



THE SPLEEN

Imaging Techniques

- CT and US: major techniques.
- Radioisotopes: → confirm the presence of functioning splenic tissue.
- MR: CT is more important

Anatomy

- **Functions:** **Formation** Fetus Blood - **Sequestration** Aged Cells – **Reservoir** R.B.Cs
- **Site:** left upper quadrant /below the diaphragm /, posterior & lateral to the stomach.
- **Surfaces:** 2 surfaces / **Diaphragmatic** → smooth and convex,
 - **Visceral**: Have concavities for the stomach, kidney, colon, and pancreas.

➔ **Size:**

Average dimensions
(in adults)
Length : 12 cm,
Width : 7 cm,
Thickness : 3 to 4 cm.

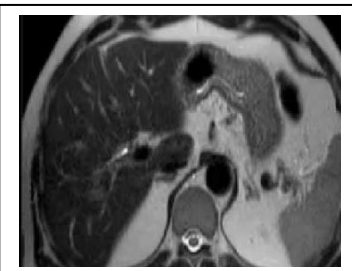
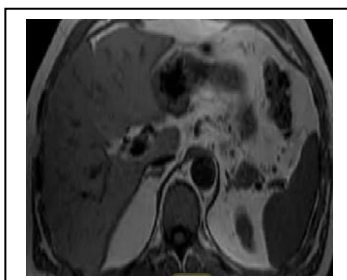
CHILD	
6 + 1/3 Age in Years	
ADULT	
Splenic / Renal index	not > 1.2
length x width x thickness	120 : 480 Cm ³

Blood supply:

- The splenic artery and vein → course through the pancreas to the splenic hilum, → divide into multiple branches.
- *Splenic arteries*: are end arteries (no anastomoses or collaterals) → Occlusion of the splenic A. or its branches → **infarction**.
- Variable rates of blood flow through splenic parenchyma → Heterogenous Enhance "Transient pseudomasses" @ early arterial phase both CT and MR .

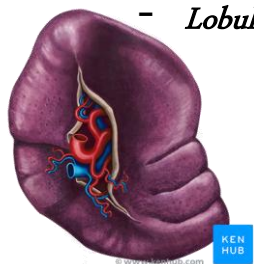
Imaging:

- On all imaging studies, → **homogeneous** appearance.
- CE CT and MR → lesions are best demonstrated .
- On **noncontrast CT**, *spleen density* is less than or equal *normal liver*.
- On MR: intensity
 - T₁WIs → is **lower** than hepatic parenchyma.
 - T₂WIs → **higher** than liver parenchyma on.





- **Arterial phase** → irregular defects in parenchymal enhancement .
- **One or 2 minutes later**, → the entire spleen is homogeneously enhanced on both CT and MR.
- **Lobulations and clefts** : **common** / must not be mistaken for *masses* or *splenic fractures*.

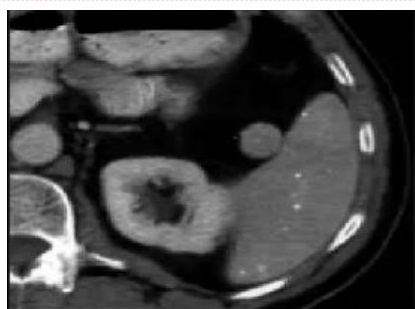


Congenital variations

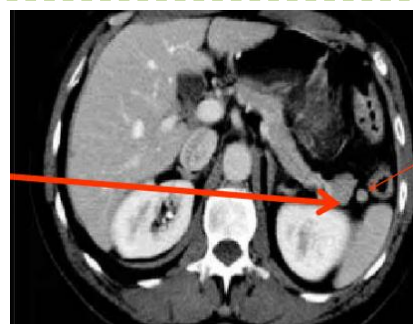
- **Accessory spleen**
10-30% may enlarge dramatically after splenectomy
- **Polysplenia / asplenia**
multiple spleens + cardiovascular and visceral anomalies
- **Wandering spleen**
mobile or ectopic spleen
Torsion → symptoms

Accessory spleens 10% to 16% of normal individuals

- **Round masses**, Single or multiple.
- **1 to 3 cm**,
- **Same** as normal splenic parenchyma.
- Usually near the splenic hilum.



