

# Pediatric Surgery

Additional Surgical Techniques



# Lesson Objectives:

1. Identify key physiological and anatomical features in pediatric surgery
2. Discuss pediatric pathology with respect to surgery
3. Discuss key elements of case planning for pediatric surgery
4. Describe psychosocial care of the pediatric patient
5. Discuss risk reduction techniques in pediatric surgery
6. List and describe common pediatric surgical procedures

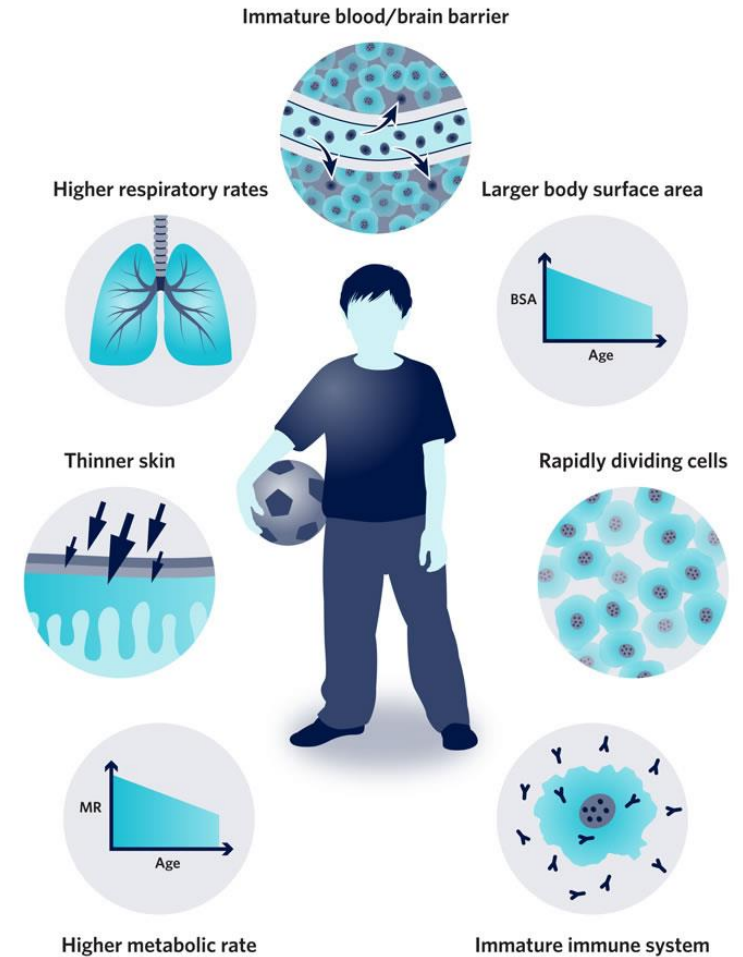
# Pediatric Surgery

- Encompasses congenital anomaly treatment, disease procedures, and trauma surgery.
- Care in pediatrics spans from neonate to late adolescence, often continuing into adulthood
- Family involvement is integral, requiring equal consideration in communication and psychosocial care.



# Physiological and Anatomical Considerations

- **Physiological Differences in Pediatric Patients:**
  - Infants and children lack developed mechanisms for maintaining temperature balance.
  - Factors like skin surface area, weight, and circulation contribute to rapid heat loss.
  - Wide temperature variation compared to adults.



# Hypothermia in Pediatric Patients

- **Hypothermia in Pediatric Patients:**
  - Risks include conduction, radiation, and anesthesia-induced effects.
  - Physiological response includes decreased metabolism, heart rate, and respiratory rate.
  - Infants lack shivering mechanism, leading to rapid hypoxemia and hypoglycemia.
- **Hyperthermia Concerns:**
  - Environmental and physiological factors contribute.
  - Misuse of warming devices and excessive covering can induce hyperthermia.

**Table 2. Differential Diagnosis for Hypothermia**

Category/ Organ System	Conditions that can result in hypothermia
Trauma	Burns, drowning, traumatic brain injury, spinal cord injury, intracranial hemorrhage, nonaccidental trauma
Toxicologic	Antidepressants, antipsychotics, barbiturates, clonidine, alcohol, opioids
Infectious disease	Sepsis, meningitis/encephalitis, brain abscess
Neurologic	Brain tumor, dysautonomia, stroke
Endocrine/metabolic	Hypoglycemia, hypothyroidism, organic acidemia, aminoacidemia, adrenal insufficiency, anemia
Cardiovascular	Cardiac arrest, cardiac arrhythmia, congenital heart disease

