National Education Curriculum
Specialty Curricula

Abdomen and Superficial Structures Including Introductory Pediatric and Musculoskeletal
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Section I: Biliary

Rationale: The sonographer is required to perform sonographic studies of the abdomen and superficial structures. Thorough knowledge of relational anatomy, physiology, pathophysiology, sonographic technique and appearances is required. The sonographer will determine normal from artifactual or pathologic conditions, modify or extend the scope of the examination as necessary, prioritize differential diagnoses and complete a technical report.

1. Describe the normal anatomy, function, sonographic technique and appearance of the biliary system
2. Correlate clinical indications and laboratory values associated with biliary disease
3. Identify biliary pathology in terms of sonographic appearances, sequelae and associated pathologies
4. Describe associated pathologies and sequelae relative to biliary disease
5. Identify normal and abnormal flow characteristics and waveforms

I. Biliary

A. Embryology

B. Anatomy

1. Gallbladder
   a. Intraperitoneal
   b. Three segments
      i) Fundus
      ii) Body
      iii) Neck
   c. Layers
      i) Epithelial
      ii) Muscular
      iii) Subserous
      iv) Serosal
   d. Relational anatomy

2. Biliary Ducts
   a. Intrahepatic
   b. Extrahepatic
      i) Common Hepatic Duct (CHD)
      ii) Cystic Duct
      iii) Common Bile Duct (CBD)

C. Variants of Normal and Congenital Anomalies

1. Gallbladder
   a. Hartman’s pouch
   b. Junctional fold
   c. Phrygian cap
   d. Septated
e. Agenesis
f. Duplication
g. Hourglass shape
h. Intrahepatic
i. Left-sided with situs inversus

D. Function
1. Gallbladder
   a. Concentrates and stores and releases bile upon the digestion of a meal
2. Biliary Ducts
   a. Transport bile
3. Laboratory Values
   a. White Blood Count (WBC)
   b. Serum bilirubin; direct, indirect and total
   c. Serum alkaline phosphatase (ALP)
   d. Liver Function Tests (LFTs)
      i) Aspartate Aminotransferase (AST)
      ii) Alanine Aminotransferase (ALT)
      iii) Prothrombin Time (PT)
      iv) Lactic Acid Dehydrogenase (LDH)

E. Indications for Examination
1. Pain
   a. Right upper quadrant (RUQ) pain, epigastric pain
   b. Right shoulder pain (with inflammation)
   c. Post prandial pain
2. Positive Murphy’s Sign
3. Nausea
4. Vomiting
5. Intolerance to Fatty Foods and Dairy Products
6. Jaundice

F. Technique
1. Patient Preparation
2. Transducer Selection
3. Patient Positioning
4. Image Optimization
5. Examination Protocol
6. Examination Limitations

G. Sonographic Findings
1. Gallbladder
   a. Anechoic lumen
   b. Thin, smooth walls
   c. Pear shape
2. Biliary Ducts
   a. Measurement
      i) Location of measurement
      ii) Anatomic variants of CBD which may confuse location of measurement
      iii) Normal diameters
         • Variation
H. Pathophysiology
1. Cholelithiasis
   a. Definition
   b. Incidence
   c. Risk factors
   d. Clinical presentation
      i) May be asymptomatic
      ii) Right upper quadrant (RUQ) pain
      iii) Nausea
      iv) Vomiting
      v) Fever
   e. Sonographic findings
      i) Acoustic shadow
      ii) Gravity dependent gallstone movement with change in body position
      iii) Floating stones secondary to an increase in the specific gravity/viscosity of the bile
   f. Correlative and/or prior imaging
   g. Treatment
2. Mirizzi Syndrome
   a. Definition
      i) Impacted stone in cystic duct, cystic duct remnant, or gallbladder neck
      ii) Partial mechanical obstruction of the CHD by compression of inflammatory reaction around impacted stone
      iii) Sequelae of jaundice, recurrent cholangitis, formation of biliary fistulas, or cholangitis cirrhosis
   b. Incidence
   c. Risk factors
   d. Clinical presentation
   e. Sonographic findings
i) 2-D
   • Pattern of ductal dilatation
ii) Doppler for vessel differentiation
f. Correlative and/or prior imaging
g. Treatment

3. Contracted or non-visualization of gallbladder
   a. Definition
   b. Incidence
   c. Risk factors
d. Clinical presentation
e. Sonographic findings
   i) Dependent on etiology
      • The “double arc” sign, or “WES” sign (W=wall, E=Echo [stone], S=Shadow)
f. Correlative and/or prior imaging
g. Treatment

4. Sludge
   a. Definition
   b. Incidence
   c. Risk factors
d. Clinical presentation
e. Sonographic findings
   i) Gravity dependent, non-shadowing echoes
   ii) Fluid-fluid level
   iii) Gallbladder isoechoic with liver when sludge fills gallbladder lumen
   iv) Atypical appearances
      • Polypoid mass or pseudotumor
      • Sludge balls
   v) Correlative and/or prior imaging
   vi) Treatment

5. Acute cholecystitis
   a. Definition
      i) Acute or chronic inflammation of gallbladder
   b. Incidence
c. Risk factors
d. Clinical presentation
   i) RUQ pain radiating to shoulder
   ii) Biliary colic
iii) Palpable RUQ mass
iv) Tender right subcostal region
v) Fever
vi) Nausea
vii) Vomiting
e. Laboratory findings
i) Elevated serum bilirubin
ii) Elevated serum transaminase
iii) Elevated alkaline phosphatase
iv) Elevated WBC
f. Sonographic findings
i) 2-D
   • Cholelithiasis
   • Diffuse gallbladder wall thickening
   • “Halo-sign” suggesting subserosal edema
   • Distended gallbladder
   • Positive sonographic Murphy’s sign
   • Sludge may be present
   • Pericholecystic fluid
ii) Doppler
   • Cystic artery and gallbladder wall flow
g. Complications
i) Empyema
ii) Emphysematous cholecystitis
iii) Gangrenous cholecystitis
iv) Perforation of the gallbladder
v) Ascending cholangitis
vi) Liver abscess
h. Correlative and/or prior imaging
i. Treatment
6. Acalculous cholecystitis
a. Definition
   i) Inflammation of gallbladder in absence of choledolithiasis
b. Incidence
c. Risk factors
d. Clinical presentation
e. Sonographic findings
i) Distended gallbladder
ii) Wall thickening
iii) Sludge
iv) Pericholecystic fluid and/or edema
f. Correlative and/or prior imaging
g. Treatment

7. Chronic cholecystitis
   a. Definition
      i) Recurrent inflammatory changes in gallbladder secondary to infection, obstruction, or metabolic disorders
   b. Incidence
c. Risk factors
d. Clinical presentation
   i) Intermittent RUQ and epigastric pain; may radiate to scapula
ii) Intolerance to fatty and fried foods
iii) Intermittent nausea and vomiting
e. Laboratory findings
   i) Elevated AST
ii) Elevated ALT
iii) Elevated ALP
   iv) Elevated WBC
f. Sonographic findings
   i) Contracted gallbladder with acoustic shadowing from cholelithiasis
   ii) Thick, echogenic gallbladder wall
   iii) Sludge may be present
   iv) Non-contractility or decreased response to cholecystokinin
g. Correlative and/or prior imaging
h. Treatment

8. Complications of surgical treatment of cholecystitis
   a. Laparoscopic versus open treatment
   b. Incidence
c. Risk factors
d. Clinical presentation
   i) Fever
   ii) New or increasing abdominal pain
   iii) Increased bile drainage
   iv) Decreasing hematocrit
e. Sonographic findings
   i) Bile leak, biloma, choleperitoneum
   ii) Subhepatic abscess
   iii) Hemorrhage
   iv) Retained stone
f. Correlative imaging
g. Treatment

9. Hydrops of the gallbladder (mucoccele)
a. Definition
   i) Abnormal accumulation of fluid within gallbladder
b. Incidence
c. Risk factors
d. Clinical presentation
   i) Asymptomatic
   ii) Palpable RUQ mass
   iii) Epigastric pain
   iv) Nausea
   v) Vomiting
e. Sonographic findings
   i) Markedly dilated gallbladder
   ii) Thin gallbladder wall
   iii) Possible stone in gallbladder neck or cystic duct
f. Correlative and/or prior imaging
g. Treatment

10. Courvoisier’s gallbladder
a. Definition
   i) Prolonged complete obstruction of the distal bile duct
b. Incidence
c. Risk factors
d. Clinical presentation
   i) Possibly asymptomatic
   ii) Palpable RUQ mass
e. Sonographic findings
   i) Markedly dilated gallbladder
   ii) Thin walls
f. Correlative and/or prior imaging
g. Treatment
11. Porcelain gallbladder
   a. Definition
      i) Rare manifestation of chronic cholecystitis
      ii) Calcification of gallbladder wall
   b. Incidence
   c. Risk factors
   d. Clinical presentation
      i) Stones
      ii) Female predominance
      iii) Associated with increased incidence of gallbladder carcinoma
   e. Sonographic findings
      i) Hyperechoic semilunar structure with posterior acoustic shadow
      ii) May simulate stone-filled gallbladder devoid of bile
      iii) Produces biconvex, curvilinear, echogenic structure with variable acoustic shadowing
   f. Correlative and/or prior imaging
   g. Treatment

12. Benign neoplasms
   a. Adenoma
      i) Definition
         • Benign epithelial tumor
      ii) Incidence
      iii) Clinical presentation
         • Asymptomatic
      iv) Sonographic findings
         • Typically small
         • Polypoid, soft-tissue masses pedunculating from gallbladder wall protruding into the gallbladder lumen
         • No acoustic shadow
         • Not gravity dependent
         • Typical location is near fundus of gallbladder
         • Doppler for any flow in stalk or mass
      v) Correlative and/or prior imaging
      vi) Treatment
   b. Adenomyomatosis
      i) Definition
         • A form of hyperplastic cholecystosis
• Proliferation of the surface epithelium of gallbladder wall with gland-like formation and diverticula of mucosa into or through a thickened muscular layer

ii) Incidence

iii) Clinical presentation

iv) Sonographic findings
• Diffuse or segmental thickening of gallbladder wall
• Intraluminal diverticula
• Cholesterol stones/crystals lodged between diverticula (Rokitansky-Aschoff sinuses)
• Comet-tail artifact

v) Correlative and/or prior imaging

vi) Treatment

c. Cholesterolosis
i) Definition
• A form of hyperplastic cholecystosis
• Types
  o Cholesterolosis (strawberry gallbladder)
  o Cholesterol polyps

ii) Incidence

iii) Risk factors

iv) Clinical presentation

v) Sonographic findings
• Non-shadowing single or multiple fixed small echodense masses projecting into lumen of gallbladder

13. Malignant neoplasms
a. Primary gallbladder carcinoma
i) Definition

ii) Incidence

iii) Risk factors

iv) Clinical presentation
• May be asymptomatic
• Loss of appetite, anorexia
• Nausea
• Vomiting
• Intolerance to fatty foods
• Belching
• Jaundice in later stage
v) **Sonographic findings**
- Wall thickening; focal or diffuse
- Intraluminal mass
- Tumor extension to liver bed
- Regional lymphadenopathy
- Ascites

vi) **Correlative and/or prior imaging**

vii) **Treatment**

b. **Metastatic disease to the gallbladder**

i) **Definition**

ii) **Incidence**

iii) **Risk factors**

iv) **Clinical presentation**

v) **Sonographic findings**
- Focal thickening of gallbladder wall
- Intraluminal mass without shadowing
- Polypoid or irregular intraluminal mass
- Indistinct walls

vi) **Correlative and/or prior imaging**

vii) **Treatment**

14. **Non-inflammatory causes of gallbladder wall thickening**

a. **Definition**

b. **Incidence**

c. **Risk factors**

d. **Clinical presentation**

e. **Sonographic findings**

i) **Diffuse thickening**
- Hypoalbuminemia – hypoproteinemia
- Ascites
- Acute hepatitis
- Congestive heart failure
- Renal disease
- AIDS
- Pancreatitis
- Cirrhosis
• Portal node lymphatic obstruction
• Sepsis
  ii) Focal thickening
  iii) Adenomyomatosis
  iv) Adenomatous polyp
  v) Cholesterol polyp
  vi) Papillary adenomas
  vii) Gallbladder carcinoma
  viii) Metastatic wall masses

I. Bile Ducts Pathophysiology

1. Ductal dilatation
   a. Definition
      i) Extrahepatic
      ii) Intrahepatic
   b. Incidence
   c. Risk factors
   d. Clinical presentation
      i) Jaundice
      ii) Elevated LFT’s
   e. Sonographic findings
      i) 2-D
         • CBD
         • CHD
         • “Parallel channel sign” or “double channel sign”
         • Intrahepatic anechoic tubular structure(s) with stellate branching pattern
         • Acoustic enhancement posterior to dilated ducts
      ii) Doppler
         • Identification of vessel versus duct
   f. Correlative and/or prior imaging
   g. Treatment

2. Choledocholithiasis
   a. Definition
      i) Bile duct stone(s)
   b. Incidence
   c. Risk factors
   d. Clinical presentation
      i) RUQ and or epigastric pain
ii) Jaundice

e. Sonographic findings
   i) Dilated duct proximal to obstruction
   ii) Dilated gallbladder
   iii) Intraductal hyperechoic focus with posterior shadowing

f. Correlative and/or prior imaging

g. Treatment

3. Cholangitis
   a. Definition
      i) Inflammation of bile ducts

   b. Types
      i) Acute bacterial
      ii) Recurrent pyogenic
      iii) HIV cholangiopathy (AIDS Cholangitis)
      iv) Sclerosing

   c. Clinical presentation
      i) Classic: fever, RUQ pain, jaundice

   d. Laboratory values
      i) Elevated bilirubin
      ii) Elevated ALP
      iii) Elevated AST
      iv) Elevated ALT
      v) Elevated WBC

   e. Sonographic findings
      i) Dilated ducts
      ii) Focal strictures
      iii) Thickened duct walls
      iv) Intraductal sludge
      v) Intrahepatic ductal stones - may not shadow
      vi) Hepatic abscess
      vii) Portal hypertension

   f. Correlative and/or prior imaging

   g. Treatment

4. Ascariasis
   a. Definition
      i) Infection by roundworm ascariasis lumbricoides

   b. Incidence
5. Hemobilia
   a. Definition
      i) Blood within biliary tree
   b. Incidence
   c. Risk factors
      i) Interventional procedures
      ii) Infection
      iii) Vascular malformation
      iv) Trauma
      v) Malignancy
   d. Clinical presentation
   e. Sonographic findings
      i) Fluid - low level echoes, gravity dependent, layering
      ii) Clot - varying echogenicity, mobile
   f. Correlative and/or prior imaging
   g. Treatment

6. Pneumobilia
   a. Definition
      i) Air (gas) within biliary tree
   b. Incidence
   c. Risk factors
      i) Biliary intervention
      ii) Emphysematous cholecystitis
      iii) Biliary/GI tract fistula
   d. Clinical presentation
   e. Sonographic findings
i) Non-dependent, hyperechoic linear structures within biliary tree
ii) Dirty shadowing and ring-down artifact
iii) Changing patient position may result in movement of bubbles
f. Correlative and/or prior imaging
g. Treatment

