

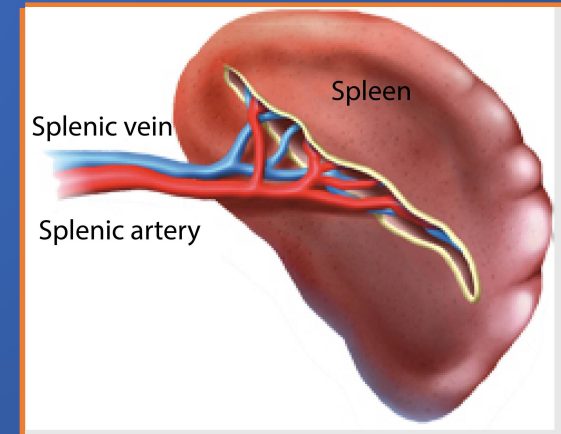
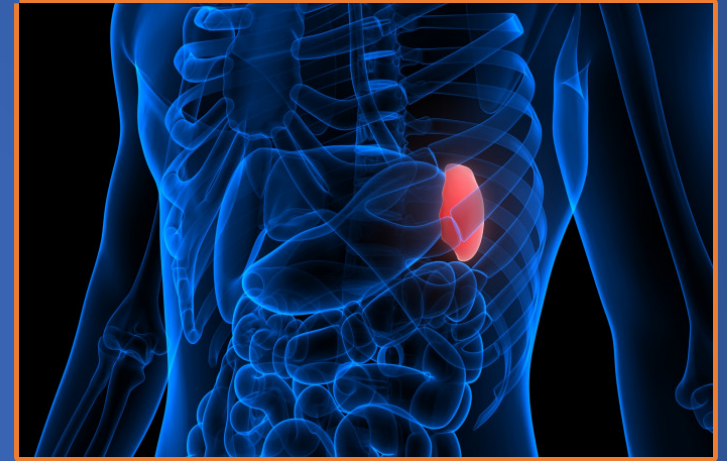
Diseases Of Spleen

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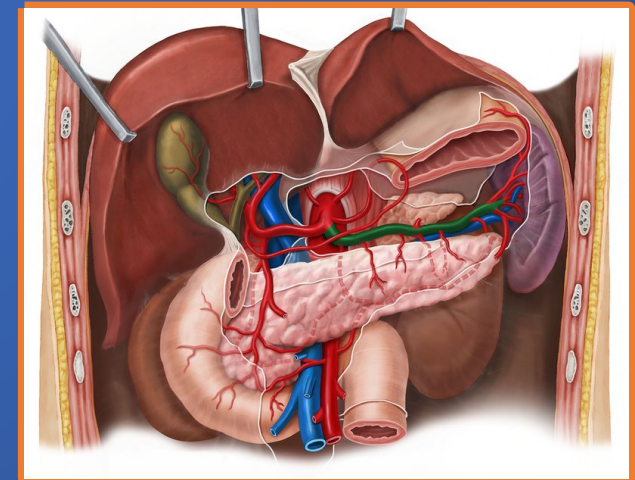
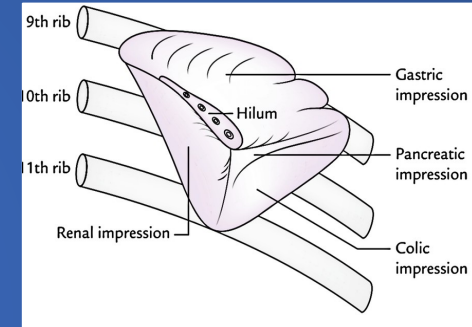


Objectives

- Anatomy & functions
- Causes of splenomegaly
- Splenic diseases of surgical interest
- Indications of splenectomy
- Hematological changes post-splenectomy
- Post-splenectomy sepsis

