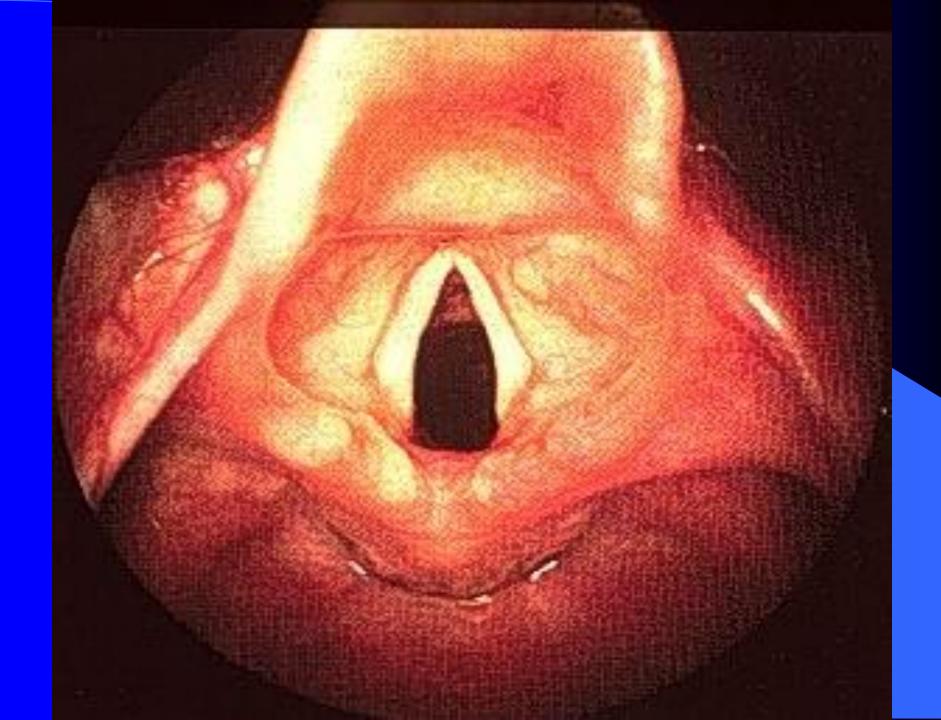
UPPER AIRWAY OBSTRUCTION



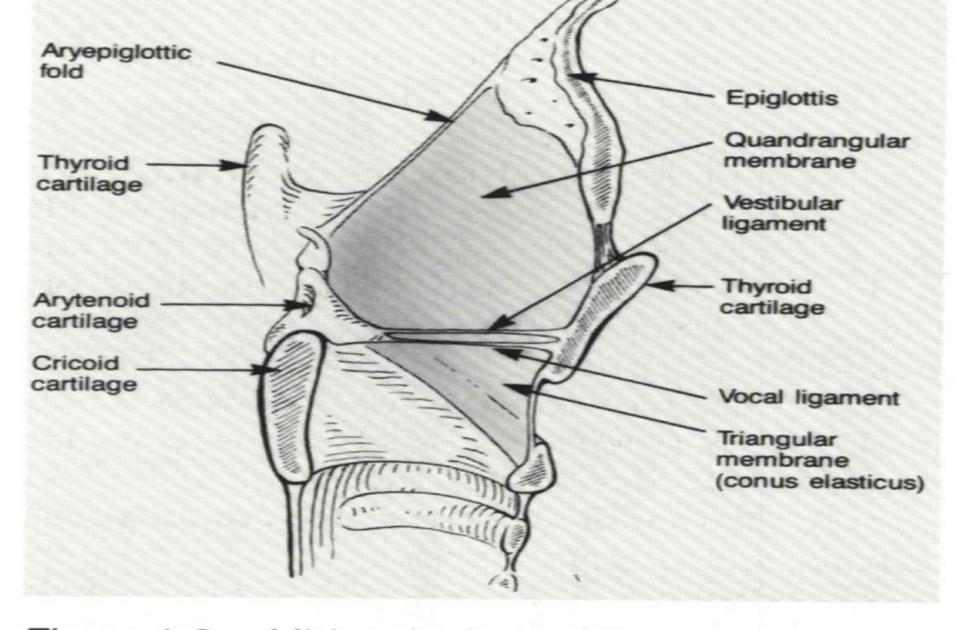


Figure 1-3. Midsagittal cut of the larynx showing the fibroelastic membrane attachments.

ANATOMY

.)

- * Infant & Pediatric larynx
 - position is higher at birth
 - epiglottis lying at the nasopharynx make the neonate an obligate nasal breather x 4-6 months
 - cartilage & soft tissue are more soft

LARYNGEAL ANATOMY in pediatric (cont.)

- Soft tissue are less adherent to the underlying cartilage
 - susceptible to collapse
 - less resistant to develop submucosal edema

Omega shape Epiglottis
Subglottis is the narrowest part of AW in children.
In adults glottis is the narrowest.

TRACHEAL ANATOMY

- consists of 16 to 20 incomplete cartilaginous rings
- the post wall is a membranous part
- length is approximately 11 cm
- diameter 19 mm male

16 mm female



TRACHEAL ANATOMY (cont.)