Ⓡ If Near Tail of Pancreas
→ Misdiagnosed
as pancreatic mass



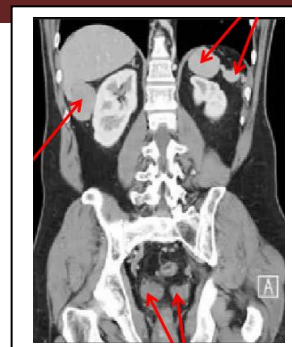
Wandering spleen

- **Laxity of the splenic ligaments**
→ spleen to be wander anywhere in the abdominal cavity.
- (+/- abnormalities of intestinal rotation,)
- **C.P.:** most cause → no symptoms. / Mass / pain ← Torsion
- **Complication** : Torsion



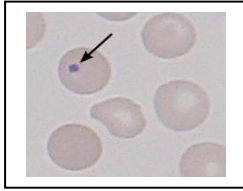
Splenosis

- **Def:** multiple **implants** of ectopic splenic tissue ,
may occur after traumatic splenic rupture.
- **Pathology** : Splenic tissue can implant anywhere in the abdominal cavity
, or (*even in the thorax if the diaphragm has been ruptured*).
 - Splenosis complicates 40% to 60% of splenic injuries.
 - Usually multiple & vary in size and shape.





- The tissue fragments enlarge over time → may simulate **peritoneal metastases**.
- After splenectomy, → seeding of splenic tissue rupture → *remaining accessory spleens* or *splenules* → may enlarge and **resume the function** of the resected spleen.



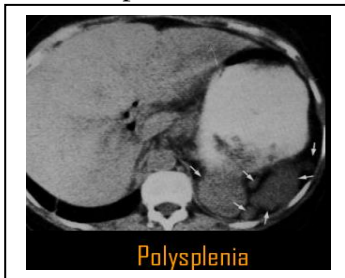
- **Howell-Jolly bodies**: bits of nuclear material seen in RBCs after splenectomy.
- *Disappearance of these Howell-Jolly bodies* → *sign of splenic regeneration*.

® **Imaging studies** → single or multiple *spleen-like* masses in the abdominal cavity + history of splenectomy.

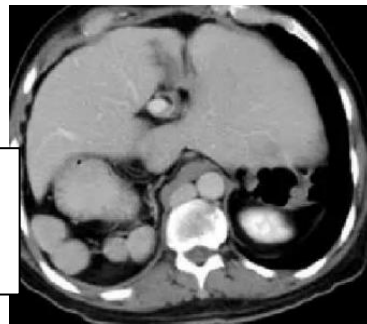
Polysplenism

Rare

- Multiple small spleens, usually located in the right abdomen + **situs ambiguous**.
- Most patients also have **cardiovascular anomalies**.



polysplenism, situs ambiguous, absent IVC and azygous continuation of the IVC.



Asplenia (Ivemark syndrome)


Mostly die before 1 year of age

- Congenital absence of the spleen.
- **Associated with**: bilateral right-sidedness + midline liver + bilateral three-lobed lungs.
- Major cardiac anomalies → in 50% of cases.



Splenomegaly

splenic length > 13 cm

Congestive	Myeloproliferative	Infection
-Portal hypertension (50% of cases) -Portal vein thrombosis 	-Lymphoma (30% of cases) -Leukemia -Polycythemia -Purpura Idiopathic thrombocytopenia -Sickle cell disease (in infants) -Spherocytosis -Thalassemia major -Hereditary Myelofibrosis	Malaria (universal in endemic areas) Schistosomiasis (endemic areas) Infectious mononucleosis Subacute bacterial endocarditis AIDS IV drug abuse
Infiltrative		
-Systemic lupus erythematosus -Amyloidosis -Gaucher disease		

CHILD

(6 + 1/3 Age in Years)

