Focal splenic lesions: Focusing on how to differentiate malignancy from benign

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Learning Objectives

 To evaluate the imaging findings of the focal splenic lesions and to characterize the features that may be helpful in the differential diagnosis of malignant and benign lesions

Background

 The mass of the spleen is often found incidentally during the evaluation of trauma and metastatic cancer. In addition, the appearance of the tumor is different from that of other organs and the characteristic features are not well known. It is difficult to distinguish between malignant and benign tumors, and unnecessary procedures such as biopsy or splenectomy are frequently performed. Therefore, we aimed to evaluate the characteristic features of CT, MR, PET, and US to distinguish between benign and malignant lesions.

Imaging Findings

- Benign
 - Congenital cyst
 - Pseudocyst
 - Lymphangioma
 - Hemangioma
 - SANT
 - Littoral cell angioma
 - Hamartoma
 - Granulomatous lesion
 - Inflammatory pseudotumor
 - Extramedullary hematopoiesis

- Malignant
 - Metastasis
 - Lymphoma
 - Angiosarcoma
 - Undiffernciated carcinoma

Benign

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Cystic lesions in spleen

 Primary (true) – epithelial lining

> Parasitic : Echinococcus Non-parasitic : Congenital Neoplastic

 Secondary (false) – no epithelial lining

> Post-traumatic Splenic infarction

Endothelial cyst



coronal

coronal

sagittal

Epithelial cyst



coronal

sagittal

Epithelial cyst



Pseudocyst



Lymphangioma

- Slow growing
- Histologic types (3):
 - simple lymphangioma
 - cavernous lymphangioma
 - cystic hygroma
- CT:
 - multiple, discrete, nonenhancing low-attenuation lesions
 - Usu. Subcapsular in location
 - Curvilinear peripheral mural calcifications
- MR: T1 high (internal bleeding, large amount of intracystic proteinaceous material)
- Angiography: Swiss cheese app. (well defined avascular lesion and stretching of intraparenchymal arterial branches)

Lymphangioma



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Hemangiolymphangioma



Hemangioma

- Most common primary benign neoplasm
- Endothelial-lined vascular channels filled with blood
- T1 low, T2 high than spleen parenchyma
- Early nodular centripetal enhancement
- Uniform delayed enhancement

Hemangioma



SANT

Sclerosing Angiomatoid Nodular Transformation

- Benign vascular lesion consisting of multiple angiomatoid nodules surrounded by dense fibrous tissue that often coalesces centrally to form a scar
- Pathogenesis: unclear
 - Hamartomatous response of the splenic red pulp to a vascular insult or stromal proliferation
 - Spectrum of IgG4-sclerosing disease
 - Final stage of inflammatory pseudotumor
 - Hamartoma
 - Organizing hematoma

- US: hypoechoic
- CT: solitary, well circumscribed, lower attenuation relative to spleen on early and portal phase, isodense on delayed image
- MR: hyperintense at the periphery, hypointense at the center, hypointense radiation bands corresponding to fibrosis, signal drop on T1 inphase sequence indicating iron deposition due to old hemorrhage, progressive enhancement
- Solitary mass, old hemorrhage, dense fibrosis, and centripetal enhancement lead to suspicion of SANT



SANT 2015



SANT



Littoral cell angioma

- Arising from littoral cell, normally line the splenic sinuses of the red pulp
- Splenomegaly
- Numerous, low-attenuating nodules on portal phase
- Histology: anastomosing vascular channels lines with tall endothelial cells and papillary fronds
- DDx: hemangioma(similar enhancement pattern of liver hemangioma), lymphoma(adenopathy or disease in other organ), disseminated infection, graulomatous disease
- Diagnosis : Biopsy in pts. Without clotting function abnormalty

Littoral cell angiosarcoma



CEUS-post

Hamartoma

- Mixture of normal splenic structures such as white and red pulp
- Commonly associated with tuberous sclerosis
- NECT: isodense to splenic parenchyma, contour distortion
- Occasionally: calcification, cystic change and fat in mass
- T1 iso, T2 heterogeneously high relative to the spleen
- Immediate diffuse enhancement on early and more uniform enhancement on delayed image

Hamartoma



Extramedullary hematopoiesis

- Compensatory response to deficient bone marrow cells
- Predominantly affect the spleen and liver
- Active lesion: T1 intermediate, T2 high , some enhancement
- Old lesion: T1, T2 low, reduced SI on inphase T1WI owing to iron deposition

Extramedullary hematopoiesis





Infection

- Infection:
 - metastatic infection (sepsis, infective endocarditis)
 - Contiguous infection (perinephric abscess, necrotic pancreatitis)
- Infarction with superimposed infection
- Trauma
- Immunodeficiency condition
- Low attenuation with rim enhancement

Abscess



Chronic granulomatous inflammation



- Spectrum of inflammatory lesions: plasma cell granuloma, plasma cell pseudotumor, myofibroblastic tumor
- Underlying inflammatory disease process
- 2 theories:
 - Primary reactive and inflammatory process
 - Low-grade neoplastic process with secondary inflammation
- Hypergammaglobulinemia and elevated inflammatory markers
- Imaging alone is of limited value in differentiating from IPT from other splenic lesions
- Nonspecific enhancement pattern because of amount of fibrosis and cellular infiltrations
- Typically a large, well circumscribed single mass in spleen







coronal

sagittal

coronal



sagittal



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3 min



10 min

HBP

In phase

Out of phase

Malignant

- Metastasis
- Lymphoma
- Angiosarcoma
- Undifferentiated carcinoma

Metastases

- Melanoma, cancer of breast, lung, ovary, stomach, and prostate
- Typically hypoechoic on US and hypodense on CT
- MR: T1 low, T2 high, diffusion restriction, various enhancement
- Capsular deposition from most commonly with gynecologic malignancy

Angiosarcoma metastases



Metastasis from intrahepatic cholangiocarcinoma



CT early a.

















ADC

Lymphoma

- Mc malignant tumor
- Secondary involvement: Hodgkin lymphoma(33%) Non-Hodgkin lymphoma(30-40%)
- Diffuse infiltration with splenomegaly, or discrete nodules or masses
- Necrosis is rare, infarction can occur
- US(hypoechoic), CT(hypodense) MR(T1 low, T2 high)
- Marked FDG avidity on FDG PET/CT

Lymphoma









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coronal

Lymphoma

 Pre











coronal



sagittal







3min



Angiosarcoma

- Most common primary malignant neoplasm of the spleen
- Arising from endothelial lining
- Well defined mass or diffusely infiltrative in appearance
- No association with chemical exposures
- Sx.: LUQ pain, anemia, thrombocytopenia
- US: splenomegaly with a heterogenous echogenic mass
- CT: enlarged spleen with area of lower and high attenuation due to acute hemorrhage or hemosiderin deposits, calcification
- MRI: T1 low , T2 high, heterogenous enhancement with multiple nodular nodular enhancing foci.
- 30% intraperitoneal rupture
- Metastasis: liver > lung, bone, BM, and lymphatic system

Angiosarcoma





coronal



Angiosarcoma





DWI



ADC





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In phase



Out of phase

3min

T1

HBP

Undifferentiated carcinoma

- Pattern-less solid, sheet-like growth of tumor cells
- Aggressive clinical course
- No nest, papillae, glands, trabeculae or spindled pattern
- No squamous or mucinous metaplasia and no/minimal neuroendocrine differentiation
- If area of differentiated component are found, the tumor is called dedifferentiated carcinoma

Undifferentiated carcinoma



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Conclusions

- Cystic lesion and wall calcification are the most likely benign findings, and malignant lesions are common in cases of diffusion restriction, spontaneous rupture, and irregular margin.
- By using up-to-date imaging techniques, it is possible to minimize unnecessary surgery or invasive procedure because of the characteristic imaging findings of focal splenic lesions.