

MANAGEMENT OF THE ANOMALOUS PEDIATRIC AIRWAY

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PEDIATRIC AIRWAY

- **Small**
- **Soft**
- **Pliable**
- **Vulnerable to minimal manipulation:**

Minor obstructive changes may lead to serious and potentially fatal conditions (reduction of neonatal glottis by 1mm decreases airway by 40%).

- **STRIDOR is the most common presentation of children with an Anomalous Airway.**

STRIDOR:

- **Harsh "musical sound"**
- **Baby's cry for help**
- **A symptom not a diagnosis**
- **Turbulent airflow through a partially obstructed airway**
- **Three types**

EVALUATION OF STRIDOR:

- **History**
- **Physical Examination**
- **Radiographic Studies**
- **Endoscopy**

ETIOLOGY OF STRIDOR:

- **Multiple:** More than 40 different etiologies in children

- **Classification:**

A. **Anatomic Region**

- Nasal and/or pharyngeal
- Laryngeal
- Tracheal and/or bronchial

B. **Underlying Cause:**

- Congenital
- Inflammatory
- Traumatic
- Neoplastic

C. **Type of Obstruction:**

- Extrinsic
- Intrinsic

MANAGEMENT OF STRIDOR

DEPENDS:

- **Severity**
- **Underlying Etiology**
- **Age**
- **Associated problems**

CLASSIFICATION:

Immediate: Epiglottitis, Foreign Body...

- **Intubation**
- **Tracheotomy**
- **Endoscopy**

Medical:

- **Observation**
- **Antibiotics**
- **Steroids**

Surgical:

- **Laser**
- **Cricoid Split**
- **Tracheotomy**
- **Laryngotracheoplasty with rib graft**

LARYNGEAL ANOMALIES

LARYNGEAL ANATOMY:

- Larynx arises from the 4th and 6th bronchial arches
- Larynx appears at 4th week of development
- Tracheoesophageal septum develops by the 6th fetal week
- Location:
 - C-4 Birth
 - C-7 Adult
- Size: Neonatal = 1/3 adult
- In infant cricoid/subglottis is the narrowest portion of the larynx
- Function:
 - Protection
 - Phonation
 - Breathing

LARYNGOMALACIA

- **Most common congenital laryngeal anomaly**
- **Stridor: worse supine, better prone**
- **Usually disappears by the age of 2 years**
- **Flabby supraglottic structures**

Diagnosis:

- **Fluoroscopy**
- **Direct laryngoscopy**

Treatment:

- **Mild-moderate:**
 - **No specific therapy:**
(Observation, early RX of URI)
 - **Resolution by 2 years**
- **Severe:**
 - **Epiglottomy**
 - **Tracheotomy (rare)**

VOCAL CORD PARALYSIS:

- Second most common laryngeal anomaly
- Frequently present at birth
- Not truly a developmental anomaly

Bilateral

- Cry may be normal
- Stridor increase with activity
- High pitched crowing
- Inspiratory stridor
- More common than unilateral

Unilateral

- Weak cry
- Mild inspiratory stridor
- Left: Thoracic and/or C/V anomalies
- Right: CNS abnormalities

Treatment:

Bilateral:

- Tracheotomy
- Nerve-muscle pedicle
- Lateralization

Unilateral:

- Observation

CONGENITAL SUBGLOTTIC STENOSIS:

- **Inspiratory and expiratory stridor**
- **Third most common laryngeal anomaly**
- **Soft tissue thickening of subglottic area**
- **Possible cyanosis**
- **Recurrent croup**
- **Possible association with other congenital lesions and syndromes**

DIAGNOSIS:

- **Radiographically: A-P and lateral neck**
- **Endoscopy:**
 - **Confirmed if a 3mm infant bronchoscope cannot be readily passed in a full-term newborn**

TREATMENT:

- **Observation**
- **Cricoid split**
- **Tracheotomy**

ACQUIRED SUBGLOTTIC STENOSIS:

- Acquired with intubation
- Tracheotomy generally required in 90% of cases
- Decreasing incidence associated with improved airway management