Surgical Anatomy

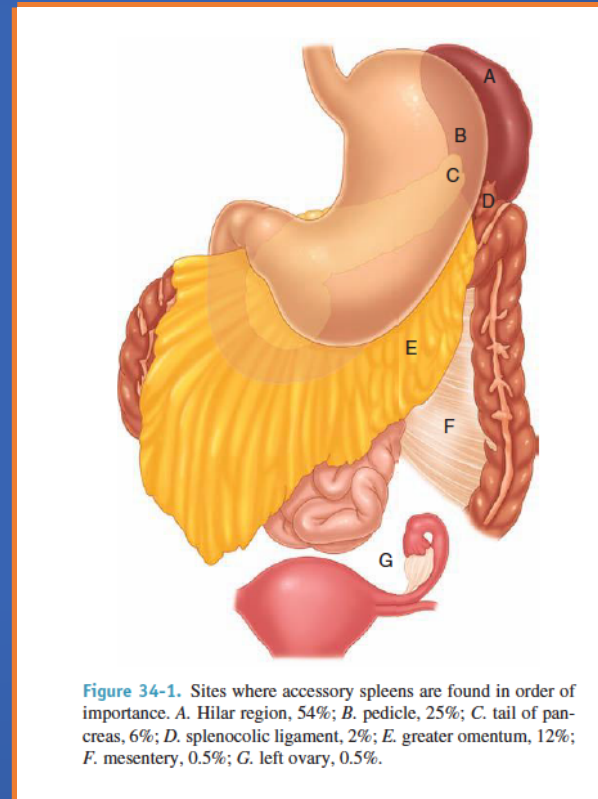
- **Convex surface & upper pole:**
 - related to diaphragm (9-11 ribs).
- **Concave surface:**
 - Fundus of stomach, tail of pancreas, & upper pole of left kidney
- **Lower pole:** rests on splenic flexure of colon



Surgical Anatomy

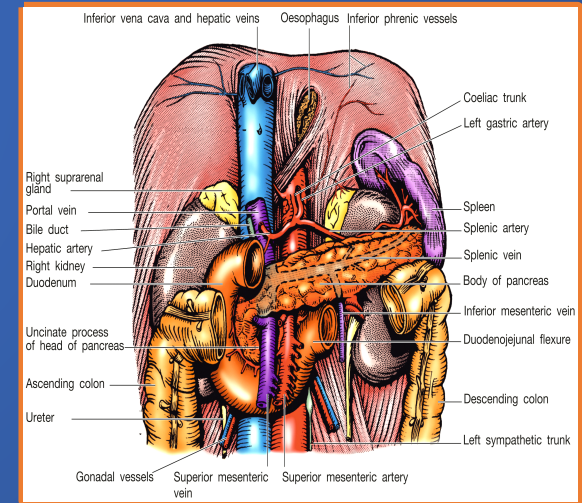
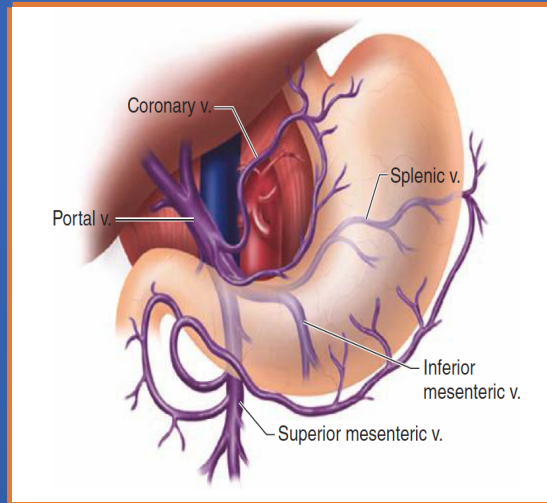
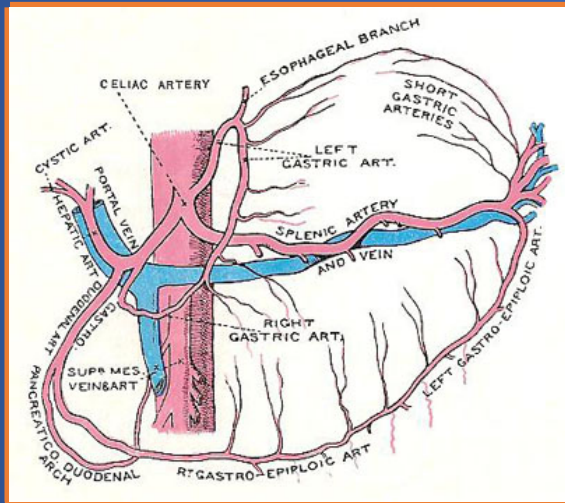
A presence of spleen