7. Malignant neoplasia
   a. Cholangiocarcinoma
      i) Definition
         - Uncommon neoplasm, typically adenocarcinoma arising from the epithelium of the bile ducts
   b. Risk factors
      i) Biliary stones
      ii) Chronic inflammation
   c. Types - defined by location
      i) Intrahepatic
         - Arise within hepatic parenchyma
      ii) Hilar (Klatskin’s)
         - Located at porta hepatis
      iii) Distal
         - Arising in distal extrahepatic ducts
d. Clinical presentation
   i) Pruritis
   ii) Jaundice
   iii) Weight loss; anorexia
   iv) Fatigue
   v) RUQ pain
   vi) Biliary colic
   vii) Palpable mass
e. Laboratory values
   i) ALP
   ii) Bilirubin-direct
f. Sonographic findings
   i) Biliary dilation (without sonographic evidence of obstruction)
   ii) Non-union of the right and left biliary ducts
   iii) Intraductal polypoid mass
   iv) Irregular soft tissue mass
   v) Soft tissue extension into ducts
vi) Tumor extension to adjacent vasculature
vii) Nodal extension to porta hepatis and preaortic regions

g. Correlative and/or prior imaging
h. Treatment

8. Metastases
a. Definition
b. Risk factors
c. Incidence
d. Clinical presentation
e. Sonographic findings
f. Correlative and/or prior imaging
g. Treatment

J. Congenital Abnormalities of Biliary System

1. Biliary Atresia
a. Definition
   i) Congenital obliteration of one or more of components of bile ducts
b. Risk factors
c. Clinical presentation
   i) Dependent on the extent of the atresia
   ii) Persistent jaundice
   iii) Portal hypertension
   iv) Cirrhosis
   v) Hepatic failure
d. Sonographic findings
   i) 2-D
      • Biliary duct dilation may or may not be present
      • Small gallbladder
      • Variable depending on disease stage
      • Portal hypertension
      • “Triangular Cord” sign
   ii) Doppler
      • Portal hypertension findings
e. Correlative and/or prior imaging
f. Treatment

2. Caroli’s disease
a. Definition
   i) Congenital biliary dilation
b. Risk factors

c. Incidence

d. Clinical presentation
   i) Pain
   ii) Fever
   iii) Intermittent jaundice

e. Sonographic findings
   i) 2-D
      • Multiple cystic structures in liver that communicate with biliary tree
      • Calculi may be present
   ii) Doppler
      • Absence of spectral Doppler or color flow in dilated structures
      • Central dot sign: small portal venous branches surrounded by dilated ducts

3. Choledochal cyst
   a. Definition
   b. Risk factors
   c. Incidence
   d. Clinical presentation
      i) Congenital dilation of common bile duct
      ii) Failure to thrive
      iii) Combination of pain, intermittent jaundice, and palpable mass
      iv) Fever
   e. Sonographic findings
      i) 2-D
         • Large, cystic mass in the porta hepatitis separate from gallbladder
         • Dilated CHD or CBD entering a cystic mass
         • Intrahepatic bile duct dilatation
         • Possible calculi
      ii) Doppler
         • Absence of spectral Doppler or color flow in dilated structures
   f. Correlative and/or prior imaging
   g. Treatment
Section II: Liver

1. Describe the normal anatomy and variants, function, sonographic technique and normal sonographic appearances of the liver
2. Evaluate the clinical indication(s) of the study and laboratory values associated with the liver pathology
3. Discuss the sonographic appearances of the various liver pathologies
4. Describe the complications of different liver pathologies and their sonographic appearance
5. Differentiate normal and abnormal flow characteristics and waveforms in the portal and hepatic venous systems and hepatic arterial system

II. Liver

A. Embryology

B. Anatomy

1. Micro
   a. Lobule

2. Macro
   a. Peritoneal
      i) Exception: bare area
   b. Glisson’s capsule
   c. Right upper quadrant (RUQ)
   d. Right and left hemiliver and sections
   e. Lobes and segments
      i) Right
         • Anterior
         • Posterior
      ii) Left
         • Medial
         • Lateral
   iii) Caudate
         • Caudate process
         • Papillary process
   f. Couinaud’s anatomy
      i) Segment I: caudate
      ii) Segment II: lateral segment of left lobe (superior)
      iii) Segment III: lateral segment of left lobe (inferior)
      iv) Segment IV: medial segment of the left lobe (quadrate)
      v) Segment V: anterior segment of the right lobe (inferior)
      vi) Segment VI: posterior segment of the right lobe (inferior)
      vii) Segment VII: posterior segment of the right lobe (superior)
viii) Segment VIII: anterior segment of the right lobe (superior)

g. Fissures
   i) Main lobar/interlobar fissure
   ii) Right intersegmental
   iii) Left intersegmental
   iv) Fissure for ligamentum venosum

h. Ligaments
   i) Coronary
   ii) Falciform
   iii) Gastrohepatic
   iv) Hepatoduodenal
   v) Left triangular (lateral)
   vi) Right triangular (lateral)
   vii) Round (ligamentum teres)

i. Vasculature
   i) Hepatic artery
      • Common
      • Proper
      • Right
      • Left
   ii) Hepatic veins
      • Left
      • Middle
      • Right
   iii) Portal veins
      • Main
      • Left
      • Right
      • Branches and tributaries
      • Collateral pathways
   iv) Normal variants

j. Relational anatomy

C. Variants
   1. Riedel’s lobe
   2. Situs inversus
   3. Diaphragmatic indentations (pseudofissures)
D. Functions
1. Formation of bile
2. Activity of reticuloendothelial tissues
3. Metabolism
4. Storage
5. Blood reservoir
6. Heat production
7. Detoxification

E. Laboratory Values for Liver Function (LFTs)
1. Aspartate aminotransferase (AST)
2. Alanine aminotransferase (ALT)
3. Bilirubin
   a. Direct (conjugated)
   b. Indirect (unconjugated/free)
4. Serum protein
5. Alkaline phosphatase (ALP)
6. Lactic dehydrogenase (LDH)
7. Prothrombin time (PT)
8. Alpha-fetal protein (AFP)

F. Indications for Examination
1. Suspicion for stenosis, thrombosis of the hepatic vasculature
2. Screening (chronic hepatitis/cirrhosis) for hepatocellular carcinoma
3. Fatigue and weakness
4. Abdominal discomfort
5. Fever
6. Abnormal liver function tests (LFTs)
7. Jaundice
8. Hepatomegaly
9. Others

G. Technique
1. Patient preparation
2. Transducer selection
3. Patient positioning
4. Image optimization
5. Image protocol

H. Normal Sonographic Findings
1. 2-D
2. Doppler
   a. Portal veins (hepatopetal, mild pulsatility)
   b. Hepatic veins (pulsatile)
   c. Hepatic arteries (low resistance forward flow)

I. Pathophysiology
   1. Diffuse disease
      a. Fatty infiltration
         i) Definition
            • Acquired, reversible condition resulting in accumulation/deposition of fatty triglycerides within liver cells
         ii) Clinical presentation
            • Asymptomatic
            • Hepatomegaly
            • Mildly elevated LFTs
         iii) Sonographic findings
            • Normal size or hepatomegaly
            • Diffuse or focal increase in liver parenchymal echogenicity
               • Focal sparing
            • Impaired visualization of hepatic vessels
         iv) Correlative and/or prior imaging
      b. Hepatitis
         i) Definition
            • Inflammation of the liver
         ii) Clinical presentation
            • Acute
               • Elevated LFTs
               • Jaundice
               • Hepatomegaly
               • Right upper quadrant pain
               • Nausea and vomiting
               • Fever
               • Other
            • Chronic
               • Variable and dependent on length and severity
iii) Sonographic findings

- Acute hepatitis
  - 2-D
    ~ Normal
    ~ Decrease in liver echogenicity
    ~ Portal vein (PV) walls may appear more echogenic relative to the liver parenchyma, “starry sky” appearance
    ~ Hepatomegaly
    ~ Associated gallbladder wall thickening

- Chronic hepatitis
  - 2-D
    ~ Increased liver echogenicity may be present
    ~ Coarse echo texture
    ~ Decreased echogenicity of portal vein walls
    ~ Portal hypertension and sequelae
    ~ Associated findings of cirrhosis
      - Doppler
        - Portal hypertension and sequelae

iv) Correlative and/or prior imaging
v) Treatment

c. Cirrhosis

i) Definition

- Irreversible and often progressive fibrosis, scarring, parenchymal necrosis, and nodular regeneration

ii) Clinical presentation

- Abnormal LFTs
- Jaundice
- Hepatomegaly
- Portal hypertension and sequelae
- Other

iii) Sonographic findings

- Early stage
  - Hepatomegaly
  - Fatty infiltration may be present
  - Surface nodularity

- Later stage
  - Smaller heterogeneous liver (anterior right and medial left sections)
o Relative caudate lobe enlargement with possible IVC compression
o Coarse liver texture with surface nodularity

• Portal hypertension and sequelae
• Ascites
• Associated hepatocellular carcinoma (HCC)
• Doppler findings

iv) Correlative and/or prior imaging
v) Treatment

2. Vascular abnormalities

a. Portal hypertension (PH)

i) Definition

• Increased pressure within the portal venous system

• Types
  o Presinusoidal
    ~ Extrahepatic presinusoidal
    ~ Intrahepatic presinusoidal
    ~ Extrahepatic postsinusoidal
  o Intrahepatic

• Clinical presentation
• Sonographic findings
  o 2-D
    ~ Increased portal vein diameter
    ~ Ascites
    ~ May have splenomegaly
  o Doppler
    ~ Hepatopetal or slow hepatofugal flow
    ~ Low velocity continuous flow
    ~ Bidirectional flow
    ~ Recanalized umbilical vein

• Correlative and/or prior imaging
• Treatment
  o Intrahepatic portosystemic venous shunt
    ~ Definition
    ~ Surgical
    ~ Transjugular intrahepatic portosystemic shunts (TIPS)
  o Sonographic findings
~ 2-D
A. Echogenic stent - most commonly placed between right portal vein and right hepatic vein
~ Doppler
A. Evaluate for stent patency and direction
B. Obtain stent and portal vein velocities, determine flow direction in the right and left portal vein branches
C. Left portal vein flow is hepatofugal, towards stent
D. Compare velocities to post placement velocities

b. Portal vein thrombosis
i) Definition
ii) Clinical presentation
iii) Sonographic findings
   • 2-D and Doppler
iv) Correlative and/or prior imaging

c. Portal vein varix, hepatic artery aneurysm
i) Definition
ii) Clinical presentation
iii) Sonographic findings
   • 2-D and Doppler
iv) Correlative and/or prior imaging
v) Treatment

d. Budd-Chiari syndrome
i) Definition
   • Obstruction of hepatic vein(s) outflow
      o May or may not involve IVC
   • Etiology (primary or secondary)
      o Congenital
      o Obstructing membranes, webs
      o Tumor extension from primary hepatic carcinoma
      o Polycythemia vera
ii) Clinical presentation
   • Right upper quadrant pain
   • Ascites
   • Hepatomegaly
iii) Sonographic findings
   • 2-D
3. Glycogen storage disease (Type I)
   a. Definition
   b. Clinical presentation
   c. Sonographic findings
   d. Correlative and/or prior imaging
   e. Treatment

4. General characteristics of focal versus diffuse liver abnormalities
   a. Sonographic findings
      i) Intrahepatic versus extrahepatic

5. Cysts
   a. Definition
      i) Fluid-filled mass with epithelial lining
   b. True liver cysts versus acquired
   c. Autosomal dominant polycystic liver disease
   d. Clinical presentation
   e. Sonographic findings
   f. Correlative and/or prior imaging
   g. Treatment

J. Infections
   1. Pyogenic (bacterial) abscess
      a. Definition
         i) Localized collection of pus
      b. Clinical presentation
         i) Fever (varied)
         ii) Pain
         iii) Pleuritic pain
         iv) Nausea
         v) Vomiting
         vi) Diarrhea
vii) Leukocytosis
viii) Elevated liver function tests (LFTs)
ix) Anemia
x) Other
c. Sonographic findings
i) Commonly located in right lobe
ii) Typically solitary
iii) Typical round or ovoid shape
iv) Irregular, poorly defined walls
v) Variable echogenicity, may appear solid
vi) Possible acoustic enhancement
vii) Microbubbles, air
viii) Calcification, if healed
d. Correlative and/or prior imaging
e. Treatment
i) Aspiration to determine organisms responsible for abscess
ii) Drainage
iii) Antibiotics

2. Amebic (parasitic) abscess
a. Definition
i) Localized cavity caused by parasite Entamoeba histolytica
b. Clinical presentation
i) Abdominal pain
ii) Diarrhea
iii) Melen
iv) Leukocytosis
v) Typically abnormal LFTs
c. Sonographic findings
i) Echogenicity varies over time
ii) Typical peripheral location contiguous with liver capsule
iii) Lack of significant wall echoes
iv) Round or oval configuration
v) Distal acoustic enhancement
d. Correlative and/or prior imaging

3. Fungal infection
a. Definition
i) Candidiasis (c. albicans)
b. Clinical presentation
   i) Increased risk for immunocompromised patients

c. Sonographic findings
   i) Possible hepatomegaly
   ii) “Wheel within a Wheel” type lesions early
   iii) Multiple “bull’s-eye” target lesions
   iv) Uniform hypoechoic lesions
   v) Echogenic foci with variable posterior acoustic shadowing in late stage

d. Correlative and/or prior imaging

e. Treatment

4. Schistosomiasis
   a. Definition
      i) Granulomatous portal vein reaction to parasitic invasion
      ii) Common parasitic infection associated with contaminated water

   b. Clinical presentation
      i) Pain
      ii) Fever
      iii) Abnormal WBC and LFT laboratory values
      iv) Presinusoidal portal hypertension

c. Sonographic findings
   i) 2-D
      • Enlarged portal vein tracts with echogenic lumen
      • Hepatomegaly in acute phase
      • Decreased size of liver with progression
   ii) Doppler
      • Evaluate for patency of portal vein

d. Correlative and/or prior imaging

e. Treatment

5. Echinococcal cyst
   a. Definition
      i) Hydatid disease
      ii) Parasitic infection

   b. Clinical presentation
      i) Abdominal pain
      ii) Fever
      iii) WBC and LFT

   c. Sonographic findings
Abdomen and Superficial Structures Including Introductory Pediatric and Musculoskeletal

i) Variable in appearance
   - Simple cyst(s) with fine textured particles within
   - Floating membranes, water lily sign
   - Cysts with daughter cyst appearance
   - Densely calcified mass
d. Correlative and/or prior imaging
e. Treatment