# Physiological and Anatomical Considerations

- **Hemostasis in Pediatric Surgery:**

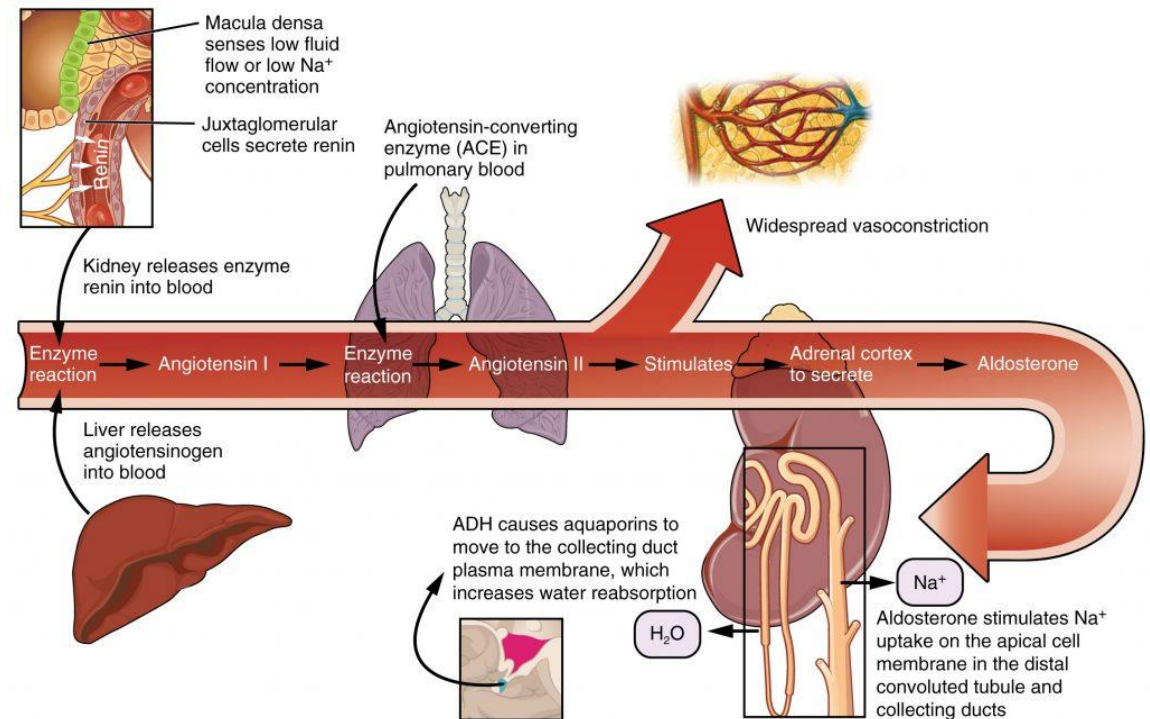
- Critical due to limited blood reserve in children.
- Monitoring blood loss and preparing for emergencies is essential.

- **Anatomical Considerations in Airway Management:**

- Differences in tongue size, airway structure, and thoracic wall rigidity.
- Infants are obligate nose breathers and have a shorter trachea compared to adults.

# Fluid Balance

- Infants can become dehydrated rapidly, so maintaining the correct amount and types of fluids is important
- Even small amounts of fluid to an adult can have large changes in children





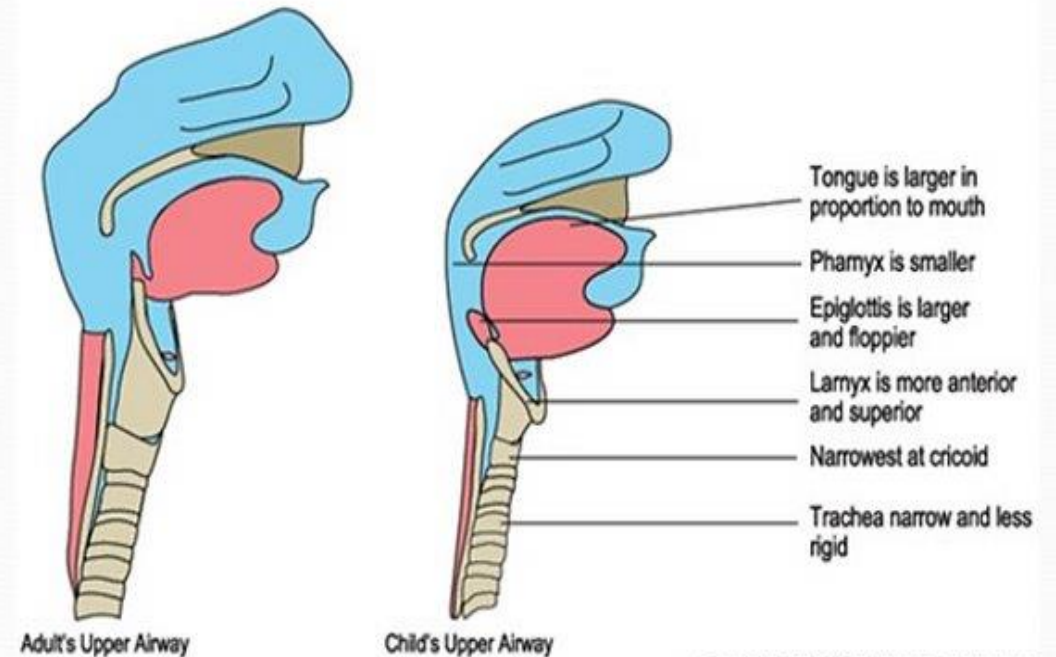
# Pediatric Airway Features

Difference exists between adult and pediatric respiratory systems in terms of structure of airways

## Features of pediatric airway:

- Large tongue
- Obligate nose breathers
- Shorter trachea
- More delicate airway
- Weak thoracic wall
- Lower residual lung capacity

