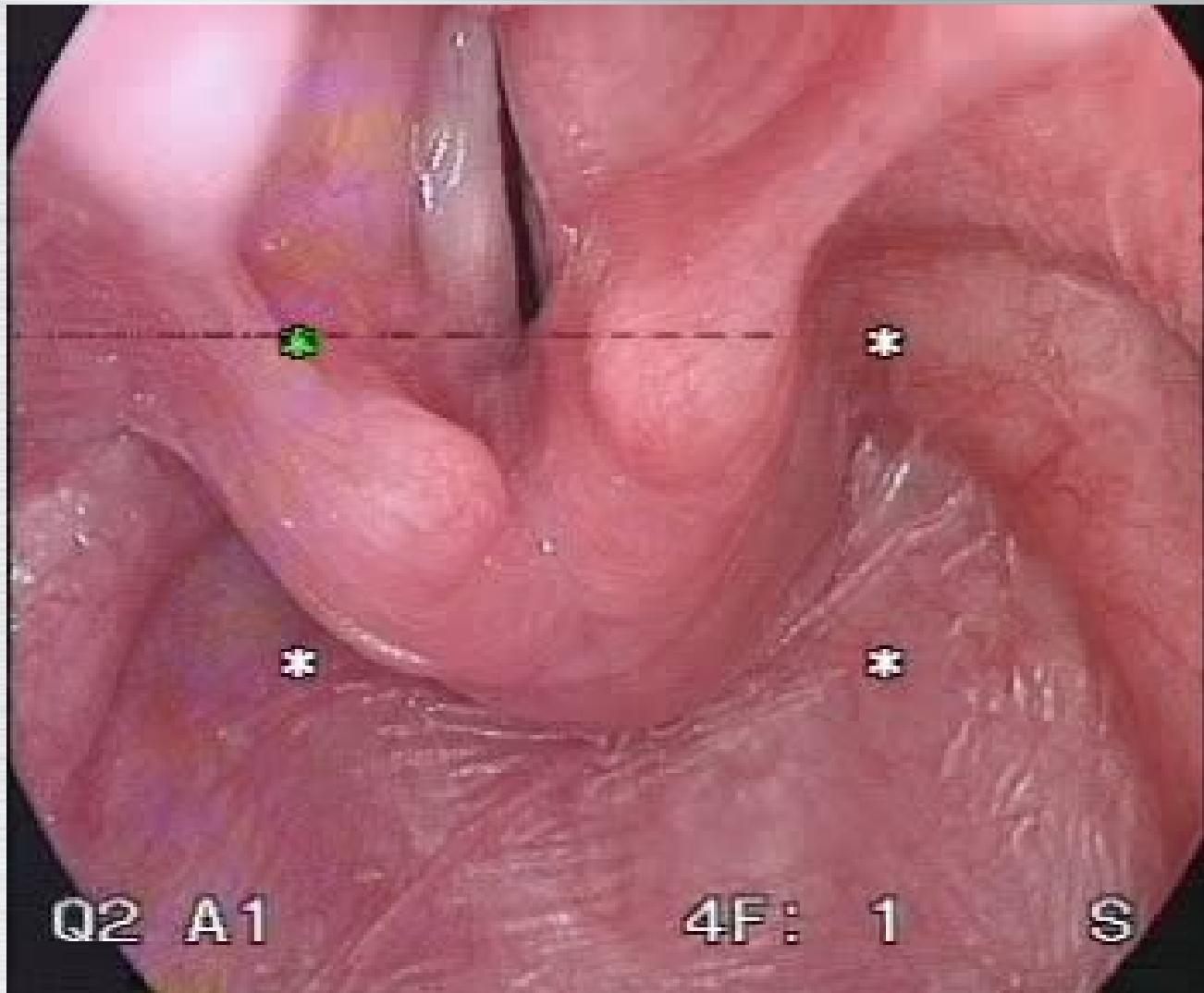
- Sheffield snip

- NOT LASER!!



Laryngomalacia

Vocal cord palsy



Vocal cord palsy

? second most common
cause of neonatal stridor

However, often more an
incoordination

Within 1st month and often
with **first breath**

Bilateral paralysis: Stridor,
cyanosis, apnea

Unilateral paralysis:
dysphonia.

Both: feeding problems,
sternal recession

Treatment

Conservative or Tracheostomy

Also:

Lateralisation: open or endoscopic

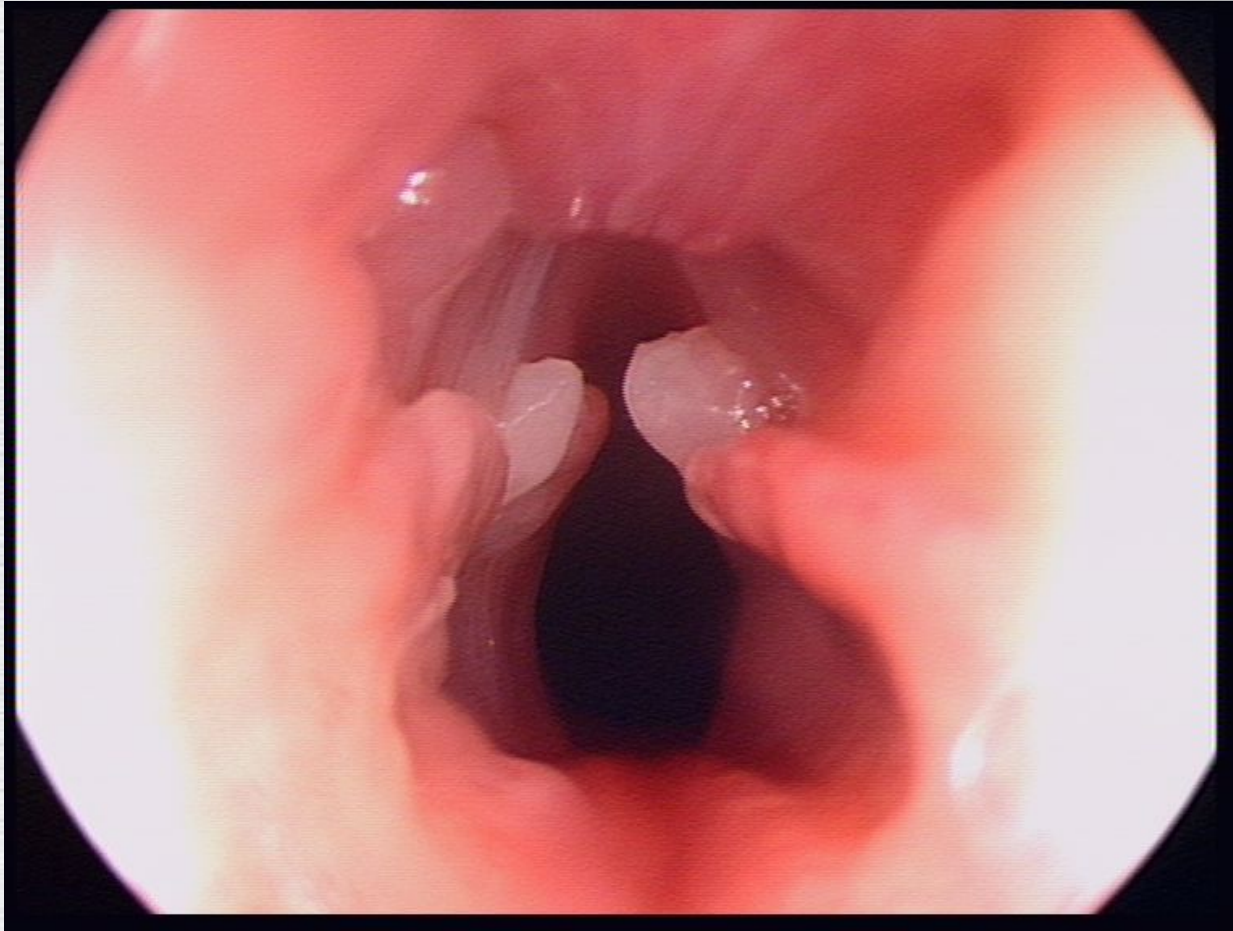
Laser arytenoidectomy

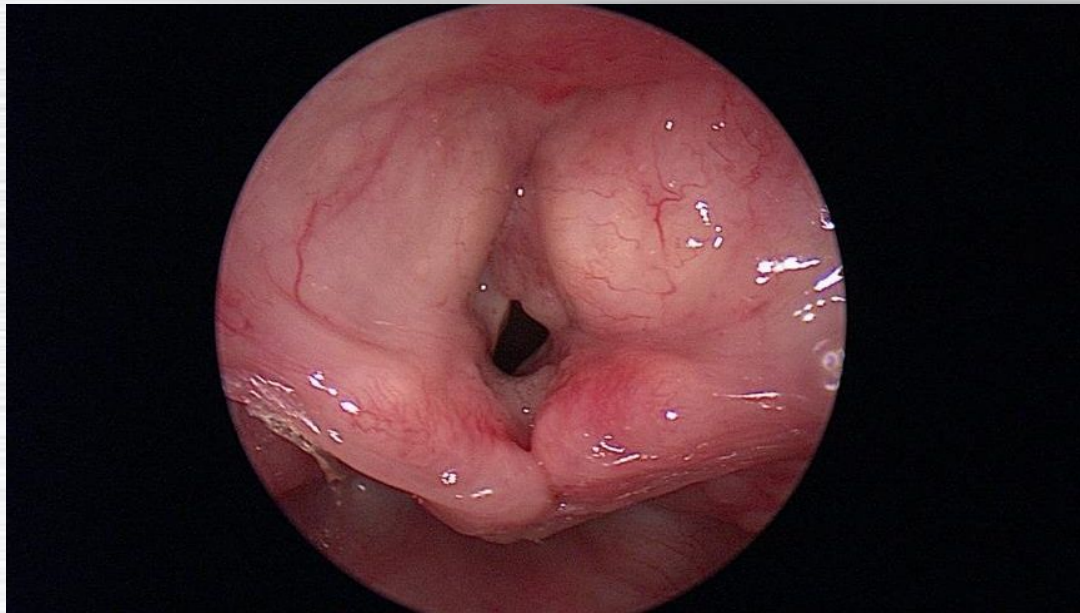
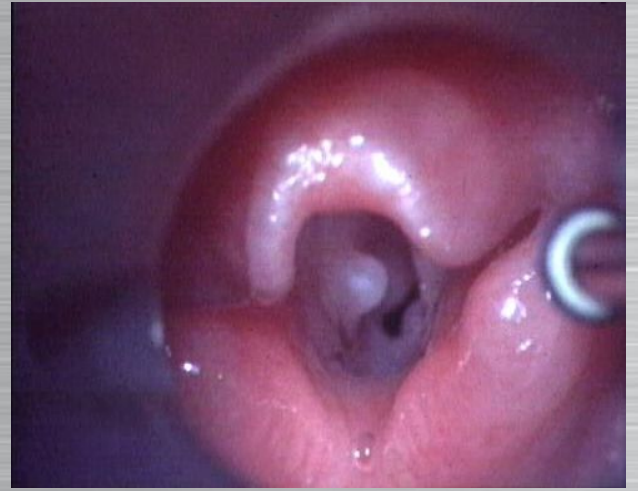
Posterior cricoid graft ? endoscopic

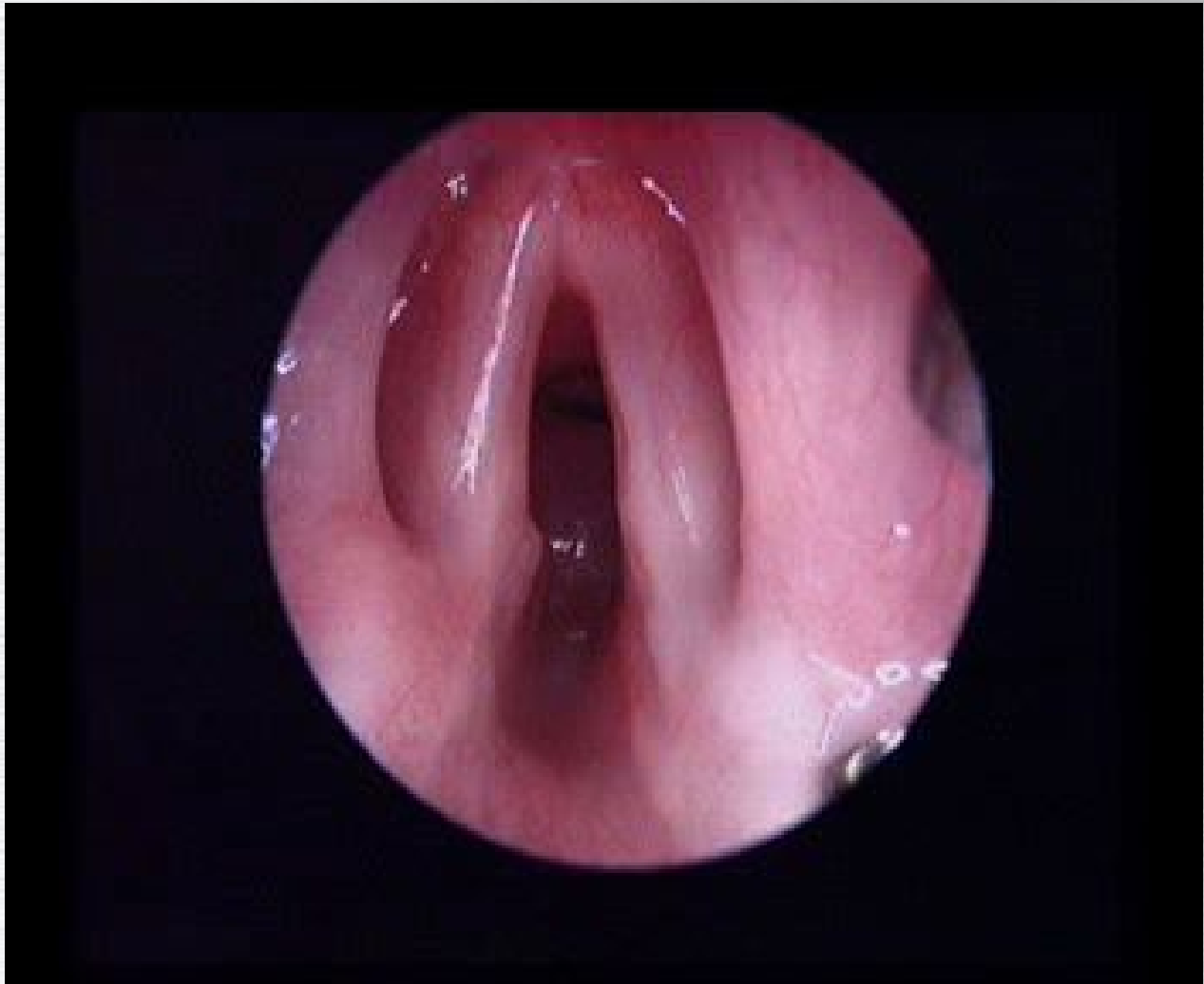
Re-inervation

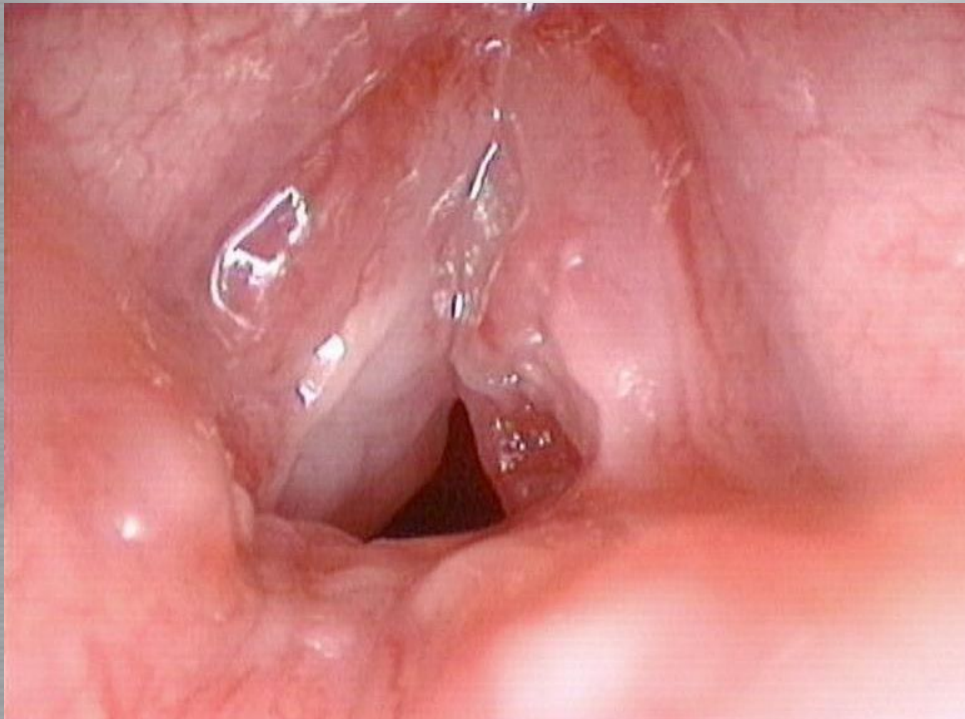
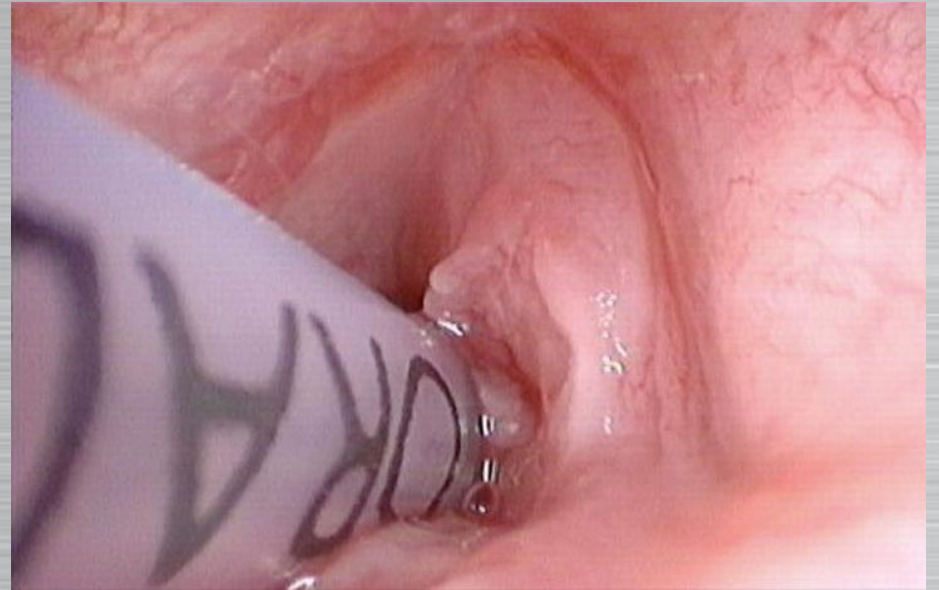
Voice vs airway