- **Accessory spleen:**
 - (10-20%)
 - Mostly hilum, may be anywhere



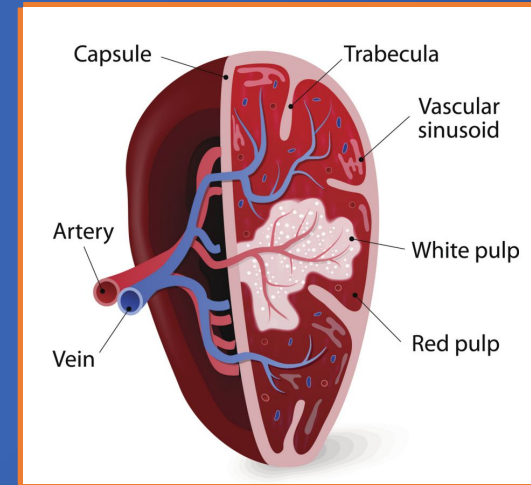
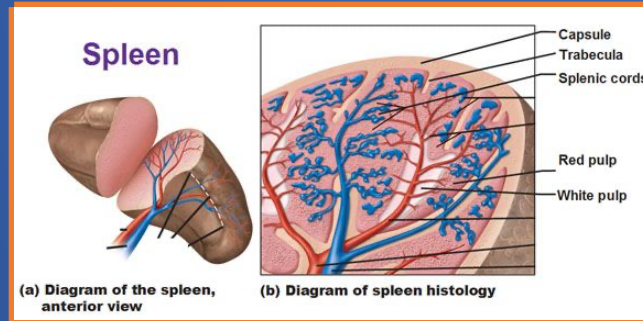
Surgical Anatomy

- **Splenic artery:** branch from celiac axis
- **Splenic vein:** joins SMV to form portal vein



Surgical Physiology

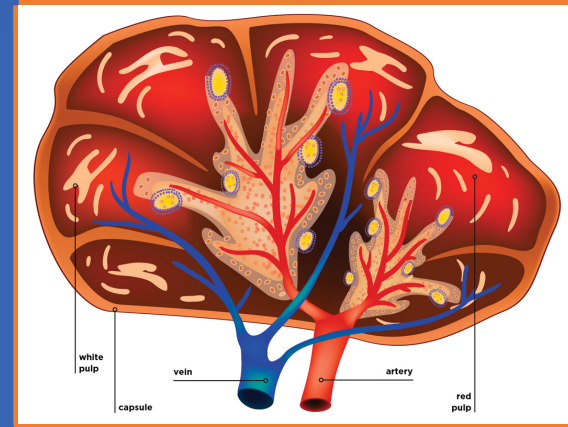
- Highly vascularized (5% CO).
- Largest **filter** of blood & a **lymphoid** organ
- Composed of **red & white** pulp.



Surgical Physiology

- Red pulp:

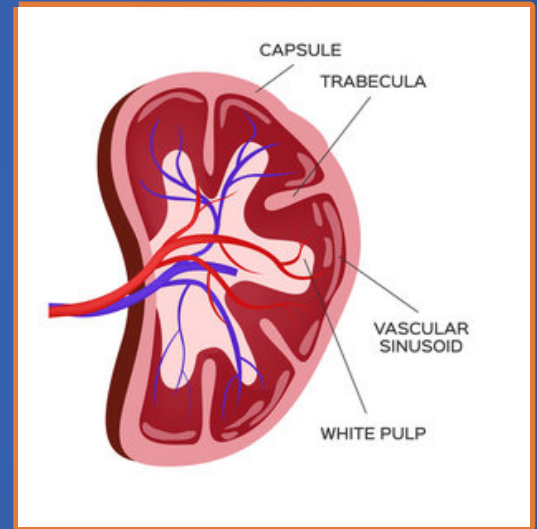
- Made up of **sinusoids**
 - Filters old RBC
 - Phagocytose
- Iron transported back to bone marrow for new RBC
- RBC & Platelets: 1% & 20-30% respectively are sequestered
- **(Howell-Jolly bodies)**: Post-splenectomy- mis-shapen RBC with nuclear remnants seen in circulation



Surgical Physiology

• White pulp:

- largest aggregation of lymphoid tissue
- Composed of **lymphoid follicles** (Malpighian bodies), lymphocytes (T & B), macrophages, & plasma cells
- Site of antigen presentation & antibody production



Immunological function

- Largest aggregation of lymphoid tissues
- Promotion of cell mediated & humoral immunity
- Antigens engulfed by macrophages for antibody production- immunoglobulin (IgM)
- Production of **opsonins, properdin** from lymphocytes
 - Binds to macrophage & leukocyte
 - Promote phagocytosis and bacteriocidal activity
- **Splenectomy** impairs immunological responses