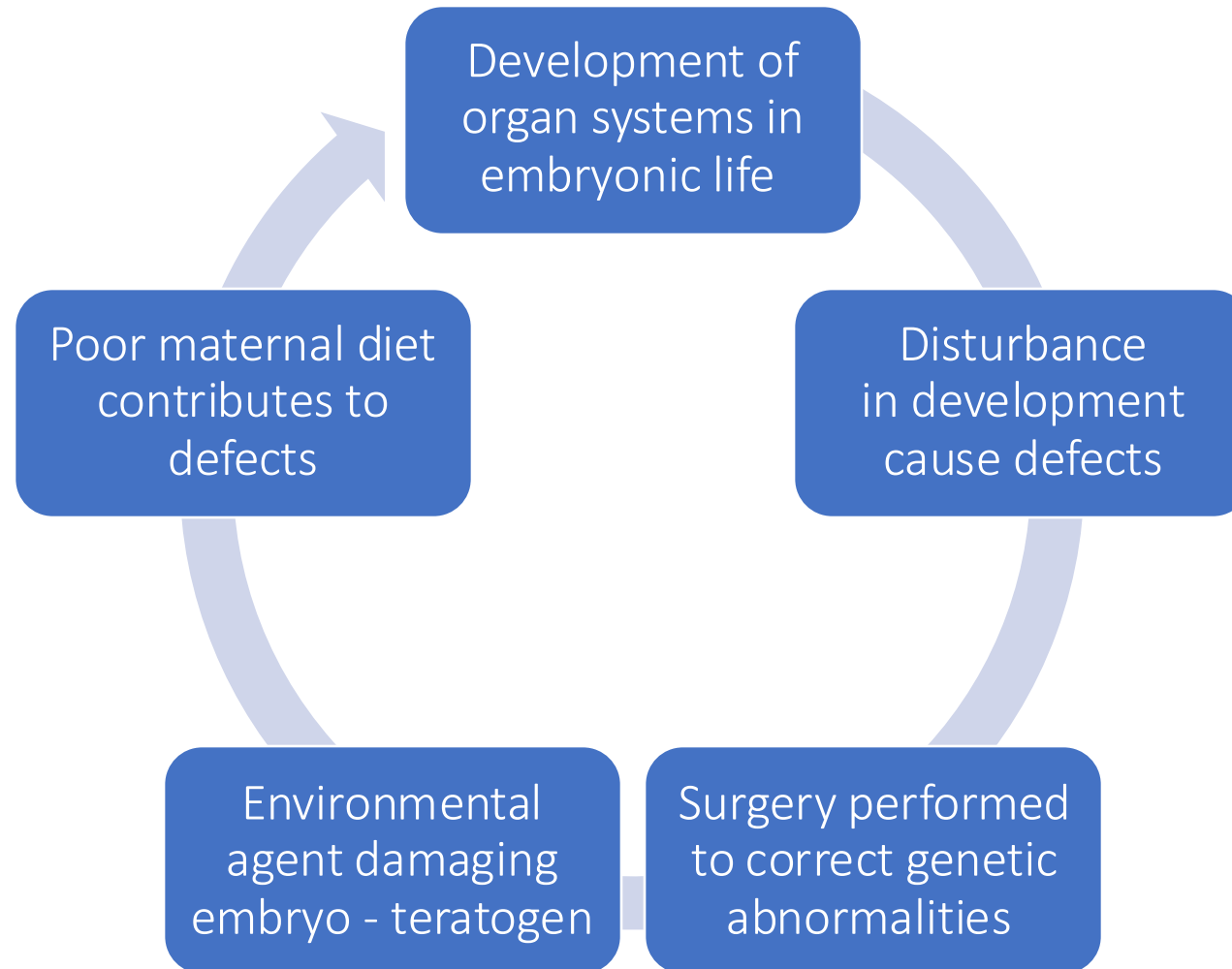
## Airway anatomy differences



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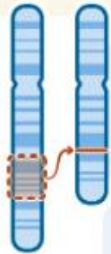
# Pathology



# Genetic Abnormality

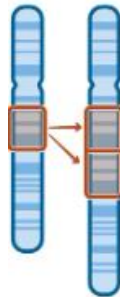
## BACKGROUND

- \* CHANGES in CHROMOSOME STRUCTURE or NUMBER
- \* NUMERICAL (ANEUPLOIDIES)
  - ↳ CHANGE to NUMBER of CHROMOSOMES
- \* STRUCTURAL
  - ↳ CHANGE to SPECIFIC PART of CHROMOSOME



### DELETION

- \* PORTION of CHROMOSOME DELETED
- ~ CRI-DU-CHAT SYNDROME



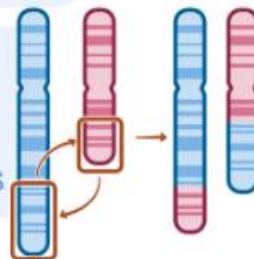
### INVERSION

- \* GENETIC MATERIAL FLIPPED in OPPOSITE DIRECTION
- ~ DO NOT OFTEN RESULT in DISEASE



### DUPLICATION

- \* PART of CHROMOSOME DUPLICATED
- ↳ EXTRA GENETIC MATERIAL
- ~ CHARCOT-MARIE-TOOTH DISEASE TYPE I



### TRANSLOCATION

- \* PIECE of ONE CHROMOSOME has BROKEN OFF & ATTACHED to ANOTHER CHROMOSOME
- ~ POTENTIAL CAUSE OF TRISOMIES

## COMMON DISORDERS

- \* TRISOMY 21
  - ↳ DOWN SYNDROME
- \* TRISOMY 18
  - ↳ EDWARDS SYNDROME
- \* TRISOMY 13
  - ↳ PATAU SYNDROME
- \* KLINEFELTER SYNDROME
  - ↳ GENOTYPE XXY
- \* TURNER SYNDROME
  - ↳ GENOTYPE X