Prolonged Intubation

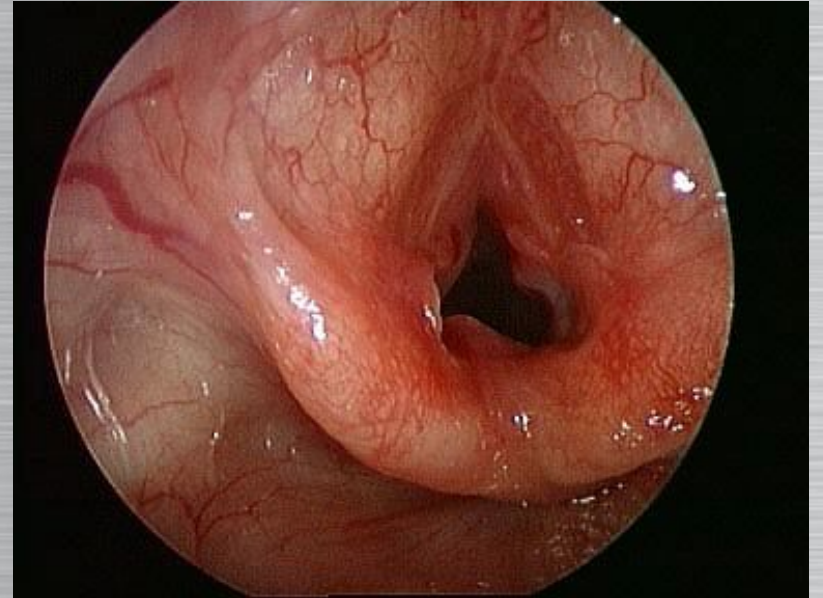




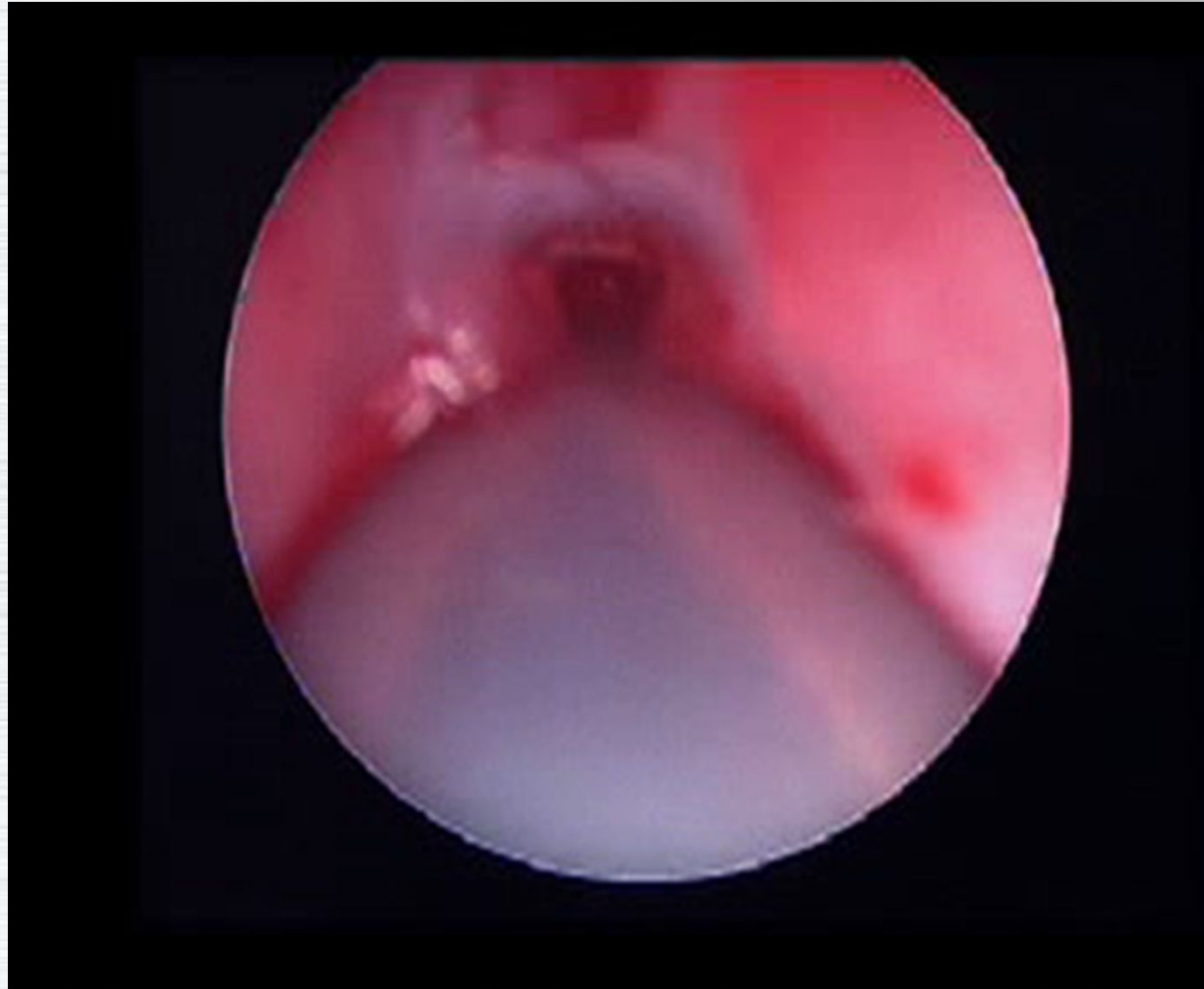




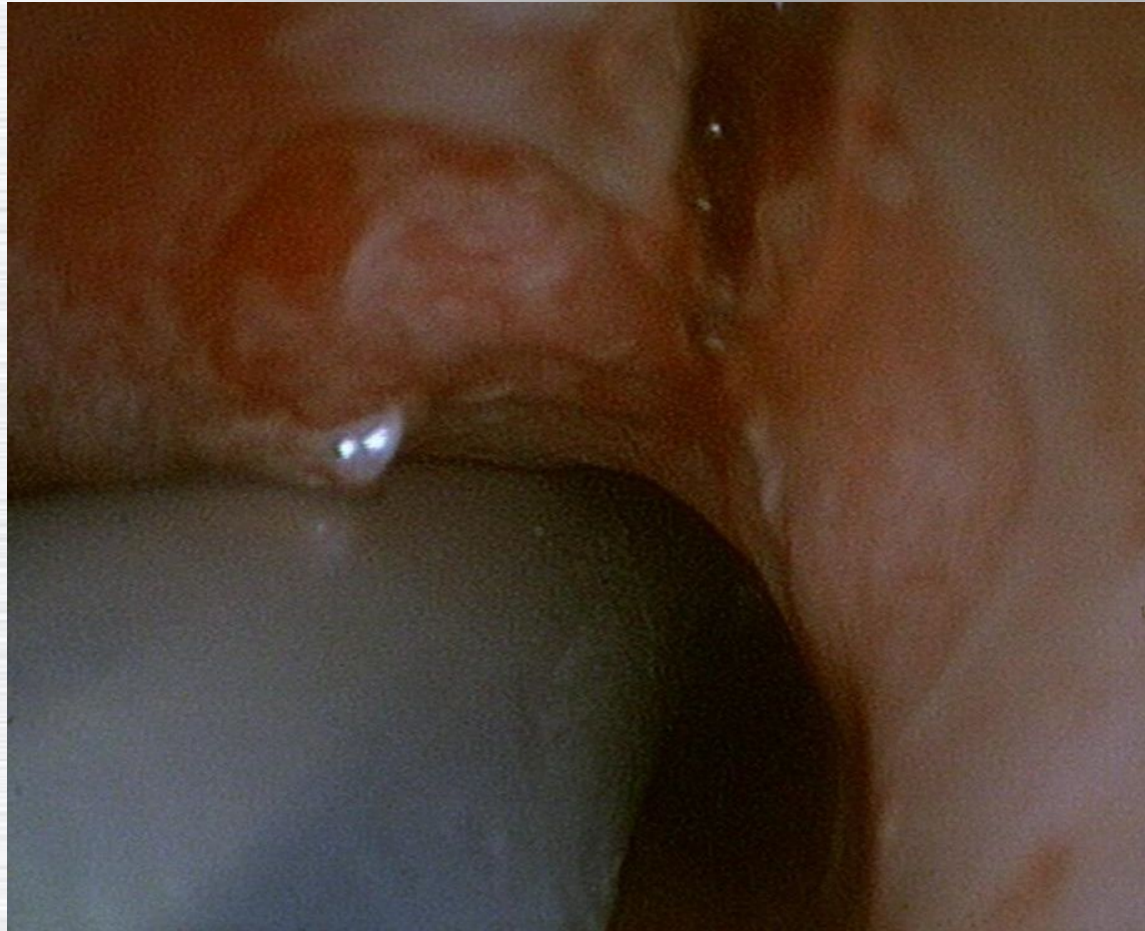
Subglottic edema - endoscopic



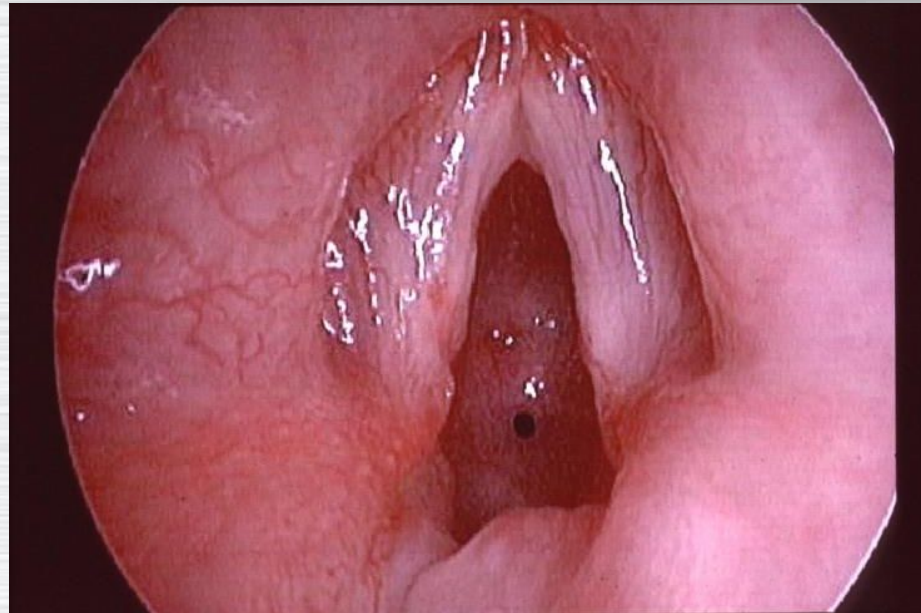
Endoscopic procedure Cut edges of cricoid

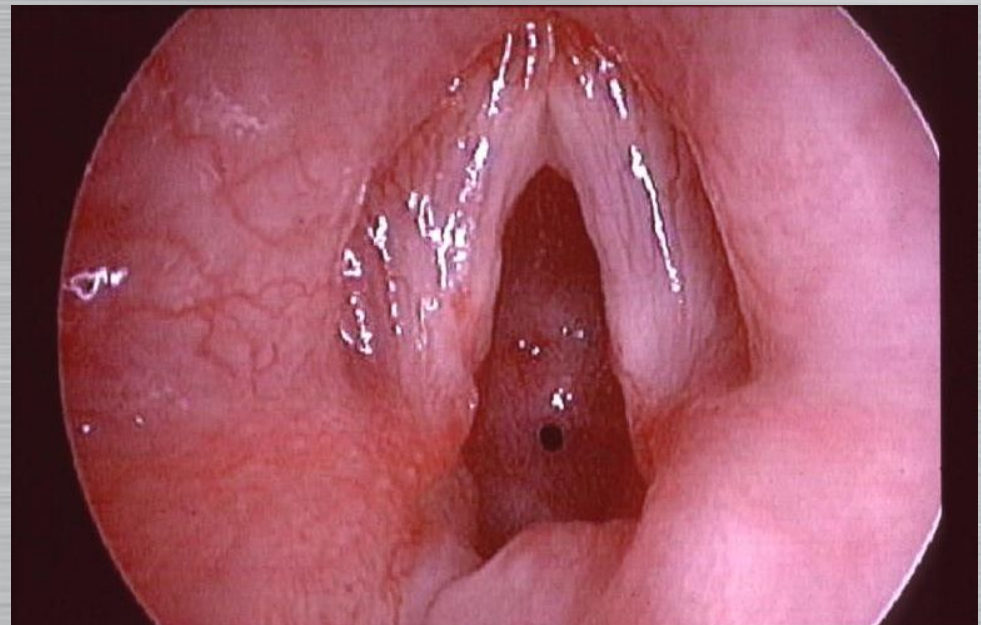
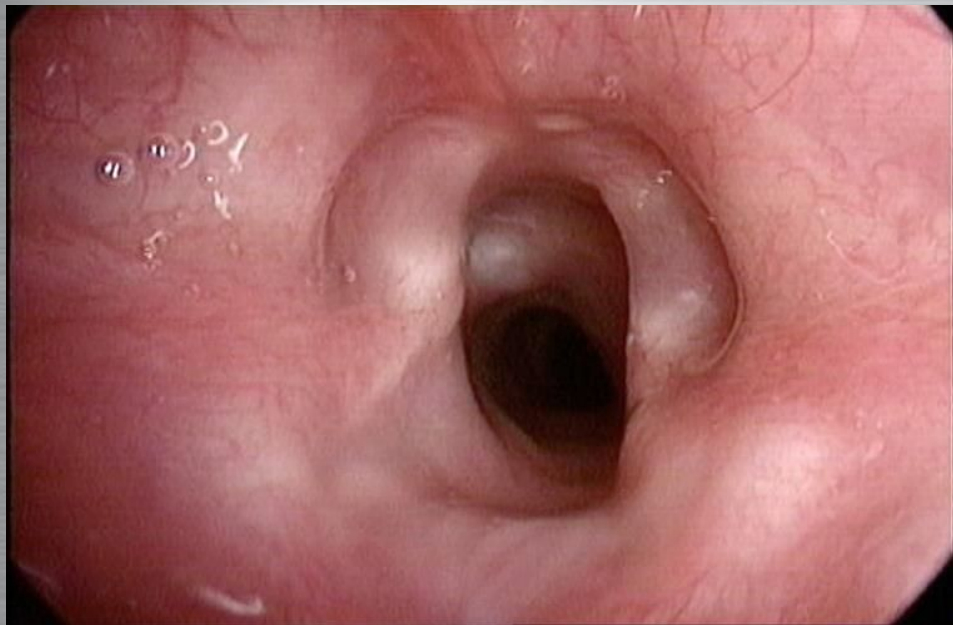
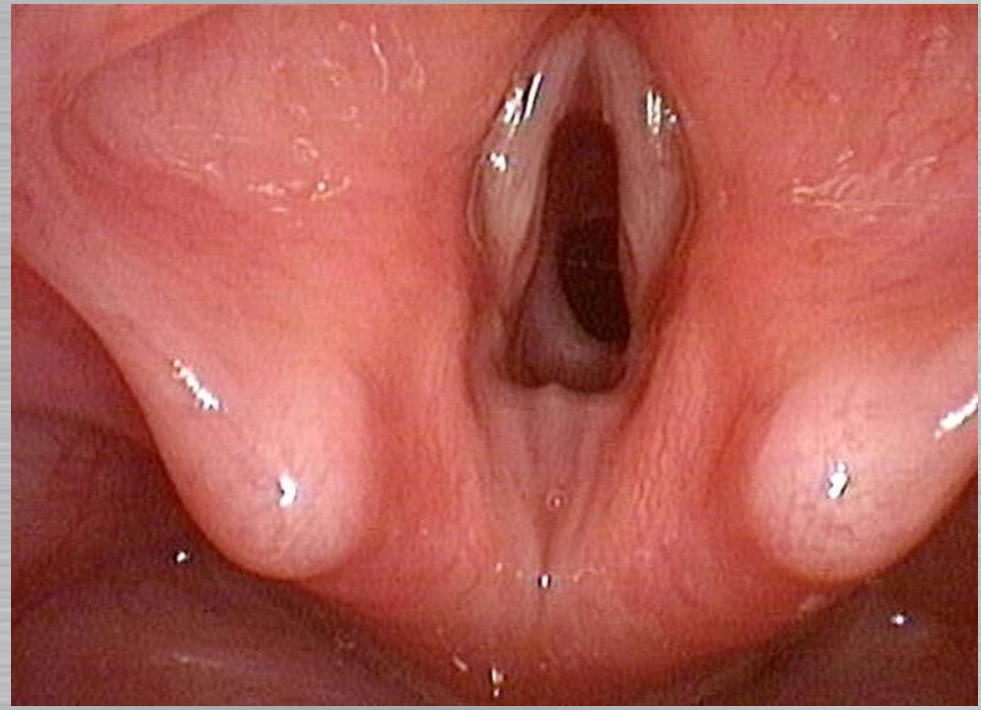
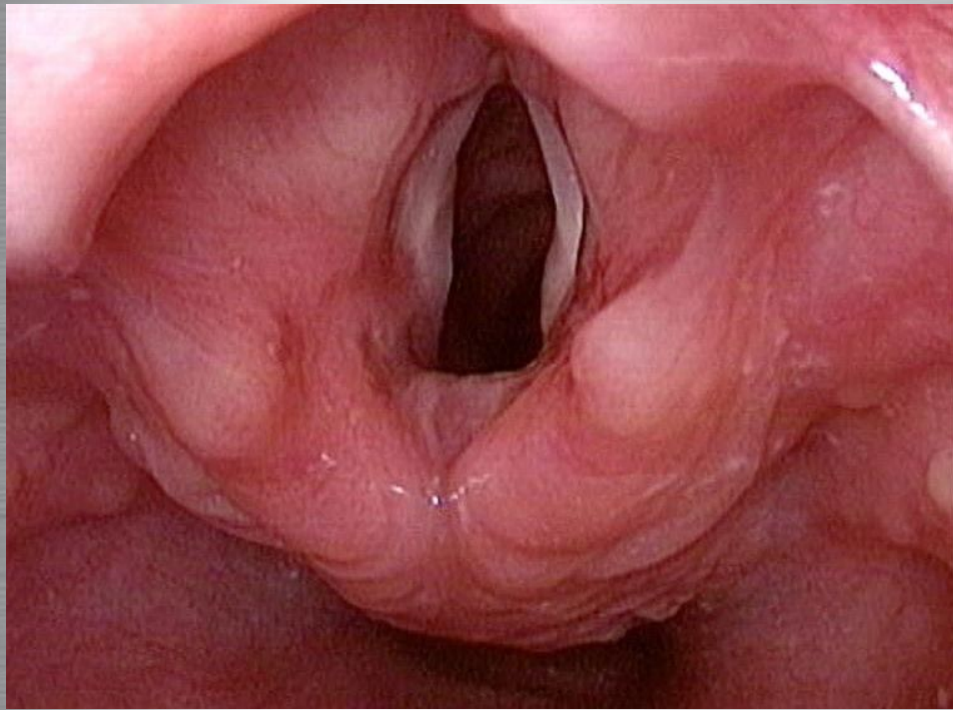