K. Benign Neoplasma
1. Adenoma
  a. Definition
     i) Benign neoplasm of normal or slightly atypical hepatocytes frequently containing areas of bile stasis and focal hemorrhage or necrosis
  b. Incidence
     i) Less common
c. Risk factors
   i) Oral contraceptives
   ii) Anabolic steroids
   iii) Associated with Type I glycogen storage disease
d. Clinical presentation
   i) Asymptomatic
   ii) Palpable mass
   iii) Severe right upper quadrant pain with bleeding or infarct within lesion
   iv) Shock due to rupture of lesion and hemoperitoneum
e. Sonographic findings
   i) 2-D
      • Mass of variable echogenicity
      • Similar to focal nodular hyperplasia, heterogeneous if intratumoral bleeding
      • Usually seen in right lobe
   ii) Doppler
      • Peritumoral flow
      • Intratumoral flow
f. Correlative and/or prior imaging
g. Treatment
2. Cavernous hemangioma
   a. Definition
      i) Congenital vascular malformation; consists of large network of vascular spaces
   b. Incidence
i) Most common benign tumor of liver, may be multiple

c. Risk factor
   i) Female predominance

d. Clinical presentation
   i) Usually asymptomatic
   ii) Large lesions can be associated with hemorrhage

e. Sonographic findings
   i) Usually round, echogenic, homogeneous, well defined
   ii) Larger lesions may be more complex
   iii) More common in right lobe
   iv) Usually solitary
   v) May demonstrate acoustic enhancement

f. Correlative and/or prior imaging

g. Treatment

3. Focal nodular hyperplasia (FNH)

a. Definition
   i) Rare, benign neoplasm composed of abnormally arranged hepatocytes, Kupffer cells, bile duct elements, and fibrous connective tissue

b. Incidence
   i) Female predominance
   ii) Less than 40 years of age

c. Risk factors
   i) Oral contraceptives

d. Clinical presentation
   i) Usually asymptomatic

e. Sonographic findings
   i) 2-D
      • Variable echogenicity, homogeneous
         o Isoechoic most common
      • Prevalence to right lobe
   ii) Doppler
      • Hypervascular with a stellate arterial pattern

f. Correlative and/or prior imaging

g. Treatment

4. Hemangioendothelioma

a. Definition
   i) Benign vascular liver tumor
b. Incidence  
i) Most common benign vascular liver tumor in infancy  
c. Risk factors  
d. Clinical presentation  
i) Hepatomegaly  
ii) Congestive heart failure  
iii) Cutaneous hemangioma  
e. Sonographic findings  
i) 2-D  
• Variable echogenicity  
• Hepatomegaly  
ii) Doppler  
• Increased flow in mass  
f. Correlative and/or prior imaging  
g. Treatment  
i) May regress spontaneously  

L. Malignant Neoplasms  
1. Hepatoblastoma  
a. Definition  
i) Germ cell tumor  
b. Incidence  
i) Most common malignant liver tumor in childhood  
c. Risk factors  
i) Beckwith-Wiedemann syndrome  
ii) Hemihypertrophy  
iii) Sporadic aniridea  
d. Clinical presentation  
i) Increased girth  
ii) Hepatomegaly  
iii) Weight loss, nausea, and vomiting  
iv) Precocious puberty  
v) Elevated serum alpha-fetoprotein (AFP)  
e. Sonographic findings  
i) 2-D  
• Heterogeneous or complex  
• Poorly marginated  
• Calcifications may be identified
ii) Doppler
   - Neovascular flow evaluation
f. Correlative and/or prior imaging
g. Treatment
2. Hepatocellular carcinoma (HCC) – hepatoma
   a. Definition
      i) Primary liver malignancy
   b. Incidence
      i) Most common primary malignancy
c. Risk factors
   i) Cirrhosis
   ii) Hepatocarcinogens in food
   iii) Chronic Hepatitis B and Hepatitis C
   iv) Metabolic disorders
d. Clinical presentation
   i) Palpable mass
   ii) Rapid liver enlargement
   iii) Unexplained mild fever
   iv) Symptoms associated with cirrhosis
   v) Elevated serum α-fetoprotein
e. Sonographic findings
   i) Variable in pattern
      - Solitary
      - Multiple
      - Dominant mass with satellite lesions
      - Diffuse parenchymal infiltration
   ii) Variable echogenicity
   iii) Hepatomegaly
   iv) Associated ascites
   v) Portal/hepatic vein invasion
   vi) Occasional bile duct invasion
f. Correlative and/or prior imaging
g. Treatment
3. Hemangiosarcoma
   a. Definition
      i) Malignant neoplasm comprised of endothelial and fibroblastic tissue and surrounds vessels
b. Incidence
   i) Rare

c. Risk factors
   i) Advanced age
   ii) Exposure to carcinogens

d. Clinical presentation

e. Sonographic findings
   i) Large mass
   ii) Mixed echogenicity

f. Correlative and/or prior imaging

g. Treatment

4. Metastasis
   a. Definition
      i) Dissemination of tumor cells to liver from primary malignant neoplasm

   b. Incidence
      i) Common site for metastatic change
      ii) Most common liver malignancy

   c. Risk factors
      i) Common primary carcinomas to metastasize to liver
         - Gastrointestinal
         - Breast
         - Lung

   d. Clinical presentation
      i) Hepatomegaly
      ii) Jaundice
      iii) Pain
      iv) Weight loss
      v) Abnormal laboratory tests

   e. Sonographic findings
      i) Variable in pattern
         - Hypoechoic
         - Hyperechoic
         - Target lesion
         - Diffusely heterogeneous
         - Calcifications
         - Cystic degeneration

   f. Correlative and/or prior imaging
g. Treatment

M. Trauma

1. Definition
   a. Hematoma may be due to trauma, post biopsy or rupture of a neoplasm such as an adenoma, metastatic choriocarcinoma, or giant cavernous hemangioma, hepatocellular carcinoma, angiosarcoma

2. Categories
   a. Rupture of the liver and capsule
   b. Separation of the capsule and the subcapsular hematoma
   c. Central rupture of the liver

3. Clinical presentation
   a. Pain
   b. Hypotension
   c. Possible decreased hematocrit

4. Sonographic findings
   a. Typically posterior segment of right lobe
   b. Varied appearance depending on the age of the hematoma
   c. Subcapsular - potential fluid bordering the liver
   d. Capsular rupture - free fluid in peritoneal cavity

N. Benign Causes of Hyperechoic Foci in the Liver Region

1. Calcification
   a. Old granulomatous disease
   b. Healed abscess or hematoma
   c. Posterior acoustic shadowing

2. Air in bile ducts
   a. Post-operative biliary surgery: endoscopic retrograde cholangiopancreatography (ERCP) with sphincterotomy, patulous sphincter of Oddi, biliary-enteric fistula
   b. Cause “comet-tail,” ring-down artifact
   c. May move with change in patient position
Section III: Pancreas

1. Describe the normal anatomy, function, sonographic technique and appearance of the pancreas
2. Correlate clinical indications and laboratory values associated with pancreatic disease
3. Discuss pancreatic pathology in terms of sonographic appearances, sequelae and associated pathologies

III. Pancreas

A. Embryology
   1. Ventral and dorsal buds

B. Anatomy
   1. Retroperitoneal
   2. Head
   3. Neck
   4. Uncinate
   5. Body
   6. Tail
   7. Vascular landmarks
   8. Size
   9. Ducts
   10. Shape

C. Relational Anatomy

D. Normal Variants
   1. Annular Pancreas
   2. Ectopic Pancreatic Tissue
   3. Pancreatic Divisum

E. Function
   1. Exocrine Function
      a. Secretions
   2. Hormone Secretion
      a. Hormones
         i) Cholecystokinin
         ii) Gastrin
         iii) Secretin
   3. Endocrine function
      a. Islet cells of Langerhans
         i) Alpha cells secrete glucagon
         ii) Beta cells secrete insulin
iii) Delta cells secrete somatostatin

F. Indications for Examination
1. Severe Epigastric Pain
2. Abdominal Pain Radiating to Back
3. Abdominal Distension with Hypoactive Bowel Sounds
4. Acute Pancreatitis
5. Chronic Pancreatitis
6. Biliary Disease
7. Abnormal Pancreatic Enzymes
8. Pancreatic Neoplasm

G. Laboratory Values
1. Amylase
   a. Serum
   b. Urine
2. Glucose
3. Lipase

H. Normal Sonographic Findings
1. Homogeneous, Coarse Echotexture
2. Echogenicity Should Be Compared to Liver
3. Size Varies with Shape

I. Technique
1. Patient Preparation
2. Transducer Selection
3. Patient Positioning
4. Image Optimization
5. Image Protocol

J. Pathophysiology
1. Congenital Diseases
   a. Cystic fibrosis
      i) Definition
         • Autonomic recessive exocrine gland disorder
         • Distention of ductules and acini with degeneration and replacement by small cysts
      ii) Incidence
      iii) Risk factors
      iv) Clinical presentation
      v) Sonographic findings
Poor visualization
- Hyperechoic gland compared to normal
- Associated liver changes
- Associated gastrointestinal (GI) tract change
- Associated gallbladder changes

vi) Correlative and/or prior imaging
vii) Treatment

2. Inflammation
a. Acute pancreatitis
i) Definition
ii) Incidence
iii) Risk factors
- Cholelithiasis
- Pancreatic hypersecretions
- Alcohol intake
- Duodenal reflux
- Blunt abdominal trauma
- Surgery and/or interventional procedures
- Metabolic disorders
- Viral and parasitic infections
- Drug exposure
- Pancreatic carcinoma
- Hereditary (idiopathic pancreatitis)

iv) Clinical presentation
- Severe pain
- Constant and intense pain radiating to back
- Fever
- Sweating
- Nausea
- Vomiting
- Gaseous distention
- Ileus
- Acute respiratory distress syndrome (ARDS)
- Acute tubular necrosis

v) Laboratory values
- Serum and urine amylase
vi) Sonographic findings
- Normal
- Diffuse enlargement with loss of normal sonographic texture
- Focal enlargement
- Echogenicity as compared to liver
- Poor visualization of the splenic vein
- Dilatation or compression of pancreatic duct

vii) Complications
- Pseudocyst formation
- Phlegmon
- Abscess
- Hemorrhage
- Biliary and duodenal obstruction
- Fluid collections
- Edema
- Acute peritonitis
- Thrombosis of the portal-splenic veins and SMV
- Left pleural effusion

viii) Correlative and/or prior imaging

ix) Treatment

3. Hemorrhagic pancreatitis
   a. Definition
   b. Risk factors
   c. Incidence
   d. Clinical presentation
      i) Severe abdominal pain
      ii) Nausea
      iii) Vomiting
      iv) Abdominal distention and varying degrees of ileus
   e. Complications
   f. Laboratory findings
      i) Increased serum amylase and serum lipase
      ii) Decreased hematocrit and serum calcium
g. Sonographic findings
   i) Acute, initial changes with progression of disease
h. Correlative and/or prior imaging
i. Treatment
4. Phlegmonous pancreatitis
   a. Definition
      i) Severe complication of acute pancreatitis
   b. Incidence
c. Risk factors
d. Clinical presentation
e. Sonographic findings
   i) Hypoechoic
f. Correlative and/or prior imaging
g. Treatment
5. Abscess
   a. Definition
   b. Incidence
c. Risk factors
d. Clinical presentation
e. Sonographic findings
   i) Smooth walls with little or no internal echoes
   ii) Irregular walls with increased internal echoes
   iii) Ranges from completely echo free to echo dense
f. Correlative and/or prior imaging
g. Treatment
6. Pseudocyst
   a. Definition
      i) Fluid collections that arise from inflammatory processes, necrosis, or hemorrhage
   b. Incidence
      i) Complication of pancreatitis or trauma
c. Risk factors
d. Clinical presentation
e. Sonographic findings
   i) Usually anechoic with acoustic enhancement
   ii) Variable
      • Complex
      • Debris with the cystic component
7. Chronic pancreatitis
   a. Definition
      i) Relapsing pancreatitis
      ii) Irreversible damage to the gland due to progressive destruction
   b. Incidence
      i) Male predominance
   c. Risk factors
      i) Alcoholism
      ii) Hypercalcemia
      iii) Hyperlipidemia
      iv) Medications
   d. Clinical presentation
      i) Persistent epigastric and radiating back pain
      ii) Nausea
      iii) Vomiting
      iv) Flatulence
      v) Weight loss
      vi) Diabetes mellitus (later stages)
   e. Laboratory values
      i) Steatorrhea
      ii) Abnormal glucose-tolerance test
   f. Sonographic findings
      i) 2-D
         • Heterogeneous
         • Increased echogenicity and attenuation
         • Calcifications
         • Ductal dilatation
         • Irregular contour to the gland or pancreatic duct
         • Focal or diffuse enlargement of gland
            o Complications
               ~ Pseudocyst
               ~ Dilatation of the biliary system
               ~ Associated pancreatic carcinoma
ii) Doppler
   - Possible thrombosis of the splenic vein and/or portal vein

  g. Correlative and/or prior imaging
  h. Treatment

K. Neoplasia
   1. Cystadenoma/cystadenocarcinoma
      a. Definition
         i) Microcystic adenoma
         ii) Macrocystic adenoma
      b. Incidence
         i) Female predominance
      c. Risk factors
      d. Clinical presentation
      e. Sonographic findings
         i) Microcystic
         ii) Macrocystic
         iii) Associated findings
      f. Correlative and/or prior imaging
      g. Treatment

   2. Adenocarcinoma
      a. Definition
      b. Incidence
         i) Most common malignancy of pancreas
         ii) Male predominance
      c. Risk factors
      d. Clinical presentation
         i) Late onset of symptoms
         ii) Weight loss
         iii) Painless jaundice
         iv) Abdominal pain with progression of disease
         v) Back pain
         vi) Anorexia
         vii) Nausea
         viii) Vomiting
         ix) Generalized malaise
         x) Weakness
         xi) New onset of diabetes
e. Sonographic findings
   i) 2-D
      • Enlargement
      • Mass with irregular borders
      • Enlarged pancreatic duct
      • Biliary duct dilatation: secondary to involvement of pancreatic head
      • Enlarged gallbladder (Courvoisier’s sign)
      • Metastasis to liver
      • Ascites
      • Regional lymph node involvement with SMA displacement
      • Encasement of mesenteric vessels
      • Other
   ii) Doppler
      • Portal vein
      • Splenic vein
f. Correlative and/or prior imaging
g. Treatment

3. Islet cell tumors
a. Definition
   i) Functional or nonfunctional; either benign or malignant neoplasms
      • Insulinoma
      • Gastrinoma
      • Other
b. Incidence
c. Risk factors
d. Clinical presentation
e. Laboratory values
f. Sonographic findings
   i) Typical locations
   ii) Size
   iii) Associated echogenicities
g. Correlative and/or prior imaging
h. Treatment
Section IV: Renal and Lower Urinary Tract

1. Describe the normal anatomy and variants, function, sonographic technique and normal sonographic appearances of the kidneys and lower urinary tract
2. Correlate the clinical indications and laboratory values associated with urinary tract disease
3. Describe urinary tract pathology in terms of sonographic appearances
4. Describe associated pathologies and sequelae relative to urinary tract disease
5. Identify normal and abnormal flow characteristics and waveforms
6. Discuss current diagnostic Doppler criteria