ADULT

Splenic / Renal index

not > 1.2

length x width x thickness

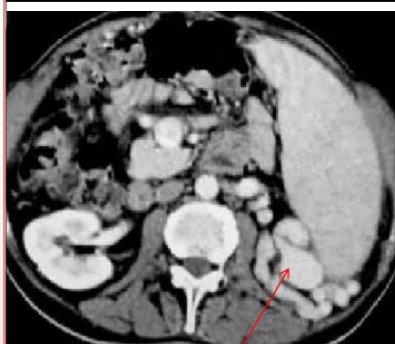
120 : 480

• Diagnosis:

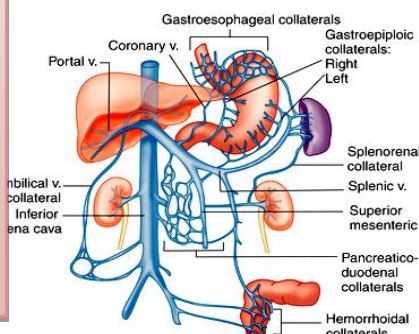
- MR → no significant benefit in the differential diagnosis of splenomegaly.
- Mild to moderate splenomegaly is seen with → portal hypertension, AIDS, storage diseases, collagen vascular disorders, and infection.
- More marked splenomegaly → usually associated with lymphoma, leukemia, infectious mononucleosis, hemolytic anemia, and myelofibrosis.

Porto systemic collateral vessels

- A WV = abdominal wall varices
- CV = coronary venous collateral vessels
- EV = esophageal varices
- GRV = gastro renal shunt
- IMV = inferior mesenteric vein
- LPV = left portal vein
- LRV = left renal vein
- MV = mesenteric varices
- OMV = omental varices
- PEV = Para esophageal varices
- PSV = perisplenic varices
- PUV = Para umbilical collateral vessels
- PV = portal vein
- RGV = retro-gastric varices
- RPPV = retroperitoneal-Para vertebral varices
- SMV = superior mesenteric vein
- SRV = splenorenal shunt
- SV = splenic vein.



Dilated porto-systemic collaterals
In portal hypertension



Sever Para esophageal varices



CYSTIC LESIONS OF THE SPLEEN

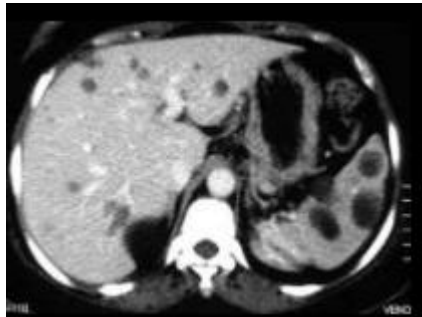
- Usually discovered accidentally
- Large or complicated cysts produce symptoms
- Complications : rupture , infection ,hemorrhage



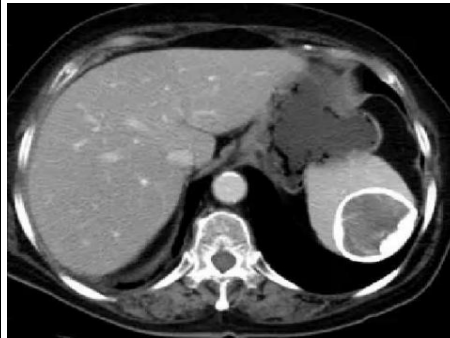
SPLENIC CYSTS

PRIMARY	SECONDARY
True Epithelial Lining	No Epithelial Lining "False"
Parasitic "Hydatid"	Post Traumatic (80%)
Cong. Simple	Infectious
Epidermoid 5%	Infarction

Multiple Simple Hepatic & splenic Cysts

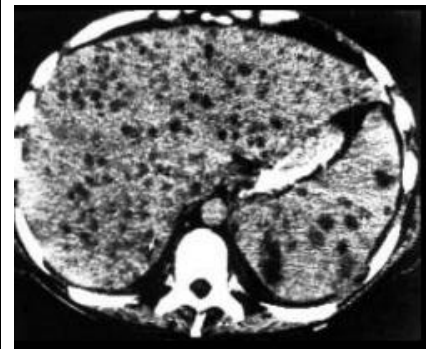


Hydatid Cyst



Abscess

"Enh. Margins & Air less common"