Cricoid Split: open procedure

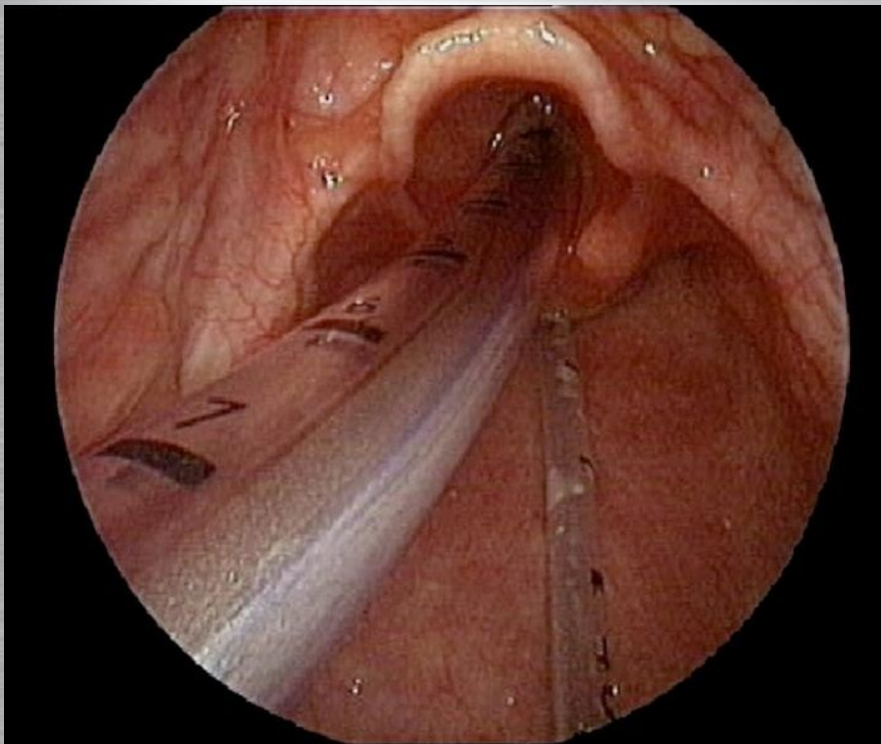








Subglottic stenosis





Staging-Sizing using ET tube

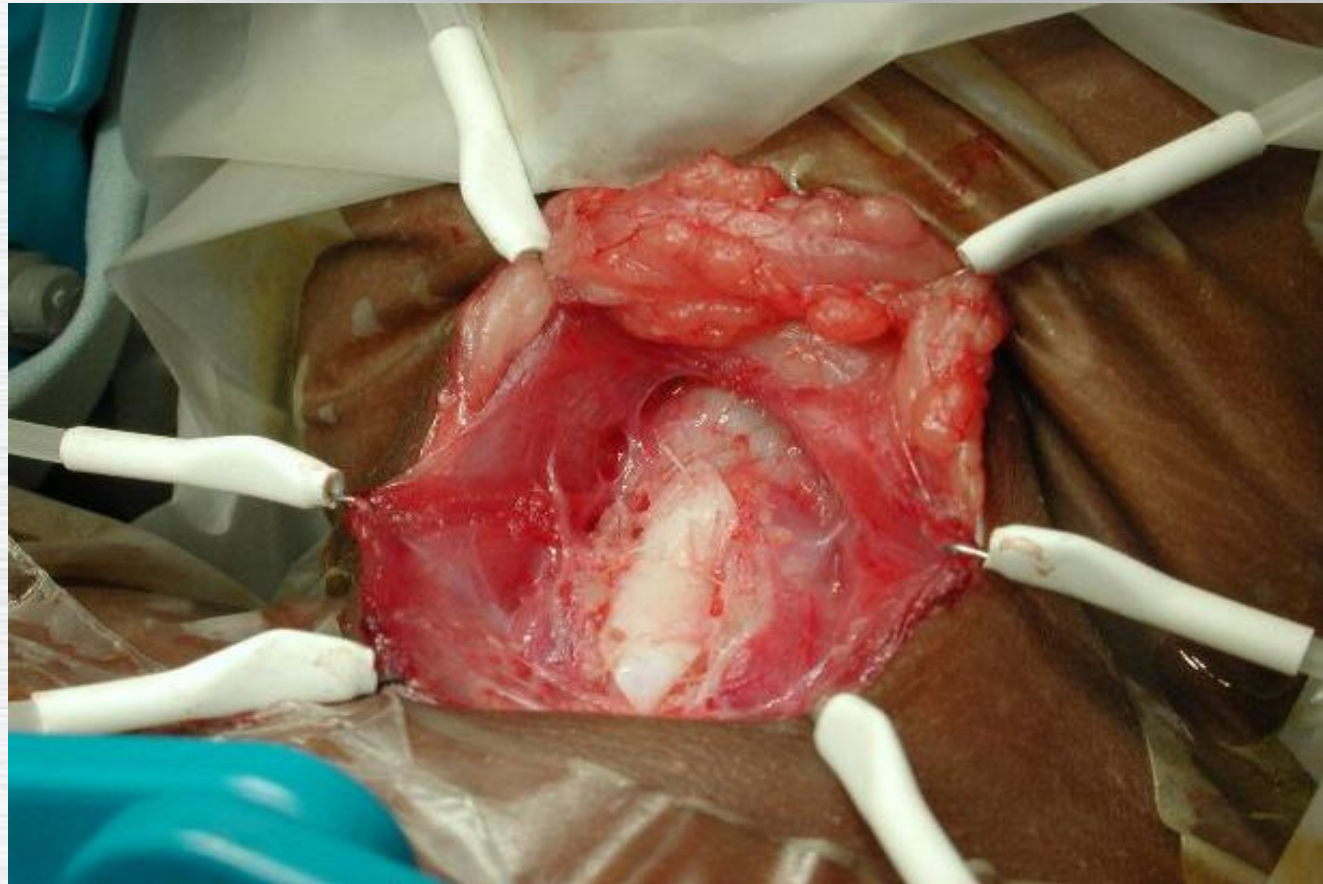


Classification	From	To
Grade I	 No Obstruction	 50% Obstruction
Grade II	 51% Obstruction	 70% Obstruction
Grade III	 71% Obstruction	 99% Obstruction
Grade IV	No Detectable Lumen	

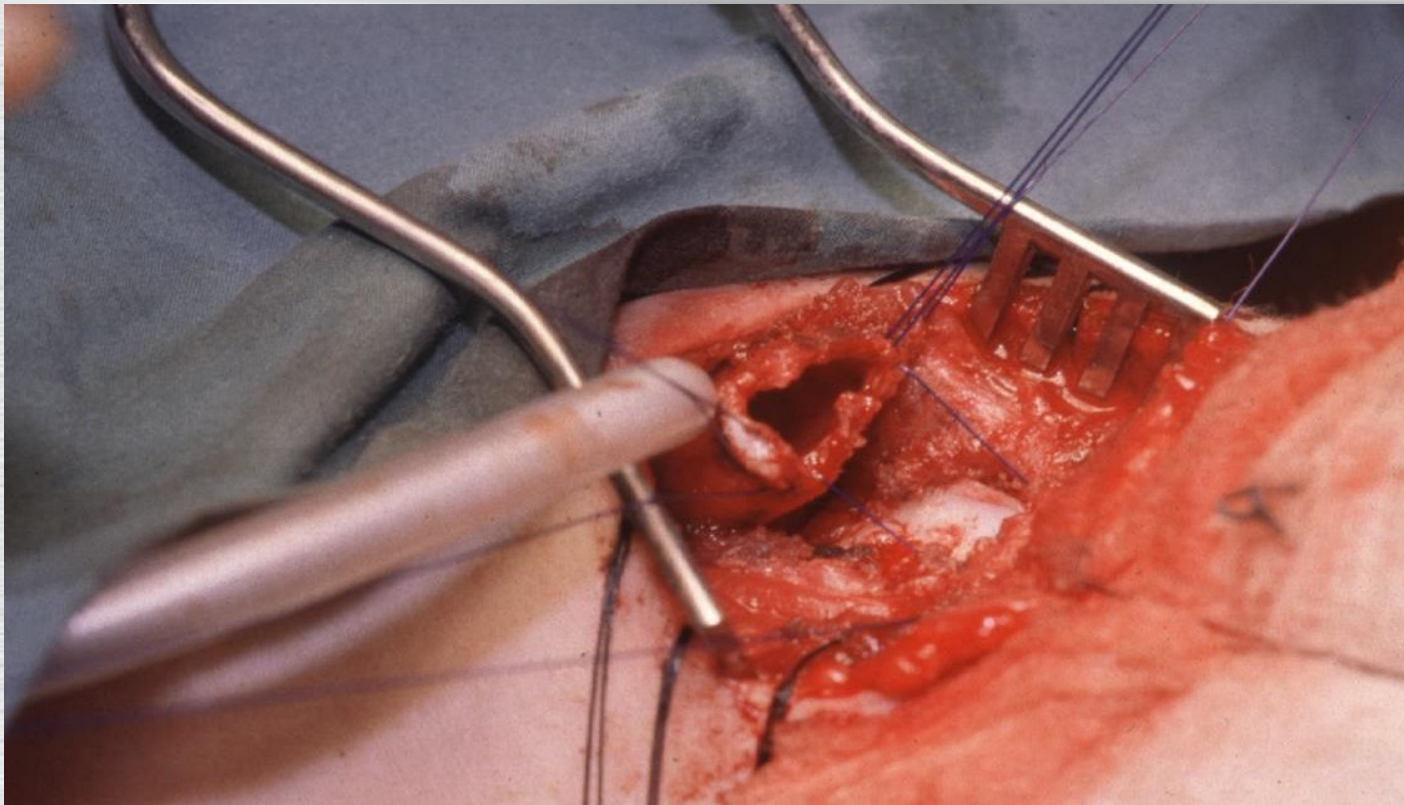
My current guidelines

Grade		
I	Conservative	
II	Endoscopic if soft	LTR once established
III	LTR	CTR if severe and clear of cords
IV	LTR	CTR if clear of cords/ t tube