IV. Renal and Lower Urinary Tract

A. Embryology

B. Anatomy

1. Micro
   a. Nephrons

2. Macro – Kidneys
   a. Paired
   b. Retroperitoneal
   c. Encapsulated
      i) Fibrous renal capsule
         • True capsule
      ii) Adipose capsule
         • Perirenal fat
      iii) Renal fascia
         • Gerota’s

d. Structure
   i) Cortex
   ii) Medulla
   iii) Pyramids
   iv) Calyces
   v) Pelvis

3. Vascular Anatomy
   a. Arterial
      i) Main renal artery
      ii) Segmental
      iii) Interlobar
      iv) Arcuate
      v) Interlobular arteries
vi) Arterioles
b. Venous
   i) Main renal vein
   ii) Segmental
   iii) Interlobar
   iv) Arcuate
   v) Interlobular
4. Macro - Lower Urinary (LU) Tract
   a. Retroperitoneal or extraperitoneal
   b. Ureters
   c. Urinary bladder
      i) Wall layers
         • Serosa
         • Muscle
         • Mucosa
      ii) Trigone
d. Urethra
5. Relational Anatomy
C. Variants
1. Renal Structure
   a. Hypertrophied column of Bertin
   b. Dromedary hump
   c. Junctional parenchymal defect (JPD)
   d. Persistent fetal lobulation (PFL)
   e. Extrarenal pelvis
   f. Horseshoe kidney
2. Renal Number
   a. Bilateral renal agenesis
   b. Unilateral renal agenesis
   c. Supernumery kidney
3. Renal Position
   a. Ectopic
   b. Crossed fused renal ectopia
4. Renal Hypoplasia
5. Renal Malrotation
6. Duplex Collecting System
7. Congenital Renal Dysplasia
a. Dysplastic renal hypoplasia
b. Unilateral multicystic dysplastic kidney

8. Congenital Renal and Medullary Cystic Disease
   a. Infantile polycystic kidney disease (IPCKD), autosomal recessive polycystic kidney disease (ARPKD)
   b. Adult polycystic kidney disease (APCKD), autosomal dominant polycystic kidney disease (ADPKD)
   c. Unilateral multicystic disease
   d. Medullary cystic disease

9. Lower Urinary Tract
   a. Bladder diverticulum
   b. Bladder duplication
   c. Bladder extrophy
   d. Posterior urethral valves (PUV)
   e. Urachal anomalies
   f. Ureterocele

D. Function
   1. Excretion of waste products of metabolism; urine production
   2. Renin-angiotensin system influences blood pressure, blood volume, and intake/excretion of salt and water
   3. Regulation of acid-base balance
   4. Regulation of serum electrolytes
   5. Renal production or erythropoietin
   6. Convey urine to bladder
   7. Reservoir for urine
   8. Convey urine from the body

E. Laboratory Values
   1. Serum Creatinine
   2. Blood Urea Nitrogen (BUN)
   3. Urinalysis

F. Indications
   1. Urinary tract infection (UTI)
   2. Palpable mass
   3. Elevated serum creatinine, BUN levels
   4. Severe flank pain
   5. Hematuria
      a. Proteinuria
   6. Oliguria
7. Azotemia
8. Assessment of mass visualized on radiography, CT scans
9. Assessment of non-visualization of kidney on intravenous pyelogram (IVP)
10. Assessment of enlarged or small kidney
11. Hypertension
12. Acute renal injury
13. Trauma or post-surgical complication

G. Sonographic Technique
1. Patient preparation
2. Transducer selection
3. Patient positioning
4. Imaging optimization
5. Imaging protocol

H. Normal Sonographic Findings
1. Measurements
   a. Length
   b. Width
   c. Thickness
   d. Cortical thickness
   e. Relative renal size
      i) Pediatric
      ii) Solitary kidney
2. Renal cortex
   a. Homogeneous
   b. Low to medium level echogenicity
   c. Hypoechoic or isoechoic to liver
3. Renal medulla
   a. Relatively hypoechoic to cortex
   b. Hyperechoic interfaces at corticomedullary junction (CMJ)
4. Renal sinus
   a. Hyperechoic to renal cortex
5. Doppler
   a. Renal arteries
   b. Renal veins
6. Ureters
   a. Not routinely visualized
   b. Anechoic tubular structure
7. Bladder
   a. Anechoic lumen
   b. Wall thickness varies with degree of distention
   c. Presence of ureteral jets on Doppler

8. Urethra
   a. Anechoic tubular structure

I. Pathophysiology
   1. Inflammatory Conditions
      a. Acute pyelonephritis (APN)
         i) Definition
            • Acute inflammation of the tubulointerstitial renal tissue
         ii) Clinical presentation
            • Flank pain
            • Fever
         iii) Sonographic findings
            • Normal appearance
            • Focal or diffuse
               o Increase in size
               o Areas of increased or decreased echogenicity
               o Absence of perfusion on Doppler
            • Loss of corticomedullary definition
            • Compression of renal sinus
            • Unilateral or bilateral
         iv) Correlative and/or prior imaging
         v) Treatment
      b. Emphysematous pyelonephritis
         i) Definition
            • Life-threatening infection of renal parenchyma with gas formation in tissue or collecting system
         ii) Clinical presentation
            • Fever
            • Flank pain
            • Dehydration
            • Acidosis
            • Electrolyte imbalance
         iii) Sonographic findings
            • Gas within parenchyma
• Presence of air within the collecting system
d) Correlative and/or prior imaging
e) Treatment
c. Chronic pyelonephritis
i) Definition
• Interstitial nephritis resulting from ongoing or recurring urinary tract infections (UTIs)
ii) Clinical presentation
• Asymptomatic
• Renal failure
iii) Sonographic findings
• Unilateral or bilateral focal areas of cortical thinning and increased echogenicity
d) Correlative and/or prior imaging
e) Treatment
d. Acute tubular necrosis (ATN)
i) Definition
• Damage and/or necrosis to the renal tubules bilaterally
ii) Clinical presentation
• Acute kidney injury
• Oliguria
• Uremia
• Electrolyte imbalance
iii) Sonographic findings
• Normal
• Bilateral enlargement
• Increased parenchymal echogenicity
d) Correlative and/or prior imaging
e) Treatment
e. Pyonephrosis
i) Definition
• The presence of pus in the urinary collecting system
ii) Clinical presentation
• Fever
• Flank pain
iii) Sonographic findings
• Hydronephrosis
• Mobile echogenic particles in collecting system

iv) Correlative and/or prior imaging
v) Treatment

f. Acute glomerulonephritis
i) Definition
• Accumulation of inflammatory elements or necrosis of glomeruli

ii) Clinical presentation
• Hematuria
• Hypertension
• Azotemia

iii) Sonographic findings
• Normal to enlarged kidneys
• Varying echogenicity of cortex
• Normal medulla

iv) Correlative and/or prior imaging
v) Treatment

g. Candidiasis
i) Definition
• Systemic fungal infection, occurring in immuno-compromised patient

ii) Clinical presentation
• Worsening of existing symptoms in chronically ill patients

iii) Sonographic findings
• Multiple focal abscesses throughout parenchyma
  • Focal masses of varying echogenicity
  • Acoustic enhancement
• Focal echogenic mass(es) within collecting system

iv) Treatment

h. Schistosomiasis
i) Definition
• Parasitic infestation of schistosome organism

ii) Clinical presentation
• Hematuria

iii) Sonographic findings
• Normal kidneys
• Urinary bladder wall thickening
• Focal bladder wall granuloma
• Calcifications of bladder wall
• Hydrenephrosis

iv) Treatment

i. Urinary tuberculosis

i) Definition
   • Infestation to the kidney of mycobacterium tuberculosis from an extraurinary source

ii) Clinical presentation
   • Asymptomatic
   • Frequency
   • Nocturia
   • Dysuria
   • Gross or microscopic hematuria
   • Sterile pyuria

iii) Sonographic findings
   • Acute
     o Focal lesions of varying echogenicity
   • Chronic
     o Varying appearances of scarring and obstruction resulting from chronic fibrotic changes
     o Decreased size
     o Increased echogenicity with shadowing (urothelial calcification)
     o May be associated ureteral stricture or small bladder volume

iv) Correlative and/or prior imaging
v) Treatment

j. Abscess

i) Definition
   • Focal collection of inflammatory and necrotic debris

ii) Clinical presentation
   • Fever
   • Flank pain
   • Localized tenderness

iii) Sonographic findings
   • Varying echogenicity
   • Acoustic enhancement
   • Shadowing
   • Central fluid with surrounding hyperemia on Doppler
iv) Correlative and/or prior imaging
v) Treatment

2. Acute kidney injury/chronic renal failure (formerly renal failure)
   a. Definition
      i) Impairment of kidney function resulting in inability of kidney(s) to maintain normal function
   b. Types
      i) Prerenal
      ii) Renal
      iii) Postrenal
   c. Clinical presentation
      i) Oliguria
      ii) Increased BUN and creatinine
      iii) Proteinuria
      iv) Anemia
      v) Hypertension
      vi) Uremia
   d. Sonographic findings
      i) Normal
      ii) Hydronephrosis
         • Acute non-obstructive
           o Normal or increased size
           o Increased parenchymal echogenicity
         • Chronic
           o Normal or decreased size
           o Increased parenchymal echogenicity
      iii) Increased size with acute
      iv) End-stage
         • Decreased size
           o Parenchymal thinning
         • Increased parenchymal echogenicity
         • Loss of parenchymal differentiation
   e. Treatment

3. Renal masses
   a. Benign renal cysts
      i) Cortical cyst
         • Definition
Benign fluid-filled mass of unknown pathogenesis

- Complications
  - Hemorrhagic
  - Infection

- Clinical presentation
  - Asymptomatic
  - Pain
  - Palpable mass
  - Fever
  - Elevated WBC

- Sonographic findings
  - Usually cortical in location
  - Simple
    - Anechoic
    - Smooth imperceptible walls
    - Lack of wall calcifications, nodules, or septations
  - Distal acoustic enhancement
  - Complex
    - Varying echogenicity
    - Septations
    - Layering
    - Wall calcifications

- Correlative and/or prior imaging

- Treatment

b. Parapelvic cyst
  i) Definition
    - Fluid-filled mass of lymphatic origin located in renal hilus
  ii) Clinical presentation
    - Asymptomatic
    - Hypertension
    - Hematuria
    - Hydronephrosis
  iii) Sonographic findings
  iv) Renal sinus location
  v) Anechoic
  vi) Well-defined walls
vii) Acoustic enhancement
viii) May have internal echoes
ix) Correlative and/or prior imaging
x) Treatment

4. Cystic disease
   a. Multicystic dysplastic kidney disease (MDKD)
      i) Definition
         • Developmental anomaly resulting in multiple renal cysts of varying sizes and
           fibrosis of renal parenchyma
      ii) Clinical presentation
         • Unilateral
         • Palpable flank mass
         • Asymptomatic
         • Bilateral
         • Incompatible with life
      iii) Sonographic findings
         • Multiple cysts of varying sizes, arranged randomly
         • No normal parenchyma
         • Late stage
           o Usually decreases in size
           o Parenchymal thinning
           o Loss of parenchymal differentiation
         • Contralateral kidney enlargement
      iv) Correlative and/or prior imaging
      v) Treatment
   
   b. Autosomal recessive polycystic kidney disease (ARPKD), formerly infantile polycystic kidney
disease (IPKD)
      i) Definition
         • Autosomal recessive inherited disorder characterized by abnormal proliferation and
           dilatation of renal tubules resulting in multiple microscopic or tiny cysts
      ii) Clinical presentation
         • Incompatible with life
         • Renal failure
         • With hepatic fibrosis – cirrhosis
      iii) Sonographic findings
         • Bilateral enlargement
         • Increased parenchymal echogenicity
Abdomen and Superficial Structures Including Introductory Pediatric and Musculoskeletal

- Loss of corticomedullary differentiation

c. Autosomal dominant polycystic kidney disease (ADPKD), formerly adult polycystic kidney disease (APKD)
   i) Definition
      - Autosomal dominant inherited disorder with bilateral development of parenchymal cysts
   ii) Clinical presentation
      - Asymptomatic (early)
      - Usually bilateral presenting in adulthood
      - Hypertension
      - Renal failure
      - Increase incidence of renal calculi and infection
         o Obstruction
            ~ Pain
         o Infection
            ~ Fever
            ~ Pain
   iii) Sonographic findings
      - Bilateral renal enlargement
      - Multiple cysts of variable size
      - Cysts of varying echogenicity
      - Acoustic enhancement
      - Calculi
   iv) Associated findings
      - Multiple cysts associated with other organs
      - Berry aneurysm

d. Medullary cystic disease
   i) Definition
      - Autosomal inherited disorder
         o Autosomal dominant: adult onset with rapid progression, uremia and death
         o Autosomal recessive: juvenile onset
   ii) Clinical presentation
      - Salt loss
      - Anemia
      - Azotemia
      - Polyuria
   iii) Sonographic findings
• Bilateral
• Increased medullary echogenicity
• Loss of parenchymal definition
• End-stage renal failure
  o Decreased renal size
  o Increase parenchymal echogenicity
  o Loss of differentiation

iv) Treatment

e. Medullary sponge kidney

i) Definition
• Bilateral, dysplastic cystic dilation of medullary and papillary portions of collecting tubules

ii) Clinical presentation
• Asymptomatic
• With complications
  o Obstruction
    ~ Pain
  o Infection
    ~ Fever
    ~ Pain

iii) Sonographic findings
• Increased medullary echogenicity
• Calculus
• Medullary nephrocalcinosis is often present

iv) Treatment

f. Acquired cystic kidney disease (ACKD)

i) Definition
• Patients undergoing dialysis develop multiple cysts in native and allograft kidneys

ii) Clinical presentation
• Asymptomatic
• Increased risk of tumor development

iii) Sonographic findings
• Multiple cysts
• Varying sizes
• Variable echogenicity
• Bilateral
5. Benign solid renal tumors
   a. Angiomyolipoma - renal hamartoma
      i) Definition
         • Benign tumor composed of fat, blood vessel, and smooth muscle tissue
      ii) Clinical presentation
         • Female predominance
         • Asymptomatic
         • Hematuria
         • Hypertension
         • Severe flank pain
         • Risk of hemorrhage > 4 cm size
      iii) Sonographic findings
         • Variable echogenicity
            • Hyperechoic lesion in renal parenchyma most common
            • Variable echogenicity if hemorrhage
         • Variable size
      iv) Correlative and/or prior imaging
      v) Treatment
   b. Adenoma
      i) Definition
         • Benign tumor derived from glandular epithelium
      ii) Clinical presentation
         • Asymptomatic
         • Painless hematuria
      iii) Sonographic findings
         • Well-defined
         • Isoechoic or hyperechoic cortical mass
      iv) Correlative and/or prior imaging
      v) Treatment
   c. Oncocytoma
      i) Definition
         • Epithelial cell tumors
      ii) Clinical presentation
• Asymptomatic
• Male predominance in later decades of life