## REDUCE RISK

- \* EATING HEALTHY
- \* ABSTAINING from SMOKING & ALCOHOL
- \* TAKING PRENATAL VITAMINS
- \* GENETIC COUNSELING



**Watch the video "Preventing Genetic Defects  
Video?" to gain insights into prevention  
mechanisms**

# Preventing Genetic Defects Video

- [Click Here](#) to watch the video

# Preventing Genetic Defects Video

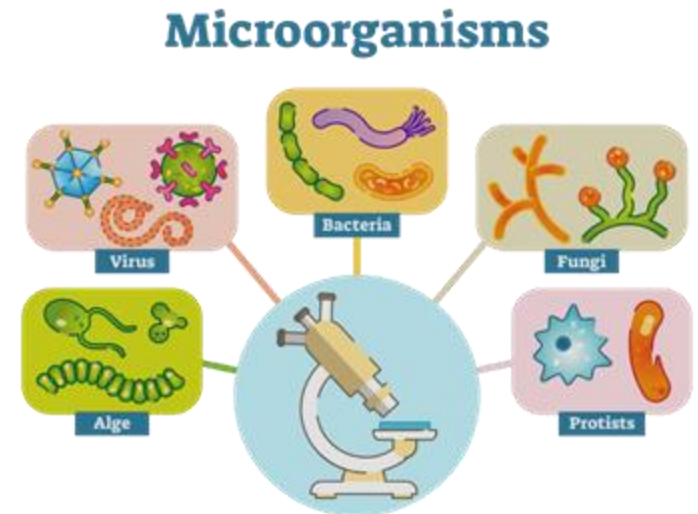
## Summary of the Video:

- Breakthrough genetic technologies offer hope for babies with genetic abnormalities.
- Scientists estimate 1 in 200 babies suffer from single gene disorders, impacting families profoundly.
- Genetic testing advancements allow pre-implantation screening for thousands of diseases, offering potential life-saving options for parents.

# Microorganisms Responsible for Genetic Abnormalities

The following infectious microorganisms cause significant congenital abnormalities (listed in parentheses):

- *Toxoplasma gondii* (cerebral calcifications, microcephaly, heart defects)
- Rubella virus—the causative agent of measles (cataract, glaucoma, deafness, heart defects, retinal defects)
- Cytomegalovirus (hydrocephalus, deafness)
- Herpes virus (microcephaly, microphthalmia, retinal defect)
- Varicella-zoster virus—the causative agent of chickenpox (muscle atrophy, mental retardation)
- *Treponema pallidum*—the causative agent of syphilis (hydrocephalus, deafness, bone defects)





# Psychological Care

- Psychosocial care involves child, caretakers, and perioperative staff
- Goal: develop strategies for perioperative experience
- Consider child's developmental stage, emotional state, and social support
- Preoperative visit/counseling involving patient, caretakers, child life specialist
- Goal: reduce anxiety, allow questions before surgery
- Greatest fears: fear of unknown, separation, mutilation

**Watch the "Communicating with Pediatric Patients"  
Video**

# Communicating with Pediatric Patients Video



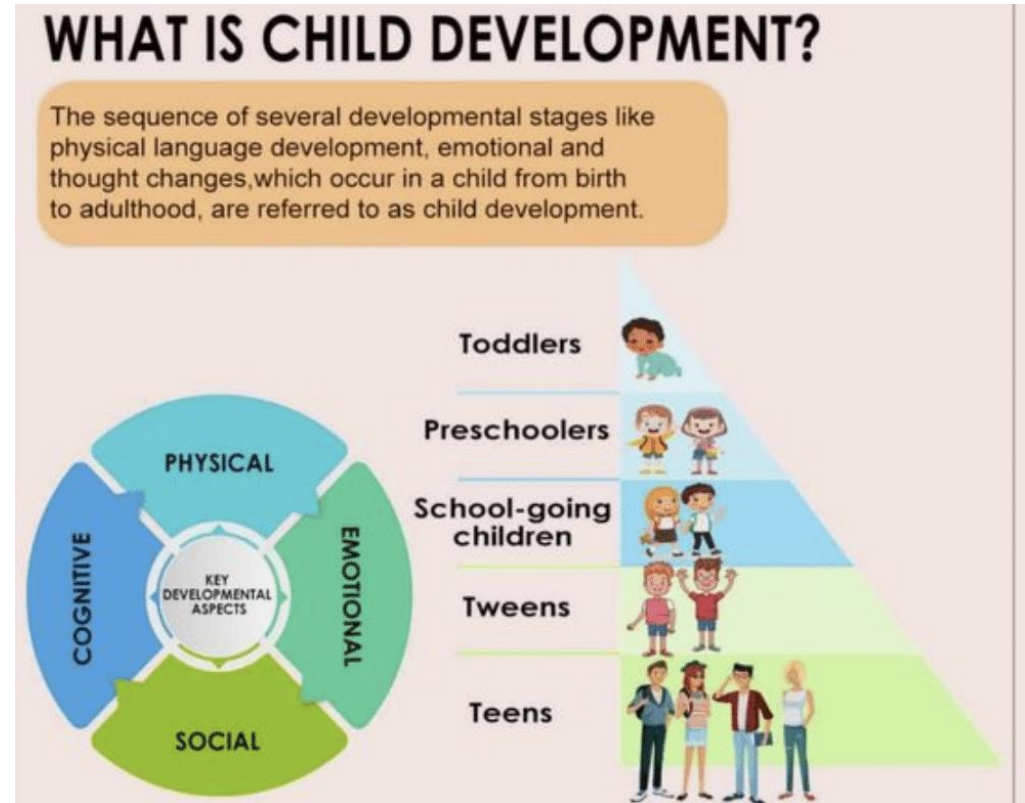
# Communicating with Pediatric Patients Video