Pediatric trachea:

Diameter: At Birth 6 mm

6 mons 7.2 mm

1 year 7.8 mm

4 years 11 mm

TABLE 91-2. APPROXIMATE SIZE OF ENDOTRACHEAL AND TRACHEOSTOMY TUBES FOR INFANTS AND CHILDREN

Age	ID Size (mm)	
Premature		
< 1000 gm	2.5	
1000-2500 gm	3.0	
Neonate-6 months	3.0-3.5	
6 months 1 year	3.5-4.0	
1-2 years	4.0-5.0	
Beyond 2 years	Age in years + 16	

Signs of Airway Obstruction

- Stridor is harsh high pitched musical sound produced by turbulance of air flow through a partial obstruction of the AW
- Stridor is a vary important sign of UAW obstruction
- It indicates pathologic narrowing of AW
 - potential resp obstruction
 - even death

Other signs of UAW obstruction

- flaring of the nasal alae
- retraction of the neck, intercostal and abdominal muscles
- Dyspnea
- Tachypnea
- Restlessness
- Cyanosis
- Subcutaneous emphysema

```
Glattle and Subplottic Commond
Sagralaryegeal
                                                                              Inflammatery
Congenital
                                                                              Viral croup
Natal obstruction, e.g., chostnal atresia, escephalocele
                                                                              Bacterial laryogeroscheitis, including dipleherta
Maconglossis, e.g., cretinien, hosphosgeoms, hemorgiona
                                                                              Tulk resolonie*
Facul sheletal anomalies—Picere Bohoa anomalad
                                                                              Surcoshe
                        Ciril culate
                                                                              Fungil, e.g., coccideddonormos
                         Trencher Collins syndrome
                                                                              Allergie referoa.
                         Missingsothia
                                                                              Koodermodynia bullene
Facial edema (fact: presunstation)
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Contraital cysts-Thyrogloud duct
                                                                               Mesplastic
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                 Rannla
                                                                                        Grandar cell myoblastonar
Lingual thymnal
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Admotomillar hypertrophy
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Scorer resulfoliarial transma
                                                                                           Lymphomerous
Neoplastic
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e.g., Bhalalossonarcoma
 Neurolopic
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 Subscrate selectrony pursescephalitis (Dosevos disease)
                                                                               Underent, v.g., post ductor legation postiatubation<sup>et</sup>
Postoperative
                                                                               Bilateral, e.g., mesingossekseele, hydrocephalus, vostoruloperitoneal
 Tongue obstruction
                                                                                                shoot failure
 Supraglottio
                                                                                              arthrogrygwais multiples congrunts'
Campenital
                                                                                              lacreshing abductor stical cord paralysis's
 Attendia
Web
                                                                                Neuroma<sup>b</sup>
                                                                                Treamatic
Langomalacia
 Cleft lervna-Cleft asytenoid
                                                                                Foreign body!
              Extension cleft.
                                                                                Inrobation-Eventurion of ventricle
                                                                                            Crumbrian things
 Interpretational System"
                                                                                            Showing.
 Ventriculor based hypertrophy**
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                Lateral
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 Showney will thereon
                                                                                 Postsurgical-trachesocrophageal fistola repris-
 Vocal road paralysis-Undateral
                       Bilderal (Arnold-Chiari syndrous)
```

Cost-Theroid cartilage forminal cyst*

Diagnostic assessment HISTORY

- Time of onset
- Possible trauma
- Characteristic of cry
- Relation of airway problem to feeding, position
- History of previous intubation
- Questions about possible aspiration of FB

HISTORY (cont.)

If stridor is present since birth:

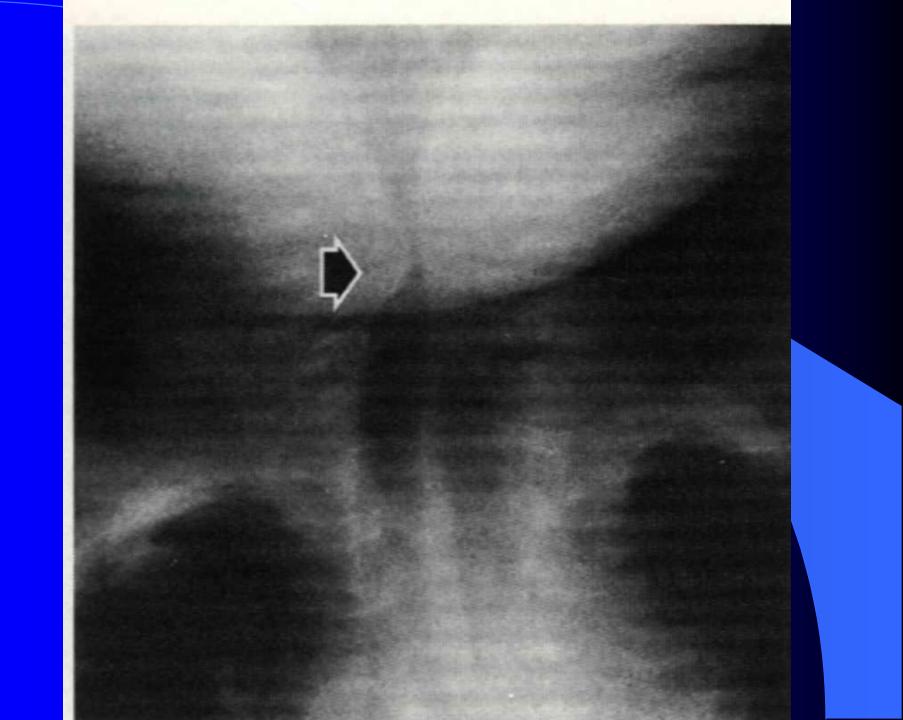
- * conginetal laryngomalacia
- * subglottic stenosis
- * vocal cord paralysis
- * vascular rings

If onset of stridor is gradual and progressing:

- * subglottic hemangioma appear between 1-3 months of age
- * papilloma of the larynx appear at 6 months of age