Splenic Sarcoidosis D.D. Metastases
Abscesses
LymphomaMultiple Splenic Calcifications
(TB or Histoplasmosis)Multiple Fungal Abscess
"Immunocompromised"

MULTIPLE SMALL CYSTS

LIVER
Metastases
Abscesses
Simple

SPLEEN
FUNGAL
ABCESS



SPLENIC TRAUMA

(Blunt / Penetrating)

- Commonest Abdominal organ to be injured. +/- Other organs Injury

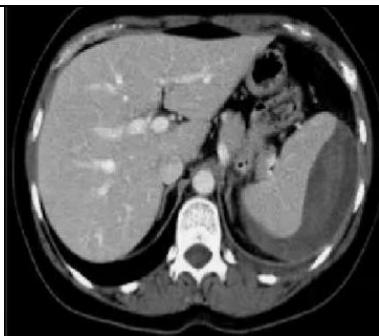
- CT → Best Modality to Diagnose. " accuracy 98% "

- Staged by Rule of 3

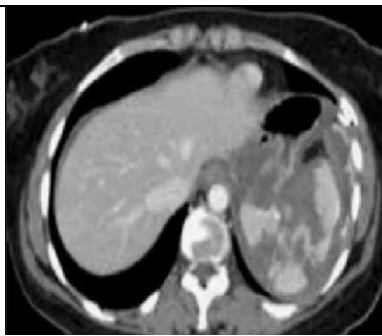
I:II → Heal in 4 m

III → " " 6 m

I	< 1 cm	Superficial laceration	subcapsular hematoma
II	1: 3 cm	Deep laceration	subcapsular hematoma
III	>3cm	Deep laceration	subcapsular hematoma
IV		Fragmentation >3 pieces	no enhancement



Subcapsular Hematoma



Shattered Spleen



Deep Laceration



Splenic hematoma



treated conservatively
over 2 weeks

SPLENIC INFARCTION

- Single or multiple **wedge shaped** hypodensities based to the splenic capsule +/- Ca

- Causes include :

- Embolic in cardiovascular diseases

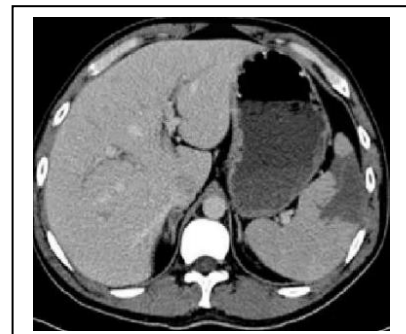
(endocarditis , AF, Local thrombosis)

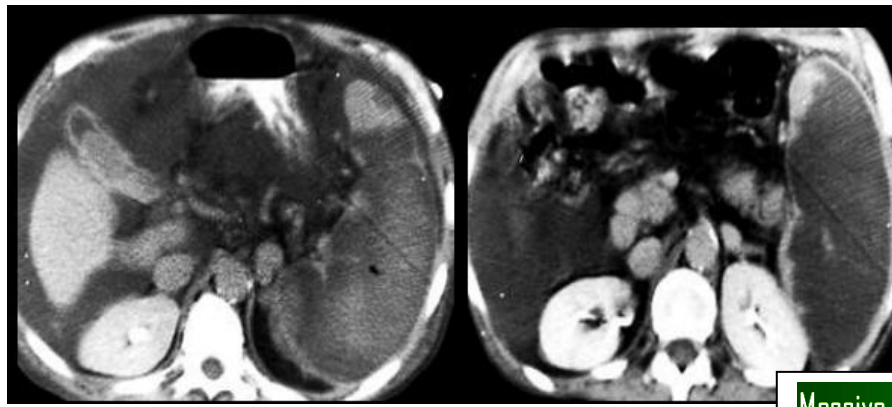
- splenic Torsion ← wandering spleen –

- pancreatitis

- AIDS,

- sickle cell disease (when Chronic → Fibrosis → Volume loss)

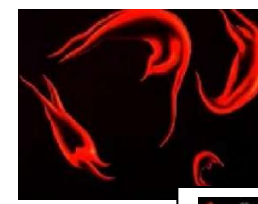




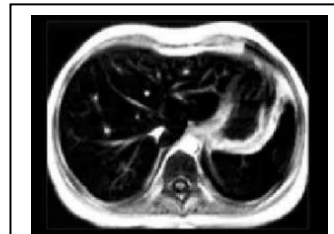
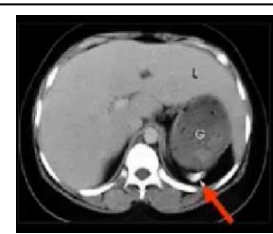
Massive infarctions

SICKLE CELL DISEASE

- Hereditary (autosomal recessive)
- formation of abnormal haemoglobin (haemoglobinopathy)
 - Abnormal shape RBCs "Sickle like" → Aggregation in Blood vessel
- At small arteriole → **INFARCTION**
- At splenic vein → blood trapping → **SQUESTRATION / RUPTURE**
- At splenic A. → **AUTO-SPLENECTOMY**
- Abnormal iron Accumulation "Haemosedrin" → **2RY HAEMOSEDROSIS**



- Splenic infarction
- Splenic sequestration
- Splenic rupture
- Hemosiderosis
- Auto-splenectomy



Infarction	SQUESTRATION	AUTO-SPLENECTOMY	2ry Haemosedrosis
Peripheral wedge shape Hypodensity	Infant or child Enlarged heterogenous spleen +/- Rupture	Total gradual Infarction → Small calcified spleen	iron overload disorder → the accumulation of hemosiderin "Diffuse loss of signal in MRI & increase density in CT.

Sickle cell disease

63Y F with sickle cell disease presented with acute abdominal pain. **CT** showed partially calcified spleen with large hypodense area of infarction. Islands of non calcified splenic parenchyma are seen. **MRI** showed low signal of the liver and spleen due to hemosiderosis with high signal liquefied infarct inside

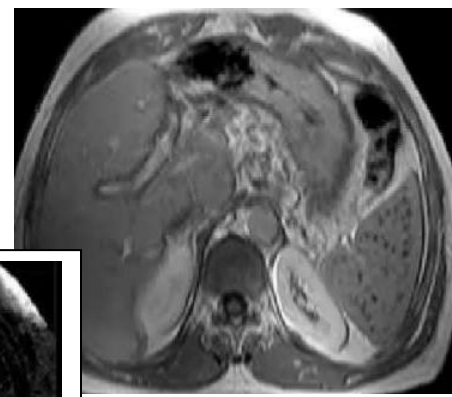
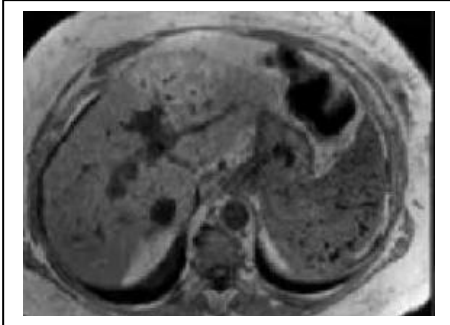




GAMNA GANDY BODIES

Only seen in MRI

- Multiple small Foci of low signal
- = Organizing foci of haemorrhage
- Seen 10 % of portal hypertension / Not related to spleen size.
- Only seen by MRI "Not by CT"



NEOPLASMS OF THE SPLEEN

Any solid lesion in the spleen is considered malignancy until proved

Normal spleen imaging → Not exclude Metastasis

- Primary : Rare
- Secondary: Uncommon



Hemangioma

-Commonest benign (14%)

May involve whole spleen → Hemangiomatosis



Lipoma

Well defined – Fat density



Angiosarcoma

Mass heterogeneous /Rare – Primary



Mets

Uncommon

Heterogeneous spread

Breast, lung, ovary, stomach, melanoma

* Calcification is rare

LYMPHOMA

The most common malignant lesion of the spleen

Splenic involvement in
HD = 27% / NHL = 35%

Lymphoma "4 patterns"

