

Congenital Diaphragmatic Hernia

Anatomy

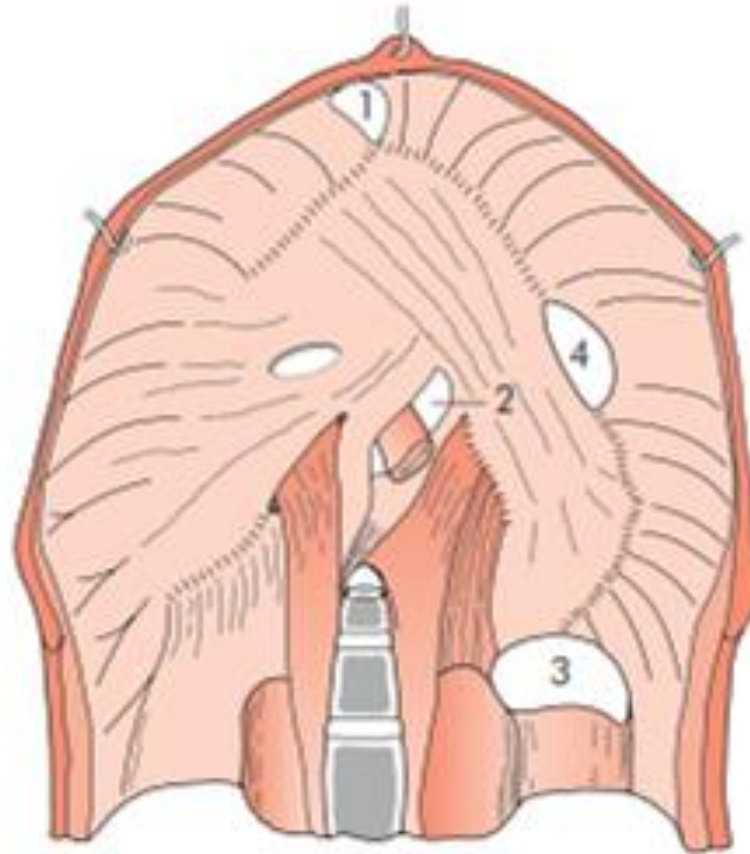
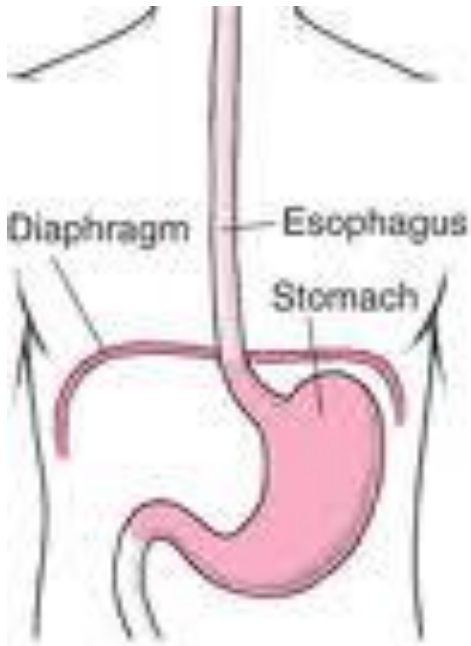
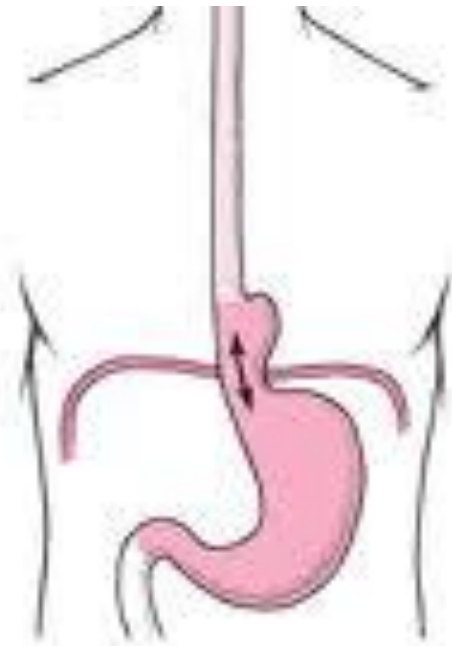


Figure 52.30 Diagram of sites of hernias. The usual sites of congenital diaphragmatic hernia: 1, foramen of Morgagni; 2, oesophageal hiatus; 3, foramen of Bochdalek (pleuroperitoneal hernia); and 4, dome