Causes of splenomegaly

- **Clinically palpable spleen enlarged 3 times**

- **Infective**: TB, abscess, HIV, malaria, schistosomiasis, hydatid cyst

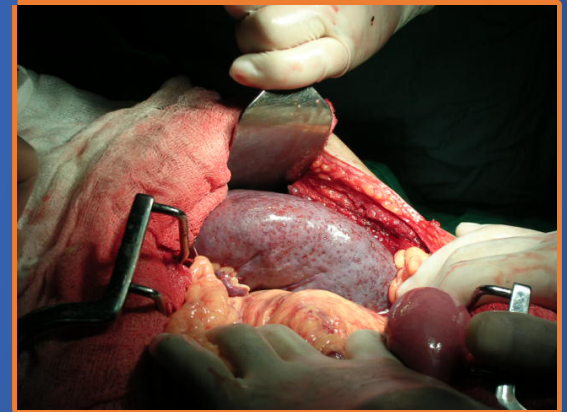
- **Blood disease**: ITP, **Hereditary spherocytosis**, autoimmune haemolytic anemia, thalassaemia, **sickle cell disease**, polycythemia, **leukaemia**

- **Metabolic**: Gaucher's disease , amyloidosis

- **Circulatory**: Portal hypertension, infarction

- **Nonparasitic Cysts**: Congenital/ acquired

- **Neoplasms**: Hodgkin's, other lymphoma, myelofibrosis, angioma

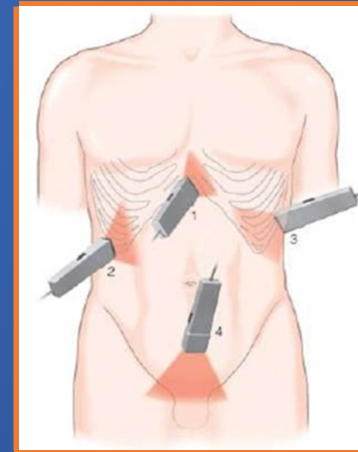


Indications of splenectomy

- **Splenic trauma**- most common indication, hemodynamic instability
- **Purpuras**: Idiopathic thrombocytopenic purpura (ITP)
- **Haemolytic anaemias**: **Hereditary spherocytosis**, Acq. haemolytic anaemia.
- **Hypersplenism**
- **Left sided portal hypertension**
- **Myelofibrosis**
- **Tumours**: Lymphomas, haemangioma
- **Cyst of spleen**
- **Splenic infarct**
- **Abscess**
- **Splenic artery aneurysm**

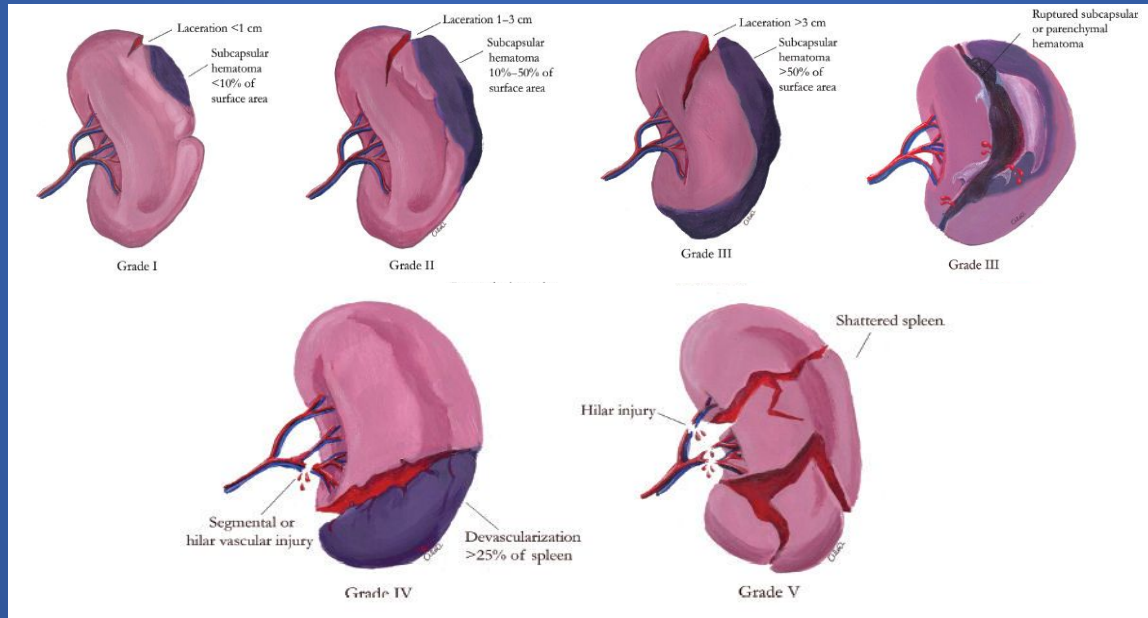
Splenic Injury

- **Aetiology: Blunt/ Penetrating injury**
- Blunt trauma: most frequently injured organ
- **Injury to left side-** chest, flank, or abdomen
- **Left lower chest & upper abdomen:**
 - **Pain, Bruising, Tenderness**
- **Diagnosis:**
 - **FAST- unstable patients**
 - **FAST+ CT-stable patients**