iii) Sonographic findings
• Solid
• Variable in size
• Variable in echogenicity
• Difficult to differentiate from adenocarcinoma

iv) Correlative and/or prior imaging

v) Treatment

d. Mesoblastic nephroma (fetal renal hamartoma)
i) Definition
• Pediatric benign tumor composed of mesoderm tissue - benign counterpart of Wilms’ tumor (nephroblastoma)

ii) Clinical presentation
• Palpable flank mass
• Hypertension

iii) Sonographic findings
• Varying echogenicity
• Possible areas of cystic degeneration
• May image calcifications - hyperechoic, shadowing foci

iv) Correlative and/or prior imaging

v) Treatment

6. Malignant renal tumors

a. Renal cell carcinoma (RCC) – hypernephroma

i) Definition
• Renal parenchymal carcinoma, consisting of tubular cells

ii) Clinical presentation
• Pain
• Hematuria
• Palpable mass

iii) Sonographic findings
• 2-D
  • Solid mass of varying echogenicity
  • Appearance is variable ranging from a solid mass to a multiocular cyst
  • Often at least partially exophytic
  • Calcifications
  • Echogenic mass(es) in renal vein, IVC, and contralateral kidney
iv) Doppler
   • Hypervascular, with high systolic and diastolic arterial flow
v) Correlative and/or prior imaging
vi) Treatment

b. Transitional cell carcinoma (TCC)
i) Definition
   • Primary malignant epithelial tumor originating in urinary collecting system
ii) Clinical presentation
   • Painless hematuria
   • Hydronephrosis
iii) Sonographic findings
   • “Faceless” kidney sign
     o Obliteration of the normal renal sinus appearance
   • Isoechoic or hypoechoic mass within collecting system
     o Renal pelvis
     o Ureter
     o Bladder
       ~ Wall thickening
         A. Focal, irregular
         ~ Intraluminal mass
   • Pelvocaliectasis
   • Hilar adenopathy
iv) Correlative and/or prior imaging
v) Treatment

 c. Wilm’s tumor – nephroblastoma
i) Definition
   • Pediatric malignant mixed tumor composed of embryonal elements
ii) Clinical presentation
   • Fever
   • Hematuria
   • Hypertension
   • Palpable flank mass
iii) Sonographic findings
   • Most commonly is homogeneous and echogenic
   • May have hypoechoic or cystic appearance
   • Large in size
• Calcifications
  • Increased risk with horseshoe kidney
  iv) Correlative and/or prior imaging
  v) Treatment
d. Metastases to the kidney
  i) Definition
  • Growth of malignant cells distant from primary tumor
  ii) Clinical presentation
  • Asymptomatic
  • Known pathology follow-up
  iii) Sonographic findings
  • Varies from hypoechoic to hyperechoic masses
  • Single or multiple masses
  • Diffusely hypoechoic
  iv) Correlative and/or prior imaging

7. Renal calcification
   a. Calculi
      i) Definition
      • Focal concentrations of calcium, uric acid, or cystine in renal parenchyma or collecting system
      ii) Clinical presentation
      • Asymptomatic
      • Acute pain
      • Hematuria
      iii) Sonographic findings
      • Hyperechoic foci
      • Acoustic shadowing
      • Obstructive hydronephrosis
      • Mobile, dependent foci in bladder hydronephrosis
      • Twinkle artifact
      • Echogenic focus in dilated distal ureter on bladder view
      iv) Correlative and/or prior imaging
      v) Treatment
   b. Nephrocalcinosis
      i) Medullary
      • Definition
• Formation of aggregates of calcium in renal tubules
  • Clinical presentation
    o Asymptomatic
    o Renal failure
    o Hyperparathyroidism
  • Sonographic findings
    o Increased echogenicity of pyramids
    o Normal cortical echogenicity
    o Distal acoustic shadowing
  • Correlative and/or prior imaging
  • Treatment

ii) Cortical
  • Definition
    o Diffuse of focal calcium deposition in renal cortex
  • Clinical presentation
    o Asymptomatic
  • Sonographic findings
    o Focal or diffuse increase in cortical echogenicity
    o Distal acoustic shadowing
    o Decrease in size
  • Correlative and/or prior imaging
  • Treatment

8. Obstructive uropathy
   a. Hydronephrosis
      i) Definition
        • Dilation of renal pelvis and calyces due to obstruction of outflow of urine from either congenital, intrinsic, or extrinsic cause
      ii) Etiology
        • Congenital
        • Intrinsic
        • Extrinsic
      iii) Types/grading
      iv) Sonographic findings
        • 2-D
          o Mild, moderate, or severe degrees of pelvocaliectasis
          o Hydroureter
          o Cortical thinning
9. Trauma
   a. Definition
      i) Assault to urinary tract from blunt force, penetrating trauma, or rupture of neoplasm resulting in possible
         - Rupture of kidney and capsule
         - Subcapsular hematoma
            o Page kidney
               ~ Extrinsic compression of the renal parenchyma
         - Parenchymal laceration
         - Urinoma
   b. Clinical presentation
      i) Abdominal pain
      ii) Decreased hematocrit
      iii) Hematuria
      iv) Oliguria
      v) Hypertension
   c. Sonographic findings
      i) 2-D
         - Sonographically occult is common
         - Diffuse heterogeneous parenchymal pattern
         - Focal mass
            o Varying echogenicity
         - Free abdominopelvic fluid
         - Loculated abdominal fluid collection
            o Anechoic
            o Septations
            o Debris
      ii) Doppler
         - Avascular areas
   d. Correlative and/or prior imaging
   e. Treatment
10. Infarction
a. Definition
   i) Necrosis of tissue due to occlusion of arterial blood supply

b. Clinical presentation
   i) Asymptomatic
   ii) Localized pain

c. Sonographic findings
   i) Focal or diffuse
   ii) Acute
      • Hypoechoic
   iii) Chronic
      • Decreased size
      • Capsular retraction or focal scarring
      • Increased echogenicity

d. Correlative and/or prior imaging

11. Vascular disorders

   a. Renal artery stenosis (RAS)
      i) Definition
         • Luminal narrowing of renal artery
      ii) Clinical presentation
         • Hypertension
      iii) Sonographic findings
         • 2-D
            o Normal
            o Decrease in size of affected kidney
            o Increase in cortical echogenicity
         • Doppler
            o Direct
               ~ Normal
               ~ Increased renal artery ratio (RAR)
               ~ Increased peak systolic velocity
               ~ Spectral broadening
               ~ Tardus parvus waveform
            o Indirect
               ~ Decreased diastolic systolic ratio (DSR)
               ~ Delayed acceleration time
               ~ Asymmetric segmental resistive index relative to other kidney
      iv) Correlative and/or prior imaging
v) Treatment

b. Renal artery occlusion
i) Definition
   • Obstruction of arterial lumen

ii) Clinical presentation
   • Asymptomatic
   • Flank pain
   • Hypertension

iii) Sonographic findings
   • 2-D
     o Decrease in renal size after acute phase
     o Increase in cortical echogenicity
   • Doppler
     o Absence of flow distal to occlusion

iv) Correlative and/or prior imaging
v) Treatment

c. Renal vein thrombosis
i) Definition
   • Presence of obstructive or non-obstructive thrombus in renal vein

ii) Clinical presentation
   • Asymptomatic
   • Flank pain

iii) Sonographic findings
   • 2-D
     o Increase in renal size in acute phase
     o Decrease in size in late or chronic phase
     o Varying echogenicity
     o Echogenic material within lumen of vein
     o IVC involvement
   • Doppler
     o Absence of flow in renal vein – variable
     o Increased arterial resistance with or without diastolic reversal

iv) Correlative and/or prior imaging
v) Treatment

d. Other vascular disorders
i) Aneurysm
• Anechoic structure with circular flow
• May be obscured by wall calcifications

ii) Pseudoaneurysm
• To-and-fro waveform
• Circular yin-yang pattern within anechoic structure

iii) Arteriovenous (AV) fistula
• Low resistance arterial flow
• Arterialized adjacent venous flow
• Tissue reverberation of adjacent parenchyma

12. Lower urinary tract pathology

a. Cystitis
i) Definition
• Inflammation of urinary bladder

ii) Clinical presentation
• Pain
• Frequency
• Urgency
• Hematuria

iii) Sonographic findings
• Normal
• Focal or diffuse wall thickening
• Luminal echoes - layering or dependent

iv) Correlative and/or prior imaging
v) Treatment

b. Bladder wall thickening
i) Definition
• Increase size of bladder wall
  ○ Causes
  ~ Muscular hypertrophy
  ~ Tumor
  ~ Infection
  ~ Under-distended bladder

ii) Clinical presentation
• Asymptomatic

iii) Sonographic findings
• Focal or diffuse increase in wall size
iv) Treatment

c. Lower tract calculi
   i) Definition
      • Presence of focal calcification in urinary collecting system
   ii) Clinical presentation
      • Asymptomatic
      • If obstructing
         o Pain
         o Hematuria
   iii) Sonographic findings
      • Focal mass of increased echogenicity
      • Distal acoustic shadowing
      • Obstructive hydronephrosis
      • Mobile if in bladder
   iv) Correlative and/or prior imaging
   v) Treatment

d. Hematoma
   i) Definition
      • Localized collection of blood within LU tract
   ii) Clinical presentation
      • Asymptomatic
   iii) Sonographic findings
      • Low-level echoes
      • Focal (mass-like)
      • Layering, gravity dependent echoes
   iv) Treatment

e. Neoplasia
   i) Adenoma
      • Definition
         o Benign glandular epithelial tumor
      • Clinical presentation
         o Asymptomatic
         o Painless hematuria
      • Sonographic findings
         o Focal luminal mass
         o Irregular wall thickening
o Varying echogenicity
  o Varying size
• Correlative and/or prior imaging
• Treatment

ii) Transitional cell carcinoma
• Definition
  o Primary malignant epithelial tumor originating in urinary collecting system
• Clinical presentation
  o Asymptomatic
  o Painless hematuria
• Sonographic findings
  o Focal luminal bladder mass
  o Irregular bladder wall thickening
  o Echogenic ureteral mass
  o Varying echogenicity
  o Obstructive hydronephrosis
• Correlative and/or prior imaging
• Treatment

iii) Metastases
• Definition
  o Growth of malignant cells distant from primary site - usually direct extension to urinary bladder
• Clinical presentation
  o Asymptomatic
  o With obstruction
    ~ Pain
    ~ Oliguria
• Sonographic findings
  o Focal luminal bladder mass
  o Irregular bladder wall thickening
  o Echogenic ureteral mass
  o Varying echogenicity
  o Obstructive hydronephrosis
• Treatment
Section V: Spleen

1. Describe the normal anatomy, function, sonographic technique and appearance of the spleen
2. Correlate clinical indications and laboratory values associated with splenic disease
3. Describe splenic pathology in terms of sonographic appearances, sequelae and associated pathologies
4. Describe associated pathologies and sequelae relative to splenic disease
5. Identify normal and abnormal flow characteristics and waveforms
6. Discuss current diagnostic Doppler criteria

V. Spleen

A. Embryology

B. Anatomy
   1. Location
   2. Organ of the lymphatic and reticuloendothelial systems
   3. Size and weight
   4. Blood supply and drainage
   5. Relational anatomy
      a. Peritoneal

C. Variants of Normal
   1. Aplasia
   2. Hypoplasia
   3. Accessory spleens
   4. Wandering spleen

D. Functions
   1. Filter
   2. Reservoir
   3. Destruction of red blood cells
   4. Formation of bile pigments
   5. Destruction of microorganisms
   6. Immunity
   7. Production of lymphocytes
   8. Production of plasma cells
   9. Erythropoiesis in the fetus
   10. Sequestration of platelets

E. Laboratory Values
   1. White blood count (WBC)
      a. Leukocytosis
      b. Leukopenia
   2. Red blood count (RBC)
a. Decreased RBC
b. Increased RBC

3. Hematocrit
   a. Increase
   b. Decrease

4. Platelet count
   a. Thrombocytosis
   b. Thrombocytopenia

F. Indications
   1. Palpable spleen
   2. Abnormal laboratory tests
   3. LUQ pain
   4. Infectious diseases
   5. Sickle cell disease

G. Scanning Technique
   1. Patient preparation
   2. Transducer selection
   3. Patient positioning
   4. Image optimization
      a. Homogeneous low to medium level echoes
      b. Isoechoic, or slightly more echogenic than normal liver
   5. Image protocol

H. Pathophysiology
   1. Congestive splenomegaly
      a. Causes
         i) Heart failure
         ii) Cirrhosis
         iii) Portal hypertension
         iv) Portal/splenic vein thrombosis
         v) Cystic fibrosis
         vi) HIV/AIDS
      b. Clinical presentation
      c. Sonographic findings
         i) Variable echogenicity
         ii) Increased size
      d. Correlative and/or prior imaging
   2. Treatment
I. Infection/Inflammation

1. Systemic
   a. Types
      i) Mononucleosis
      ii) Tuberculosis
      iii) Histoplasmosis
      iv) Schistosomiasis
      v) Sarcoidosis
      vi) Candidiasis
   b. Clinical presentation
      i) Varies
   c. Sonographic findings
      i) Variable dependent on type
      ii) Normal to increase in size
   d. Correlative and/or prior imaging
   e. Treatment

2. Focal
   a. Types
      i) Echinococcus
      ii) Abscess
   b. Clinical presentation
      i) LUQ pain
      ii) Fever
   c. Sonographic findings
      i) Focal mass of varying echogenicity
      ii) Irregular borders
      iii) Normal to increase size
      iv) Ill-defined, thickened borders
      v) Possible distal acoustic enhancement
      vi) Shadow from presence of gas in the abscess
   d. Correlative and/or prior imaging
   e. Treatment

J. Neoplasia

1. Benign
   a. Hamartoma
      i) Definition
      ii) Clinical presentation
iii) Sonographic findings
   • Focal well-defined mass
   • Variable echogenicity

iv) Correlative and/or prior imaging
v) Treatment

b. Hemangioma
i) Definition
ii) Clinical presentation
iii) Sonographic findings
   • Typically small
   • Focal
   • Well-defined borders
   • Variable echogenicity

iv) Correlative and/or prior imaging
v) Treatment

2. Malignant
   a. Leukemia
      i) Definition
      ii) Clinical presentation
      iii) Sonographic findings
         • Marked splenomegaly
      iv) Correlative and/or prior imaging
      v) Treatment

   b. Lymphoma
      i) Definition
      ii) Clinical presentation
      iii) Sonographic findings
         • Focal mass
            o Typically hypoechoic mass
            o Poorly defined margins
         • Diffuse splenomegaly
      iv) Correlative and/or prior imaging
      v) Treatment

   c. Hemangiosarcoma
      i) Definition
      ii) Clinical presentation
      iii) Sonographic findings
• Focal well-defined mass
• Variable echogenicity
• Variable size
  iv) Correlative and/or prior imaging
  v) Treatment
d. Metastasis
  i) Definition
  ii) Clinical presentation
  iii) Sonographic findings
    • Multiple focal masses
      o Variable size and echogenicity
  iv) Correlative and/or prior imaging
  v) Treatment