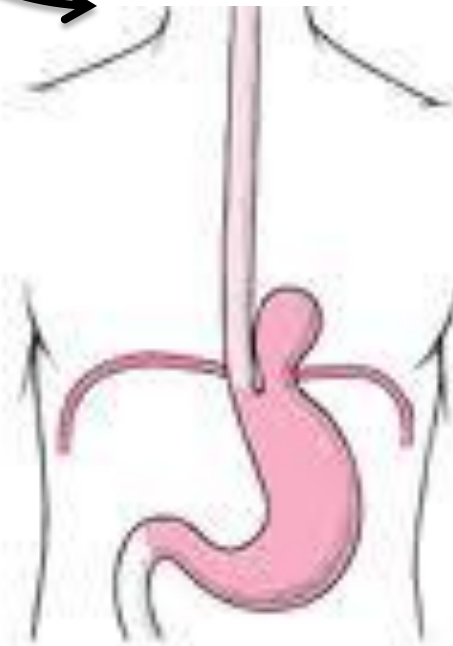
Hiatus Hernia



Normal Esophagus and Stomach

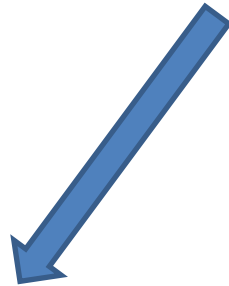


Sliding Hiatus Hernia



Paraesophageal Hiatus Hernia

Congenital diaphragmatic hernia

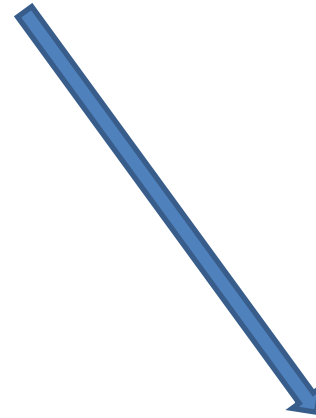


Bochdalek Hernia

postrolateral

85-90%

80-90% Lt. side



Morgagni Hernia

retrosternal

2-6%

Pathophysiology:

- Pulmonary hypoplasia.
- Pulmonary hypertension
- Persistent fetal circulation



C/F:

➤ Prenatal:

US



24-26 wk 50-60%
polyhydraminous
Echogenic chest mass
bowel loops in the chest
Mediastinal shift
Intrathoracic liver

➤ Postnatal :

Immediately after birth

low Apgar score

tachypnea

grunting

cyanosis

Cry  worsening due to more swallowed air.

10% discovered outside the neonatal period.

Later in life *incidentally*

recurrent chest infection

Late presentation has good prognosis due to minimal or absent pulmonary hypoplasia and hypertension.

O/E:

- Scaphoid abdomen.
- Increased antero-posterior diameter of the chest.
- Decreased air entry on the affected side with decreased tidal volume.
- Bowel sound may be heard in the chest.
- Shifting of cardiac impulse to the right.



Diagnosis:

-- CXR

- Air & fluid filled loops of bowel in the chest.
- Paucity of intestinal gas in the abdomen.
- Mediastinal shift (heart shifted to the right).
- Tip of NG tube seen in the chest.



-- Barium study



Treatment:

➤ Resuscitation

NG tube

Bag mask ventilation is contraindicated

Endotracheal intubation + oxygen.

Watch O₂ saturation (pulse oxymetry)

Urinary catheter

Arterial & venous line

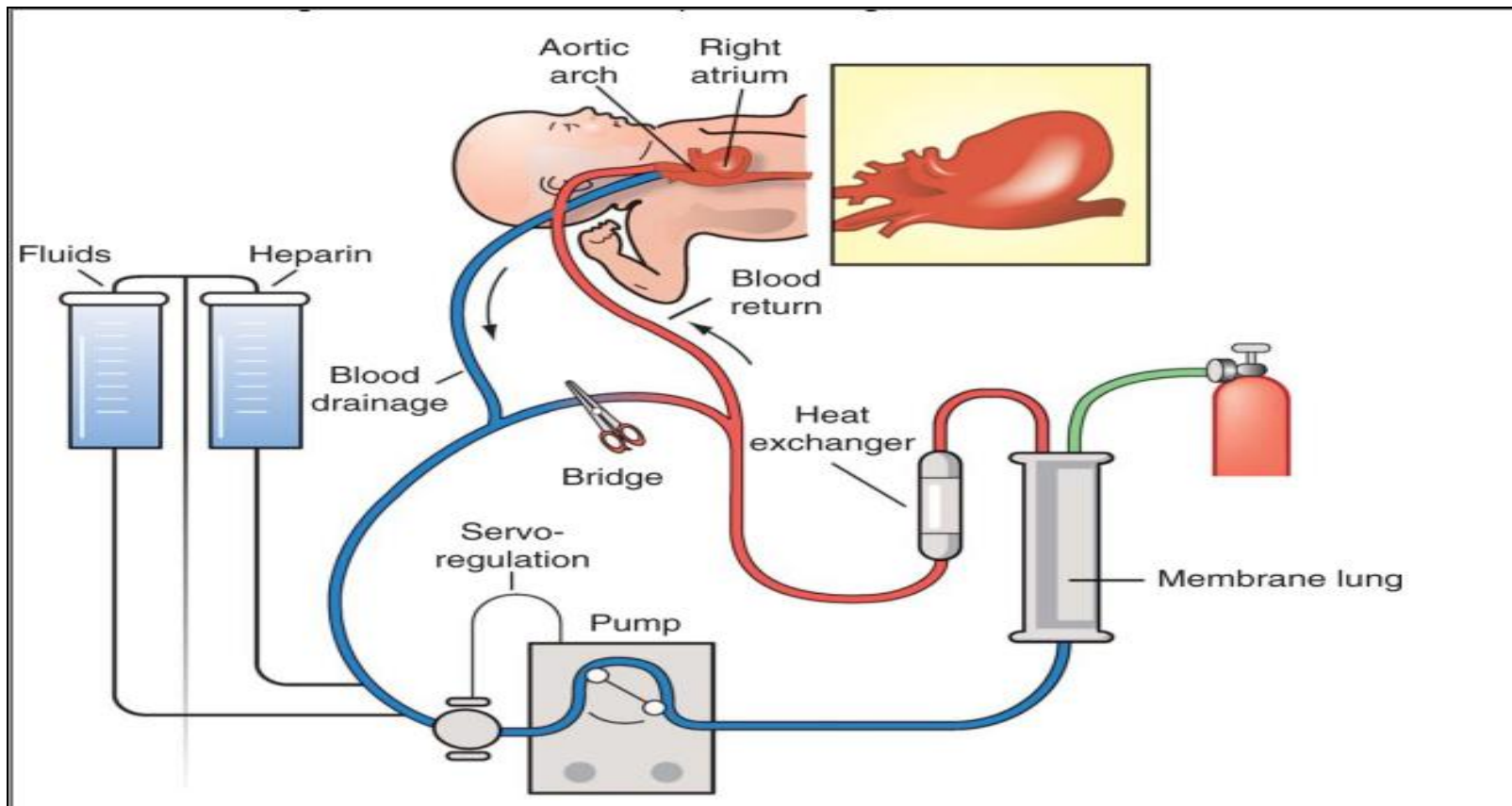
Fluid therapy

Echo study

Refer to pediatric surgical center

Therapeutic options for stabilization prior to surgery

- mechanical ventilation: high frequency oscillatory ventilation(HFOV).
- inhaled nitric oxide (NO).
- surfactant therapy.
- ECMO (extra corporeal membrane oxygenation).



➤ SURGERY:

delayed repair

transabdominal X transthoracic

subcostal incision

Reduction of hernial contents

1^o closure

Prosthetic mesh

ventral hernia for big defects

Fetal intervention:

1. Tracheal occlusion.

2. Prenatal steroid.

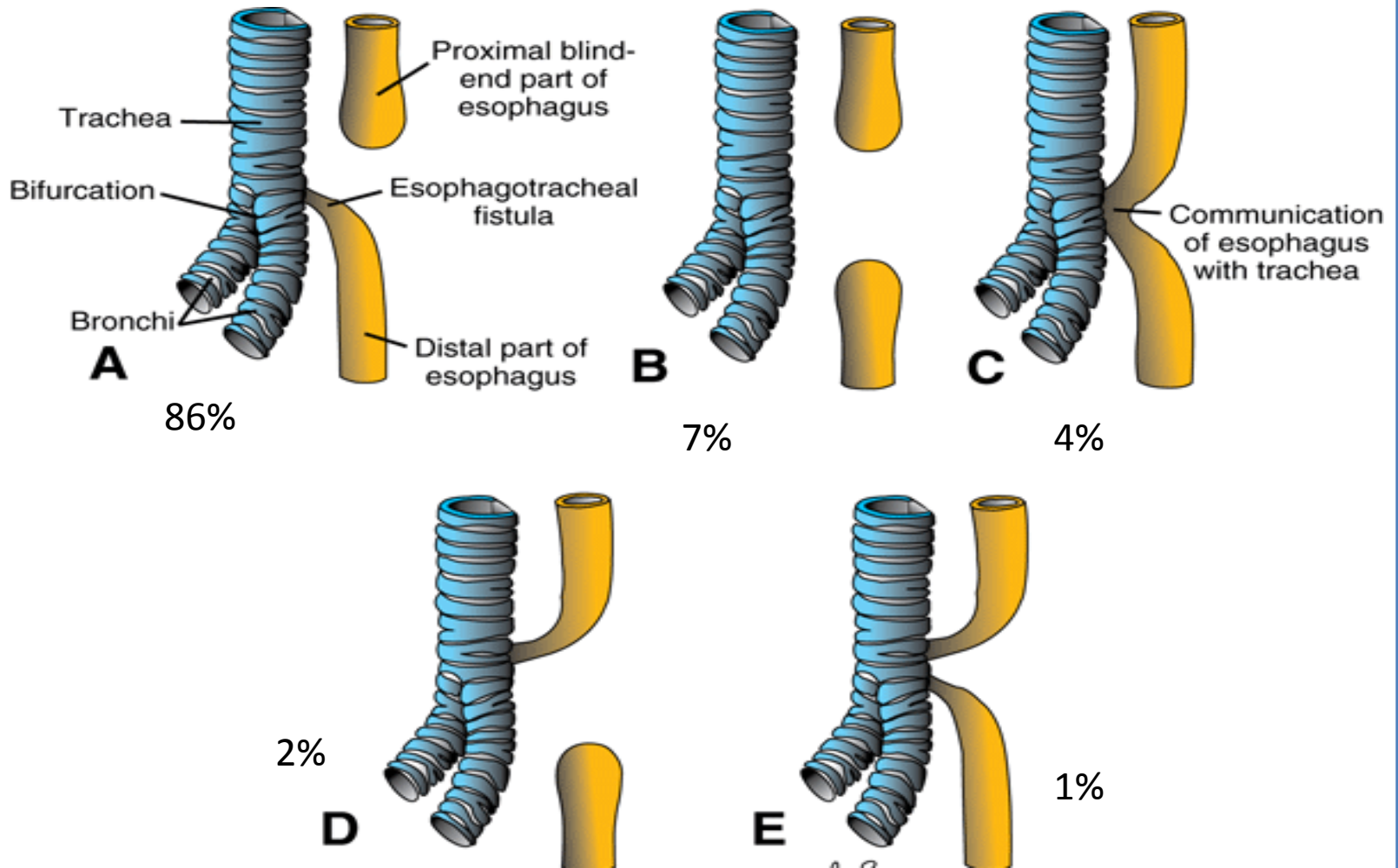
3. Intrauterine surgical repair.

Esophageal Atresia

+ / -

Tracheoesophageal Fistula

Classification



Pathophysiology :-

- Blind end esophageal pouch = pooled secretion.
- Fistulous connection with trachea = aspiration.
- Disordered peristalsis in lower esophagus = GERD.
- Tracheomalacia = respiratory obstruction.

Presentation :-

Polyhydraminous.

Frothy secretion.

Respiratory distress.

- Resistance to NG tube insertion.
- Plain X-ray.







Initial management :-

Suctioning.

Positioning.

Vascular line.

Endotracheal intubation.

Incubator , warm , humidity.

Check for associated anomalies.

Referral to pediatric surgical center.

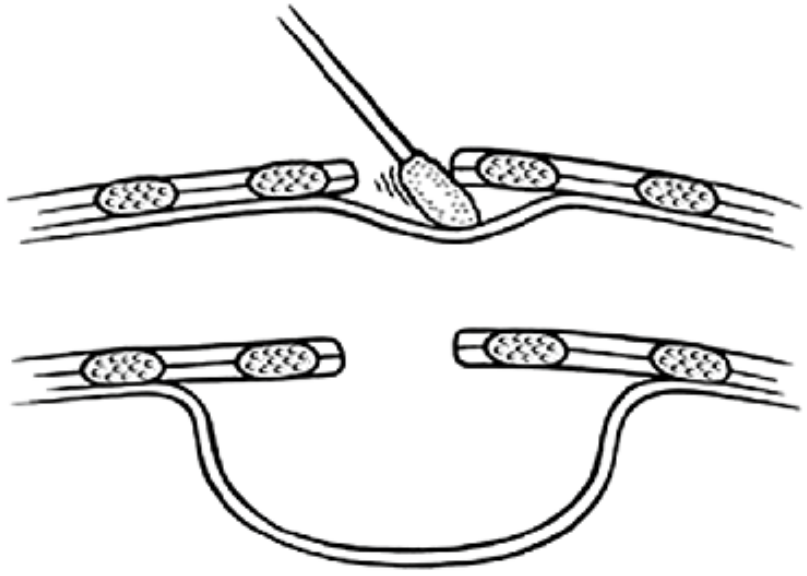
Operative management :-

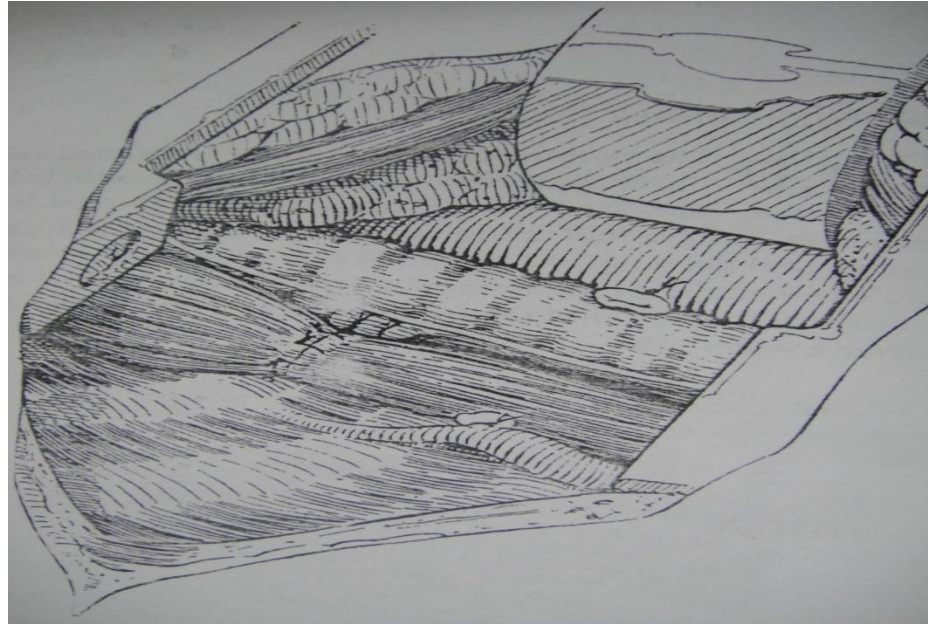
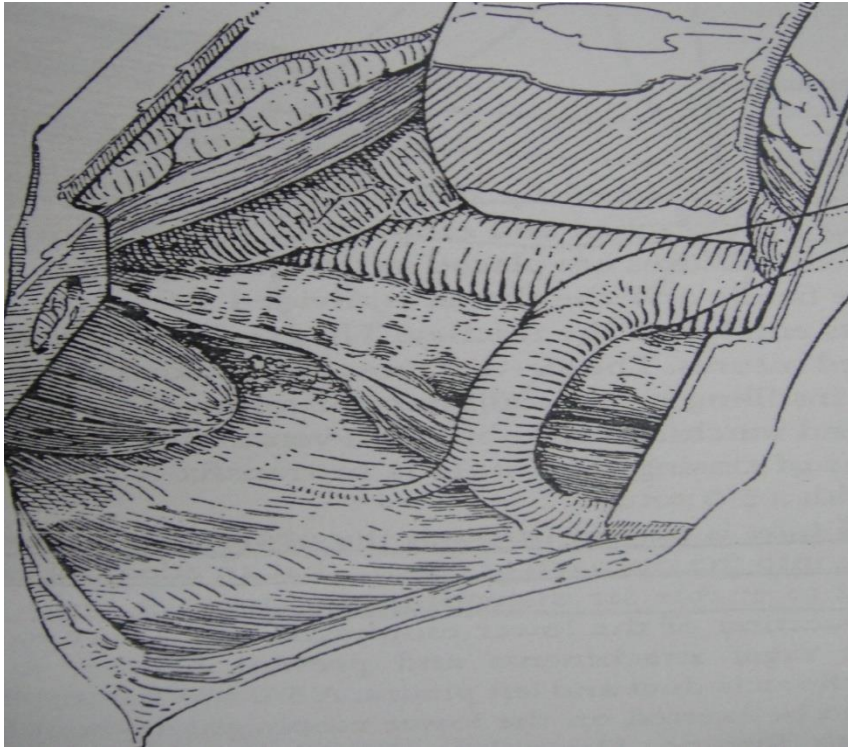
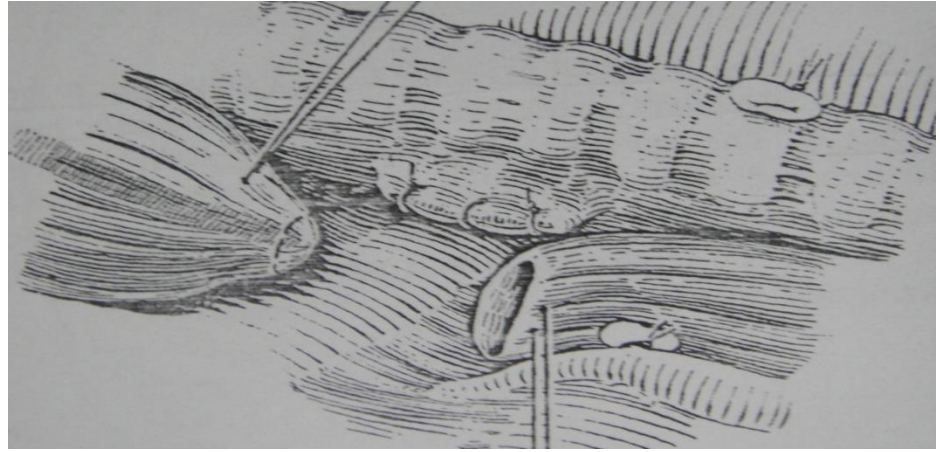
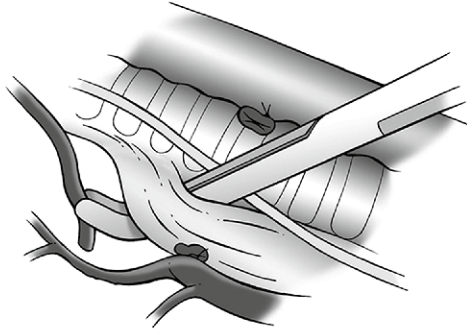
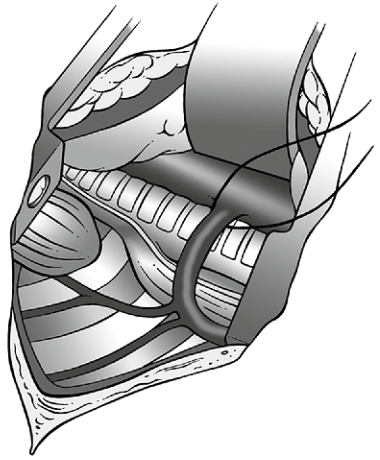
Pre-op. Echo study.

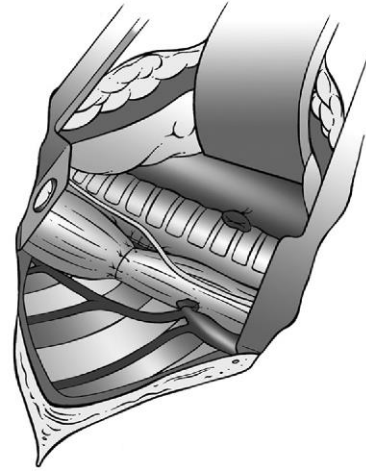
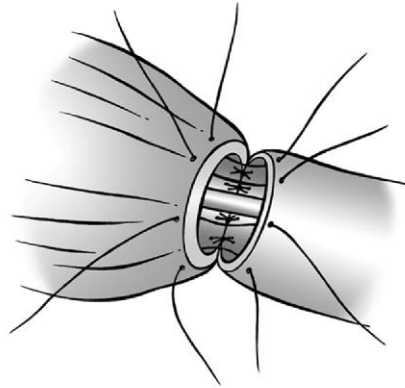
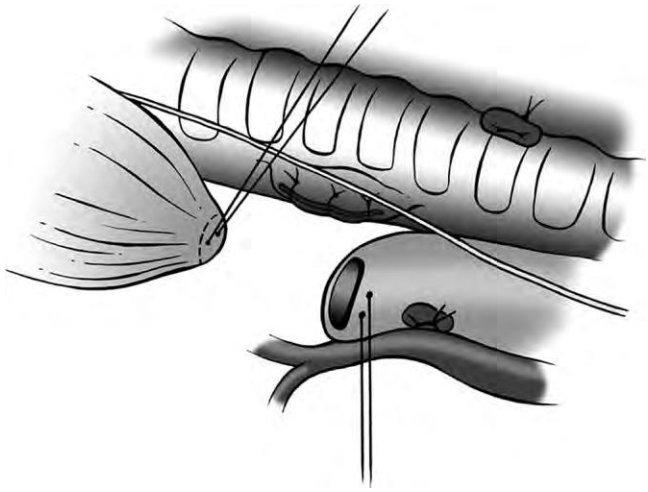
Intra-op. Esophagoscopy.

Intra-op. Bronchoscopy.

thoracotomy.







Childhood tumors

The commonest malignancies in childhood are :-

1- leukemia.

2- CNS tumors.

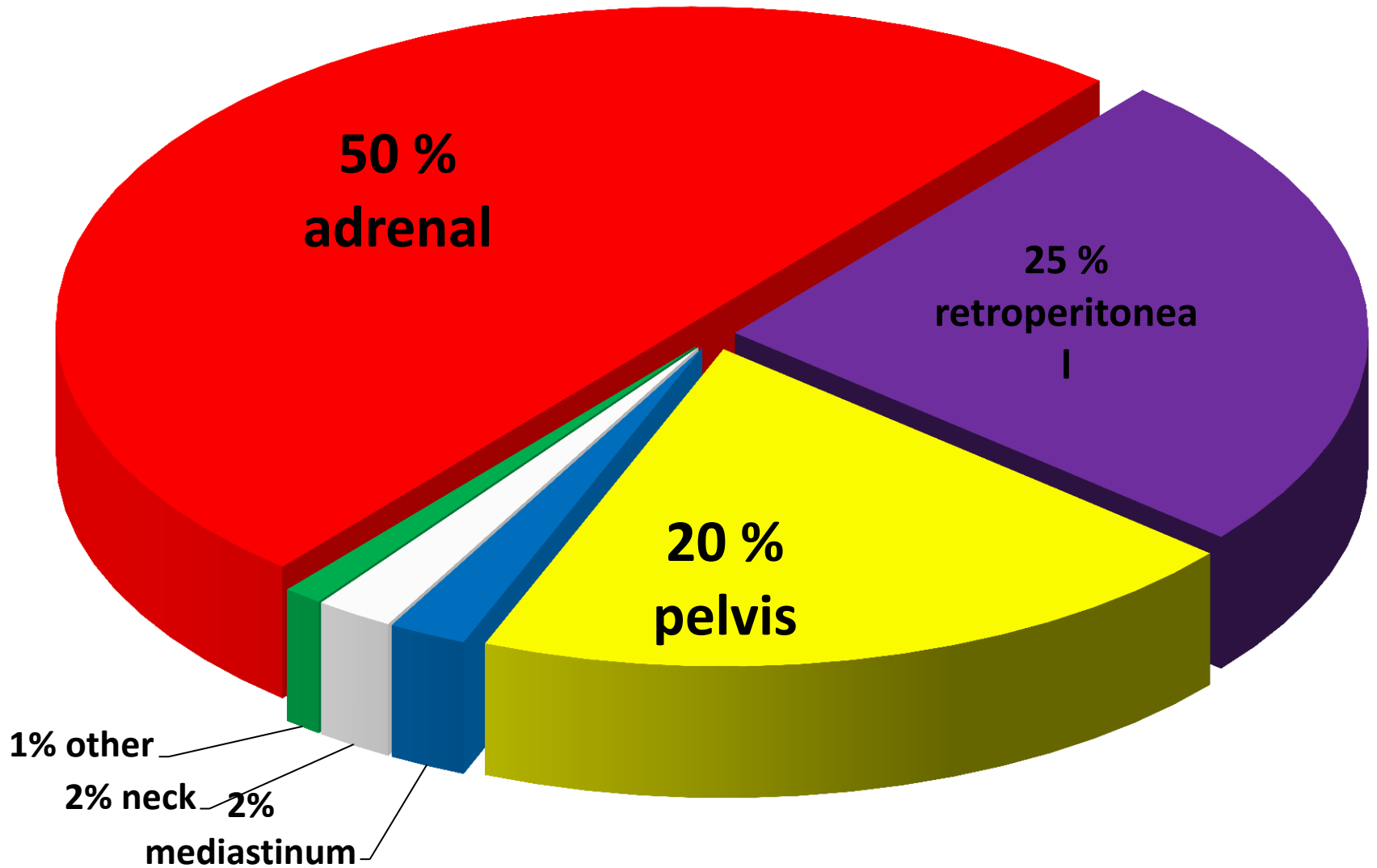
3- lymphoma.

4- neuroblastoma.

5- Wilm's tumor. (nephroblastoma).

6- the other are retinoblastoma, hepatic tumor, bone tumor, soft tissue tumor.

Neuroblastoma



Presentation :-

Patients with NB usually present with signs and symptoms that reflect the primary site and extent of disease, although localized disease is often asymptomatic.

- Fever , weight loss , anorexia , sweating.
- Skeletal involvement (bone and joint pain).
- Orbital sign (ecchymosis and proptosis).”black eye”
- Horner syndrome.
- Pelvic involvement.
- Cervical involvement (dysphagia)

Investigation

❑ *24 hr urinary catecholamine metabolites (VMA & HVA)*

❑ *Basic evaluation. (CBC, biochem., RFT, LFT, LDH, ...)*

❑ *Imaging.*

❑ *Biopsy.*

Ultrasound.

Abdominal X- ray.

CXR.

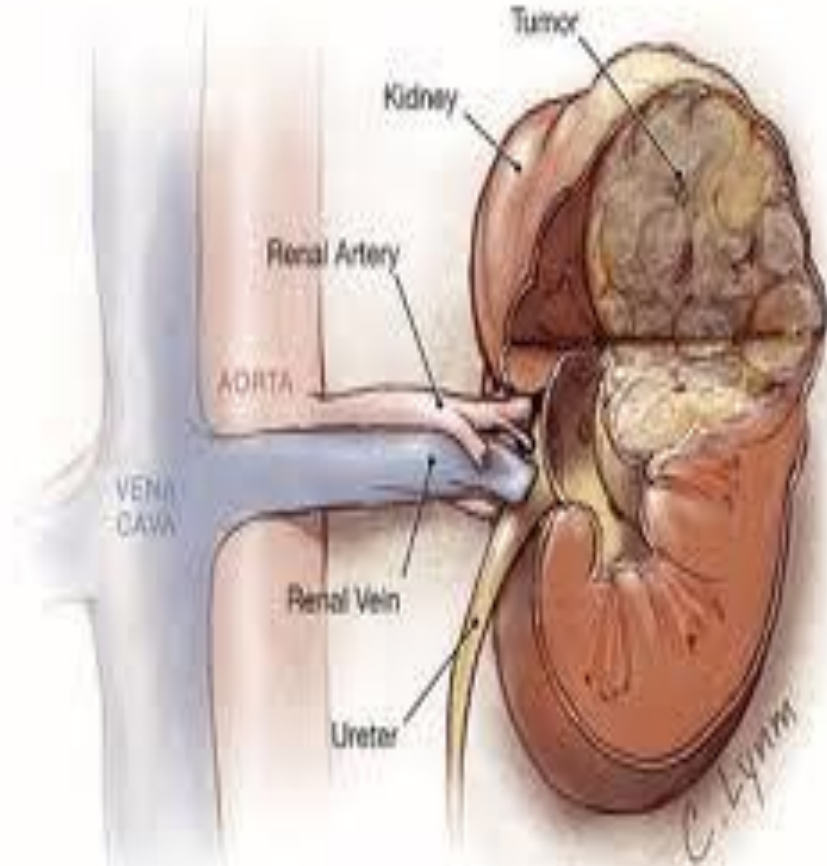
CT , MRI.

MIBG scan.

TREATMENT

- MDT
- Chemotherapy.
- Surgical intervention.
- Radiotherapy.

Nephroblastoma (Wilms' tumor).



Clinical presentation :-

- **80%** *asymptomatic flank mass.*
- *Hematuria.*
- *Fever.*
- *Hypertension.*

TREATMENT

- MDT
- CHEMOTHERAPY
- SURGERY

Sacrococcygeal teratoma

Teratos = monster.

Onkoma = swelling.

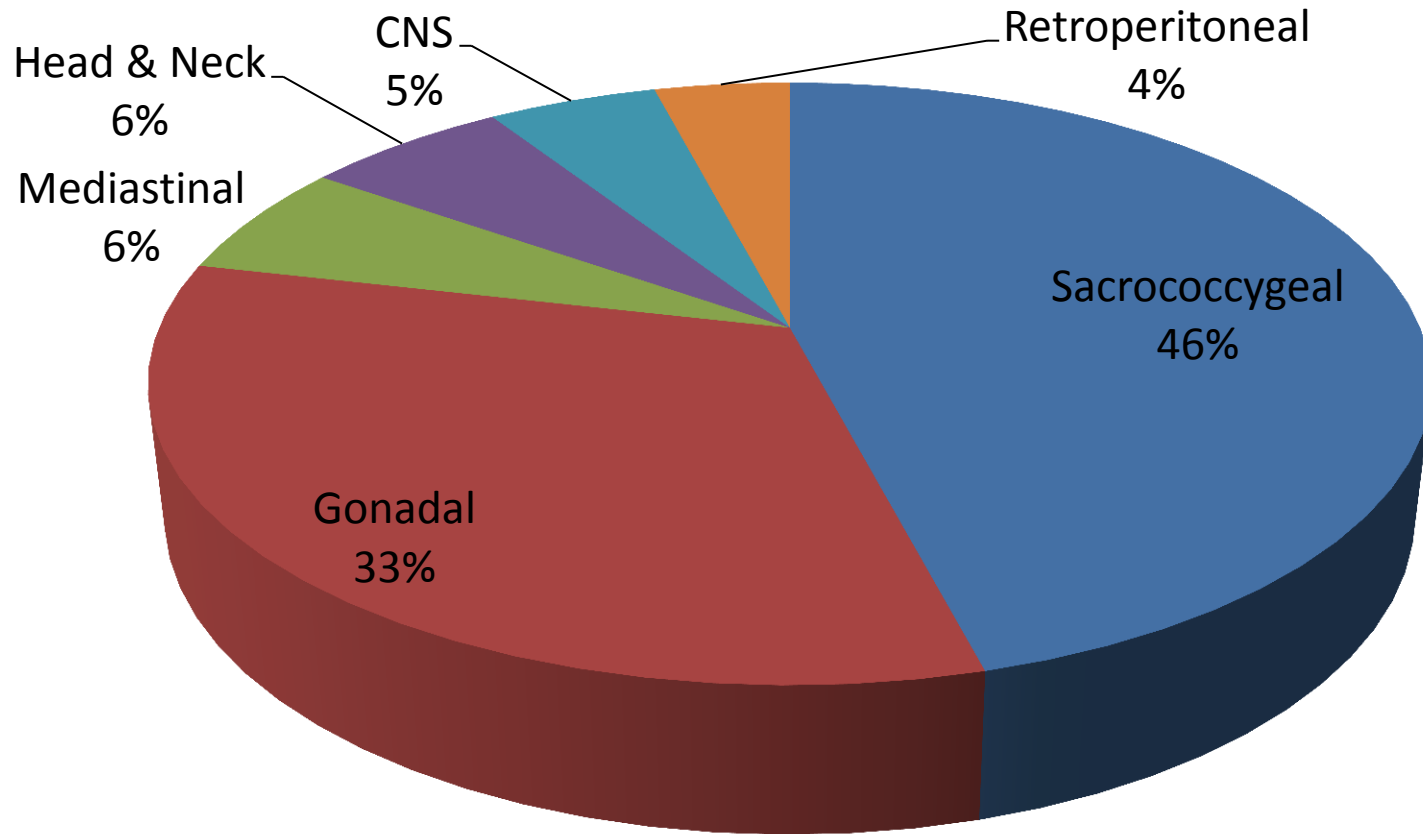
The tumor is composed of multiple tissues foreign to the site of origin.



©2007 Marcos Antonio Velasco Sanchez

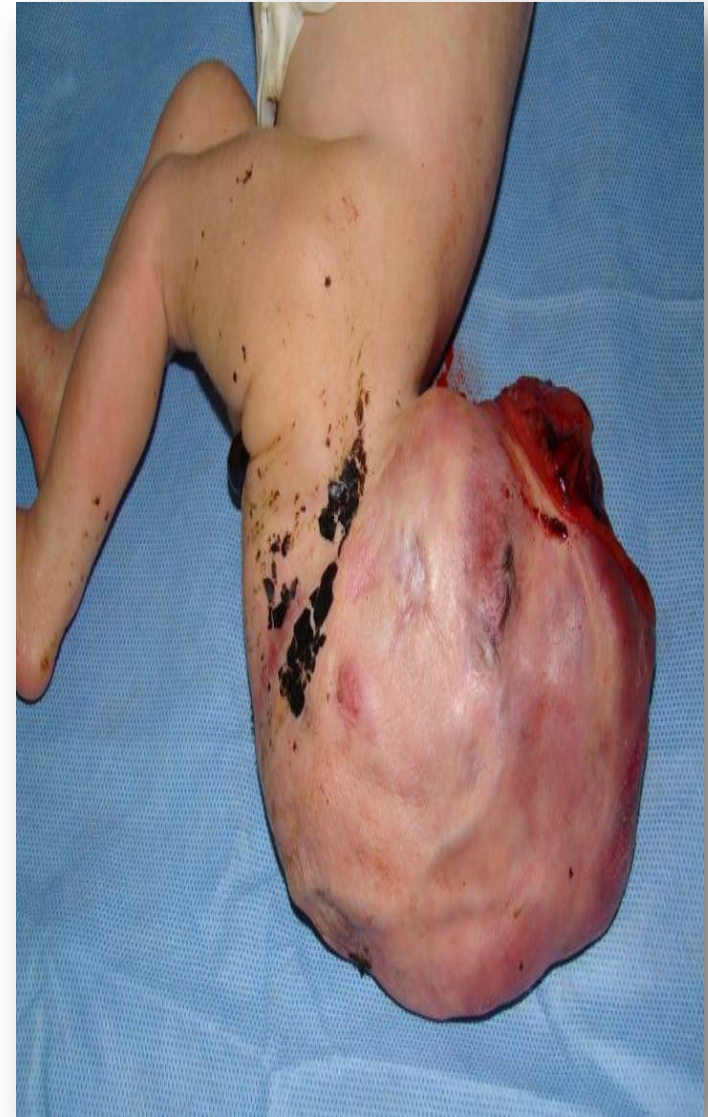


Teratoma



Clinical presentation :-

- ✓ **Visible caudal mass.**
- ✓ **Constipation.**
- ✓ **Urine retention.**
- ✓ **Leg edema.**
- ✓ **High output cardiac failure.**
- ✓ **Intrapartum complications.**



Surgery

- ***Risk of malignancy.*** 10% birth-----75% 1 year
- ***En bloc excision with the coccyx.***
- ***Serum tumor markers.*** Carcinoembryonic Ag., HCG, AFP