RADIOLOGIC EVALUATION

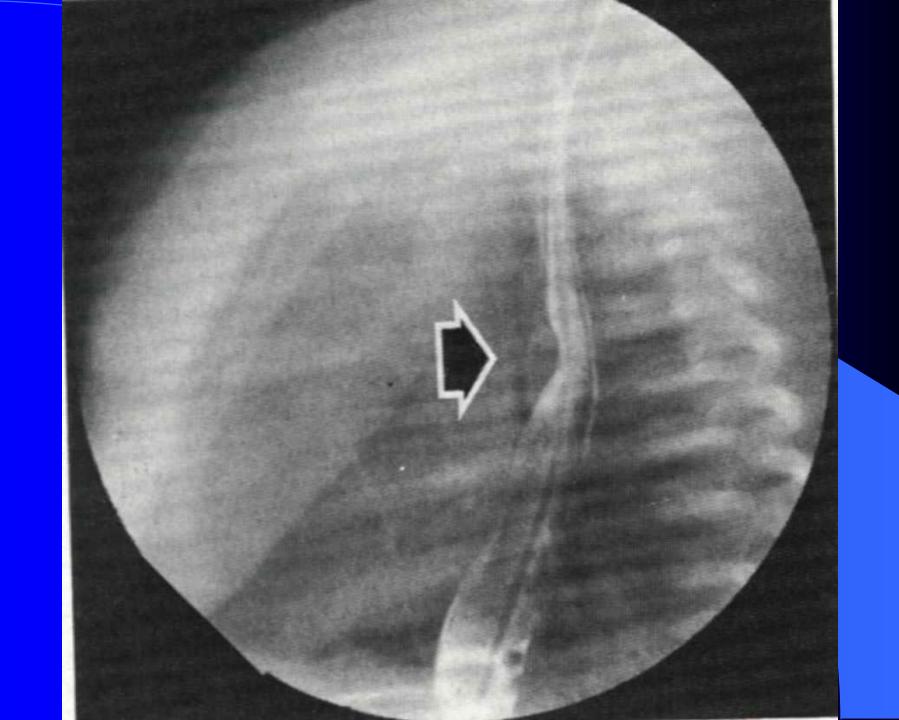
- Indicated for patient without respiratory distress
 - * Plain views soft tissue neck A.P. lateral
 - chest
 - * Mobile pharyngeal tissue may bulge during expiration in normal infants
 - * High-kilovoltage technique (croup series) AP view assesses subglottic region
 - * Flouroscopy: dynamic AW changes



RADIOLOGIC EVALUATION (cont.)

- * Barium swallow
 - Assess swallowing
 - R/O presence of vascular rings
- * CT scan.} good in evaluating mediastinum
- * MRI.





ENDOSCOPIC EVALUATION

- Mirror Examination: is not endoscopic.
 In older children and adults can provide information about hypopharynx and larynx.
- Telescopic Examination
 - * Fibroeptic Endoscopy excellent to assess the movement of VC.
 - * Rigid bronchoscopy done under GA
 - may enable removal of FB
 - assess the AW down to the main stem bronchi

Diagnostic assessment

- Flow volume loop
- ABG
 - late indicator of AWO
 - should not be used routinly to assess degree of obstruction

THERAPEUTIC OPTIONS

1. Observation/Medical Support

- a. ICU
- b. Airway team availability
- c. Oxygenation
- d. Steroids
- e. Antibiotics

(Cont. Therapeutic options)

- 2. Heimlich maneuver
- 3. N. P. Airway
- 4. Oral Airway
- 5. Esophageal airway
- 6. Transoral intubation

Cont. Therapeutic Options

- 7. Nasal intubation
- 8. Flexible fibroptic intubation
- 9. Transtracheal jet ventilation
- 10. Cricothyroidotomy
- 11. Tracheostomy

Table 131-1. Advantages and disadvantages of various types of airways

Airway	Indications	Advantages	Disadvantages
Nasopharyngeal	Obtundation with man- dibutar retrusion or pharyngeal collapse	Simple Maintains airway Facilitates suctioning Well-tolerated in alert patients	Requires normal ventilatory drive Can cause epistaxis
Oral	Obtundation Obstructed or injured na- sal airway	Simple Maintains airway Facilitates suctioning	Easily dislodged Requires normal ventilatory drive Poorly tolerated by alert patients
Oral intubation	Need for controlled ventu- lation Failure of simple mea- sures Aspiration control	Allows full control of ventilation Prevents aspiration	Requires expertise and proper equipment Potential injury to larynx and pharynx
Blind nasal	Potential cervical spine injury with need for airway control Massive oral cavity injury precluding oral intuba- tion	Can be performed without hy- perextending neck Eliminates need for laryngeal visualization	May aggravate airway problem if poorly executed May cause epistaxis Requires prior experience
Fiberoptic nasal	General airway control Potential cervical spine injury Massive oral cavity injury	Altows laryngeal visualization and placement of tube under direct visualization Less chance for error compared with blind nasal intubation	Difficult to see if excess secre- tions or bleeding present Requires added equipment and expertise
Transtracheal (jet ventilation)	Failure of above methods	Very rapid control of oxygena- tion Easily performed	Requires special equipment Requires training Potential submucosal inflation
Cricothyroidotomy	Failure of intubation No laryngeal injury	More rapid than tracheotomy Good airway control	Surgical procedure Can cause both acute and chronic laryngeal injury
Tracheostomy	Laryngeal trauma Oropharyngeal obstruction not controlled by intubation	Avoids entry into larynx	Time consuming Anatomically more complex than cricothyroidotomy Numerous potential complica- tions

SURGICAL TECHNIQUES

1. TRANS TRACHEAL NEEDLE VENTILATION

- where immediate ventilation is required
- can support ventilation for several hours
- Technique
 - 12, 14 or 16 gauge cannula
 - high press ventilation system (50 p.s.I.)
 attached