K. Cysts
  1. Types
    a. True cyst
      i) Congenital
    b. Post-traumatic
    c. Infection
    d. Parasitic
    e. Neoplastic
  2. Clinical presentation
  3. Sonographic findings
    a. Variable, related to types and cause
      i) Well-defined walls
      ii) Anechoic, hypoechoic
      iii) Posterior enhancement
      iv) Internal echoes
      v) Fluid-fluid level
      vi) Wall thickening
      vii) Possible wall calcification
  4. Correlative and/or prior imaging
  5. Treatment

L. Blood Disorders
  1. Types
    a. Hemolytic anemia
    b. Extramedullary hematopoiesis
i) Myelofibrosis

2. Clinical presentation

3. Sonographic findings

4. Correlative and/or prior imaging

5. Treatment

M. Storage Disease

1. Types
   a. Gaucher disease
   b. Diabetes mellitus
   c. Niemann-Pick disease
   d. Amyloidosis
   e. Histiocytosis
   f. Hemochromatosis

2. Clinical presentation

3. Sonographic findings

4. Correlative and/or prior imaging

5. Treatment

N. Infarction

1. Definition
   a. Occlusion of segment of splenic artery
   b. Embolic source

2. Risk factors

3. Clinical presentation
   a. Usually asymptomatic
   b. Sudden onset of LUQ pain

4. Sonographic presentation
   a. Wedge-shaped lesion
      i) Well-defined borders
      ii) Base toward subcapsular surface of spleen
   b. Variable echogenicity
      i) Time and age of infarction
   c. Possible small spleen

5. Correlative and/or prior imaging

6. Treatment

O. Splenic Sequestration Crisis in Sickle Cell Disease
1. Definition
2. Clinical presentation
3. Sonographic findings
4. Correlative and/or prior imaging
5. Treatment

P. Trauma
1. Definition
2. Clinical presentation
3. Sonographic findings
   a. Hematoma
      i) Subcapsular or intraparenchymal
   b. Free or focal accumulation of blood in intraperitoneal or retroperitoneal cavity
   c. Variable echogenicity
      i) Age and progression
   d. Other
      i) Enlarged spleen
      ii) Irregular borders
      iii) Left pleural effusion
4. Correlative and/or prior imaging
5. Treatment

Q. Congenital Disorders
1. Asplenia syndrome
   a. Definition
      i) Congenital absence of spleen
   b. Associated with bilateral right-sidedness
      i) Visceral heterotaxy
   c. Incidence
      i) Rare
   d. Clinical presentation
      i) Impaired immune response
   e. Sonographic findings
      i) Absent spleen
      ii) Possible maldevelopments
         • Abnormal location of IVC
         • Horseshoe kidneys
         • Abnormal location of stomach
2. Polysplenia syndrome
a. Definition
   i) Accessory splenic tissue
b. Associated with bilateral left-sidedness
   i) Visceral heterotaxy
   ii) Associated with other congenital abdominal anomalies
c. Incidence
   i) Rare
d. Clinical presentation
   i) Asymptomatic
   ii) Dependent upon presence of other abnormalities
e. Sonographic findings
   i) Biliary atresia with absent gallbladder
   ii) Azygous/herniazygous continuation of IVC with interruption of hepatic segment of IVC
   iii) Genitourinary anomalies
   iv) Abnormal location of liver
f. Correlative and/or prior imaging
g. Treatment
Section VI: Adrenal

1. Describe the normal anatomy and variants, function, sonographic technique and sonographic appearance of the adrenal glands
2. Correlate the clinical indications and laboratory values associated with adrenal pathology
3. Describe adrenal pathology in terms of sonographic appearances, sequelae and associated pathologies
4. Describe associated pathologies and sequelae

VI. Adrenal

A. Adrenal Glands
   1. Embryology
   2. Anatomy
      a. Micro
         i) Cortex
            • Zona glomerulosa
            • Zona fasciculata
            • Zona reticularis
         ii) Medulla
      b. Macro
         i) Paired organs
         ii) Retroperitoneal
         iii) Arterial supply
         iv) Venous drainage
      c. Relational
   3. Variants
      a. Agenesis
   4. Function
      a. Cortex secretes steroids
         i) Mineralocorticoids
         ii) Glucocorticoids
         iii) Gonadal hormones
      b. Medulla secretes
         i) Catecholamines
            • Epinephrine
            • Norepinephrine
         ii) Endocrine glands
      c. Hyperfunction
         i) Cushing’s disease/syndrome
ii) Adrenogenital (AG) syndrome
iii) Conn’s disease
d. Hypofunction
   i) Addison’s disease
5. Laboratory values
   a. Aldosterone
   b. Cortisol
c. Adrenocorticotropic hormone (ACTH)
d. Catecholamines
e. Total metanephrines
f. Vanillylmandelic acid (VMA)
6. Indications
   a. Decreased hematocrit
   b. Hypertension
c. Virilism
7. Sonographic technique
   a. Patient preparation
   b. Transducer selection
c. Patient positioning
d. Imaging optimization
e. Imaging protocol
8. Normal sonographic findings
   a. Pediatric
      i) Cortex hypoechoic to medulla
      ii) Medulla hyperechoic to cortex
   b. Adult
      i) Not routinely visualized
9. Pathophysiology
   a. Neoplasia
      i) Adenoma
         • Definition
            o Benign adrenal tumor of epithelial origin
         • Clinical presentation
            o Asymptomatic
            o Cushing’s syndrome
            o Conn’s disease
         • Sonographic findings
Focal mass
- Variable size
- Unilateral or bilateral
- Variable echogenicity
- Correlative and/or prior imaging
- Treatment

ii) Adenocarcinoma
- Definition
  - Primary adrenal malignant tumor of epithelial origin
- Clinical presentation
  - Asymptomatic
  - Cushing’s syndrome
  - Adrenogenital (AG) syndrome
  - Precocious puberty
  - Conn’s syndrome
- Sonographic findings
  - Variable size
  - Variable echogenicity
  - Complex
  - Calcifications
- Correlative and/or prior imaging
- Treatment

iii) Neuroblastoma
- Definition
  - Pediatric sarcoma arising in adrenal medulla
  - Can also originate in autonomic nervous system
- Clinical presentation
  - Asymptomatic
  - Palpable mass
  - Weight loss
  - Pallor
  - Fever
  - Hypertension
  - Tachycardia
  - Sweats
- Sonographic findings
  - Focal mass
iv) Pheochromocytoma

- **Definition**
  - Chromaffin cell tumor originating in adrenal medulla or chromaffin tissue of sympathetic paraganglia

- **Clinical presentation**
  - Hypertension
  - Headache
  - Palpitations
  - Tachycardia
  - Excessive perspiration

- **Sonographic findings**
  - Focal mass
  - Unilateral or bilateral
  - Variable in appearance

- **Correlative and/or prior imaging**

- **Treatment**

v) Metastases

- **Definition**
  - Growth of malignant cells distant from primary site

- **Clinical presentation**
  - Asymptomatic
  - Addison’s disease

- **Sonographic findings**
  - Focal mass
  - Unilateral or bilateral
  - Varying echogenicity
  - Varying size

- **Correlative and/or prior imaging**

- **Treatment**

b. Inflammation/Infection

i) Abscess

- **Definition**
• Focal collection of pus from disintegration of tissue
• Clinical presentation
• Fever
• Pain
• Sonographic findings
• Focal mass
• Variable echogenicity
• Acoustic enhancement
• Fluid-fluid levels
• Hyperechoic foci with shadowing
• Correlative and/or prior imaging
• Treatment

ii) Infectious disease

• Tuberculosis
  o Chronic, granulomatous infection by mycobacterium tuberculosis
• Histoplasmosis
  o Infection caused by inhalation of fungal spores histoplasma capsalutum
• Cytomegalovirus
  o Infection by species-specific herpes-type virus
• Clinical presentation
  o Addison’s disease
• Sonographic findings
  o Acute
  o Bilateral enlargement
  o Inhomogeneous
  o Chronic
  o Calcification
• Correlative and/or prior imaging
• Treatment

c. Other

i) Hemorrhage

• Types
  o Spontaneous
  o Posttraumatic
  o Neonatal
• Clinical presentation
- Decreased hematocrit
- Abdominal pain
- Asymptomatic

• Sonographic findings
  - Appearance dependent upon age of bleed

• Correlative and/or prior imaging

• Treatment

ii) Cyst

• Types
  - Endothelial
  - Pseudocyst
  - Epithelial
  - Parasitic

iii) Clinical presentation

• Asymptomatic

iv) Sonographic findings

• Round
• Anechoic
• Acoustic enhancement
• Wall calcification

v) Correlative and/or prior imaging

vi) Treatment
Section VII: Abdominal Vasculature

1. Describe the normal anatomy and variants, function and sonographic appearance of the major abdominal vasculature
2. Understand the clinical indication of the study and laboratory values associated with vascular disease.
3. Recite the protocol for sonographic assessment of the abdominopelvic vasculature
4. Describe vascular pathology in terms of sonographic appearances, sequelae and associated pathologies
5. Identify normal and abnormal flow characteristics and waveforms
6. Discuss current diagnostic Doppler criteria

VII. Abdominal Vasculature

A. Embryology

B. Anatomy

1. Vessel wall layers
   a. Tunica intima
   b. Tunica media
   c. Tunica adventitia

2. Arterial anatomy
   a. Abdominal aorta
      i) Anterior branches
         • Celiac axis/trunk
            o Splenic
            o Left gastric
            o Hepatic artery
               ~ Common hepatic
               ~ Proper hepatic
                  • Right hepatic (cystic)
                  • Left hepatic
               ~ Gastroduodenal
         • Superior mesenteric artery (SMA)
         • Inferior mesenteric artery (IMA)
      ii) Lateral branches
         • Right renal
         • Left renal
            o Left gonadal
         • Right gonadal
         • Right common iliac
            o Right internal iliac
            o Right external iliac
• Left common iliac
  o Left internal iliac
  o Left external iliac

C. Arterial Variants
1. Most common
   a. Multiple renal arteries
   b. Replaced right hepatic artery
   c. Common trunk of SMA/celiac artery

D. Functions
1. Arteries
2. Veins
3. Capillaries

E. Laboratory Values
1. Hematocrit
2. White blood count (WBC)

F. Indications
1. Pulsatile abdominal mass
2. Abdominal pain
3. Abdominal bruit
4. Hemodynamic compromise of lower extremities
5. Swelling of the lower extremities

G. Sonographic Technique
1. Patient preparation
2. Transducer selection
3. Positioning
4. Image optimization
5. Image protocol

H. Normal Arterial Sonographic Findings
1. Measurements
   a. AP aorta – outer to outer wall (sagittal) and transverse (right to left) – outer to outer wall (coronal)
   b. Proximal, mid, distal aorta
   c. Proximal common iliac
2. Parallel walls
3. Pulsatility
4. Anechoic lumen
5. Doppler
   a. Aorta
i) High resistance proximal to renal arteries
ii) Triphasic distal to renal arteries

b. Celiac
i) Low resistance
c. Splenic
i) Low resistance
ii) Turbulent distally
d. Hepatic
i) Low resistance
e. SMA
i) Pre-prandial
   • High resistance
ii) Post-prandial
   • Low resistance
f. Renals
i) Low resistance

I. Arterial Pathophysiology
1. Atherosclerosis
   a. Definition
      i) Type
   b. Complications
      i) Thrombus
      ii) Embolus
      iii) Stenosis
      iv) Occlusion
      v) Aneurysm

2. Aortic ectasia/arteriomegaly
   a. Definition
      i) Diffuse, mild enlargement of aorta without distal tapering
   b. Clinical presentation
      i) Asymptomatic
   c. Sonographic findings
      i) Diffuse enlargement of aorta
   d. Correlative and/or prior imaging

3. Aneurysm
   a. Definition
      i) Abnormal dilation of segment of arterial wall
b. Types
   i) Fusiform
   ii) Bulbous
   iii) Saccular
   iv) Dumbbell
   v) Mycotic

c. Abdominal aortic aneurysm (AAA)
   i) Focal or diffuse dilation of abdominal aorta
   ii) Clinical presentation
      • Asymptomatic
      • Palpable, pulsatile abdominal mass
      • Abdominal, back, or leg pain
      • Abdominal bruit
   iii) Sonographic findings
      • 2-D
         o Increased size of aorta (focal or diffuse)
         o Types
         o Location
         o Measurement technique
         o Irregular walls
         o Calcifications
         o Thrombus
      • Doppler
   iv) Correlative and/or prior imaging
   v) Treatment
      • Surgical
      • Endovascular stent graft

d. Other arterial abdominal aneurysms
   i) Iliac
   ii) Splenic
   iii) Hepatic

4. Aortic dissection
   a. Definition
      i) False lumen between intima and media of vessel wall
      ii) Most common is origination in thoracic aorta
   b. Clinical presentation
      i) Abdominal pain
c. Sonographic findings
   i) Flow within wall and lumen of vessel
   ii) Mild to no dilation of the vessel
   iii) Thin, echogenic linear structure in true lumen (intimal flap)

5. Aortic rupture
   a. Definition
      i) Disruption of the aortic wall, typically infrarenal and left lateral location
   b. Clinical presentation
      i) Abdominal pain
      ii) Loss of consciousness
      iii) Decreased vital signs
      iv) Hypotensive
   c. Sonographic findings
      i) Para-aortic hematoma, typically left flank
      ii) Varying echogenicity
      iii) May displace left kidney

6. Pseudoaneurysm
   a. Definition
      i) May be due to infection (mycotic), post procedure, trauma
      ii) May result from puncture of two or three layers of vessel wall
      iii) Does not have a true wall (three layers)
      iv) Allows blood to escape into surrounding tissues
      v) Most common cause is arterial catheterization or trauma
   b. Clinical presentation
      i) Focal abdominal pain
      ii) Pulsatile mass
      iii) Bruising
   c. Sonographic findings
      i) Focal mass of varying echogenicity
      ii) Identification of narrow connection (neck) between vessel and mass
      iii) Doppler
         • To-and-fro spectral appearance in neck
         • Yin-yang color appearance in flow lumen
   d. Treatment
      i) Ultrasound-guided thrombin
      ii) Ultrasound-guided compression
      iii) Surgery
7. Renal artery stenosis
   a. Definition
      i) Elevation in blood pressure due to decreased renal blood flow
   b. Clinical presentation
      i) Hypertension
   c. Sonographic findings
      i) Elevated velocity in the main renal artery
      ii) Parvus/tardus pattern in the intrarenal arteries

8. Renal transplants
   a. Sonographic assessment
      i) 2D
      ii) Doppler
   b. Rejection
   c. Acute tubular necrosis