- **Summary of Video:**

- Pediatric patients will often be nervous about going to the doctor
- Find ways to relate to patients and help them feel at ease
- Communicate on their level
  - Each developmental stage will communicate different
- Explains things in advance and as they happen to help them feel at ease
- Use simple terms or analogies they can relate to
- Engage parents: children will trust their parents and they can help them warm up

# Developmental Stages of the Child

- Infants: Birth to 12 months
- Toddlers: 13 months to 2 years
- Preschoolers: 3 to 5 years
- Middle Childhood: 6 to 11 years
- Early Adolescence: 12 to 14 years
- Late Adolescence : 15 to 17 years



**Watch the "Growth and Development of Infants, Toddlers, Preschoolers and School Aged Children" video to gain insights into the development stages of the child and the physio-social changes associated with each stage**



## Growth and Development Video



# Case Planning

Anesthesia

Safety of the pediatric patient

Transport of the Pediatric Patient

Positioning the Patient

Electrosurgery

Instruments

Sponges

Sutures

# Anesthesia

(Slide 1 of 2)

- General anesthesia often necessary for pediatric patients due to airway control needs and inability to cooperate.
- Variances in anesthesia approach due to physiological and anatomical differences from adults.



# Anesthesia

(Slide 2 of 2)

## Preoperative Preparation:

- Psychological preparation involves family engagement and addressing child's fears.
- Assessment includes health history, medications, allergies, and lab tests.
- Parental presence during induction increasingly accepted and may reduce preoperative medication need.

## Anesthesia Induction and Maintenance:

- Various induction methods based on age, IV access, and preferences.
- Inhalation anesthesia commonly used for induction and maintenance.
- Continuous monitoring and IV access secured during maintenance phase.

## Emergence and Recovery:

- Recovery influenced by administered drugs, with opioids and sedatives potentially prolonging emergence.
- Post-anesthesia complications include nausea, vomiting, respiratory depression, and delirium in pediatric patients.

# Pediatric Safety

- Maintain a safe environment
- Collaborate with other perioperative team members
- Address specific developmental stages



# Transport and Positioning of Pediatric Patient

## Transport

- Neonates and Infants: Transport Method
- Young Children: Transport Procedures
- Patients with Monitoring Devices: Special Considerations
- Challenges with Inpatient Toddlers: Safety Measures

## Positioning

- Principles of Safe Patient Positioning
- Importance of Appropriate Padding
- Considerations for Surgical Positioning
- Avoiding Skin Injury: Tape Alternatives



# Pediatric Surgical Procedures

- Repair of a cleft lip
- Repair of a cleft palate
- Choanal atresia
- Correction of esophageal atresia and tracheoesophageal fistula
- Pyloromyotomy
- Repair of an omphalocele
- Repair of bladder exstrophy/epispadias
- Orchiopexy for an undescended testicle
- Repair of pectus excavatum
- Radical nephrectomy: Wilms tumor
- Repair of a myelomeningocele
- Correction of syndactyly

***Watch the "Pediatric Surgery" Video***

# Pediatric Surgery Video



# Pediatric Surgery Video

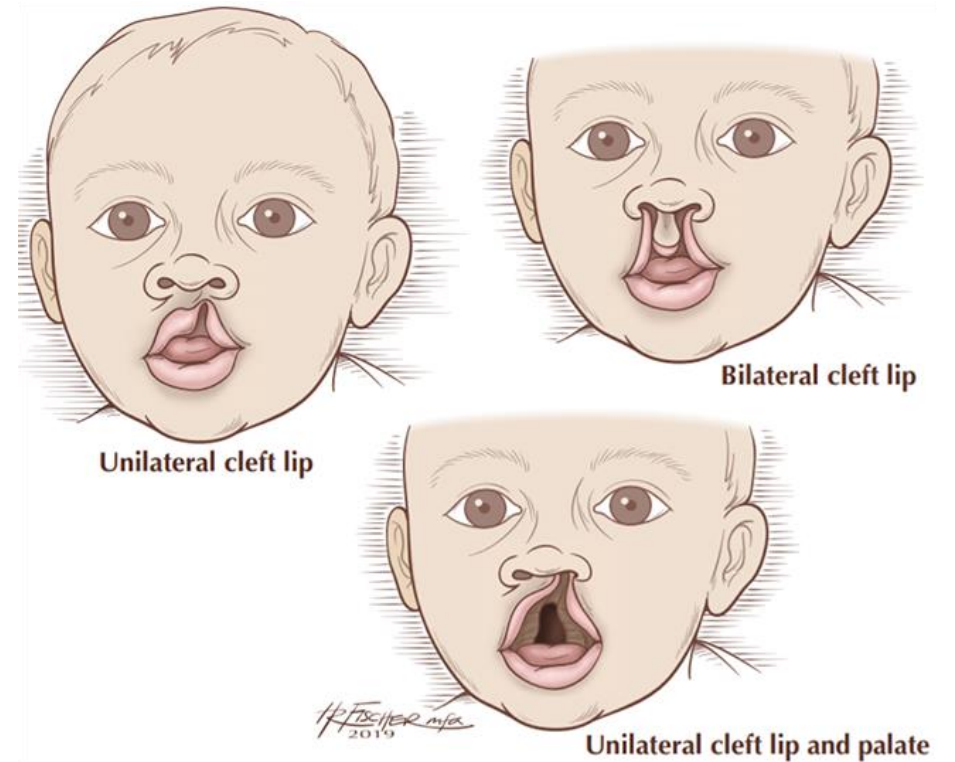
- Summary of Video:
  - Focus on individualism of pediatric patients and their needs
  - Each patient is unique
    - Developmental stage
    - Vital signs
    - Weight and Medication Dosing
    - Family Members