(Cont. Surgical Techniques)

- Complication
 - * failure to establish an AW
 - * misplaced catheter in soft tissue of the neck (esp. in children)
 - pneumo mediastinum
 - pneumothorax
 - total obstruction of the airway preventsadequate ventilation *

Cont. Surgical Techniques

3. CRICOTHYROIDOTOMY

generally for emergency UAO.
 (failed or contraindication intubation)

 elective for head and neck or cardiovascular procedures

- where access to tracheal rings is limitted

(Cont.Cricothyroidotomy)

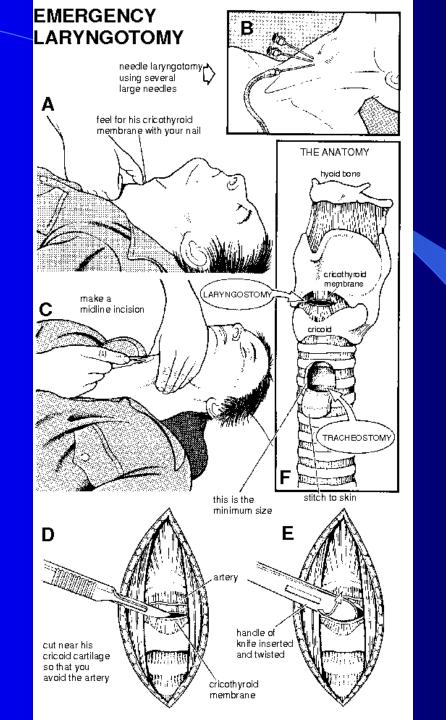
- PROCEDURE

may utilize horizontal or vertical incision

 use small trach. tube or endotracheal tube (Cont. Cricothyroidotomy)

-COMPLICATION

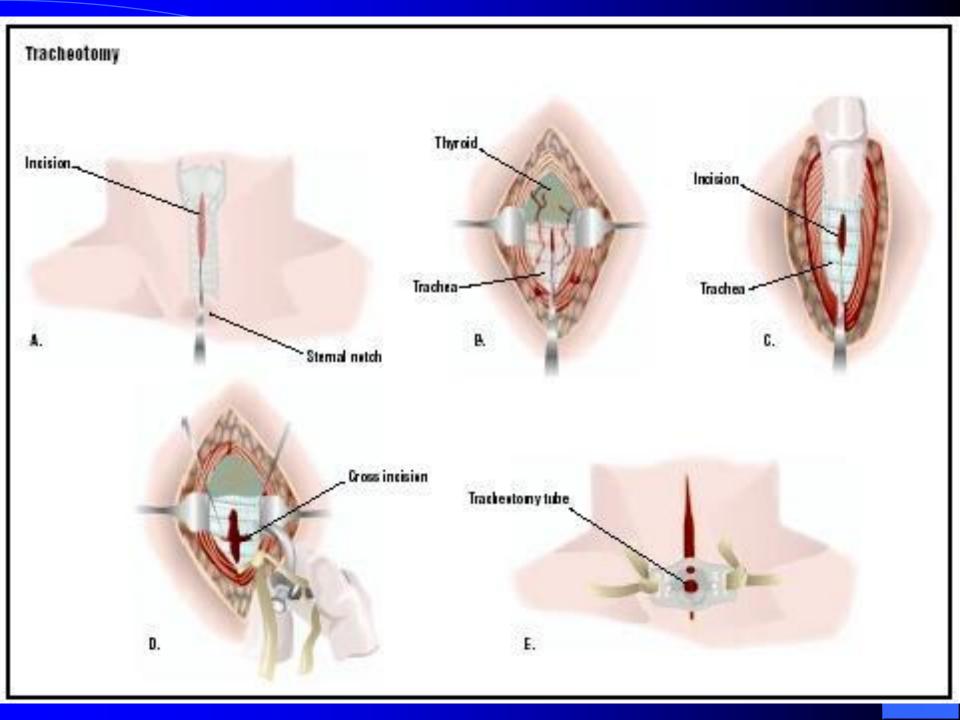
- injury of anterior jugular vein, great vessels
- injury of recurrent laryngeal nerve
- subglottic and laryngeal stenosis(especially in children)



(Cont. Surgical Technique)

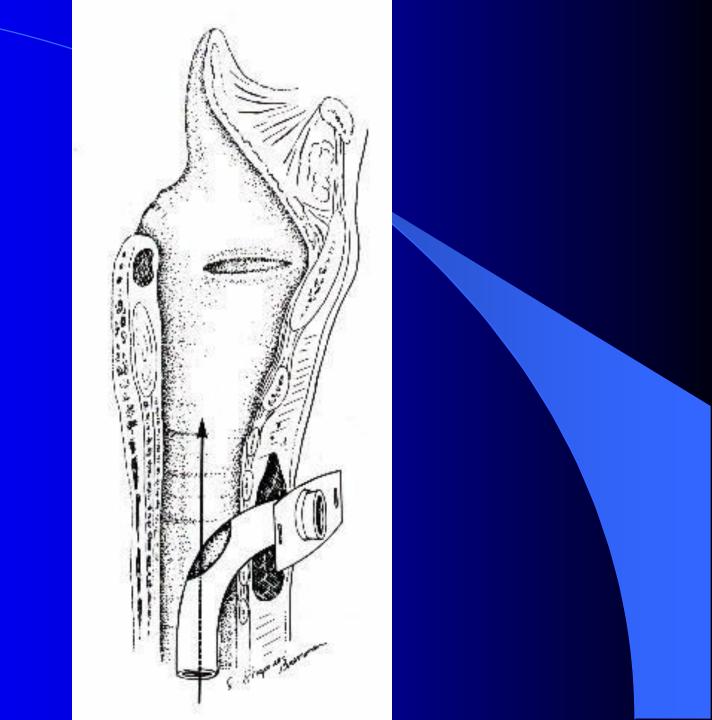
4. TRACHEOSTOMY

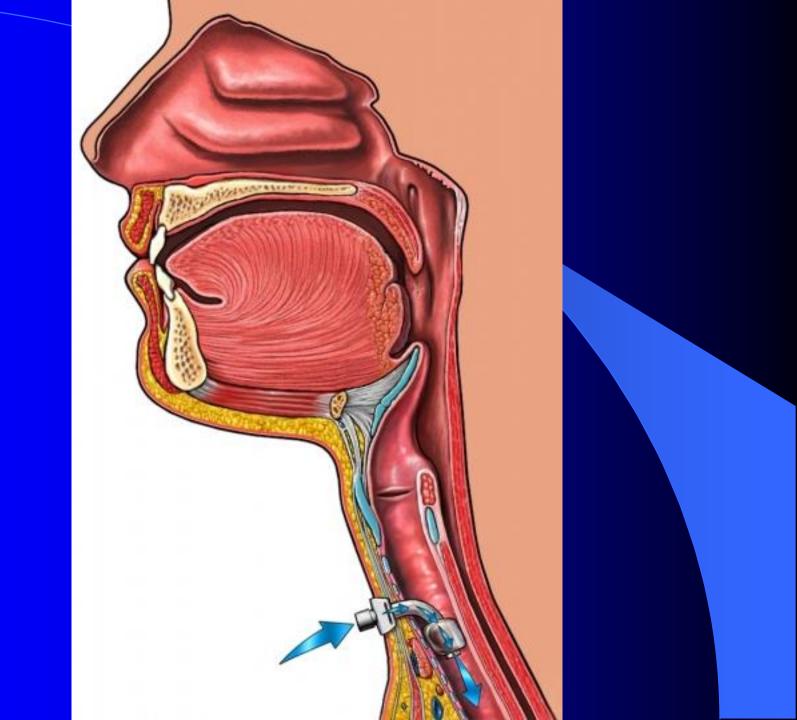
- * for emergency or elective-airway obstruction acute or chronic-airway obstruction
- * in emergency tracheostomy
 - vertical incision is preferred
 - heamostasis after establishing airway obstruction











AIRWAY EMERGENCY

A. TUMOR

- commonly tumors of aerodigestive tract or thyroid
- typically present with gradual airway obstruction
- initial management
 - 0₂, humidification
 steroids and IV antibiotics

(Cont -Airway Emergency- Tumor)

Airway stabilization

- organization between Surgeon and Anasthatist
- avoid blind attempt of intubation
- if available, fiberoptic intubation (experience)
- percutaneous jet ventilation to stabilize patient

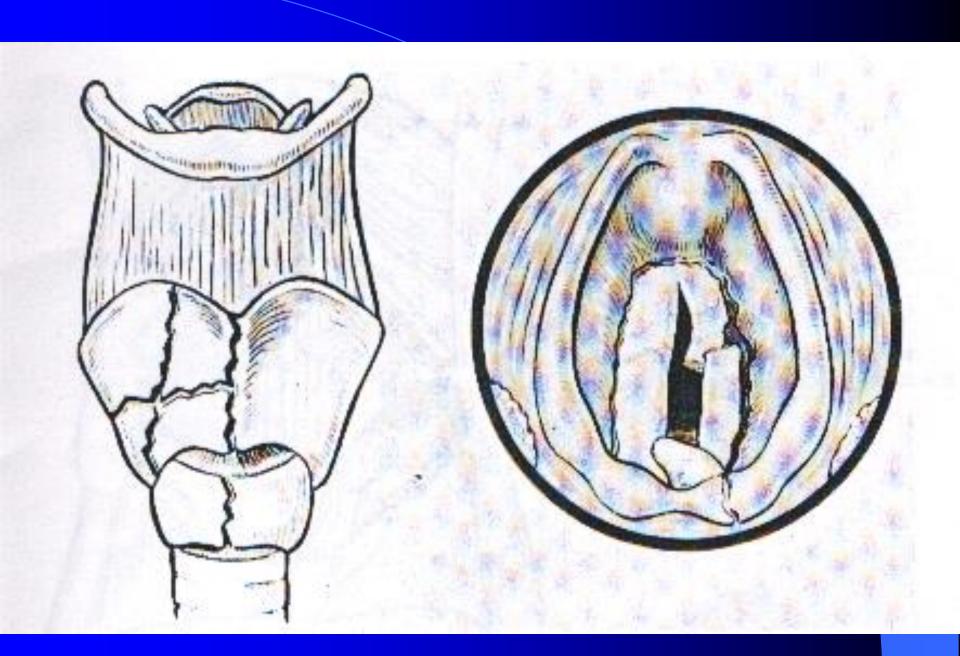
(Cont. Airway stabilization in tumor)

 elective awake tracheostomy under local anaesthesia is the safest method to secure the airway

precipitation of complete obstruction
 necessitates emergent cricothyroidotomy or tracheostomy

B. TRAUMA

- * Presenting SX
 - Hoarseness
 - Pain tenderness
 - Hemoptysis
 - Dysphagia
 - SC emphysema
 - Impaired respiration
 - Haematoma



CLASSIFICATION OF LARYNGEAL TRAUMA & TREATMENT

TYPE I

- minor endolaryngeal haematoma or laceration absence of detectable fracture of laryngeal skeleton
- Management
 - 24 / 48 hours observation in ICU
 - head of bed elevated
 - humidification & systemic steroids

(Cont. Classification of laryngeal trauma & treatment)

TYPE II

- edema, haematoma, mucosal disruption
 no exposed cartilage, no displaced fracture
- Management
- tracheostomy under local anaesthesia
- CT scan to R/O displaced fracture

(Cont.Classification of laryngeal trauma & treatment

TYPE III

- massive edema with large mucosal laceration,
 exposed cartilage, displaced fracture
 V.C. motion impairment
- Management
 - tracheostomy
 - laryngoscopy
 - exploration and repair

(Cont.-Classification of laryngeal trauma and treatment

TYPE IV

- same as III but more severe
- Management
 - explore and repair
 - require endolaryngeal stent

(Cont.Airway Emergency

C. BURN PATIENT

- airway management is controversial

- considering the choice of airway

(Cont.-Burn Patient-Considering the choice of AW)

Oral or nasal endotracheal tube

- May exacerbate existing thermal injury
- Inadvertent extubation is a potential disaster
- When facial grafting is necessary
 tube and ties will limit the access
- Tube obstruction occur more frequent

(Cont.-burn patient-considering the choice of AW)

► Tracheostomy

Reported to have higher mortality rate
 as a result of infectious complication
 (pulmonary sepsis, necrotizing tracheitis, mediastinitis)

- Bleeding, pneumothorax, trancheal stenosis

(Cont.-burn patient-considering the choice of AW)

Tracheostomy cont

- Edema of the neck results in
 - difficult procedure
 - inadvertent removal of the tube

- Cricothryroidotomy, may establish the airway more easily

(Cont. Burn patients)

- Stabilization of airway is indicated for thermal injury of trachea, and extensive burns of the face or oropharynx. Where impending UAWO necessitates intubation

- Intubation for assisted ventilation is required for inhilation injury with:
 - changes in ABG, O₂ sat, and increase CO₁

(Cont. Burn patients)

Once decision of intubation is made:

- ET should be attempted initially
- if necessary, leave X 3-4 wks
- utilize this time for grafting neck burns
- shift to tracheostomy after that if necessary

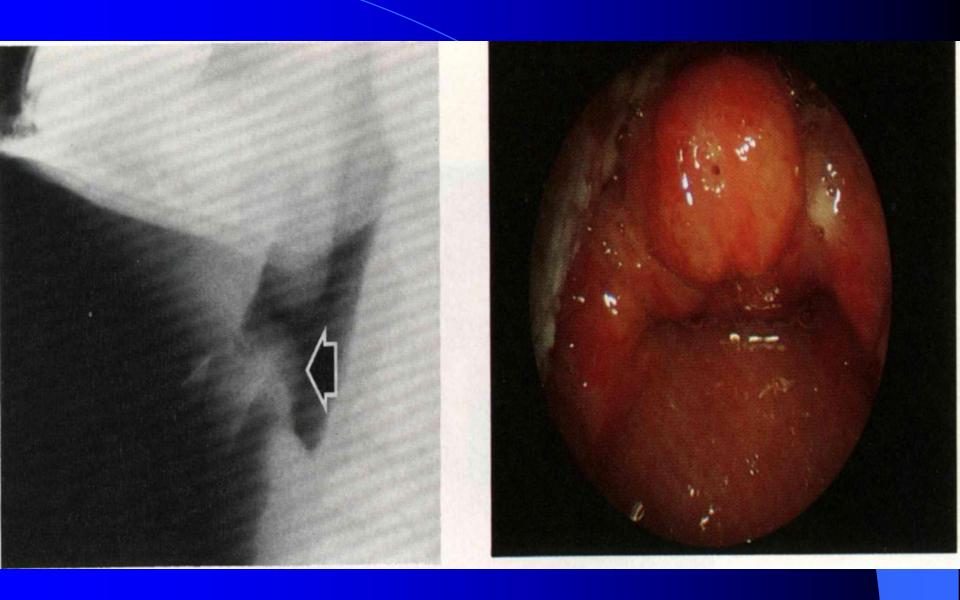
D. SUPRAGLOTTITIS / EPIGLOTTITIS

-PAEDIATRIC

- sudden onset
- rapidly progressive course
- high fever, respiratory distress
- drooling, painful swallowing

ADULT

- dysphagia, severe sore throat
- fever, stridor, voice change



MANAGEMENT

- CHILDREN

- secure airway

ET tube

tracheostomy

- ADULT

- frequently observed in an ICU
- may need intubation

FOREIGN BODY

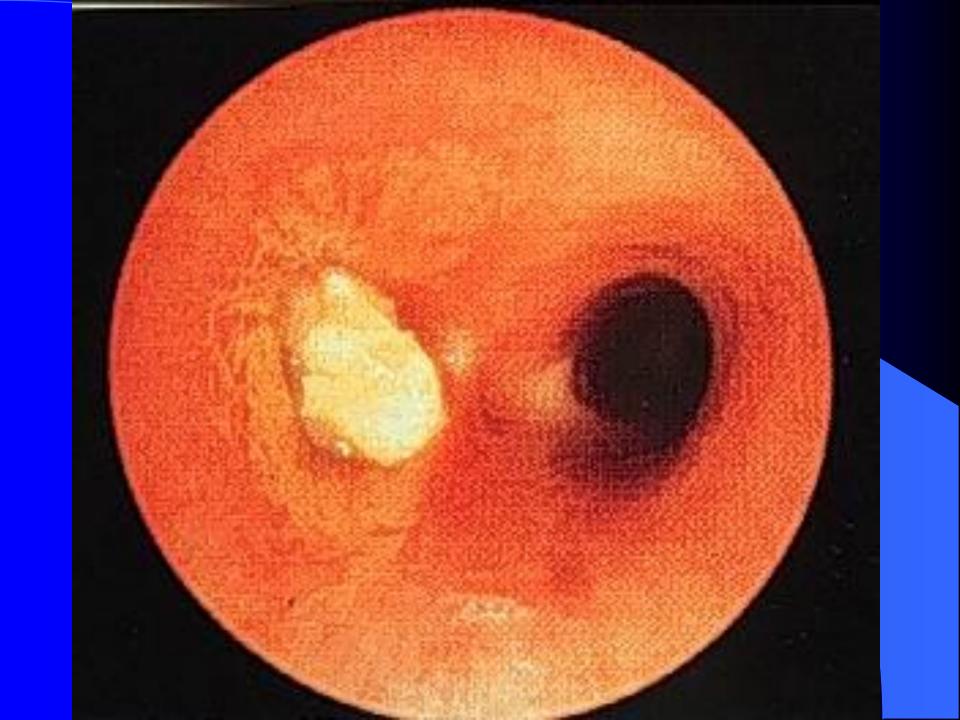
- Death from FB aspiration in USA is about 3000 per year for all ages
- Complete AW obstruction may be recognized in the conscious child as sudden resp. distress
 - inability to speak or cough

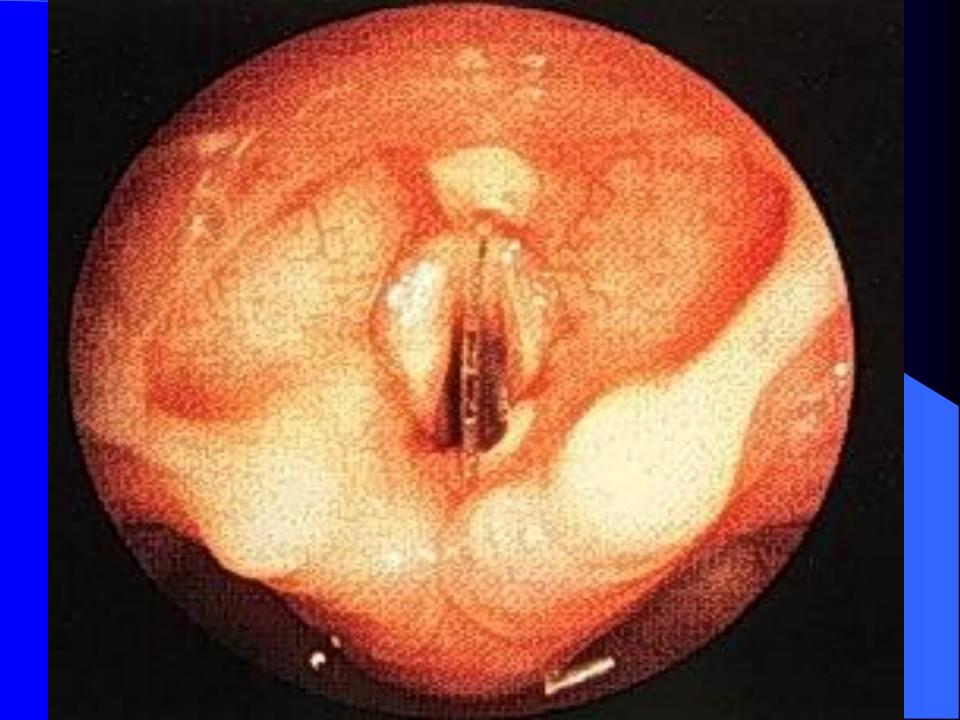
FORIGN BODY (cont.)

- Types of FB
 - * vegetable matter are the most common in the children's AW
 - * metal
 - * plastic

FB (cont)

- Location of FB in the AW
 - * commonly the final destination is one of the main bronchi right more common than left
 - * Larynx -sharp objects
 - * Trachea if there is narrowing in the trachea





FOREIGN BODY (cont.)

PRESENTATION

- Usually coughing, choking, gaging and wheazing
- No symptoms or signs
 Mimik different acute or chronic disease of lungs
 e.g. croups, bronchial asthma.
- . Stage of complications





FOREIGN BODY (cont.)

DIAGNOSIS

- Radiologic extended soft tissue neck}
 - PA, lateral chest

most efficacious

- It can demonstrate FB
 - Emphysema, atelectasis of the lung

FOREIGN BODY (cont.)

- Management:
 - * Endoscopic removal is both diagnostic and theraputic.

LARYNGOMALACIA

- Accounts for 60% of laryngeal problems in newborn
- Due to falcidity or incoordination of supra laryngeal cartilages which are pulled inside the lumen during inspiration leading to UAW obstruction
- Characterized by stridor in the first few weeks

LARYNGOMALACIA (cont.)

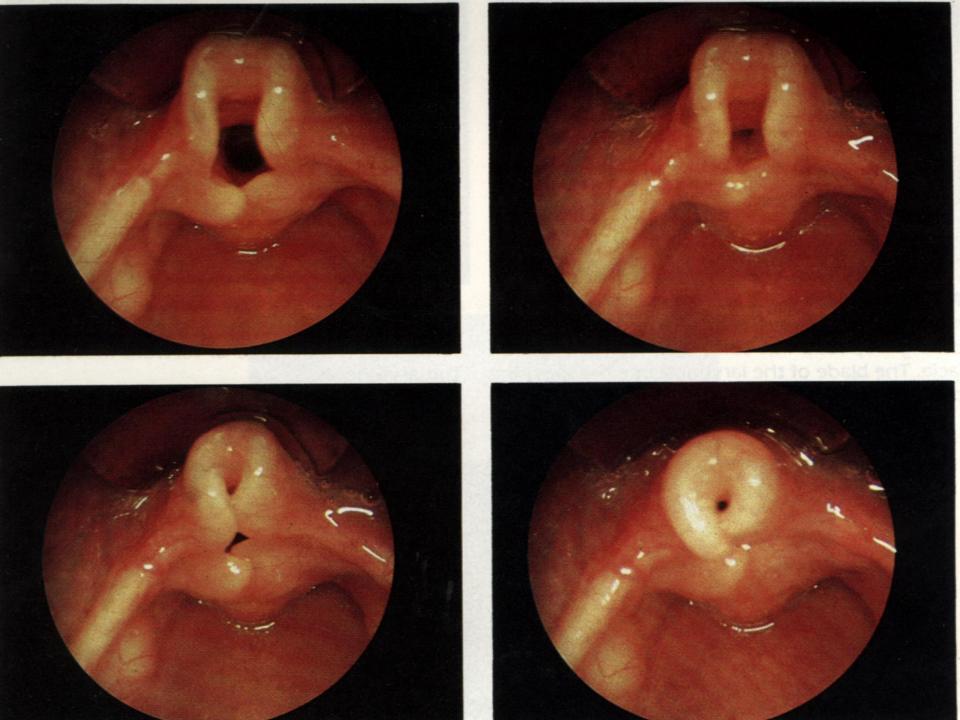
Cause is unknown:

- * edema of the Aryepiglottic folds and loose suspension of the epiglottis.
- * embryologically rapid growth of the third branchial arch causes the epiglottis to curl open itself forming an omega shape.
- * Neurologic immaturity of Brainstem & vagus infolding of the aryteroids in the AW

LARYNGOMALACIA (cont.)

Diagnosis:

- * can only be confirmed by direct observation of movement of supraglottis during respiration
- * fibroeptic evaluation is the most appropriate method of visualization
 - * Radiologic evaluation by (high voltage X-ray PA lateral) may help in excluding the presence of associated AW problem: e.g. SGS.
 - Innominate artery compression.



COMPLICATIONL of laryngomalacia

- Feeding difficulty
- Failure to thrive

LARYNGOMALACIA

Treatment:

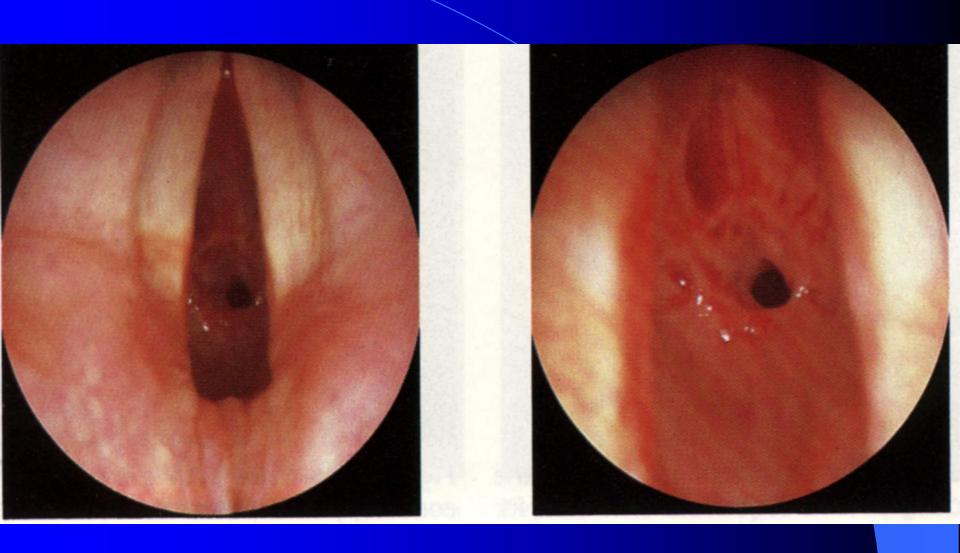
- Reassurance
- Infant can outgrow this problem
- Tracheostomy for severe cases.
- Epiglottoplasty for severe cases

SUBGLOTTIC STENOSIS

It is a narrowing of the subglottis.
 in newborn SG diameter of less than 3.5 mm.

Two types of Subglottic Stenosis:

- * congenital
- * acquired the commonest



Subglottic stenosis presentation

- Mild cases may present as recurrent croup secondary to URTI
- Generally present with symptoms and signs of URT obstruction.

SGS EVALUATION

- Plain film of the neck (high KV)
- MRI for difficult cases
- confirm the diagnosis by endoscopy

SGS MANAGEMENT

- * Endotracheal intubation} emergency situations
 - * Tracheotomy
 - * Cricothyroidotomy
 - * Endoscopic Techniques
 - dilatation
 - laser

SGS MANAGEMENT (cont.)

- Open Surgical Technique
 - * Cricoid split
 - * Laryngotracheoplasty + Rib graft + stent
 - * Recection and primary anastomosis

Choanal atresia

- Uncommon anomaly 1/5000-8000
- Unilateral: present late
- Bilateral: birth emergency
- Mixed bone-membranous CA account for 90%
- Remaining bony CA

Association of CA

CA may be associated with other anomalies in 20-50% of cases
 CHARGE
 VATER

craniofacial anomalies

Examination of CA

- Infants; failure to pass# 6- 8 catheter
 - pyriform aperture stenosis (1 CM)
 - choanal atresia (3.5 cm)
- Fiberoptic nasoscope



Surgical repair of CA

- Many approaches;
 - transpalatal
 - transnasal
 - transantral
 - trans-septal
- Endoscopic repair outcome is variable
- Success rates reported to range between 20-80%