J. Venous anatomy
   1. Inferior vena cava (IVC)
      a. Tributaries
         i) Right and left common iliacs
         ii) Ascending lumbar
         iii) Right gonadal
         iv) Renal
             • Right
             • Left
                 o Left gonadal
         v) Hepatic
             • Right
             • Middle
             • Left

   2. Hepatopetal venous system
      a. Main portal
         i) Tributaries
         ii) Splenic
         iii) Inferior mesenteric (IMV)
         iv) Superior mesenteric (SMV)
         v) Coronary (left gastric)
         vi) Pyloric (right gastric)
      b. Branches
i) Right portal
   - Anterior
   - Posterior

ii) Left portal
   - Medial
   - Lateral

3. Variants/anomalies
   a. Accessory hepatic veins
   b. Transposition of the IVC
   c. Partial transposition
   d. Situs inversus
   e. Asplenia syndrome
   f. Azygous continuation of the IVC
   g. Retro-aortic left renal vein

4. Indications
   a. Known pathology
   b. Follow up studies
   c. IVC filter
   d. Swelling of trunk, lower extremities

5. Sonographic technique
   a. Patient preparation
   b. Transducer selection
   c. Patient positioning
   d. Imaging optimization
   e. Imaging protocol

6. Normal sonographic findings
   a. Cava and tributaries
      i) Anechoic lumen
      ii) Varying luminal size
   b. Portal
      i) Anechoic lumen
      ii) Portal vein walls hyperechoic to liver parenchyma

7. Normal Doppler appearance
   a. IVC
      i) Pulsatile Doppler pattern by diaphragm and phasic Doppler pattern in distal abdomen
   b. Hepatic veins
      i) Pulsatile
c. Renal veins
   i) Monophasic

d. Splenic vein
   i) Hepatopetal

e. Portal vein
   i) Hepatopetal flow

8. Venous pathophysiology
   a. IVC
      i) Intraluminal thrombus
         • Definition
            o Most common intraluminal finding in IVC
            o Extension of thrombus from tributary vein – renal, hepatic, iliac, gonadal, adrenal
         • Clinical presentation
            o Asymptomatic
            o Swelling/edema of lower extremities
         • Sonographic findings
            o 2-D
               ~ Intraluminal mass of varying echogenicity; if malignant – lumen enlargement
               ~ Luminal stenosis or occlusion
            o Doppler
         • Correlative and/or prior imaging
         • Treatment

      ii) Intravascular tumor
         • Definition
            o Most common - caval thrombus due to tumor; renal cell carcinoma, hepatocellular carcinoma, breast and adrenal carcinoma
            o Primary caval tumors are rare with leiomyosarcoma
         • Sonographic findings
            o 2-D
               ~ Vessel enlargement
               ~ Intraluminal mass of varying echogenicity
            o Doppler
               ~ Signal from tumor mass
         • Correlative and/or prior imaging
         • Treatment

      iii) Arteriovenous (AV) Shunts
         • Definition
o Abnormal connection between artery and vein

- Types
- Indications
  o Presence of bruit or thrill associated with dilated pulsatile mass
  o Low back and abdominal pain
  o Swelling of lower trunk and extremities
  o Renal A-V fistulas
  o Hematuria
  o Hypertension
  o Cardiomegaly
  o Heart failure (rare)
  o Abdominal bruit
- Doppler findings
  o Pulsatile, arterialized venous waveform
  o Increased systolic and diastolic arterial velocities proximal to fistula (decreased RI)
  o Reverts to normal diastolic flow distal to fistula
  o Perivascular tissue vibration
- Correlative and/or prior imaging
- Treatment

b. Portal hypertension (PH)
  i) Definition
  o Increased resistance to flow or increased flow in portal system
    o Prehepatic
    o Intrahepatic
    o Posthepatic
  ii) Clinical presentation
    o Asymptomatic
    o Symptomatic
      o Hematemesis, varices
      o Abdominal distention, ascites
      o Splenomegaly
      o Hepatic failure
      o Jaundice
  iii) Sonographic findings
    o 2-D
      o Increase in portal vein diameter
      o Splenomegaly
o Presence of portosystemic collaterals
o Ascites
o Variable hepatic appearance
• Doppler
  o Hepatofugal portal flow
  o Enlarged tortuous hepatic artery with high velocities
iv) Correlative and/or prior imaging
c. Portal vein thrombosis
  i) Presence of thrombus in portal system, may be benign or malignant
  ii) Clinical presentation
  iii) Sonographic findings
    • 2-D
      o Intraluminal thrombus of variable echogenicity
      o Presence of collaterals (cavernous transformation)
      o Presence of flow in the thrombus (malignant)
    • Doppler
      o Lack of flow in portal vein
      o Enlarged hepatic artery
iv) Correlative and/or prior imaging
v) Treatment
    • Transjugular intrahepatic portosystemic shunt (TIPS)
      o Pathophysiology
      o Sonographic assessment
d. Cavernous transformation of the portal vein
  i) Definition
  ii) Clinical presentation
  iii) Sonographic findings
    • 2-D
      o Multiple anechoic tubular channels in porta hepatis indicating presence of collaterals
    • Doppler
      o Multiple small portal venous-like vessels in porta hepatic which may extend into the liver
iv) Correlative and/or prior imaging
v) Treatment
Section VIII: Gastrointestinal Tract (GI)

1. Describe the normal anatomy and variants, function, sonographic technique and normal sonographic appearances of the GI tract
2. Correlate the clinical indications and laboratory values associated with GI tract disease
3. Describe GI tract pathology in terms of sonographic appearances
4. Describe associated pathologies and sequelae relative to GI tract disease

VIII. Gastrointestinal Tract

A. Embryology
B. Anatomy

1. Wall
   a. Constant with layers; listed from the outer layer to inner layer
      i) Serosa – outermost
      ii) Muscularis propria
      iii) Submucosa
      iv) Mucosa
         • Muscularis mucosa
         • Lamina propria
         • Epithelial layer
   b. Specific morphologic features
      i) Stomach
         • Rugae
      ii) Small intestine
         • Valvulae conniventes
      iii) Large intestine
         • Haustra

2. Macro
   a. Esophagus
   b. Stomach
      i) Parts
         • Fundus
         • Body
         • Pylorus
      ii) Openings
         • Cardiac
         • Pyloric
      iii) Curvatures
Abdomen and Superficial Structures Including Introductory Pediatric and Musculoskeletal

- Greater
- Lesser

iv) Surfaces
  - Anterior/superior
  - Posterior/inferior

c. Small intestine
  i) Duodenum
     - Superior/first part
     - Descending/second part
     - Transverse/third part
     - Ascending/fourth part
  ii) Jejunum
  iii) Ileum
     - Ileocecal valve

d. Large intestine/Colon
  i) Cecum
  ii) Appendix
  iii) Ascending colon
  iv) Transverse colon
  v) Descending colon
  vi) Sigmoid colon
  vii) Rectum
  viii) Anal canal

3. Relational anatomy

C. Variants
   1. Duplication cyst

D. Function
   1. Esophagus
   2. Stomach
      a. Storage of food
      b. Mixing of food with gastric secretions until chyme is formed
      c. Permits food to empty slowly into duodenum
   3. Small intestine
      a. Absorption of food
      b. Break down of starch, protein and fats
      c. Emulsifies fats
   4. Colon
a. Intestinal walls reabsorb liquid
b. Wastes formed into solid feces

E. Laboratory Values
1. WBC
2. Hematocrit
3. Hemoglobin

F. Indications
1. Pain
2. Vomiting
3. Bleeding
4. Fever
5. Leukocytosis
6. Palpable mass
7. Distension related to bowel obstruction

G. Sonographic Technique
1. Patient preparation
   a. Six hour fast is recommended
2. Transducer selection
3. Patient positioning
   a. Supine
   b. Decubitus
4. Imaging optimization
   a. High contrast grey scale
   b. Color Doppler
5. Imaging protocol
   a. Survey should include the entire region of the bowel with methodical approach to avoid omission of regions.

H. Normal Sonographic findings
1. The ‘gut signature’ (from lumen outward)
   a. Hyperechoic mucosal layer
   b. Hypoechoic intramural layer
   c. Hyperechoic submucosa
   d. Hypoechoic muscularis propria
   e. Hyperechoic serosal layer
2. Compressibility
   a. Normal bowel is compressible
   b. Thickened or infiltrated bowel is generally not
3. Measurements
   a. Wall
   b. Appendix
4. Doppler

I. Pathology
   1. Inflammation
      a. Crohn’s disease
         i) Definition
            • Manifestation of inflammatory bowel disease (IBD) consisting of chronic,
              relapsing bouts of inflammation – most commonly involving terminal portion of
              the ileum
         ii) Complications
            • Fistula formation
            • Abscess
            • Perianal ulcerations/fistulae
            • Luminal narrowing
            • Stricture
            • Perforation
            • Fistula formation
            • Inflammatory mass
         iii) Clinical presentation
            • Pain
            • Fever
            • Weight loss
            • Diarrhea
         iv) Sonographic findings
            • Related to the abnormal or involved bowel
              o Target pattern
              o Thickened bowel wall
              o Thickened bowel loops
              o Non-compressibility of bowel loop
              o Increased echogenicity of mesentery – creeping fat
              o Increased blood flow as assessed on color Doppler imaging
              o Lymphadenopathy
            • Related to complications
              o Abscess/phlegmon
              o Fistula
Dilated fluid filled bowel proximal to a site of stricture or luminal compromise, suggesting incomplete bowel obstruction

v) Correlative and/or prior imaging
vi) Treatment

b. Appendicitis
i) Definition
   • Inflammation usually caused by obstruction
ii) Complications
   • Rupture
   • Abscess
iii) Clinical presentation
   • RLQ pain
   • Fever
iv) Sonographic findings
   • RLQ target pattern
   • Increased size of appendix
   • Increased size of appendiceal wall
   • Non-compressibility
   • Pericecal fluid
   • Abscess
   • Appendicolith
   • Fecalith
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   • Increased size of appendiceal wall
   • Non-compressibility
   • Pericecal fluid
   • Abscess
   • Appendicolith
   • Fecalith
• Non-compressibility of wall
• Pericolonic fat changes
  o Poorly defined hyperechoic zones
• Abscess
  o Localized fluid collections
  o Acoustic enhancement
• Fistula
  o Linear echogenic tracts connecting bowel to other structures
• Correlative and/or prior imaging
• Treatment
d. Ischemic bowel disease
  i) Definition
    • Inadequate blood supply resulting in vascular compromise to bowel
  ii) Clinical presentation
    • Abdominal pain
    • Weight loss
    • Post-prandial pain
  iii) Sonographic findings
    • Stenosis of arterial supply
    • Occlusion of arterial supply
    • Gut wall thickening
  iv) Correlative and/or prior imaging
  v) Treatment

2. Neoplasia
   a. Adenocarcinoma
      i) Definition
        • Malignant primary neoplasia derived from glandular tissue
          o Most common location is the colon; rectum and rectosigmoid
      ii) Clinical presentation
        • Dependent on location
        • Asymptomatic
        • Obstruction
        • Anemia
        • Change in bowel habits
        • Weight loss
      iii) Sonographic findings
• Most common appearance is that of a mural mass
• Thickened bowel wall
• Target appearance or pseudokidney
• Hypoechoic mass
• Loss of continuity of wall layers

iv) Correlative and/or prior imaging
v) Treatment

b. Leiomyoma
i) Definition
• Benign tumor composed of smooth muscle cells – stomach is most common GI location
ii) Clinical presentation
• Obstruction
• Weight loss
iii) Sonographic findings
• Thickened bowel wall
• Target pattern
• Variable echogenicity
iv) Correlative and/or prior imaging
v) Treatment

c. Leiomyosarcoma
i) Definition
• Malignant sarcoma containing smooth muscle cells
• Most commonly referred to as gastrointestinal stromal tumor or GIST
ii) Clinical presentation
• Obstruction
• Weight loss
iii) Sonographic findings
• Target pattern
• Thickened bowel wall
• Mural mass
• Variable echogenicity
  o Complexity related to the tendency of necrosis
iv) Correlative and/or prior imaging
v) Treatment

d. Lymphoma
i) Definition
   • Neoplastic primary of lymphoid tissue – may be primary to GI tract or metastatic

ii) Clinical presentation
   • Dependent on location
   • Asymptomatic
   • Obstruction
   • Weight loss

iii) Sonographic findings
   • Thickened bowel wall, with a strong tendency to hypoechogenicity
   • Target pattern
   • Hypoechoic mass

iv) Correlative and/or prior imaging

v) Treatment

e. Metastases
i) Definition
   • Growth of malignant cells distant from primary site

ii) Clinical presentation
   • Dependent on location
   • Asymptomatic
   • Obstruction
   • Weight loss

iii) Sonographic findings
   • Thickened bowel wall
     o May produce a focal mass or a plaque like infiltration of the bowel wall
   • Target pattern
   • Varying echogenicity

iv) Correlative and/or prior imaging

v) Treatment

3. Obstruction
a. Mechanical
i) Physical cause of obstruction of GI lumen

ii) Clinical presentation
   • Abdominal pain
   • Vomiting
   • Abdominal distention

iii) Sonographic findings
• Dilated bowel loops
• Hyperperistalsis
• Site of obstruction is marked by a caliber alteration of the dilated bowel. A mass or bowel wall thickening may be shown at this site.

iv) Correlative and/or prior imaging
v) Treatment

b. Paralytic ileus
i) Paralysis of intestinal musculature
ii) Clinical presentation
• Change in bowel habits
• Abdominal distention
• Abdominal pain

iii) Sonographic findings
• Dilated bowel loops
• Fluid-fluid level in dilated loop
• Aperistaltic bowel loops

iv) Correlative and/or prior imaging
v) Treatment

c. Intussusception
i) Definition
• Invagination of segment of bowel into adjoining section

ii) Clinical presentation
• Abdominal pain
• Abdominal distention
• Vomiting
• Stool mixed with blood and mucus

iii) Sonographic findings
• Multiple concentric rings
• Target appearance
• Pseudokidney appearance

iv) Correlative and/or prior imaging
v) Treatment

d. Volvulus
i) Definition
• Torsion of a loop of bowel

ii) Clinical presentation
• Abdominal pain
• Vomiting

iii) Sonographic findings
• Dilated bowel loops
• Swirling / whirlpool sign of mesentery

iv) Correlative and/or prior imaging

v) Treatment
e. Hypertrophic pyloric stenosis (HPS)
i) Definition
• Pediatric condition characterized by hypertrophy and hyperplasia of circular muscle resulting in elongation of pylorus and constriction of canal

ii) Clinical presentation
• Projectile vomiting
• Failure to thrive
• Age range

iii) Sonographic findings
• Target pattern
• Elongated canal
• Thick muscular wall
• Non-visualization of emptying of stomach contents into duodenum

iv) Correlative and/or prior imaging

v) Treatment
Section IX: Abdominopelvic Wall and Cavities

1. Describe the normal anatomy and variants, function, sonographic technique and appearance of the abdominopelvic wall and cavities
2. Correlate the clinical indications and laboratory values associated with abdominopelvic wall and cavity pathology
3. Describe abdominopelvic wall and cavity pathology in terms of sonographic appearances, sequelae and associated pathologies