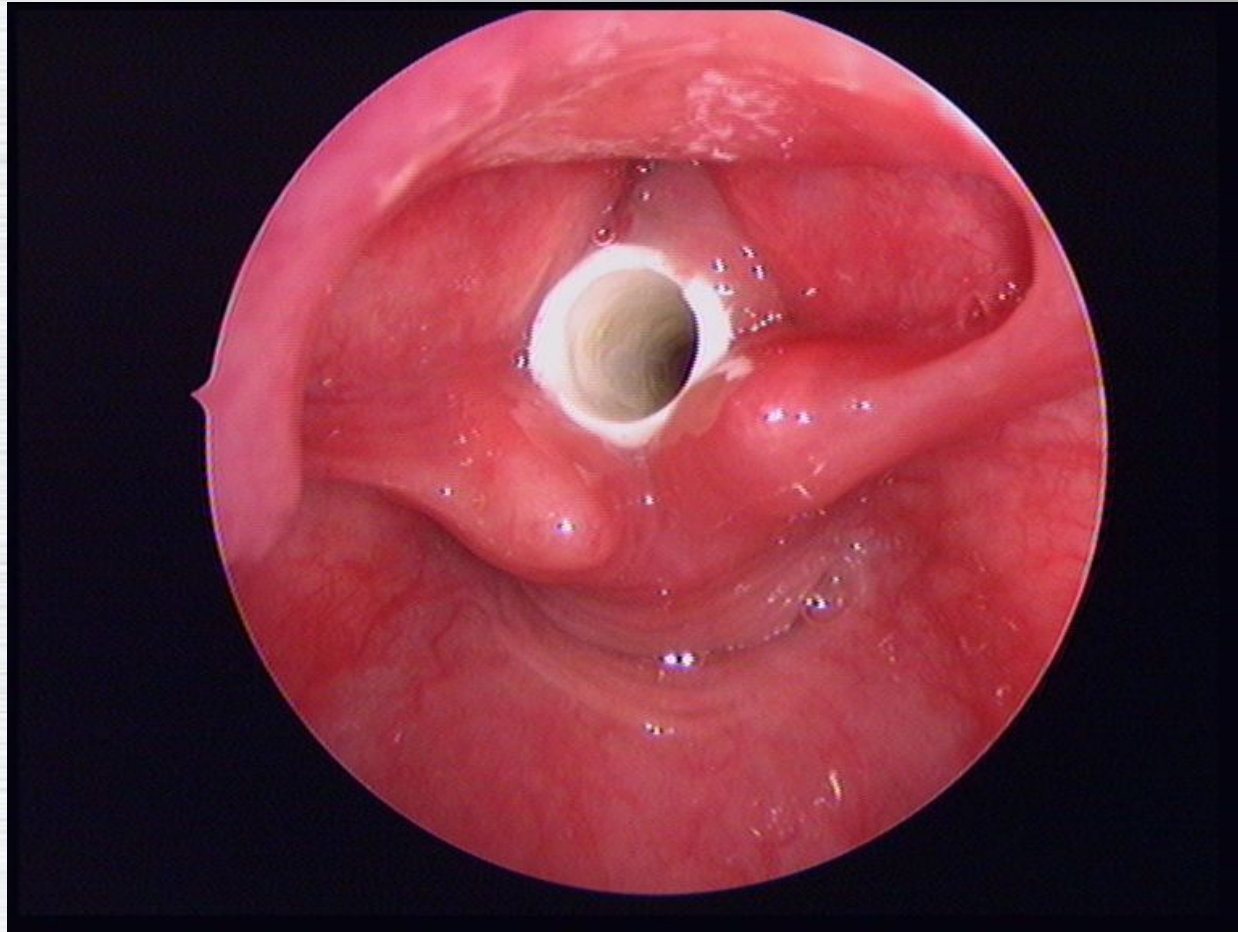
LTR Graft placement



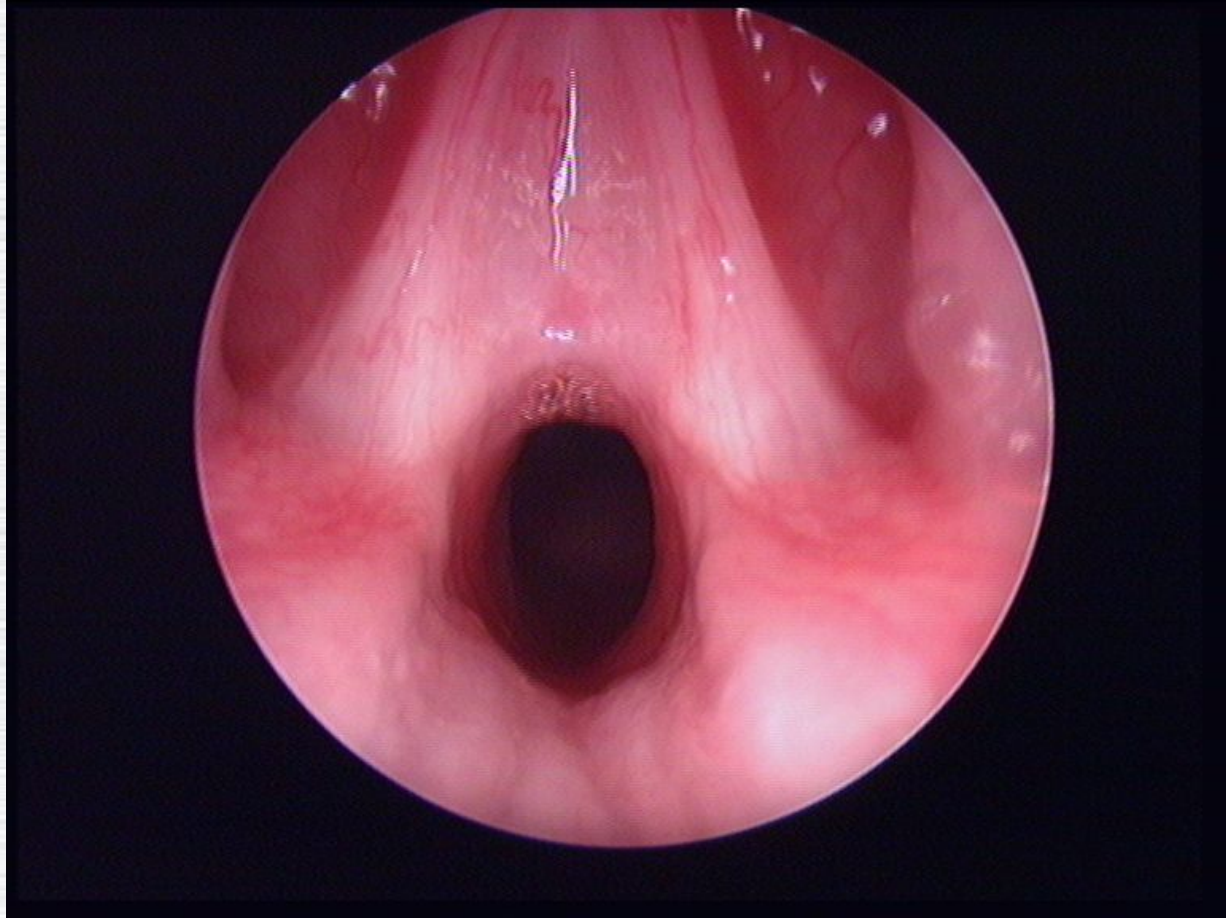
Cricotracheal resection

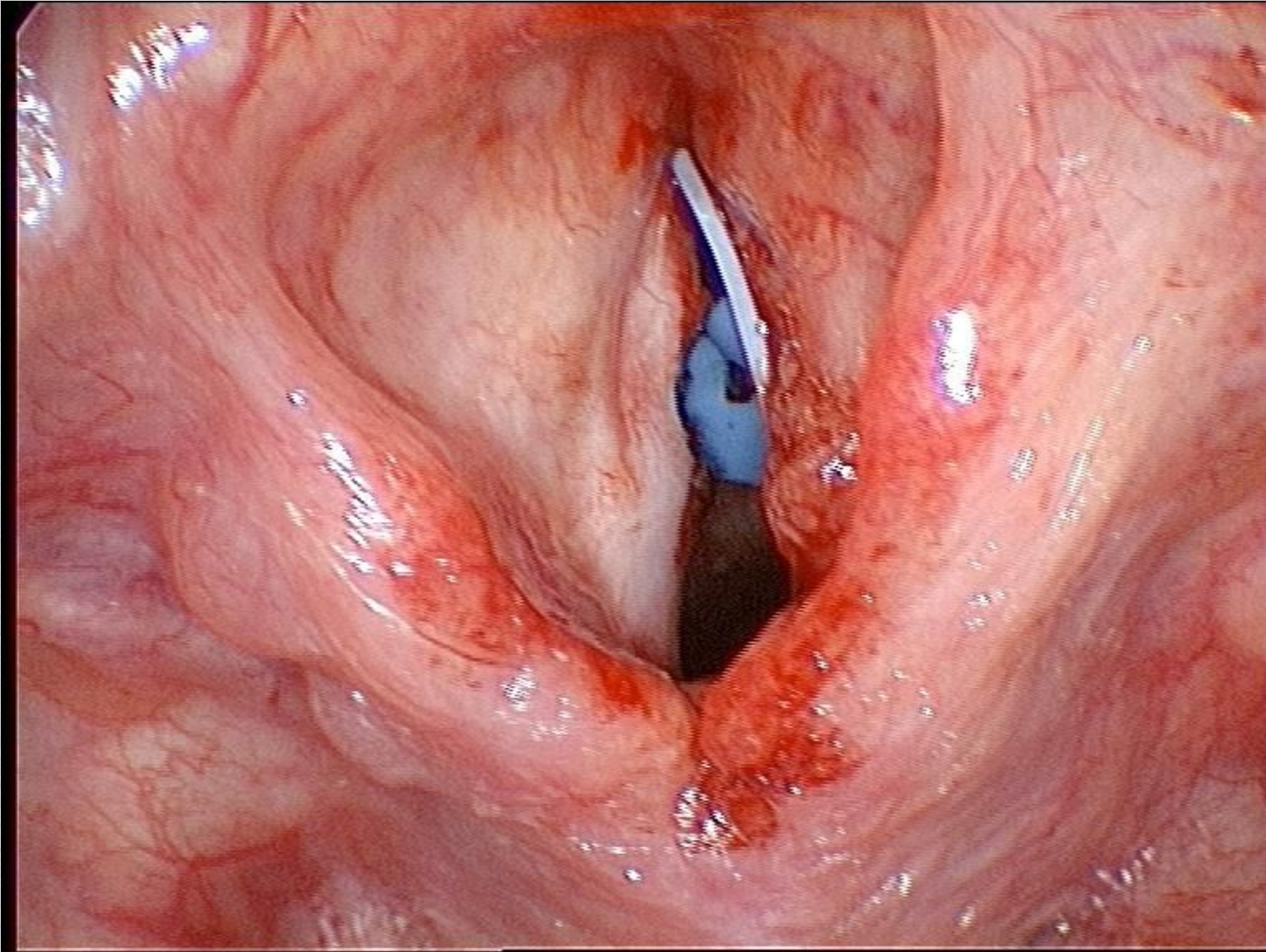


T-Tube In Situ

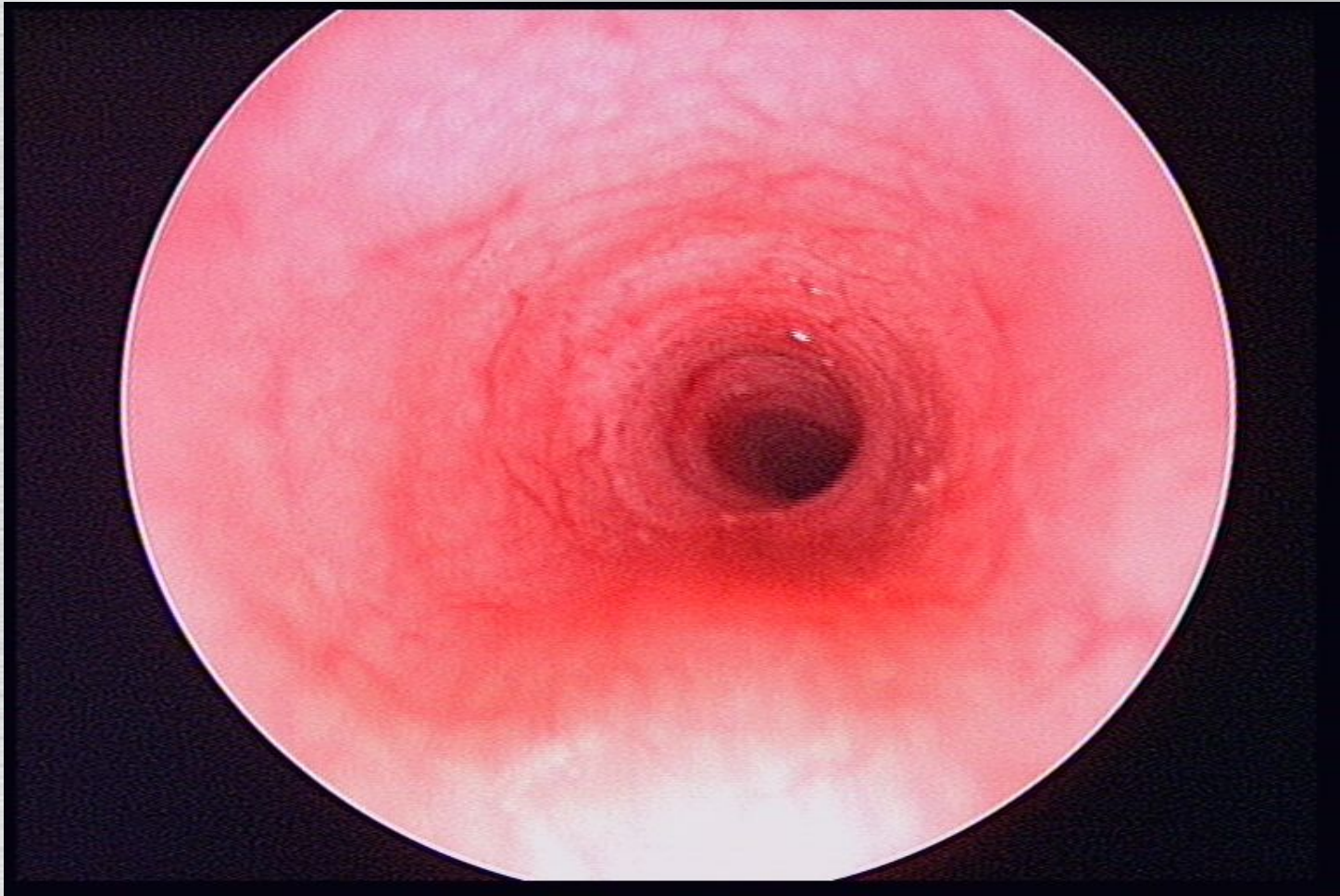


Webs

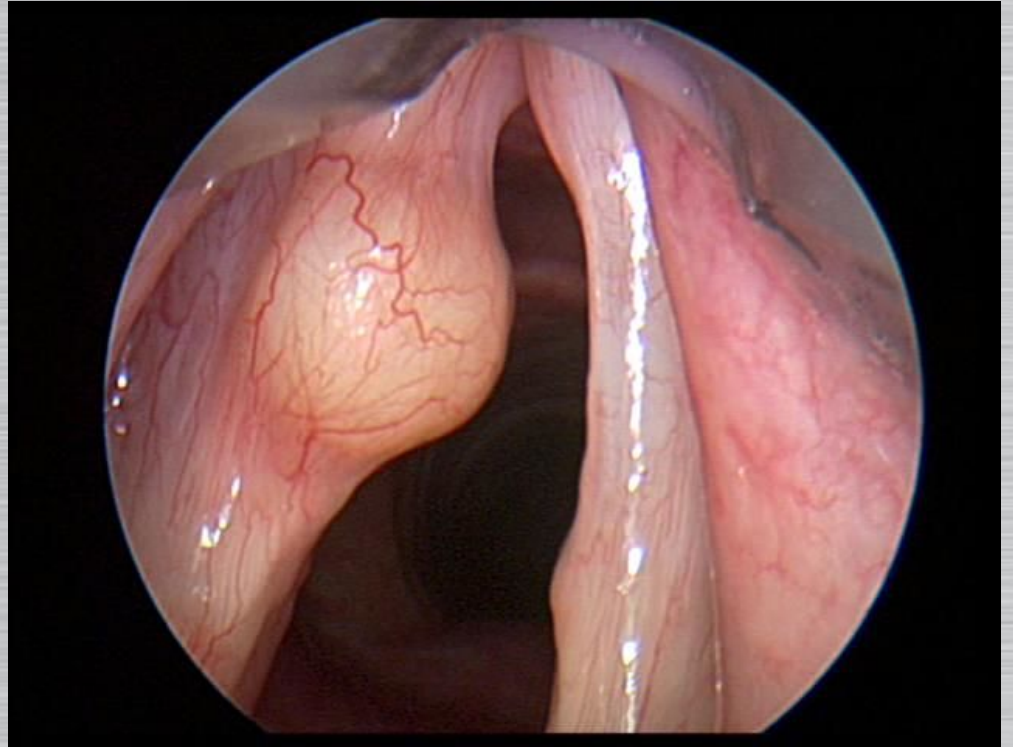
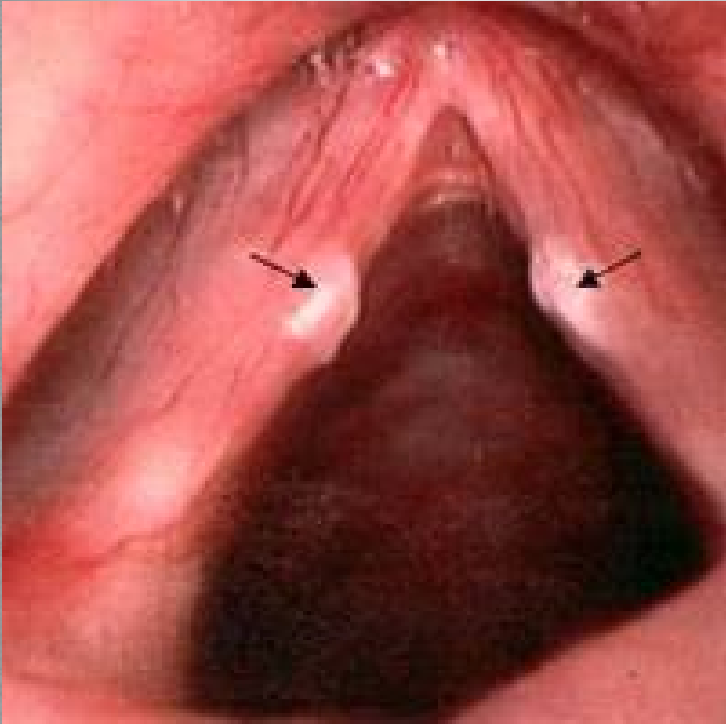




