

# ***Lymphadenopathy & Splenomegaly***

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# ***SPLENOMEGALY***

## *At a glance*

- ❑ Definition
- ❑ Pathophysiology & Epidemiology
- ❑ Etiology
- ❑ History & Ph Ex
- ❑ Differential Diagnoses
- ❑ Workup
  - Laboratory Studies
  - Imaging Studies
- ❑ Treatment & Management

## *Definition*

Splenomegaly is defined as enlargement of the spleen, measured by size or weight.

# Pathophysiology

## Anatomy

The spleen is the largest lymphoid organ in the body.

6/0 cm at age 3 months

6/5 cm at age 6 months

7/0 cm at age 12 months

8/0 cm at age 2 years

9/0 cm at age 4 years

9/5 cm at age 6 years

10/0 cm at age 8 years

11/0 cm at age 10 years

11/5 cm at age 12 years

12/0 cm at age 15 years < (females)

13/0 cm at age 15 years < (males)

## *Function of spleen*

- The filtration of defective blood cells
- Reservoir for platelets
- Produces blood components
- Immunologic function

# *Epidemiology*

A 1- to 2-cm splenic tip is palpable :

- approximately 30% of full-term neonates
- as many as 10% of healthy children overall.

# *Etiology*

Splenomegaly is often categorized in :

- Sequestration of blood cells - Such as in hemolytic conditions
- Proliferation due to infection or inflammation
- Deposition - Such as in Niemann-Pick and Gaucher disease and in some infections



# *Etiology*

- Infiltration due to granulomatous, histiocytic, lymphoproliferative , or malignant conditions
- Endowment - As caused by space-occupying lesions

# *Etiology*

Splenic masses

Hyperplasia

Malignancy

Portal venous system abnormalities

Deposition causing splenomegaly

Masses

Extramedullary hematopoiesis

Hypersplenism

Splenic sequestration

# *History*

- Duration of known enlargement of the spleen.
- Exposure to hepatotoxic agents or microorganisms resulting in hepatitis or portal hypertension.
- Abdominal trauma that may cause splenic hematoma.
- Signs of infection or known infections such as hepatitis, mononucleosis, malaria, or salmonellosis.
- Inflammatory bowel disease

# *History*

- Bone pain, fever, malaise, lethargy, pallor, bruising, weight loss, night sweats, or other findings that may indicate malignancy
- Jaundice suggestive of hepatobiliary disease
- Anemia
- Cholecystectomy
- Splenectomy - Eg, due to hemolytic anemi

# *History*

- Fever or rigors indicative of infection
- Jaundice (evidence of liver disease).
- Abnormal bleeding or bruising (hematologic malignancy).
- Family history of hemolytic anemia (thalassemia major).
- Travel to endemic areas (e.g., malaria).
- Trauma (splenic hematoma).

# *Physical Examination*

- The patient should be examined in the supine or right lateral decubitus position.
- Palpation should start at the pubis and move toward the left upper quadrant to identify the medial and inferior border of the spleen.

## *Physical Examination*

- Size of spleen; consistency, tenderness.
- Hepatomegaly.
- Lymphadenopathy.
- Fever.
- Ecchymoses, purpura, petechiae.

# *Physical Examination*

- Pruritus, pallor, icterus, exanthem, or enanthem
- Abnormal vital signs
- Ophthalmologic abnormalities (uveitis, iritis, vascular occlusion, opacification)
- Abnormal heart sounds
- Dyspnea, abnormal breath sounds
- Hepatomegaly, abdominal masses or tenderness



# *Differential Diagnoses*

- Acute Lymphoblastic Leukemia (ALL)
- Cavernous transformation of the portal vein
- Gaucher Disease
- Heart Failure, Congestive
- Hemolytic Anemia
- Hepatitis
- Hereditary Spherocytosis

# *Differential Diagnoses*

- Hodgkin Lymphoma
- Immunodeficiency disorders
- Lipid Storage Disorders
- Portal Hypertension
- Sickle Cell Disease
- Systemic Lupus Erythematosus (SLE)
- Tuberculosis (TB)

## *Workup in pediatric splenomegaly*

- Splenomegaly is usually the result of a systemic disorder rather than primary splenic disease.
- Therefore, diagnostic studies are not directed solely towards the spleen.
- The most useful initial laboratory test is the complete blood count (CBC) with manual differential and peripheral blood smear.
- This test should be performed on all patients with an enlarged spleen.

# *Laboratory Investigations*

- Evaluation for evidence of hemolytic disease
- Evaluation for infection
- Evaluation for liver disease
- *Evaluation for portal hypertension*
- *Evaluation for connective tissue disease*
- *Evaluation for infiltrative disease*
- *Lymph node biopsy*

# *Laboratory Studies*

The CBC may be revealing, as follows:

- The WBC count
- The platelet count
- Hemoglobin concentrations
- LFT

## *Workup in pediatric splenomegaly*

Ultrasonography alone is the most appropriate means of imaging the spleen in pediatric patients, s noninvasive and does no radiation.

## *Imaging studies:*

- CT scan
- Magnetic resonance imaging (MRI)
- Liver – spleen scans with  $^{99m}\text{Tc}$ -sulfur colloid

# ***Treatment & Management***

Because splenomegaly is usually the result of a systemic disease, the primary goal is treatment of the underlying process.

## ➤ ***Splenectomy***

- the benefit-to-risk ratio must be carefully determined if splenectomy is considered.



*Thank You For Your Attention*