MANAGEMENT:

- Observation
- Anterior/posterior cricoid split
- Auricular cartilage graft
- Costal cartilage graft
- Tracheotomy

SUBGLOTTIC HEMANGIOMA:

- Symptomatic after birth
- Most common neoplasm of the larynx in children
- May be associated with other hemangiomas

DIAGNOSIS:

- X-ray: Subglottic narrowing/mass
- Endoscopy

ENDOSCOPIC FINDINGS:

- **Smooth compressible mass below posterior commissure with lateral extension**
- **Covered by mucosa**
- **Rarely extends to trachea or true vocal cords**

TREATMENT:

- **Laser**
- **Steroids**
- **Tracheotomy**

LARYNGEAL WEBS:

- **Glottic web: most life threatening anomaly**
- **98% anteriorly positioned**
- **Hoarseness**
- **Aphonia**
- **Stridor**
- **Airway obstruction**

DIAGNOSIS:

- **Flexible and/or rigid endoscopy**

TREATMENT:

- **Lysis:**
 - **Laser**
 - **Knife**
- **Tracheotomy**
- **Open procedure**

LARYNGEAL CYSTS:

- **Supraglottic region**
- **First few months of life**
- **Low cuboidal epithelium**
- **Arvepiglottic folds and epiglottic: most common location**
- **Presentation: Stridor and severe upper airway obstruction**

DIAGNOSIS:

- **Radiographically**
- **Endoscopy**

TREATMENT:

- **Endoscopy: Removal of cyst roof**
- **External approach with tracheotomy: in case of recurrence**
- **Aspiration is not sufficient**

LARYNGOTRACHEOESOPHAGEAL CLEFTS

- **Incomplete closure of cricoid**
- **Incomplete closure of tracheoesophageal septum**

TYPES:

- **Type I: Cricoid and interarytenoid muscle**
- **Type II: Six proximal tracheal rings**
- **Type III: Extension to carina**

SYMPTOMS:

- **Stridor and respiratory distress**
- **Feeding problems**
- **Repeated aspiration/pneumonia**

ASSOCIATED FINDINGS:

- **Polyhydramnios**
- **Prematurity**
- **Congenital defects**

DIAGNOSIS:

- **History**
- **Radiographic studies**
- **Endoscopy**
- **Awareness of the anomaly**

CHOANAL ATRESIA

- A developmental failure of the posterior nasal cavity to communicate with the nasopharynx
- Etiology: Persistence of nasobuccal membrane
- Occurs in 1/5000 - 1/7000 live births
- Unilateral more than bilateral
- Bony vs. membranous (9:1)
- Associated abnormalities (20% - 50%) CHARGE
- Right: Left (2:1)
- Females:Males (2:1)

DIAGNOSIS:

- Clinical presentation/history
- Physical Examination
- Radiographic studies

MANAGEMENT:

- Medical: Temporary/Early
- Surgical: Definitive

MEDICAL MANAGMENT:

- Secure airway
- Adequate feeding and care

SURGICAL MANAGMENT:

- Challenging
- No one type of repair can be used in every case with complete success and no complications

TYPES:

- Transnasal (dilation, drillout, curettage, laser, endoscopic)
- Transpalatal
- Trans septal
- Transantral
- Sublabial

SURGICAL REPAIR

TRANSPALATAL

- Better visualization
- Preservation of nasal lining
- Shorter stenting period
- Higher success rate
- Older children
- Developmental effects

TRANSNASAL

- Limited visualization
- Lower success rate
- Younger children
- Quick
- Minimal blood loss

CONCLUSION

STRIDOR:

- Anomalous Airway
- "Baby's cry for help"
- Warrants immediate attention

MANAGEMENT OF ANOMALOUS AIRWAY

- Plan of action
- Orderly and Focused
 - Where
 - What
 - Severity
 - Alternate airway

LARYNGEAL MANIPULATION = EDEMA = OBSTRUCTION

TEAMWORK + ACCURATE DX + ADEQUATE RX

EQUALS

SUCCESS