# Repair of a Cleft Lip

## Closure of cleft defect in the lip

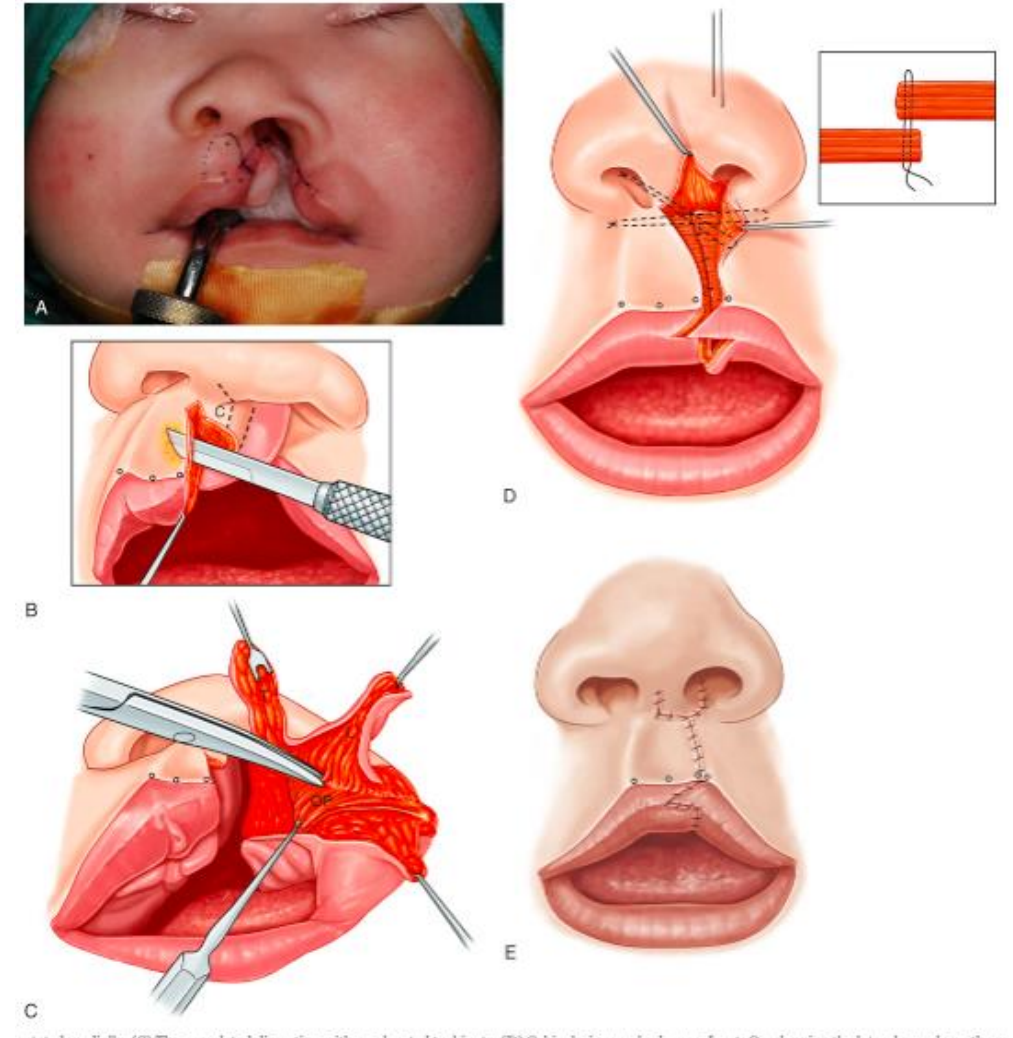
### Pathology

- Formation of philtrum and upper lip during embryonic development.
- Interruption leads to cleft lip.
- May be complete/incomplete, unilateral/bilateral.
- Often associated with clefts of the soft palate.
- Repaired in stages, initial at 10-12 weeks.
- Preoperative planning involves precise measurements.