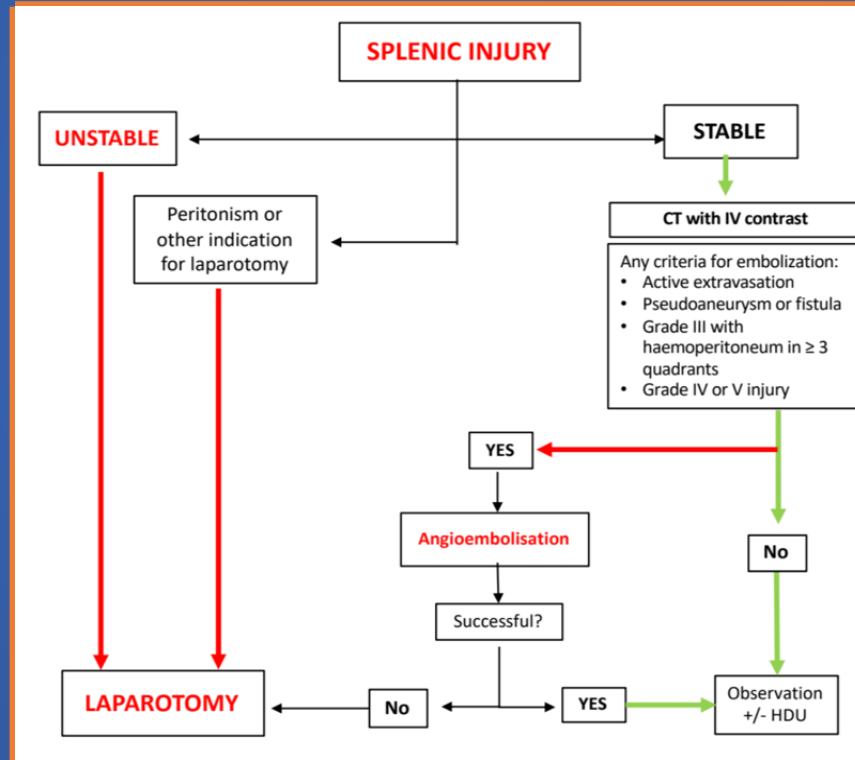


Splenic Injury

- American Association for the Surgery of Trauma (AAST) splenic injury scale



Splenic Injury



Idiopathic Thrombocytopenic Purpura ITP

- ITP in children:
 - 2-4 years age
 - Acute
 - Usually post viral,
 - Most recover without treatment.

Idiopathic Thrombocytopenic Purpura

ITP

- ITP in adults:

- Chronic
- Antibody (IgG) against platelets
- Low platelets $<50,000$ (epistaxis, GI bleeding, ecchymosis)
- Mild splenomegaly.
- Initial therapy (if bleeding)- prednisolone, platelet concentrate, & immunoglobulin.
- Splenectomy: (Commonest elective indication)
 - Persistent $<30,000$ platelet after 4-6 weeks of medical therapy .
 - Severe thrombocytopaenia- platelet concentrate given after splenic artery ligation. Long time remission in 65%
- 2nd line therapy- Rituximab (anti-CD20 monoclonal antibody)

Hereditary Spherocytosis

- **Autosomal dominant** disorder
- Increased **permeability of cell** membrane to Na, cell **swelling & fragility**
- **RBC- spherical**, fragile, trapped in spleen & destroyed.
- **Excessive haemolysis-** jaundice, anaemia, splenomegaly,
- **Pigment gallstone** formation in **30-60%**.