Tracheal Stenosis



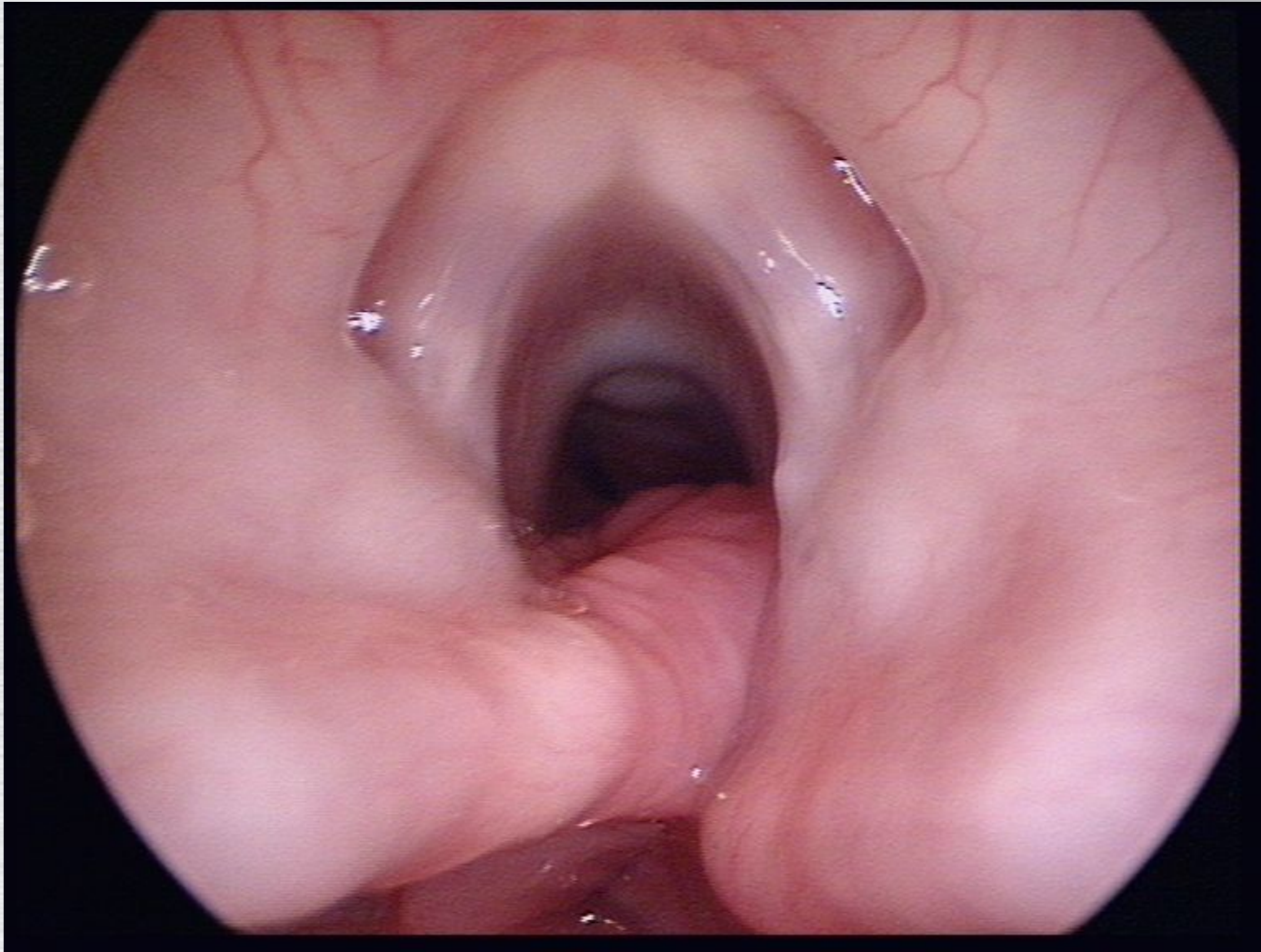
Vc nodules vs intracordal cyst



Valecullar cyst



Cleft larynx



Laryngeal Cleft



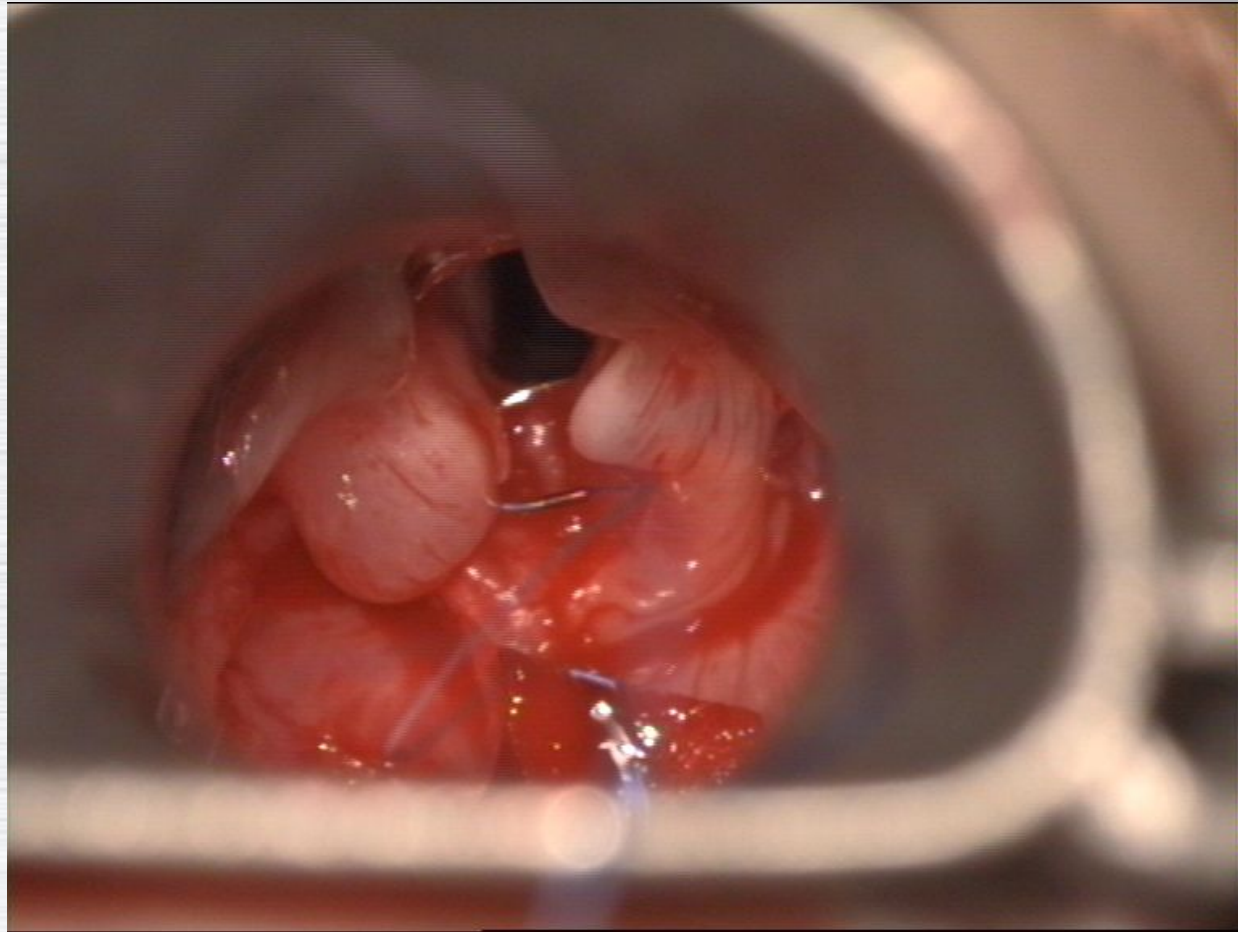
Type I
- To VC's

Type II
- Involving cricoid

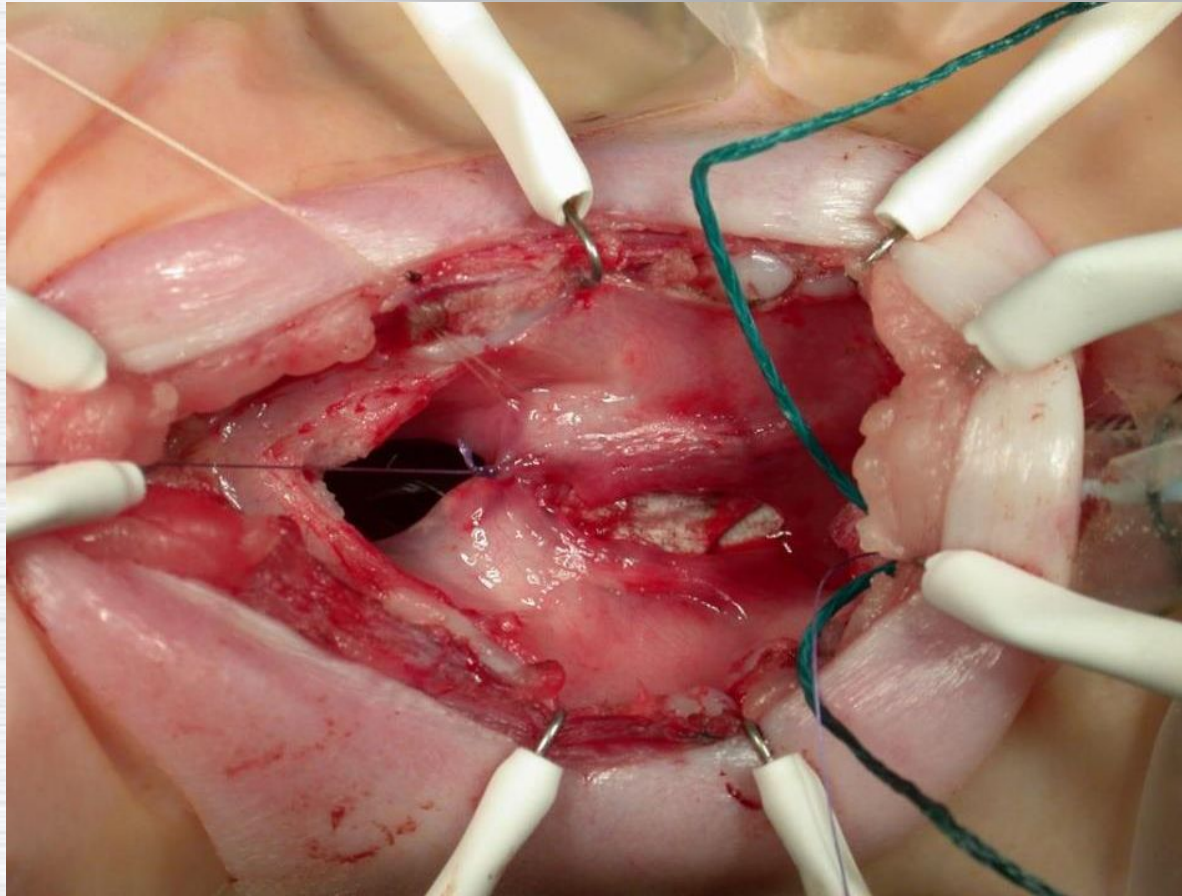
Type III
- above thoracic
outlet

Type IV
- below thoracic
outlet

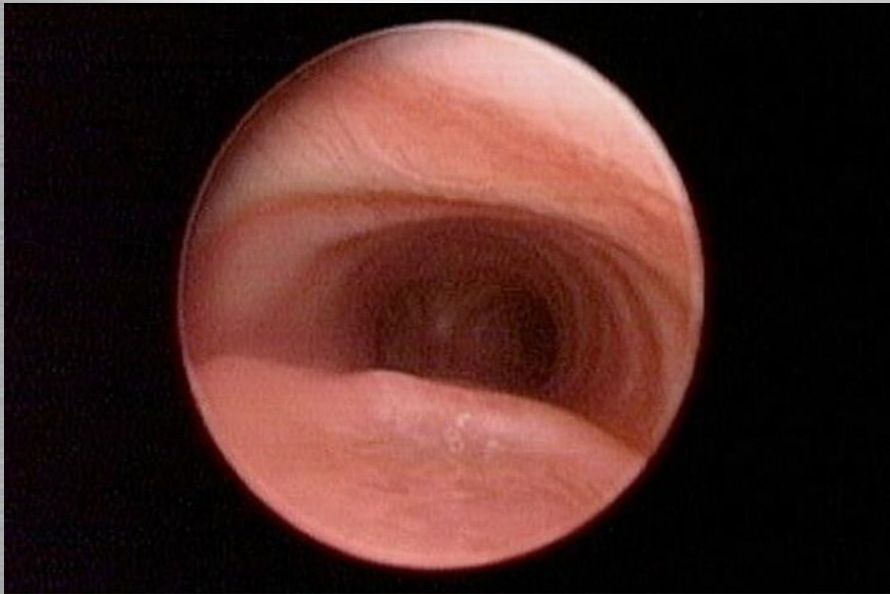
Endoscopic repair



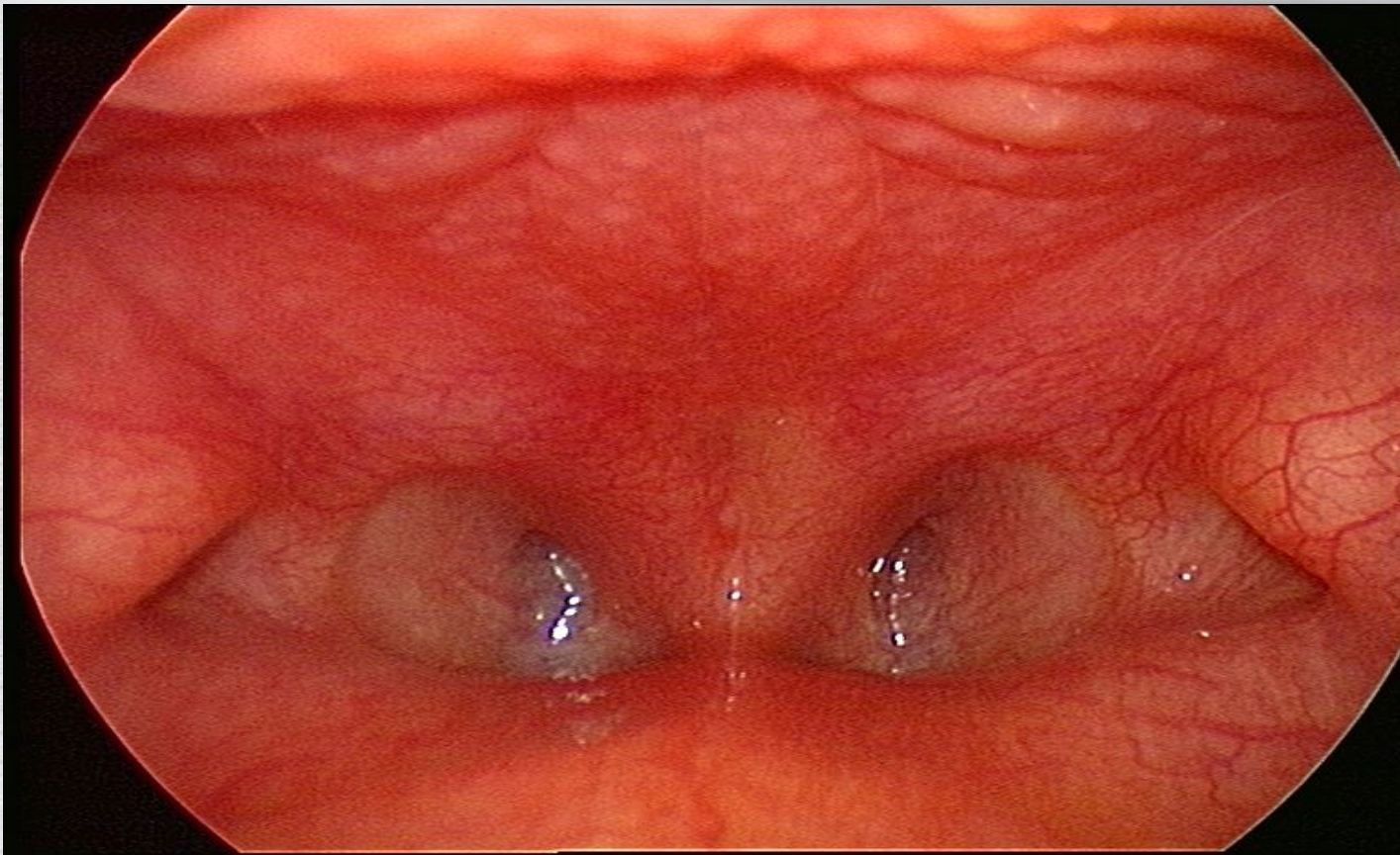
Open Repair



Trache Oesophageal Fistula



Choanal atresia



Choanal atresia

Presentation – Bilateral choanal atresia

A neonatal respiratory emergency
Immediate management with taped-in oral airway
Urgent CHARGE work-up and ?CT scan
Trans-nasal correction in first week of life



Choanal atresia

4 of the 6 'C-H-A-R-G-E' features.

C oloboma

H eart disease

A tresia choanae

R etarded growth and development

G enital hypoplasia

E ar anomolies and/or deafness

Presentation – Unilateral choanal atresia

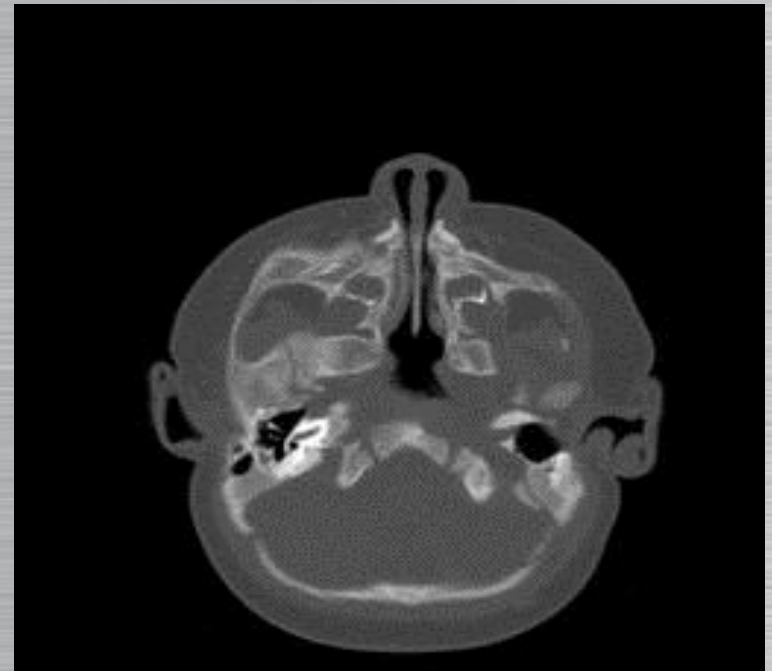
Non-urgent presentation with unilateral nasal discharge and obstruction

Routine CT scan

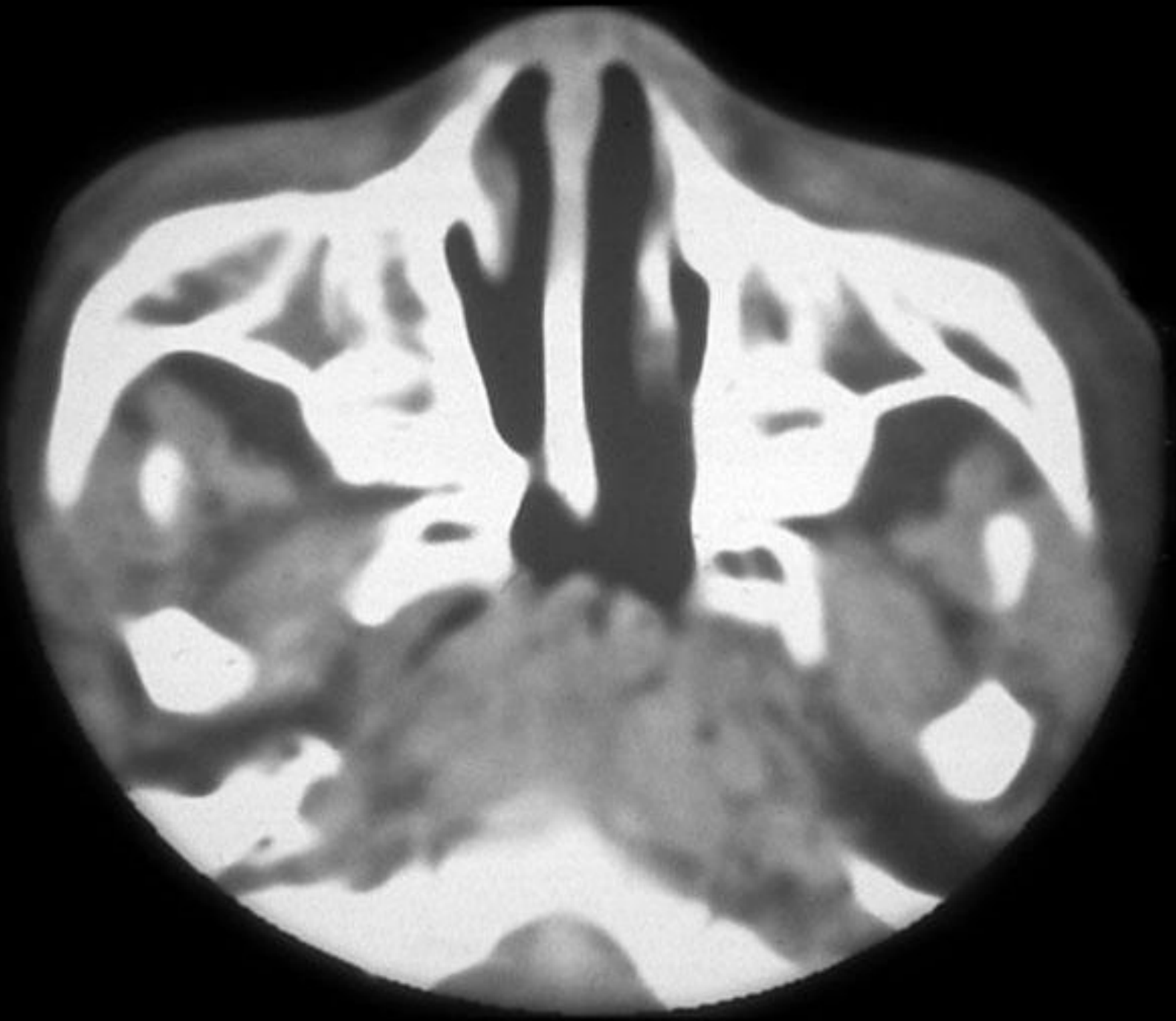
Elective correction 1-5 years

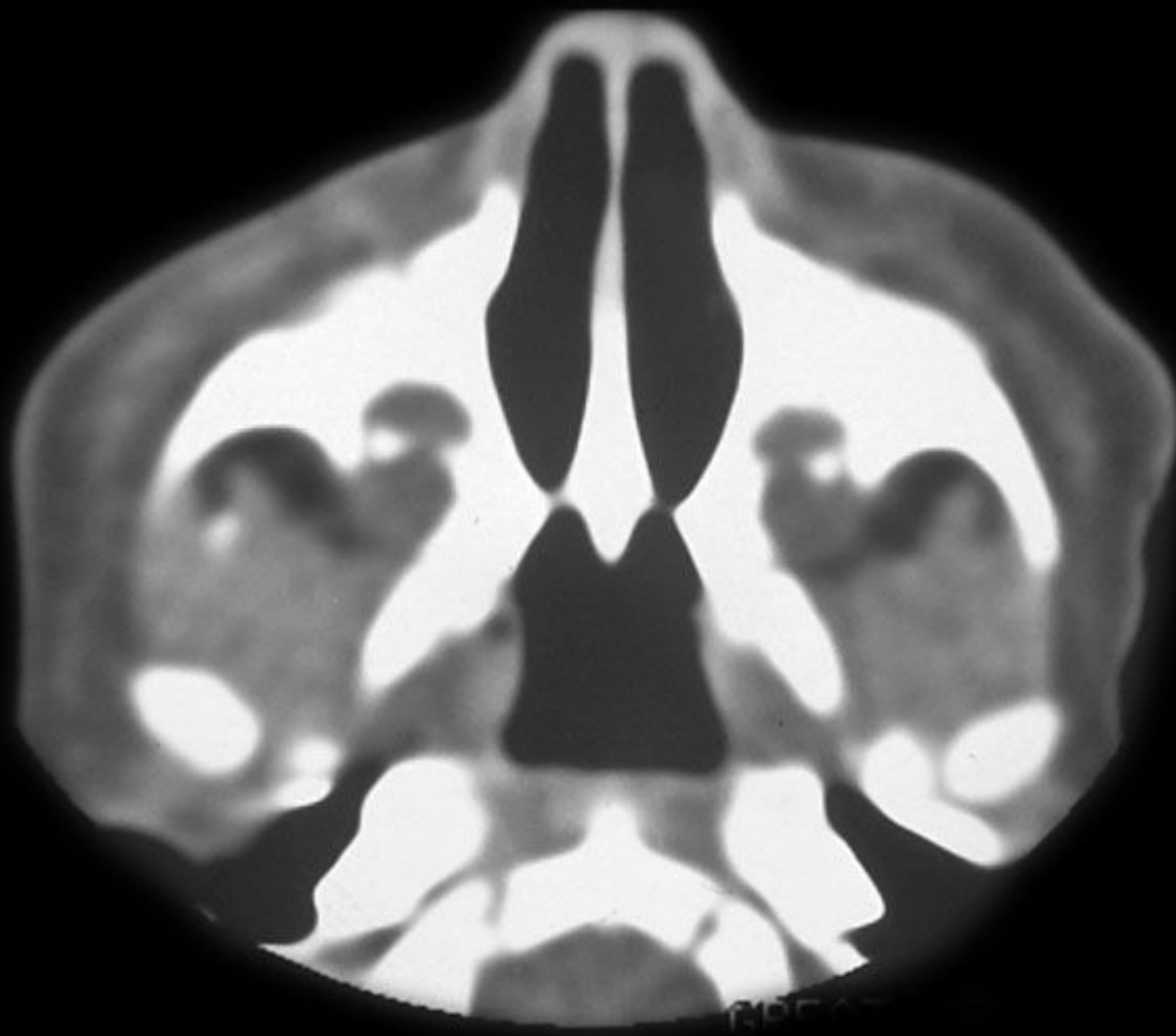
Differential diagnosis

- Neonatal rhinitis
- Masses
 - Post nasal space
 - Teratoma
 - Anterior nasal space
 - Glioma
 - *Midline nasal dermoid*
 - Meningocele
- Mid nasal and pyriform aperture stenosis
 - (Foreign body if unilateral)