IX. Abdominopelvic Wall and Cavities

A. Abdominopelvic Wall and Cavities
   1. Embryology
   2. Anatomy
      a. Abdominal wall layers
         i) Skin
         ii) Superficial fascia
         iii) Subcutaneous fat
         iv) Muscles
      b. Abdominopelvic muscles
         i) Rectus
         ii) External oblique
         iii) Internal oblique
         iv) Transversus abdominus
         v) Psoas
         vi) Quadratus lumborum
      c. Abdominal wall fascia, tendons and ligaments
         i) Linea alba
         ii) Linea semilunaris
         iii) Inguinal ligament
      d. Cavities
         i) Peritoneal layers
            • Layers
               o Parietal
               o Visceral
         ii) Peritoneal compartments
            • Lesser sac - left posterior subhepatic
               o Foramen of Winslow
            • Greater sac
o Subdiaphragmatic
   ~ Right
   ~ Left anterior – perigastric
   ~ Left posterior – perisplenic
o Subhepatic
   ~ Right
      · Hepatorenal
      · Morison’s pouch
   ~ Left anterior
      · Hepatogastric recess
o Subsplenic
o Perinephric
o Pelvic
   ~ Male
      · Rectovesicle
   ~ Female
      · Vesicouterine
      · Rectouterine (Pouch of Douglas (POD))

iii) Peritoneal reflections
    • Ligaments/mesentery
    • Falciform
    • Coronary
    • Triangular
    • Hepatoduodenal
    • Hepatogastric
    • Gastrocolic
    • Gastroplenic
    • Transverse mesocolon
    • Sigmoid mesocolon
    • Small bowel mesentery

iv) Peritoneal omenta/compartments
    • Greater
    • Lesser

v) Retroperitoneal compartments
    • Anterior pararenal
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- Perirenal
  - Anterior perirenal fascia/Gerota’s fascia
  - Posterior perirenal fascia/Zuckerkandl’s fascia
- Posterior pararenal
- Retrofascial space
  - Psoas muscle
  - Quadratus lumborum muscle

vi) Retroperitoneal organs/structures
- Diaphragmatic crura
- Vessels
  - Aorta
  - Inferior vena cava (IVC)
  - Superior mesenteric
  - Hepatic
  - Splenic
- Pancreas
  - Second, third, and fourth parts of the duodenum
- Kidneys
- Adrenals
  - Ascending, descending colon
  - Distal portion of common bile duct
  - Lymph nodes

vii) Extraperitoneal spaces
- Space of Retzius
- Rectovesicle
- Right, left paravesicle
- Presacral
- Right, left pararectal

viii) Extraperitoneal organs
- Female
  - Urinary bladder
  - Distal ureters
  - Uterus
  - Rectum
- Male
  - Urinary bladder
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- Distal ureters
- Prostate
- Seminal vesicles
- Rectum
- Posterior pararenal
- Retrofascial space
  
  ix) Thoracic cavity
  - Pleural cavities
    - Right
    - Left
  - Layers
    - Parietal
    - Visceral
  - Organs
    - Lungs
    - Heart
  
  x) Relational anatomy

3. Variants
   a. Urachal anomalies

4. Indications
   a. Fever
   b. Pain
   c. Palpable mass
   d. Bleeding
   e. Trauma
   f. Post surgery
   g. Distended abdomen
   h. Dyspnea
   i. Differentiate loculated from nonloculated intrathoracic mass
   j. Differentiate pleural from thoracic masses
   k. Delineation of opacities noted on chest x-ray

5. Sonographic Technique
   a. Patient preparation
   b. Transducer selection
   c. Patient positioning
   d. Image optimization
   e. Image protocol
6. Normal sonographic appearance
   a. Abdominal wall
      i) Skin
         • Thin, highly echogenic
      ii) Fat
         • Variable echogenicity
         • Usually hypoechoic to skin and muscle
      iii) Muscle
         • Usually hyperechoic to subcutaneous fat
   b. Peritoneal cavity
   c. Retroperitoneum
   d. Thoracic cavity
      i) The lung-air/visceral pleura interface is echogenic

7. Pathophysiology
   a. Abdominal wall
      i) Hernia
         • Definition
            o Protrusion of bowel or fat through opening in ventral wall where no muscle is present – may be acquired or congenital
         • Types/locations
            o Ventral
               ~ Umbilical
               ~ Spieghelian
               • Spieghelian hernia (inferior lumbar or incisional)
            • Sonographic technique
               o High resolution linear or curved linear array transducer
               o Fascial defect
               o Valsalva maneuver
      ii) Rectus sheath hematoma
         • Definition
            o Focal collection of extravasated blood contained within rectus sheath
         • Clinical presentation
            o Pain
            o Palpable mass
            o Decreased hematocrit
         • Sonographic findings may include
            o Focal mass
i) Varying echogenicity
   - Anechoic (early)
   - Echogenic (later)

iii) Abscess
   - Definition
     - Focal collection of pus from disintegration of tissue
   - Clinical presentation
     - Focal pain
     - Palpable mass
     - Fever
     - Elevated WBC
   - Sonographic findings may include
     - Focal well-defined mass
     - Varying echogenicity fluid-fluid levels
     - Hyperechoic foci with shadowing

iv) Neoplasia
   - Desmoid
     - Definition
       - Benign, unencapsulated fibromatous tumor arising from muscle sheath
     - Clinical presentation
       - Palpable mass
     - Sonographic findings
       - Focal mass
       - Varying echogenicity
     - Correlative and/or prior imaging
     - Treatment
   - Metastases
     - Definition
       - Growth of malignant cells from a distant primary site
     - Clinical presentation
     - Sonographic findings
       - Focal mass
       - Varying echogenicity
     - Correlative and/or prior imaging
     - Treatment

b. Abdominopelvic cavity
   i) Adenopathy (lymphadenopathy)
• Definition
  o Enlargement of lymph nodes caused by inflammatory processes, primary tumor, or metastatic spread of cancer
• Sites
  o Para-aortic chain
  o Mesenteric and celiac
  o Pelvic adenopathy
  o Iliac chain
  o Hypogastric chain
  o Intraperitoneal chain
    ~ Splenic hilum
    ~ Parapancreatic
    ~ Hepatic hilum
• Sonographic findings
  o Mass(es)
  o Homogeneous, low to medium level internal echoes
  o Para-aortic nodes may be individually enlarged or present as lobulated or sheet-like masses
  o Enlarged mantle of nodes can surround aorta and possibly IVC
  o Displacement of superior mesenteric and celiac vessels anteriorly
  o Anterior displacement of aorta
  o Compression of IVC
• Correlative and/or prior imaging
• Treatment
  ii) Retroperitoneal fibrosis
    • Definition
      o Chronic inflammatory process with fibrous tissue proliferation confined to paravertebral and central region of abdomen
    • Clinical presentation
      o Urinary obstruction
      o Nausea and vomiting
    • Sonographic findings
      o Solid, low-level echo mass anterior and lateral to the aorta
  iii) Retroperitoneal tumors
    • Primary malignant
      o Liposarcoma
        ~ Arising from fat
      o Leiomyosarcoma
Arising from smooth muscle
- Rhabdomyosarcoma

Arising from striated muscle
- Fibrosarcoma
- Arising from connective tissue

Benign retroperitoneal tumors
- Lipoma
  - Arising from fat
- Leiomyoma
  - Arising from smooth muscle
- Rhabdomyoma
  - Arising from striated muscle
- Fibroma
  - Arising from connective tissue

Sonographic findings
- Solid mass
- Variable size and appearance
- May invade surrounding tissues
- Displacement of vasculature (usually anteriorly)
- Compression of the IVC and urinary bladder
- Obstruction of the kidneys, ureters, extrahepatic bilary ducts

Correlative and/or prior imaging

Treatment
c. Retroperitoneal fluid collections
  i) Hematoma
  ii) Lymphocele
  iii) Urinoma
  iv) Abscess
  v) Clinical presentation
    - Abdominal pain
    - Fever
    - Palpable mass
  vi) Sonographic findings
    - Focal mass
    - Varying appearance
  vii) Correlative and/or prior imaging
  viii) Treatment
d. Peritoneal fluid collections
   i) Ascites
      - Excessive accumulation of serous fluid in peritoneal cavity
      - Clinical presentation
      - Sonographic findings
         o Transudative ascites
            ~ Reflection of the dynamics of the peritoneal cavity
            ~ Anechoic
            ~ Free floating bowel loops
            ~ Fluid conforms to surrounding structures
            ~ Gallbladder wall thickening
            ~ Changes with patient position and probe pressure
         o Exudative ascites
            ~ Generally a reflection of inflammation or neoplasia
            ~ Echogenic debris
            ~ Septations
            ~ Matted bowel loops
            ~ Thickened interfaces
         o Loculated
            ~ Does not conform to surrounding structures
            ~ Does not change with patient position
      - Correlative and/or prior imaging
      - Treatment
   ii) Focal fluid collections
      - Types
         o Abscess
         o Lymphocele
         o Seroma
         o Biloma
      - Clinical presentation
         o Abdominal pain
         o Fever
         o Palpable mass
      - Sonographic findings
         o Focal mass
         o Variable echogenicity
            ~ Echogenic
e. Pseudomyxoma peritonei
   i) Definition
   • Metastatic spread of malignant tumor filling peritoneal cavity with mucinous material and gelatinous ascites. Tumor implants on peritoneal surfaces and producing adhesions.
     o Most commonly secondary to malignant tumor of the appendix.
   • Clinical presentation
     o Abdominal pain
     o Abdominal distention
   • Sonographic findings
     o Complex ascites
     o Non-mobile echogenic foci within ascites
     o Scalloping of the convexities of the spleen and liver
   • Correlative and/or prior imaging
   • Treatment

f. Peritoneal carcinomatosis
   i) Definition
   • Metastatic spread with diffuse involvement of peritoneum
   ii) Clinical presentation
   • Abdominal distention
   • Abdominal pain
   iii) Sonographic findings
   • Ascites
   • Irregular masses
   • Echo poor nodules
   • Thickening of peritoneum
   iv) Correlative and/or prior imaging
   v) Treatment
g. Thoracic cavity
   i) Pleural effusion
      • Accumulation of fluid in pleural space
      • Clinical presentation
         o Dyspnea
         o Chest pain
      • Sonographic findings
         o Supradiaphragmatic fluid
         o Variable echogenicity
         o Anechoic
         o Echogenic
      • Correlative and/or prior imaging
      • Treatment
h. Tumor
   i) Metastases
      • Growth of malignant cells distant from primary site
      • Sonographic findings
         o Pleural effusion
         o Pleural thickening
         o Solid nodules in pleural cavity
   ii) Mesothelioma
      • Rare, fatal pleural tumor
      • Sonographic findings
         o Diffuse pleural thickening
         o Calcifications
         o Pleural effusion
i. Lung abscess
   i) Localized collection of pus from disintegration of tissue
   ii) Sonographic findings
      • Thick, irregular walls
      • Echogenic debris
      • Expansion of entire mass with inspiration
j. Mediastinal lymphadenopathy
   i) Enlargement of mediastinal nodes
   ii) Sonographic findings
      • Homogeneous masses
• Calcification
• Large, homogeneous solid mass

k. Pericardial effusion
   i) Accumulation of fluid between pericardial layers
   ii) Sonographic findings
      • Fluid in pericardial sac
      • Varying echogenicity

l. Atelectasis
   i) Absence of air in all or part of lung
   ii) Sonographic findings
      • Wedge-shaped, highly echogenic homogeneous mass
      • Mobile, floating lung
      • Motion of lung with respiration
      • Pleural effusion

m. Lung consolidation
   i) Lung filled with fluid and cells
   ii) Sonographic findings
      • Wedge-shaped, echo-poor mass
      • Homogeneous
      • Motion with respiration
Section X: Interventional

1. List indications for performing invasive procedures
2. Describe the sonographic technique for interventional procedures
3. Correlate clinical indications and laboratory values associated with interventional procedures
4. Describe the role of sonography in interventional procedures
5. Identify normal and abnormal flow characteristics associated with pathology
6. List various complications associated with interventional procedures

X. Interventional

A. Clinical Indications
   1. Differentiating between benign process, primary cancer, metastatic disease, or infectious process and obtaining tissue for immunohistochemistry and molecular markers
   2. Obtaining tissue in order to test for specific tumor markers/receptors that may help to tailor treatment regimen
   3. Progression of disease
      a. Hepatitis
      b. Renal failure
      c. Cirrhosis
   4. Verifying the type and presence of disease
   5. Drainage of fluid or fluid collection
      a. Therapeutic tap
      b. Diagnostic tap
   6. Distinguishing between a fluid-filled mass and a solid tumor
   7. Obtain parenchymal sample to diagnose parenchymal disease
      a. Liver
      b. Kidney
      c. Transplanted organ
   8. Determine cause of organ dysfunction or rejection
   9. Obtain fluid for cells – diagnostic tap
      a. Amniotic
      b. Ascites
      c. Pleural
      d. Pericardial
   10. Obtain malignant tissue for clinical trials

B. Consent Process
   1. Risks of procedures
   2. Benefits of Procedure
   3. Alternatives to Procedure
4. How Procedure is Performed

5. Witnessing a Consent

6. Time Out

C. Types

1. Biopsy
   a. Removal and examination of living tissue from the living body
   b. Fine needle aspiration (FNA)
      i) Obtains cells from tissue
      ii) Capillary technique
      iii) 18-25 gauge needles
   c. Suction technique
      i) 20-26 gauge needles
   d. Core
      i) Obtain core sample of tissue
      ii) 14-20 gauge needles
      iii) 1 or 2 cm long samples
   e. Needle types
      i) Spinal
      ii) Franseen
      iii) Chiba
      iv) Echogenic tips
      v) Echogenic stylets
      vi) Catheters for aspirating large volumes of fluid

2. Percutaneous aspiration and/or drainage
   a. Paracentesis
   b. Thoracentesis
   c. Amniocentesis
   d. Fluid collections
   e. Indwelling catheter
      i) Percutaneous nephrostomy
      ii) Abscess

D. Contraindications

1. Bleeding disorder
   a. Laboratory values
      i) Prothrombin Time (PT)
      ii) PTT
      iii) INR
b. Bleeding correction methods
   i) Platelet therapy
   ii) Fresh frozen plasma (FFP)
2. Anti-coagulant therapy
3. Inability to give consent
4. Inability to cooperate
E. Risk Factors of Procedure
1. Bleeding
2. Infection
3. Seeding of biopsy tract with tumor
4. Inconclusive results
5. Inadequate sample
6. Organ damage or tissue adjacent to organ
7. Collapsed lung if thoracentesis
F. Sedation
1. General anesthesia/deep sedation
   a. Pediatric patient
   b. Patient and performing team comfort
   c. Patient asleep
   d. Greatest risk to patient
   e. Usually performed by anesthesia team
2. Conscious sedation
   a. