

Upper Airway Obstruction

This lecture was taken with Dr.AL-Ammar

Special thanks to our old colleague Dr.Hz
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ψ Anatomy of the larynx and trachea:

1. Laryngeal anatomy:

The laryngeal framework is composed of a cartilage.

***three unpaired cartilages:**

- a. cricoid: the only complete ring in the airway. Unable to expand. The narrowest part of airways in children. Liable to injury.
- b. thyroid: the biggest.
- c. epiglottis.

***three paired cartilages:**

- a. arytenoids: pyramid shape. Base on the cricoid's lamina. Attached anterior to the vocal ligament. Posterior to cricoarytenoid muscle. Most important as it's responsible for movement of vocal cord. (most imp. One)
- b. corniculate.
- c. cuneiform.
- d. triticeus.

***laryngeal membranes and ligaments:**

- extrinsic: attach the larynx to other structures.
- intrinsic: between the laryngeal cartilage.
- thyrohyoid membrane.
- cricothyroid membrane: part of conus elasticus.

(cricohyroidectomy: easiest approach to emergency airway obstruction) but not in children

Tracheostomy which done in 2nd, 3rd, or 4th tracheal ring.

- quadrangular membrane.

***muscles:**

- extrinsic.
- intrinsic: all supplied by recurrent laryngeal nerve, except the cricothyroid muscle supplied by external laryngeal nerve.

***infant and pediatric larynx:**

- position is higher at birth. In adult from 2C to 3C.
- epiglottis lying at the nasopharynx makes the neonate an obligate nasal breather, for 4-6 months.
- cartilage and soft tissues are softer.
- soft tissues are less adherent to the underlying cartilage → susceptible to collapse, and less resistance to develop submucosal edema.
- subglottis is the narrowest part of airway in children. It's surrounded by cricoid cartilage.
- in adults the narrowest part of glottis.

2. tracheal anatomy:

- consists of 16-20 incomplete cartilaginous ring, the posterior wall is a membranous part.
- diameter: 19 mm in males.
16 mm in females.
- length is approximately 11 cm.

***pediatric trachea:**

- | | | | |
|-------------|-------------------|---|------------------------|
| - diameter: | -at birth- 6 mm | } | don't worry about this |
| | -6 months- 7.2 mm | | |
| | -1 yr- 7.8 mm | | |
| | -4 yrs- 11 mm | | |

- significance of knowing these diameter: in case of intubation and use of bronchoscopy.

ψ Signs of airway obstruction:

- stridor is harsh high pitched musical sound produced by turbulence of air flow → partial obstruction of AW. It can be inspiratory or expiratory depends on: site and degree of obstruction.

***stridor indicates:**

1. pathologic narrowing of AW.
2. potential respiratory obstruction.
3. even death.

- Usually: → supraglottic is expiratory.

→ Glottis and infraglottic is biphasic.

→ Lower trachea is inspiratory.

- stridor is a very important sign of upper airway obstruction.

[note: table: causes of stridor and respiratory distress originating within upper respiratory tract in children....ect.]

***Other signs of upper airway obstruction:**

- flaring of nasal alae.

- retraction of the neck, intercostals and abdominal muscles.

- dyspnoea.

- tachypnea.

- restlessness.

- cyanosis.(Sub costal emphysema.)

***history:**

- time of onset.

- characteristic of cry.

- Possible trauma.

- Hx of previous intubation.

- relation of airway to problem to feeding, position.

- question about possible aspiration of foreign body.

***if stridor present since birth:**

1. congenital laryngomalacia.

2. subglottic stenosis.

3. vocal cord paralysis.

4. vascular rings.

***if stridor is gradual and progressing:**

- subglottic hemangioma appear between 1-3 months of age.

- papilloma of the larynx appear at 6 months of age.

ψ Differential diagnosis of upper airway obstruction:

1. Congenital:
 - a. laryngeal web.
 - b. laryngomalasia.
 - c. subglottic haemangioma.
2. acquired:
 - a. inflammatory: e.g. epiglottitis, laryngobronchitis.
 - b. infectious: laryngeal diphtheria.
 - c. traumatic: corrosive inhalation.
 - d. neoplastic: papillomatosis, laryngeal cancer.
 - e. foreign body.

ψ Radiologic Evaluation:

- indicated for patients without respiratory distress.

1-plain view soft tissue of the neck.

A.P , lateral . chest (inspiratory + expiratory)

2- mobile pharyngeal tissue may bulge during expiration in normal infants.

3- high kilovoltage techniques (crowp series) AP view assesses subglottis region.

4-fluoroscopy : dynamic airway changes (it's video x-ray , without contrast)

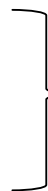
5-barium swallow

. assess swallowing (slow swallowing can lead to aspiration)

. rule out : presence of vascular rings → can affect the airway

MRI

CT scan



ARE GOOD IN EVALUATING MEDIASTINUM.

ψ ENDOSCOPIC EVALUATION:

1-minor examination : is not endoscopic in older children + adults

Can provide information about hypo pharynx and larynx.

2-telescopic examination:

Fiberoptic endoscopy: excellent to assess the movement of vocal cords.

3-rigid bronchoscopy:

-done under GA

-may enable removal of foreign body.

-assess the AW down to the main stem bronchi.

ψ Therapeutic options for upper airway obstruction:

- 1-observation / medical support: →
- a-I.C.U
 - b- SECURE AIRWAY
 - c- OXYGENATION
 - d- STEROIDS
 - e- ANTIBIOTICS

2- heimlick maneuver.

3- nasopharyngeal airway.

4- oral airway.

5- esophageal airway.

6- trans oral intubation.

7- nasal intubation.

8- flexible fibrotic intubation.

9- trans tracheal intubation.

10- cricothyroidotomy

11-

***SURGICAL techniques:**

1-transtacheal needle ventilation (for short time in emergency):

a-when immediate ventilation is required.

b-support ventilation for several hours.

c-technique:-

→ 12,14,16, gauge cannula

→ high pressure ventilation system (50 p.s.i) is attached.

d-complications:

- failure to establish airway.
- misplaced catheter in soft tissue of the neck (esp. in child coz the cartilage is soft normally and it'll compressed not penetrated) can lead to pneumomediastinum or pneumothorax.
- total obstruction of airway preventing adequate ventilation.

2-cricothyroidotomy:

-generally for ER UAWO (if intubation has failed or it's contraindications)

- elective for head & neck & c.v.s procedures.

- also used where access to tracheal rings is limited (e.g kyphosis)

*** procedures :**

a- may use horizontal or vertical incision.

b- use small tracheal tube or endotracheal tube

***complications:**

a-injury to int. Jugular vein (great vessels)

b- injury to recurrent laryngeal nerve

c- subglottic or laryngeal stenosis (esp. in child)

3-tracheostomy:

-for ER or elective upper airway obstruction , also for acute or chronic upper airway obstruction.

-in ER: → vertical incision is preferred

→ maintain homeostasis after establishing airway.

ψ Other diagnostic test :

1. flow volume loop.
2. ABG (arterial blood gases).

@late indicator of upper airway obstruction , that's why it shouldn't be used routinely or in ER to assess the degree of obstruction .

ψ AIRWAY EMERGENCY :

1)TUMOR :-

-commonly, tumours of aerodigestive tract goal to stabilization.

-organization should happen between surgeon and anaesthetist.

-avoidance of blind attempt of intubation .

- *stabilization should be either by: -*

a- elective awake tracheotomy under local Anaesthesia (preferred)(is the safest).

b- fiberoptic intubation .

NOT cricothyrotomy because it may precipitate complete obstruction.

2)TRAUMA :-

-Elective awake tracheotomy under local anaesthesia .

3)BURN PATIENT :-

-Endotracheal intubation is preferred .

****we secure airway for burn patients when there is :***

a-burn to face , oropharynx & upper airway in general .

b-intubation injury .

ψ epiglottitis (supraglottitis):-

-infection of the epiglottis .

-commonly affect children between 2-6 years.

***causes :-**

-almost always bacterial in origin (H.influenza type B account for 90% of cases, with H.influenza type B vaccine the incidence of it is less (commonest cause in children).

-other organisms: pneumococci , sterpt. Beta hemolytic .

[if you suspect patient with epiglottitis → you have to examine him under controlled situation in OR not in ER because , you may stimulate the epiglottis then the patient will collapse (die)]

***clinical feature:-**

-sudden onset and rapidly progressive course.

-high fever , respiratory distress , drooling and painful swallowing.

-may quickly progress to respiratory obstruction .

-examination of pharynx can reveal cherry red or edematous epiglottis.

-should avoid manipulation (e.g. intubations) of the pharynx → may lead to complete obstruction of the larynx .

-x-ray (lateral view) of the neck can show the thumb sign (edema of the epiglottis) if necessary.

***management:-**

-in children airway should be opened either 1)endotracheal tube.

2)tracheostomy tube.

-in adult 1)observe in ICU.

2)may need intubation.

@antibiotic should be started:-

- 1- initially: cefuroxime , cefotaxime or ceftriaxone.
- 2- Then change antibiotics according to sensitivity.

ψ foreign body:-

-death from foreign body aspiration in USA is about 3000 per year for all ages.

-complete airway obstruction may be recognized in the conscious child as sudden respiratory distress , inability to speak or breathe or cough .

***types of foreign body:-**

1-vegetable matter are the most common in the children's airway.

2-plastic.

3-metal.

***location of foreign body in the airway:-**

-commonly the final destination is one of the main bronchi (right one is more common than left because it is more straight and wider).

-larynx : for sharp objects .

- trachea : if there is narrowing in the trachea .

***presentation :-**

(acutely)1-usually coughing , choking , gagging and wheezing .

(followed by) 2- no symptoms or signs (mimic different acute or chronic disease of the lung e.g. croup, bronchial asthma .

(finally)3-stage of complication (erosion, infection, abscess)

***diagnosis:-**

- radiological : extended soft tissue in the neck , PA, lateral chest (most efficacious)

- it can demonstrate foreign body : emphysema (commonest & earliest), then atelectasis of the lung.

[do it in inspiration & expiration you will see the findings in expiration]

@ the scope the best Dx & Rx method .

***management:-**

- endoscopic removal is both diagnostic & therapeutic .

!! note: table for first aid for the choking child , recommendations of the American academy of pediatric.(long table we couldn't write it)

ψ laryngomalagia:-

- accounts for 60% of laryngeal problems in newborn.

-characterizes by stridor in the first few weeks.

-due to flaccidity or incoordination of supralaryngeal cartilage which are pulled inside the lumen during inspiration leading to upper airway obstruction.

-cause is unknown

-Edema of the aryepiglottic folds and loose suspension of the epiglottis.

-Embryologically rapid growth of the 3rd branchial arch causes the epiglottis to curl open & self forming an omega shape .

-Neurological immaturity of brainstem and vague- infolding of the arytenoids in the airway .

***Diagnosis :-**

-Can only be confirmed by direct observation of movement of subglottis during respiration .

-Fibrotic 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.: → subglottis stenosis (SGS).

→innominate artery compression .

***Complication of laryngomalacia :-**

1-failure to thrive (it's one of the high index indicate chronicity).

2-feeding difficulty .

***treatment :- “ > 95% of cases are mild “**

✓ reassurance.

✓ Infant can outgrow this problem .

✓ Tracheostomy for sever cases only in case of children (because of high mortality rate) *not good for them because high mortality rate that happen because they have small trachea , small tube can easily obstruct it so we use Tracheostomy .*

✓ Epiglottoplasty for sever cases (definitive procedure for laryngomalacia , done under general anesthesia).

ψ SUBGLOTTIC STENOSIS :

-It's a narrowing of the subglottis. In newborn the subglottis diameter of less than 3.5 mm.

-Two type of subglottic stenosis :- 1- congenital .

2-acquired . (*the most common*)

[note : the most common cause of acquired stenosis is Endotracheal intubation .]

***subglottis stenosis presentation :-**

- mild cases may present as recurrent croup (laryngotracheobronchitis) secondary of upper respiratory tract infection .

- Generally present with symptoms and sign of upper respiratory tract obstruction .

- * subglottis stenosis evaluation :-**
- 1-plain film of the neck (high KV) .
 - 2-confirm the diagnosis by endoscopy .
 - 3-MRI for difficult cases .

*** subglottis stenosis management :-**

1. Endotracheal intubation .
2. Tracheostomy (preferred) (the best) .
3. cricothyroidctomy above the stenosis, we can pass a rigid tube to bypass the stenosis, sometime we cannot bypass → Tracheostomy .

} emergency situation

4. endoscopic techniques :-

- dilatation .
- laser .

} for immediate scarring .

5. open surgical technique :-

- a-cricoid split for small children + mild stenosis .
- b-resection and primary anastomosis .
- c-laryngotracheoplast + Rib graft + (stent for 6w) .

ψ Choanal Artesia :

- uncommon anomaly .
- unilateral :- present late .
- bilateral :- birth emergency .
- mixed bone-membranous CA account 90% .
- remaining bony CA .

***associate of Choanal Artesia :-**

- associated with other anomaly in 20%- 50% :-

1. **CHARGE** (the most common)
2. **VATER**
3. **craniofacial anomaly** .

****dear friends we are so so so sorry we couldn't write symptoms of this anomaly****

„done“