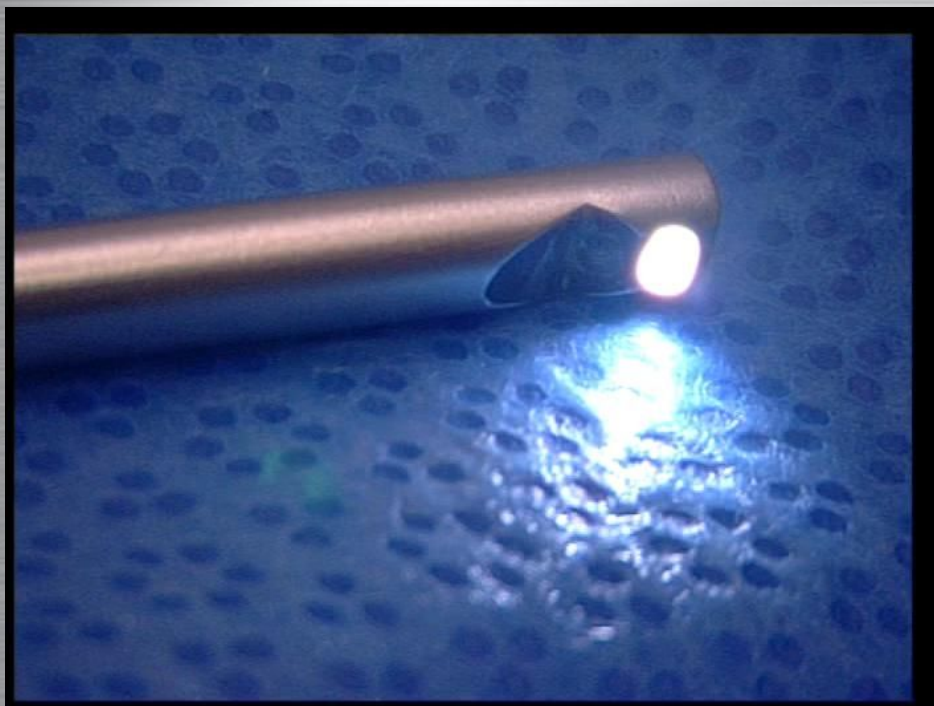


Choanal atresia

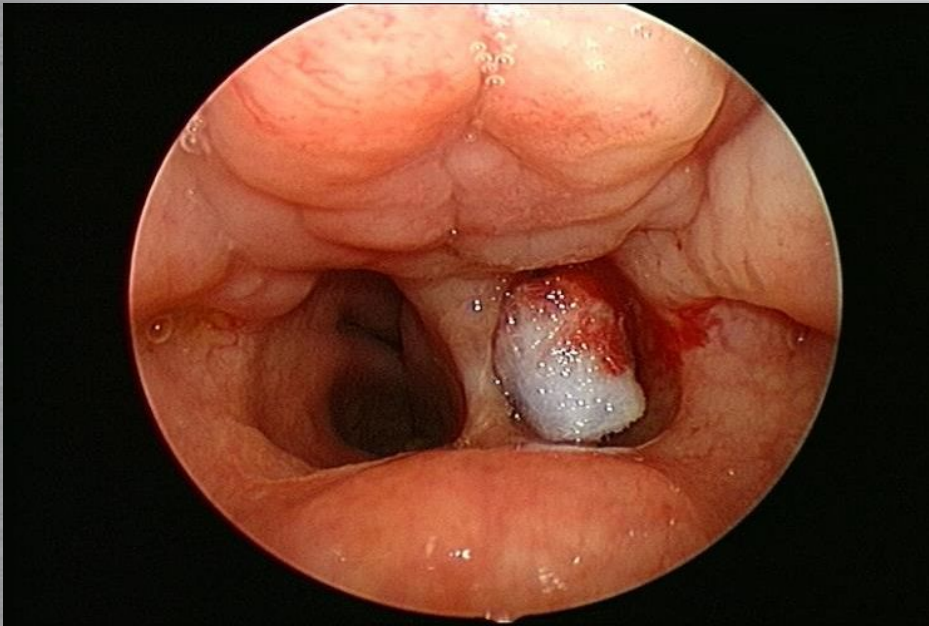




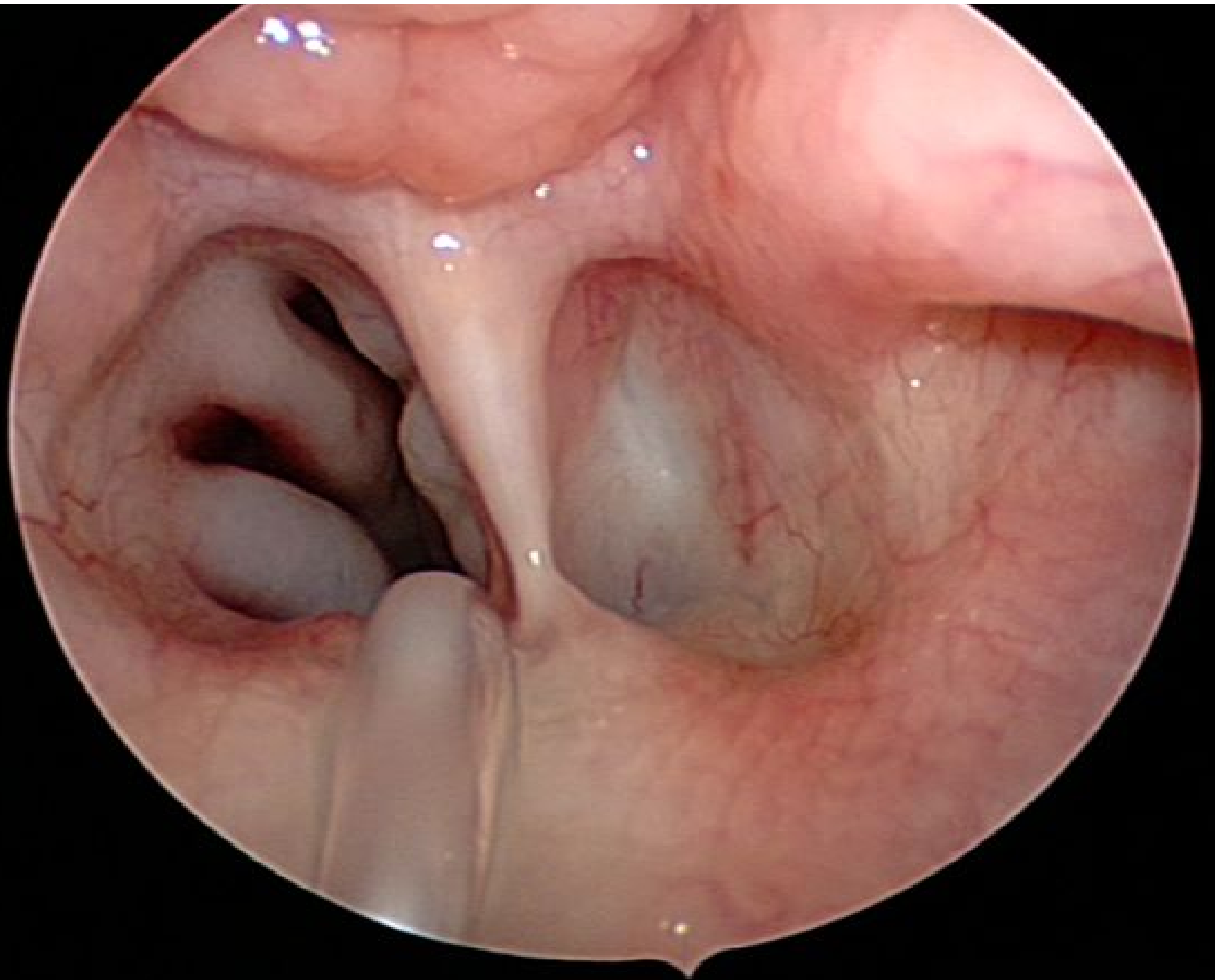
Trans nasal approach with 120° telescope



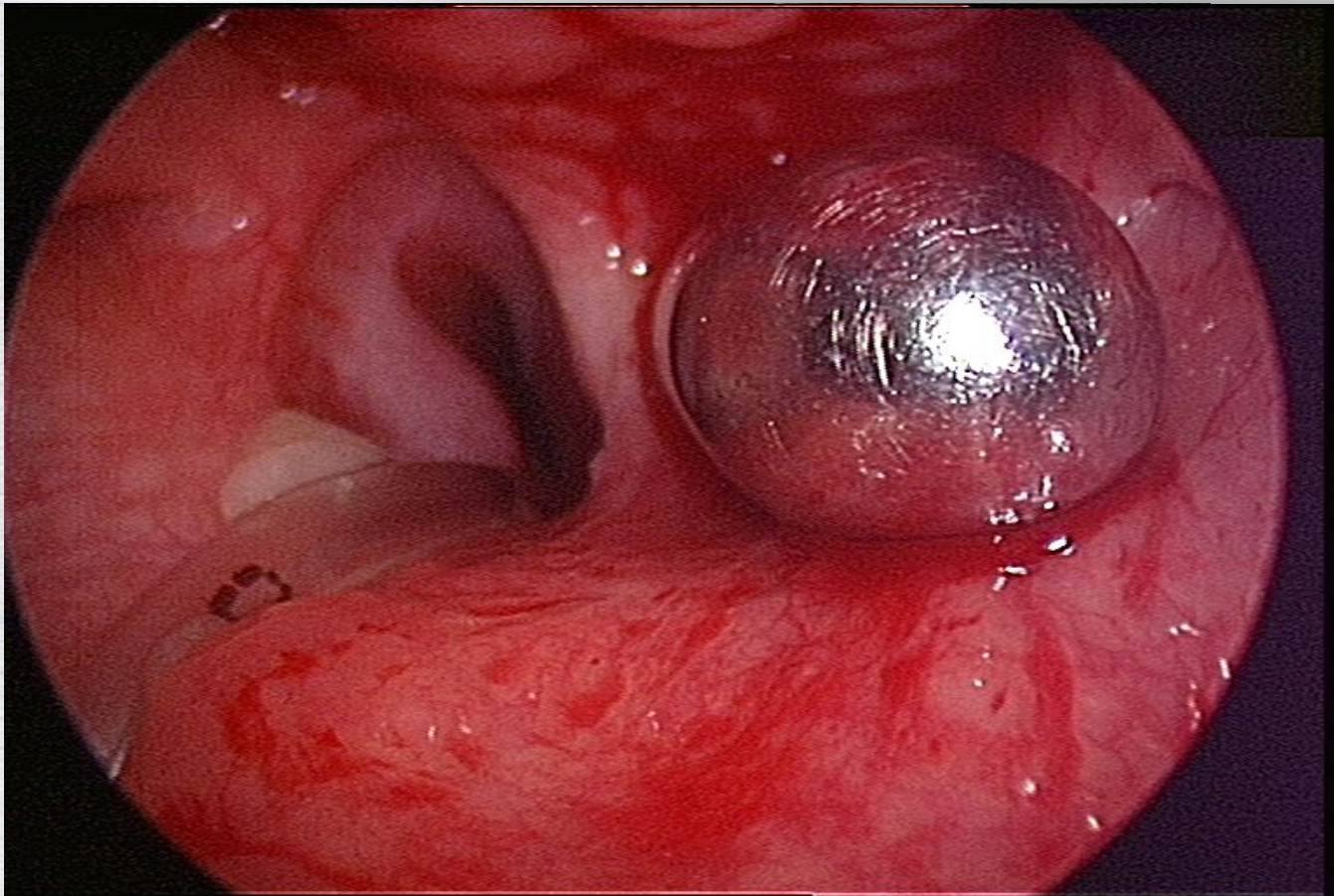
Adrenaline and dilators



Choanal atresia



Trans nasal approach with 120° telescope - dilatation

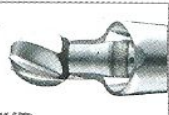


Trans nasal approach with 120° telescope - drilling



A. Pediatric Round Bur, 2.9mm

- Length: 10cm
- Straight shaft
- Application: choanal atresia
- Operating speed: up to 5,000 RPM FWD
- 5/box



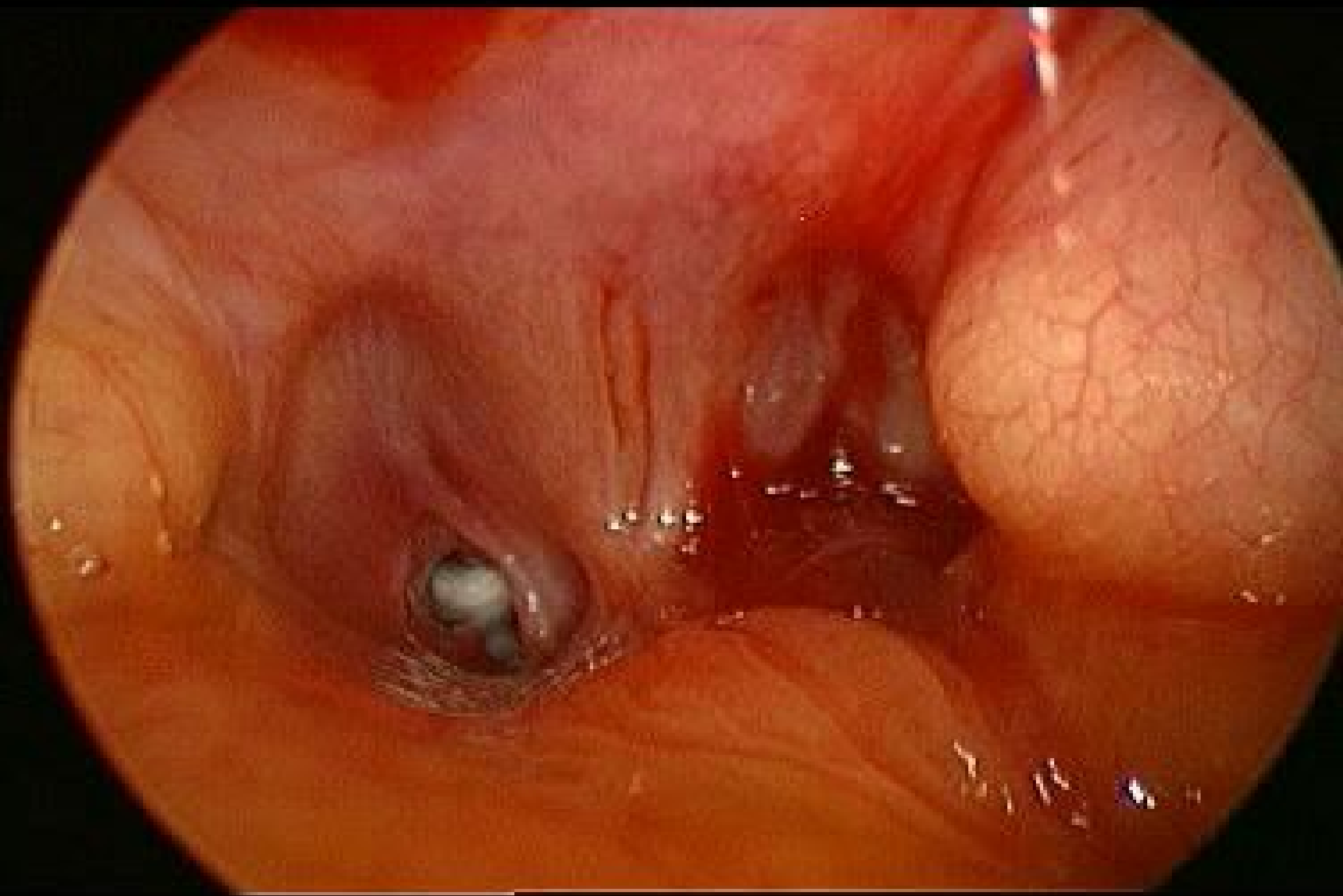
18-82960

G. Choanal Atresia Bur, High-Speed, 4mm

- Length: 13cm
- Curved shaft
- Cannulated suction bur tip
- Application: removal of vomer
- Operating speed: up to 12,000 RPM FWD
- 3/box
- Developed in conjunction with Gary Josephson, MD

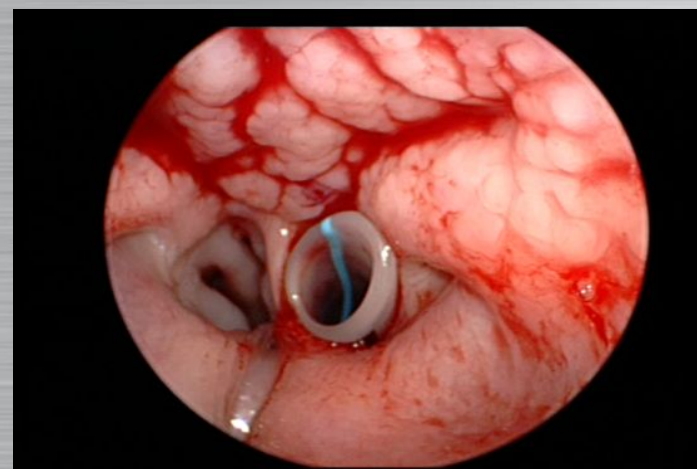


18-83673HS



Stents

- Bilateral cases
 - External bridge piece to avoid damage to columella
 - 6 weeks
 - Only dilate if needed
 - 4.5 Portex for term
- Unilateral
 - Avoid
 - Intranasal



Choanal atresia

Mitomycin

Sub-cytotoxic dose inhibits fibroblasts

2 mg/ml applied topically for 4 minutes

Topical Mitomycin as an Adjunct to Choanal Atresia Repair -

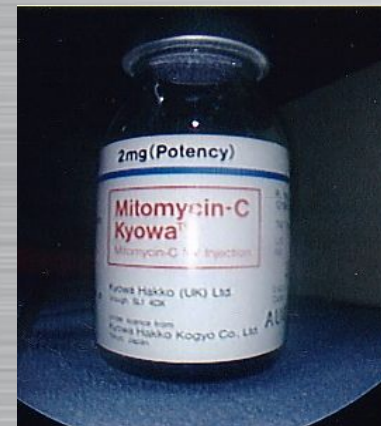
Arch Otolaryngol. 2002;128:398-400.

Surgical Management of Choanal Atresia Improved Outcome Using Mitomycin (0.4 mg/mL)

Arch Otolaryngol. 2001;127:1375-1380.