# Repair of a Cleft Lip Procedure

- **Preparation:** Patient prepped, draped; marked for incisions.
- **Incisions:** Made along vermilion border, mucosa separated from orbicularis oris muscle.
- **Z-plasty Flaps:** Created through skin, muscle, and mucosa; hemostasis maintained.
- **Closure:** Mucosa closed with interrupted absorbable sutures; muscle closed with subcuticular absorbable sutures; vermilion border and skin closed.





**Watch the "Cleft Treatment" video to  
gain insights into the procedure!**

# Cleft Treatment Video

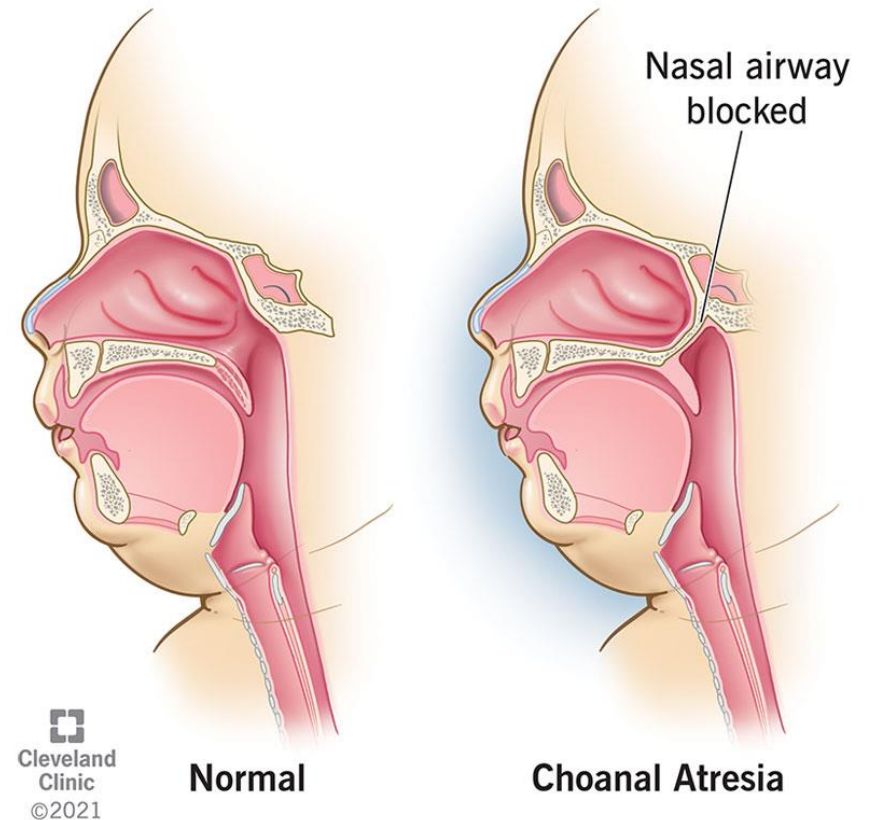


# Cleft Treatment Video

- **Summary of the video**
- Various types: Unilateral, Bilateral, Isolated Cleft Palate.
- Procedures at different ages: Lip repair at 4-6 months, Palate repair at 9 months.
- Treatment at Boston Children's Hospital: High-volume center, Multidisciplinary team available.

# Choanal Atresia

- Congenital blockage of the choana, the passage between the nasal cavity and the throat
- Surgery involves puncturing and removal of obstructing tissue
- Placement of a stent to maintain airway patency during healing
- Pediatric sinuscope for visualization and use of tissue-shaver and rongeur for obstruction removal



# Correction of Esophageal Atresia and Tracheoesophageal Fistula (EA/TEF)

- Surgical procedure to address defects in the esophagus and abnormal connections between the esophagus and trachea
- Goal: Restore continuity of the esophagus and prevent aspiration of food and saliva into the lungs
- **Pathology of EA/TEF**
  - EA: Absence or closure of a normal esophageal opening
  - TEF: Abnormal connection between esophagus and trachea
  - Associated birth defects: Imperforate anus, cardiac defects, limb deformities, neural tube defects

**Watch "Esophageal Atresia & Tracheoesophageal  
Fistula" video to gain insights into what this defect is in  
detail**

# Esophageal Atresia and Tracheoesophageal Fistula (EA/TEF) Video



# Esophageal Atresia and Tracheoesophageal Fistula (EA/TEF) Video

- **Summary of the Video**
- Introduction
- What is esophageal atresia and a tracheoesophageal fistula?
- Risk factors, signs/symptoms, diagnosis, treatment, and nursing care of esophageal atresia and a tracheoesophageal fistula.