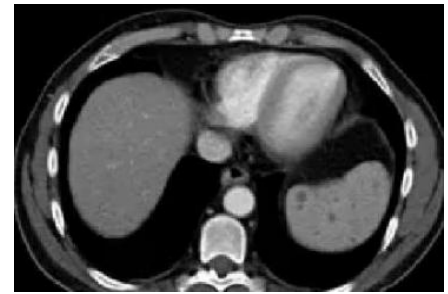
- ♦ Enlarged spleen with homogenous CT
- ♦ Solitary focal lesion
- ♦ Multiple Splenic lesions
- ♦ Diffuse infiltration



SPOTTED SPLEEN

- Multiple low attenuation nodules
- Uncommon Finding / But common in Immunosuppressed "Infectious"

Neoplasm	Infections
<ul style="list-style-type: none"> - Lymphoma "coomonest" - Metastases - Hemangioma 	<ul style="list-style-type: none"> - Fungul - Mycobacterial - Parasetic
	Sarcoid



CASES

Splenic TB 			
Lymphoma 			
			Leiomyosarcoma & Liver mets

**-Accessory Spleen****(SplenuLe)**

- 10.15% of Normal Ind
- Single or Multi
- Round
- 1.3 cm
- Same text as SPL
- Usual periHilar

POLYSPLENIA

- Rare
- Multiple small Spl tissue.

ASPLENIA

- (Ivemark Synd)
- Cong Absent spleen.
- Most die e in 1 y
- +other cong Anomalies

•WANDERING SPLEEN•

- Lax Lig
- Spleen not in Posit
- Hilum directed post

•SPLENOSIS•

- Mlti implanted splen tissue
- After Traumatic rupture
- 40.60% SPL Injury
- Multiple
- Vary size/shape

•Splenic Regeneration•

- EnLargment of access Spl after splenectomy
- > Splenic like masses
- Howell-Jolly bodies
- bits of nuclear mator seen in RBCs after splenectomy.
- Reappear of it = Splenic Regeneration.

**•Cystic Lesions•****1-Post-Traumatic:**

- 80% •False •Thick wall+/-sept
- Calc 30.40%

2-Epidermoid:

- True •Calc < 5%

3-Pancreatic

- e pancreatitis 1.5 %
- Fluid slide on tail to hilum

4-Abscess : "Ring enhance"**5- Micro Abscesses**

- Immunodef •5.20mm

6- Hydatid Cyst

- 2% of hydated cases
- Criteria: may be
 - a-wall calc
 - b-Floating shadows
 - c-Inner cysts "Diagnostic"

• Solid Lesions •**1- LYMPHOMA**

- Commonest
- HD 25% -NHL 35%
- Predispose to Infarc
- =Many patterns.
- Diffuse enlarg
- Solitary Mass
- Multi Mass
- Miliary Nodules
- Direct Invas

2- METASTASES

- Mostly Micro
- "Not imaging detectabl"
- Single/Multi mass
- Low dens
- Low T1 - Hi T2
- CaLC rare

=Melanoma->Cystic Met

NE.Normal spleen not exclude Metastases.

3-INFARCTION

- Wedge shape
- SubCapsular Defect
- Low dens
- T2 hi
- =Complication.
- Hematoma -Infection
- Rupture->Hage

4-Gammmma Gandy

- Portal hptense => Small Hags
- Best seen e MR
- Multi nodules
- LOW T1&T2 "Hemosedrin"

5-HEMANGIOMA

- Commonest 1ry -As in Liver
- Hypoechoic
- Hypodens
- Lo T1 Hi T2

6-ANGIOSARCOMA

- Very Rare -Give wide Metas
- Multi well defined nodules enh
- or Diffuse involving spleen

• Multi CALCEIFICATIONS

- TB. 1st
- Histoplasmosis
- Hemangioma
- Phlebolith
- AIDS

•Multi Small LESIONS•

- MicroAbscesses
- Lymphoma
- Sarcoidosis
- Gamna Gandy
- Metastases
- Histoplasmosis

→Sources

- **Fundamentals Sec VII -CHAPTER 28 -**
- **Primer of Diagnostic radiology 6th**
- **Lecture of Prof. Mamdouh Mahfouz**
- <https://radiopaedia.org>