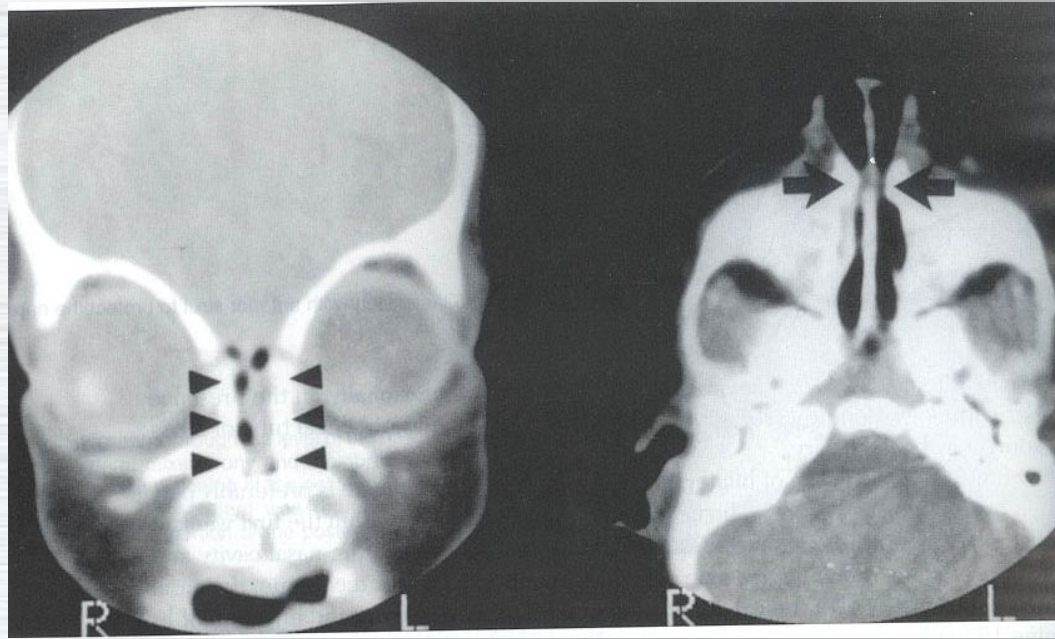
The expression of mRNA for some extracellular matrix proteins (elastase, hyaluronidase, and procollagen) was downregulated in the mitomycin test groups

Laryngoscope. 113(2):237-242, February 2003



Choanal atresia

Pyramiform Stenosis



PROGRESSIVE

Definition: JRRP

Juvenile

Mean age at diagnosis 3years

More aggressive than adult disease

Recurrent

Average lifetime procedures = 21

Respiratory

Usually larynx

If extends below larynx tends to be younger

Papillomatosis

With rare dysplasia and progression to carcinoma

HPV types

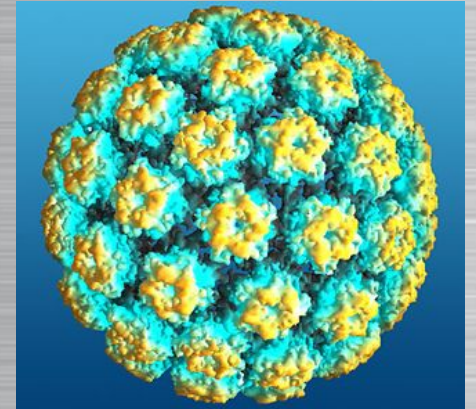
6

11



Benign genital warts/JRRP

In JRRP, 11 associated with more severe disease "Eleven Ξ Heaven"

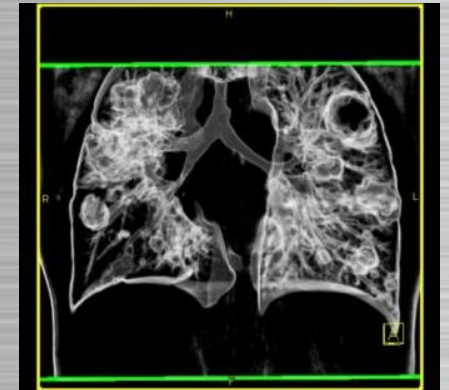


16

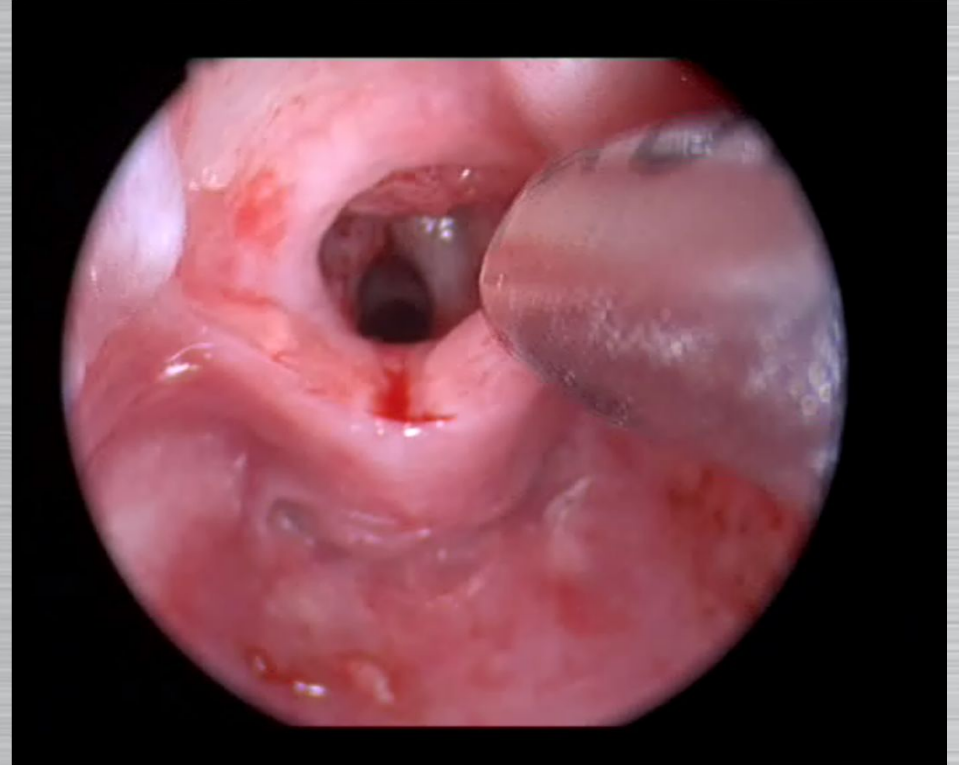
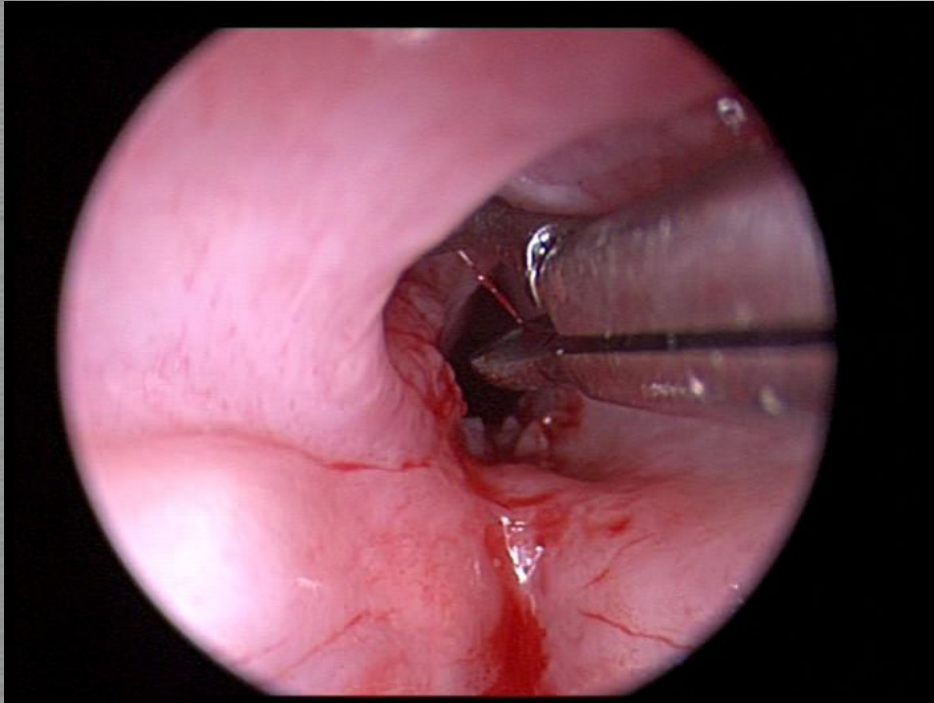
18



Cervical cancer



Juvenile respiratory papilloma

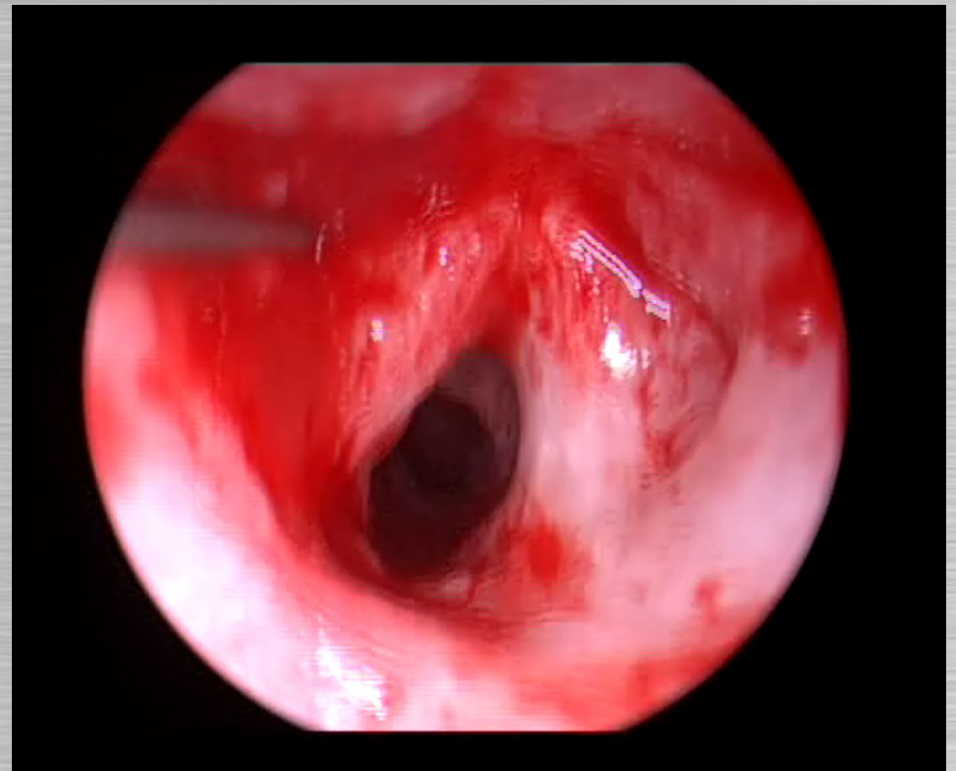


- Cidofovir

 - 5mg/ml

 - 3ml = 15mg

- Risk/Benefit



Subglottic Haemangioma



Subglottic Haemangioma

- Presentation peaks at 6 weeks

- stridor,
- feeding difficulties,
- FTT

- Usual site is left lateral subglottis

- Occasionally in trachea and glottis



Subglottic Haemangioma

- 50% of patients with SGH have associated head and neck cutaneous haemangioma

- Diagnosis at endoscopy
- Can be biopsied safely as capillary not cavernous haemangioma



Management

–*Cutaneous* usually regress by 6-8 years

–*Subglottic* involute sufficiently so that most are asymptomatic by 2-3 years

- Observation

50 - 70% will need a tracheostomy until about 18 months

Medical treatments

Surgical treatments

Management

- Interferon
- Steroids
- Propranalol
- Tracheostomy
- Laser KTP/C0₂
- Steroid injection
- Open excision
- Microdebrider

First GOS patient: SGH, trachy after debulk, July 2008



Propranolol

Week1 1mg/kg/day divided into 3 doses.

Week 2: ↑ 2mg/kg/day divided into 3 doses

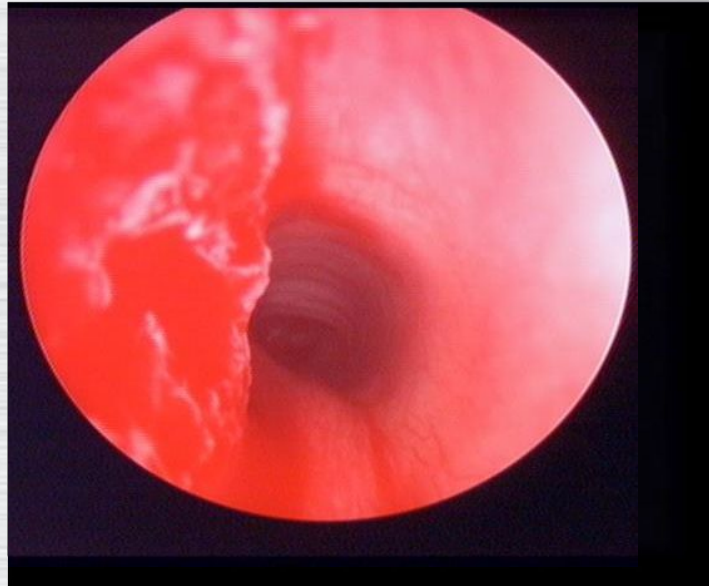
Up to 9/12 adjust by weight

From 9/12 no weight adjustment

?12/12 ½ dose every week.

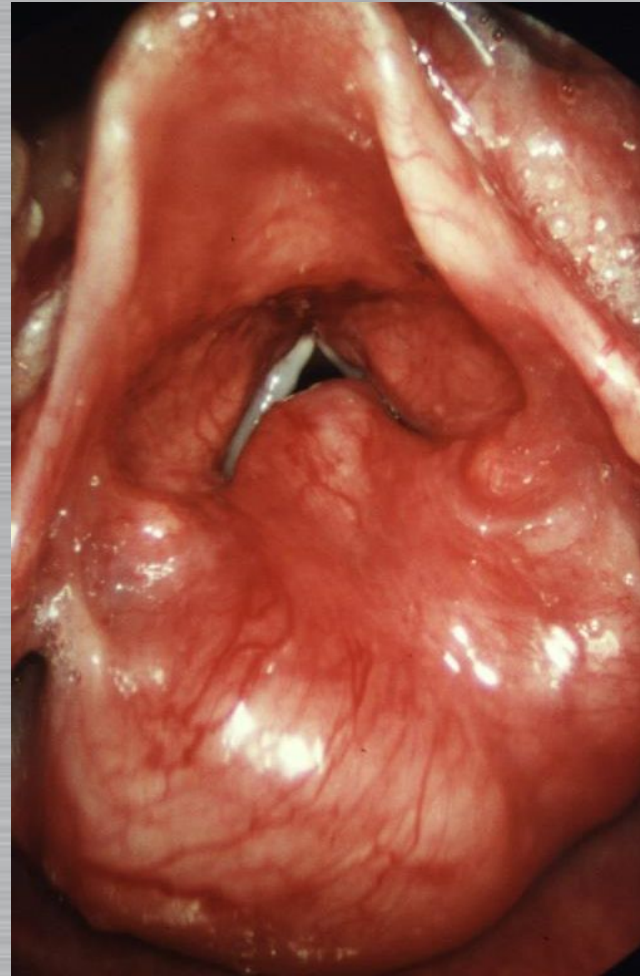
Subglottic Haemangioma

- Endoscopic resection of haemangioma

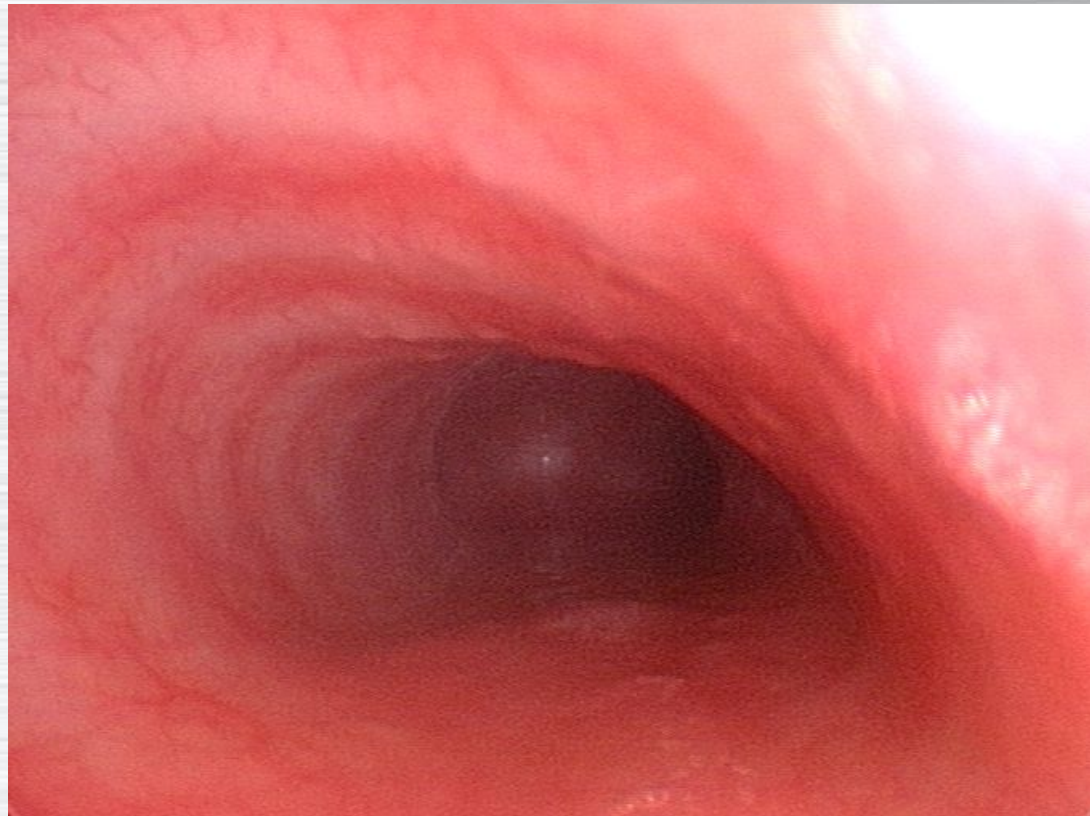


Tumours

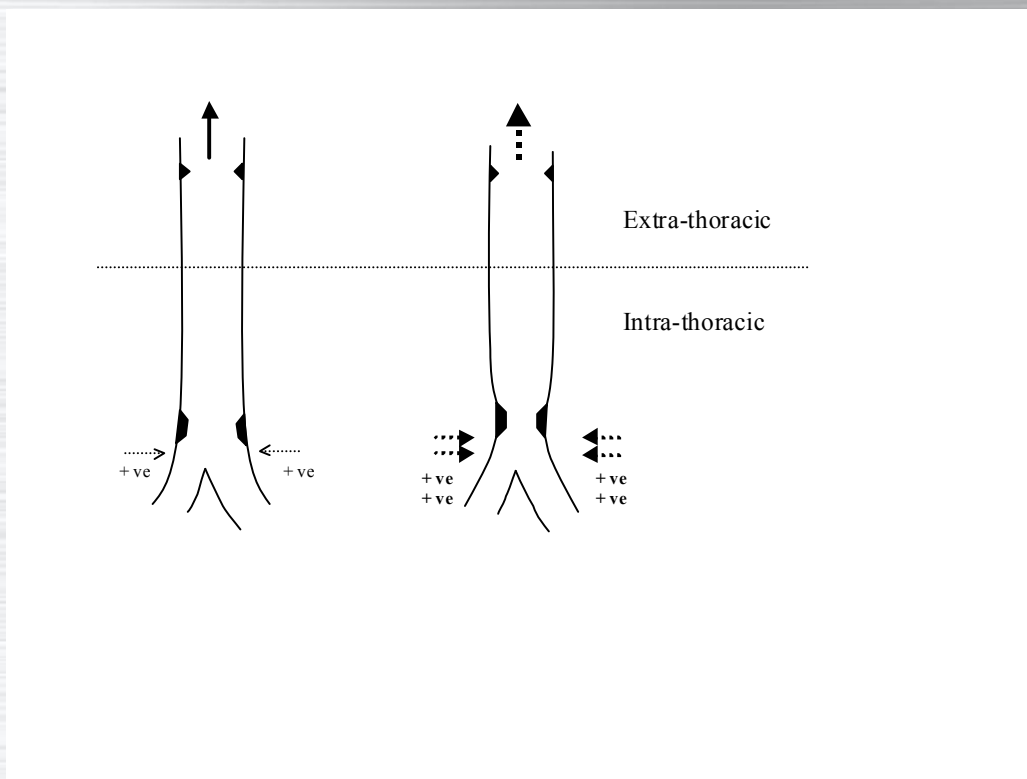
- RRP commonest
- Chondroma
- Squamous Ca 2nd RRP
- Rhabdomyosarcoma
- Lymphoma
- Vascular - disappears