Hereditary Spherocytosis

- Spontaneous **remission & relapse**.
- Haemolytic crisis needs blood transfusion
- Mild cases managed without splenectomy.
- **Mild / severe : Splenectomy** after age **6 years** (risk of **OPSI**)
- Simultaneous **Cholecystectomy** if gallstone present

Acquired Haemolytic Anaemia

- **Aetiology:** Haemolysis due to exposure to **drugs**, or immune reaction as in **SLE**, chronic lymphatic **leukaemia** or mycoplasma pneumoniae **infection**.
- **Initial treatment:** Steroid therapy.
- **Splenectomy:**
 - No response to steroid
 - Relapse on cessation of steroids

Hypersplenism

- **Splenomegaly, pancytopenia, normal bone marrow & no autoimmune disorder**
 - .
- Causes: Malaria, portal hypertension, rheumatoid arthritis, myeloproliferative disorder.
- **Sequestration & destruction**- predominantly **WBC & platelets**
- Anaemia, leukopenia & thrombocytopenia.
- Splenectomy- sometime after benefit & risk assessment.

Segmental Portal Hypertension

- (Left sided PH)
- Aetiology:
 - Thrombosis of splenic vein
 - Acute/ chronic pancreatitis,
 - Carcinoma pancreas.
- UGI bleed from **gastric varices**, hypersplenism.
- Endoscopic control of varices unsuccessful
- **Splenectomy+ ligation** of vessel on greater curvature of stomach very effective.

Proliferative disorders

- **Myelofibrosis:** Proliferation of mesenchymal elements (connective tissues) bone marrow fibrosis, spleen, liver, lymph nodes & extramedullary haemopoieses.
- Seen in over 50 age group.
- Huge splenomegaly & infarct causes discomfort.
- Splenectomy relieves symptoms.
- **Tumours:** Large haemangioma, Non-Hodgkin's lymphoma confined to spleen-splenectomy

Uncommon indications of Splenectomy

- **Cysts:**

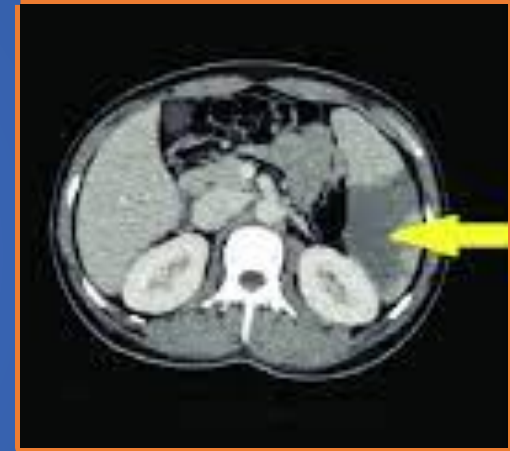
- Congenital, degenerative, hydatid disease.
- **Infarction spleen:** arterial embolism,
- Asymptomatic/ pain LUQ.
- CT: hypo-perfused area

- Splenic **abscess**

- Splenic artery **aneurysm:**

- Typically incidental finding, female, rupture during pregnancy.
- **Treatment:** proximal location – proximal & distal ligation. Distal location- proximal ligation with splenectomy

- **Part of other surgery :** Distal pancreatectomy, radical gastrectomy for ca.



Complications of Splenectomy

- Early:

- Haemorrhage (2-5%)
- Organ injury- pancreas (0-6%-open, up to 16%- laparoscopic), splenic flexure , stomach

Complications of Splenectomy

- Delayed:

- Fistula- stomach, pancreas
- Sub-diaphragmatic collection
- Left basal atelectasis & pleural effusion
- Thrombocytosis- thrombotic complications
- OPSI- H influenzae, Meningococcus
- Splenosis- splenic rupture, bag rupture in laparoscopic surg.

Effects of Splenectomy

- **RBC:** Howell Jolly bodies, erythroblasts
- **WBC:** Leucocytosis
- **Platelet:** Thrombocytosis, increased adhesiveness.
- **Immunological defects:**
 - ↓ serum IgM level
 - ↓ level of phagocyte promoting peptide
 - ↓ response to particulate antigens

Immunization

- **Prone to infection** (encapsulated bacteria- Strep pneumoniae, Neisseria meningitides, Haemophilus influenzae)
- **Elective splenectomy:** vaccination **2-3 weeks before surgery**
- **Emergency splenectomy-** vaccination **postoperatively**
- Polyvalent pneumococcal vaccine (pneumovax)



Immunization

- Not previous immunized persons:
 - Strep pneumoniae (booster dose in 8 weeks)
 - Haemophilus influenza type b (Hib)
 - Meningococcal type c
- Life long antibiotic prophylaxis:
 - Oral phenoxymethyl penicillin or erythromycin

References

- Principles and Practice of Surgery
 - Pg 229-232

Thanks