# EA/TEF Surgical Options and Post-Op Care

- **Surgical Options**

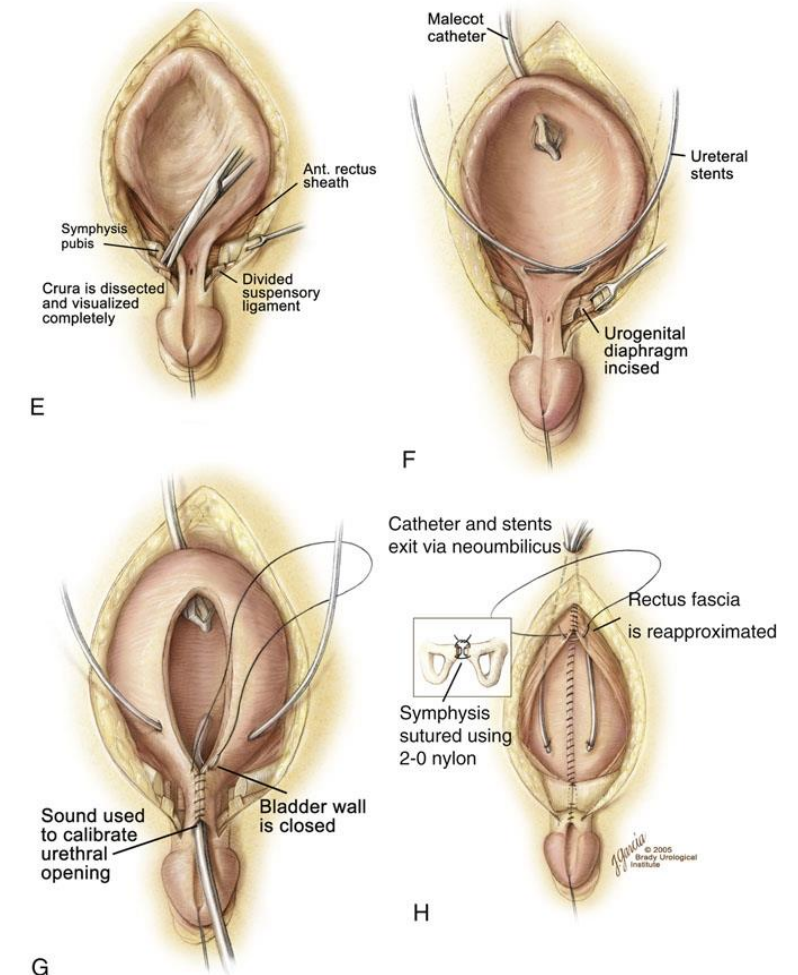
- Dependent on type and severity of defect
- Staged or delayed repair may be necessary
- Primary closure preferred when possible

- **Postoperative Care**

- Airway management
- Pharyngeal suctioning
- Monitoring for complications like infection and anastomotic leakage

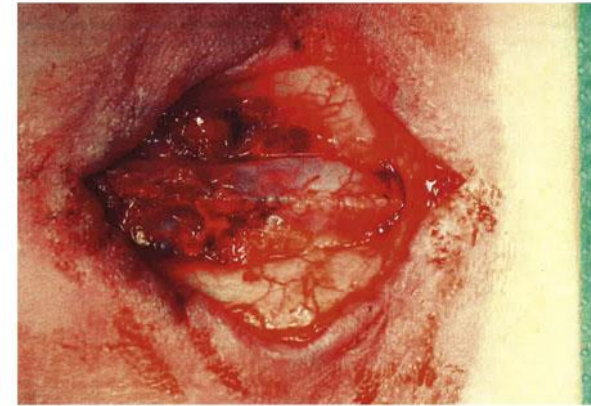
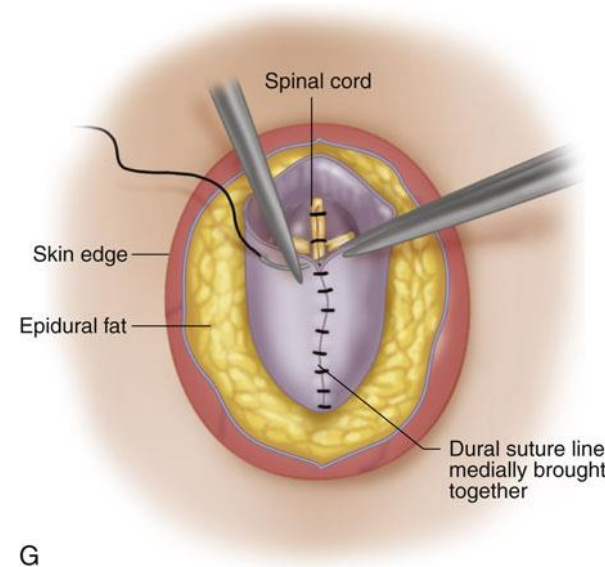
# Repair of Bladder Extrophy/Epispadias

- **Surgical goal:** Restore normal functions of lower urinary tract.
- **Pathology:** Complex congenital anomalies involving lower genitourinary tract and skeletal system.
- **Anomalies include:**
  - Bladder exstrophy (exposed bladder wall outside the body)
  - Epispadias (urethra exposed or terminates on dorsum of penis)
- **Technical Points:**
  - Reconstruction usually in three stages:
    - Closure of bladder and abdomen (24-48 hours of life).
    - Repair of epispadias (2-3 years old).
    - Achieving urinary continence (4-5 years old).



# Repair of a Myelomeningocele:

- Surgical goal: Close dural and cutaneous defect, preserving neural function.
- Immediate repair crucial for CSF leak prevention; otherwise, performed within 48 hours to prevent infection.
- Latex-free protocols often employed due to frequent surgeries and latex sensitization risk.
- Vulnerability to hypothermia requires warm solutions and delicate instrument use.
- Often accompanied by hydrocephalus, necessitating potential ventricular shunt placement.



# Read Chapter 33 from the E-book

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[Click Here](#) to access Chapter 33!

# Thank you!

Get ready for your quiz and rest of the activities now. Best of luck!



# Congratulations!

Lesson 33 is complete.