Tracheobronchomalacia

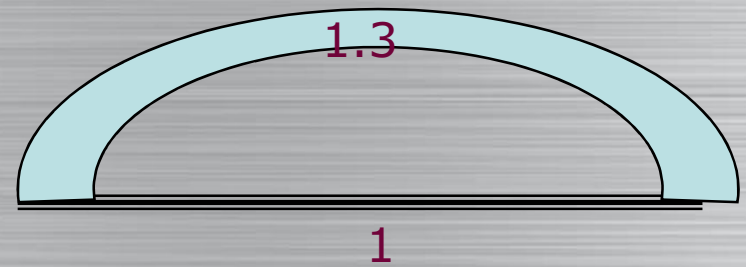
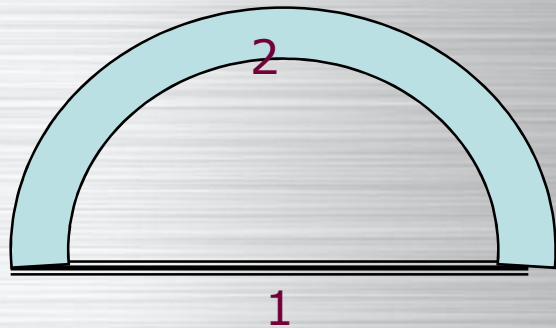


Collapse and increasing obstruction on expiration with intra thoracic obstruction



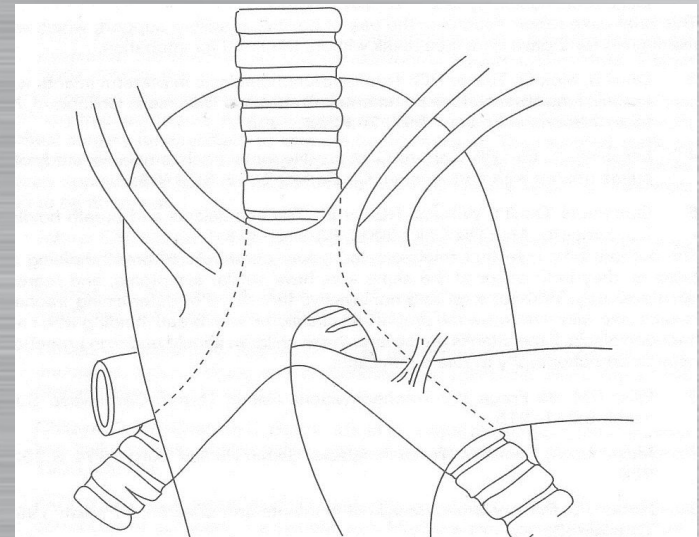
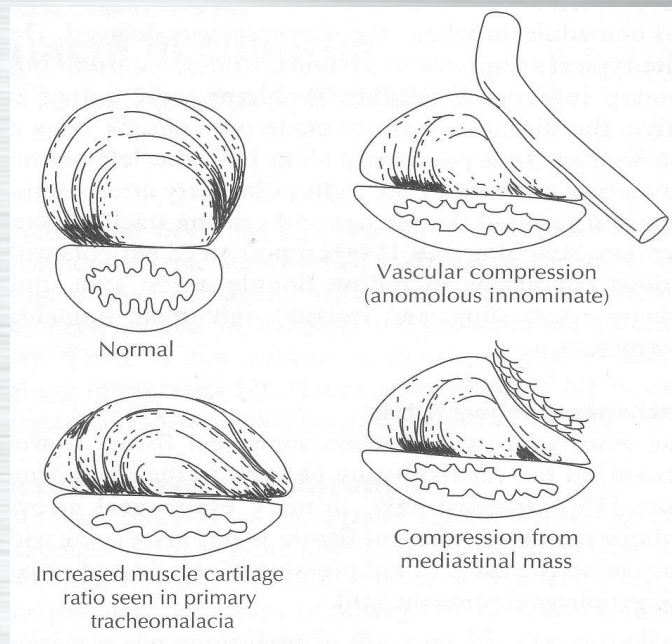
Primary Tracheobronchomalacia

Cartilage to muscle Ratio should be 2:1



Secondary Tracheobronchomalacia

- Compression



Tracheobronchomalacia Diagnosis

Cyanotic episodes

Cough

Aspiration

Expiratory

MLB

–Avoid physical or airway splinting
(underdiagnosis)

–Coughing (overdiagnosis)

•Bronchography

•stridor



Tracheobronchomalacia

- Prognosis

- Usually improves by 18 months

- Can be severe (fatal)

- Treatment

- Aortopexy

- Bronchopexy

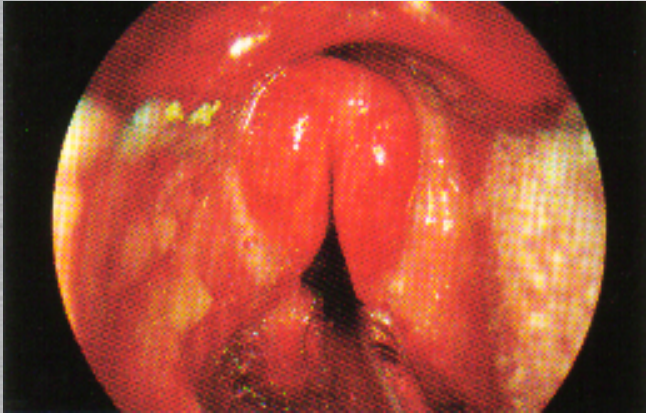
- Stents

Cystic Hygroma



ACUTE

Acute Epiglottitis



H. Infuenzae type b

Age 3 – 6 years

Sore throat, drooling, fever

Upright position, neck extended

Intubate in OT

Laryngotracheobronchitis (Croup)

Parainfluenza virus

Ages 3 months to 3 years

Barking cough

Progresses slowly

Rarely requires intubation



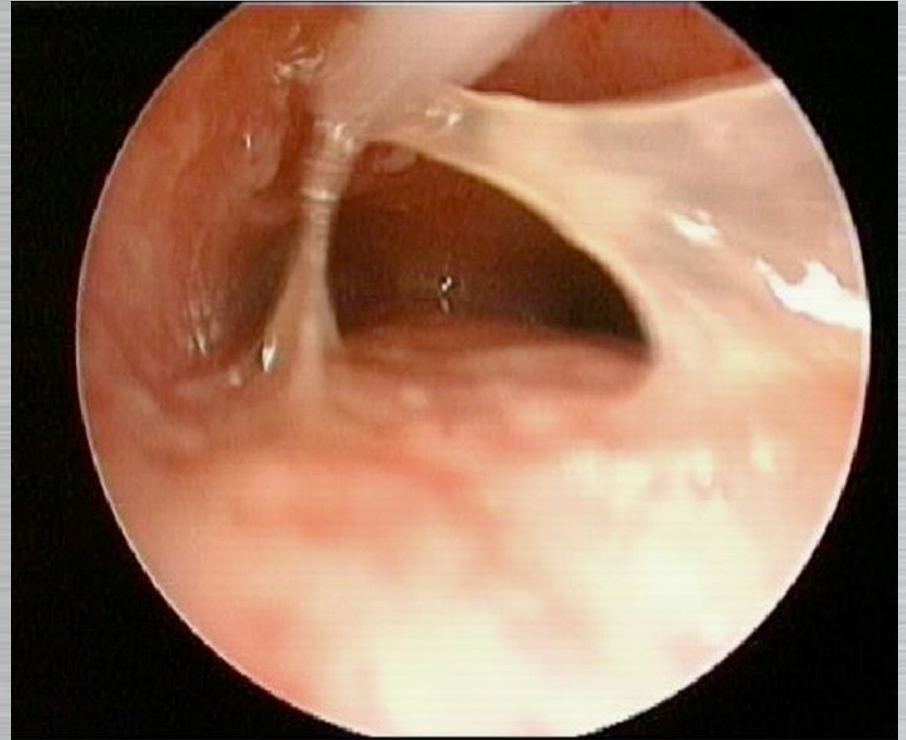
Steeple sign

Retropharyngeal Abscess

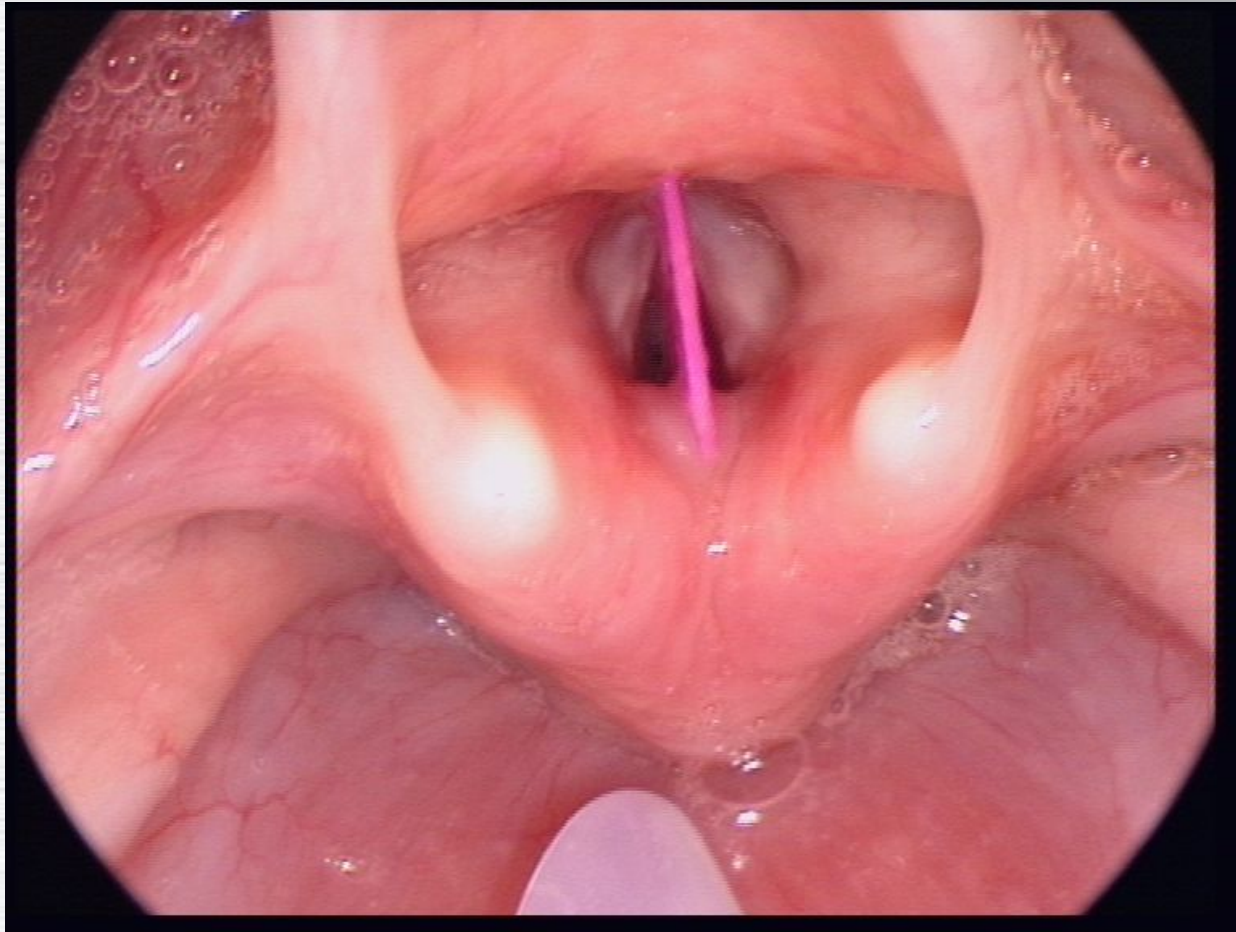


Bacterial tracheitis

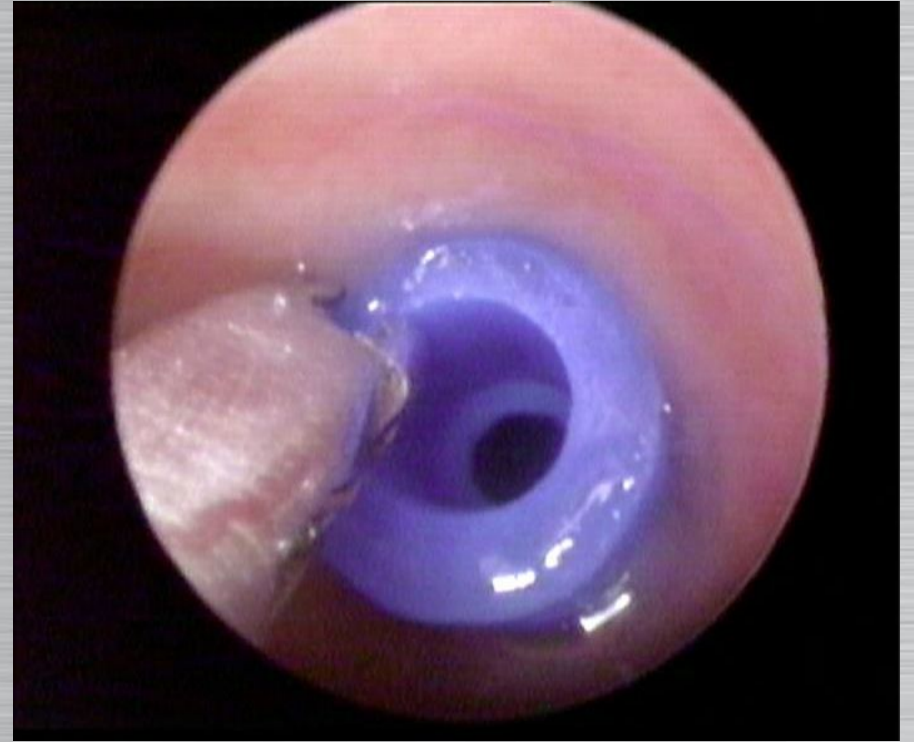
- Staph Aureus
- H. Flu

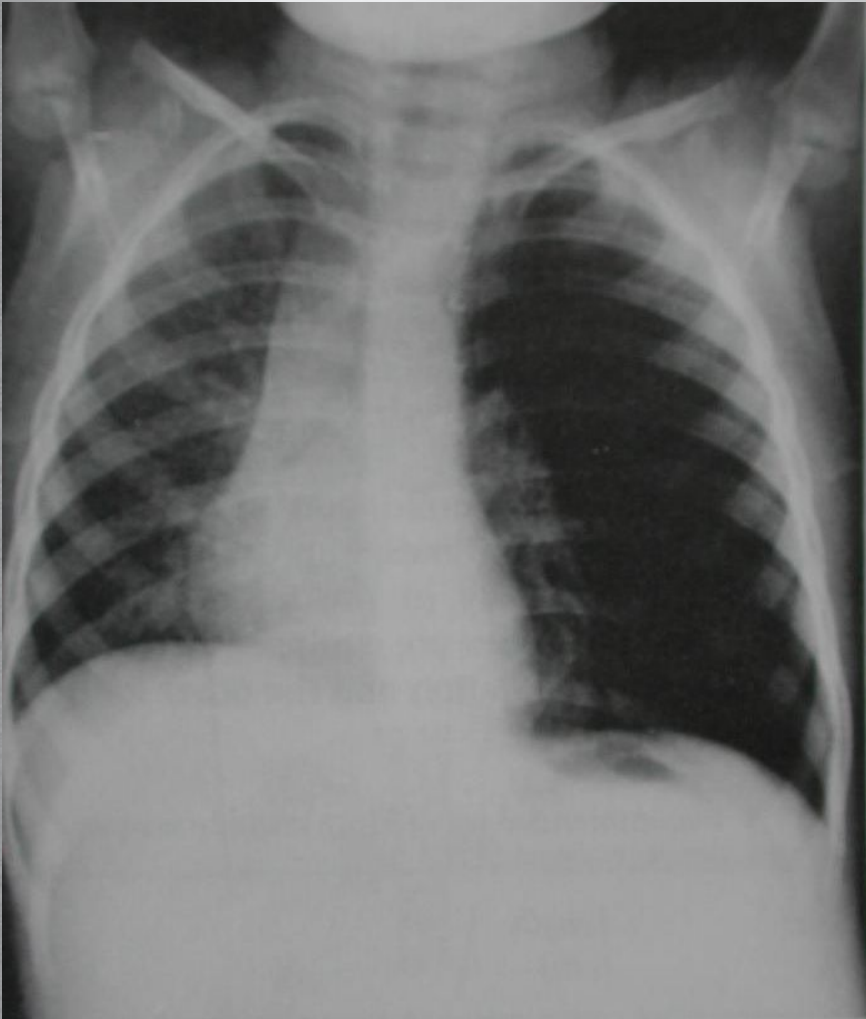


Laryngeal FB

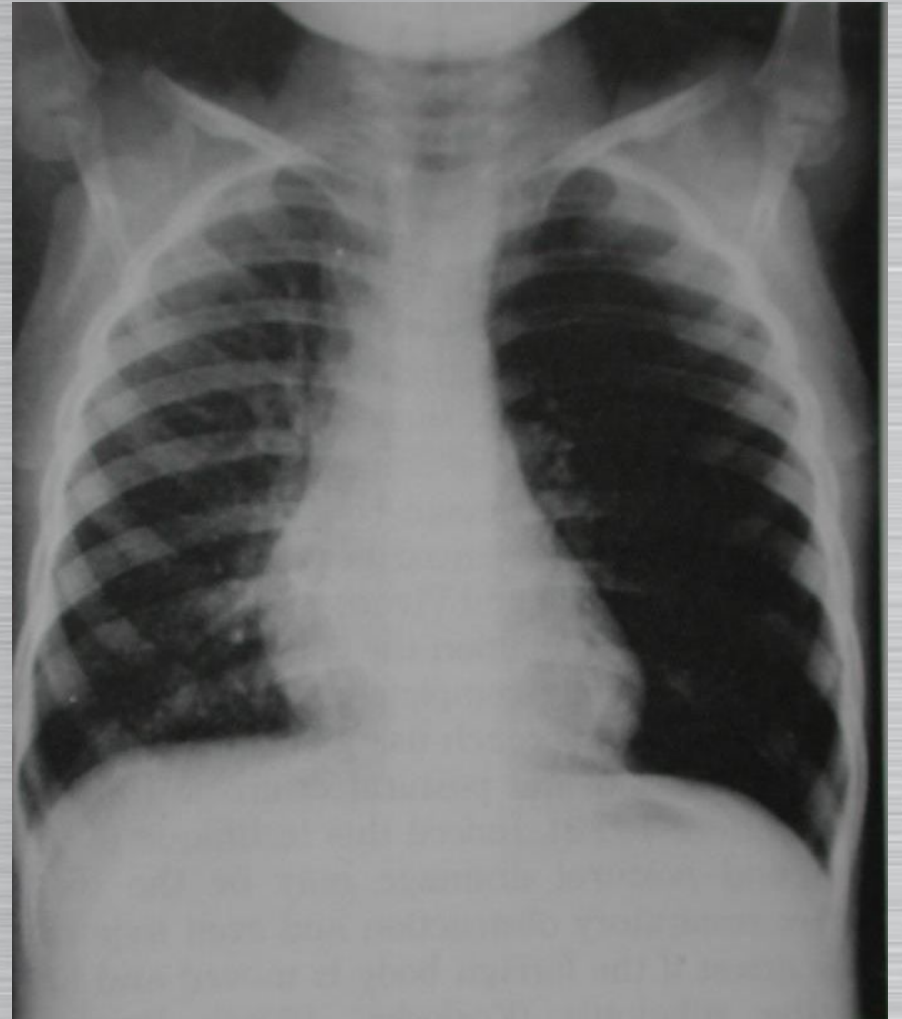


Bronchial Foreign bodies





Expiration



Inspiration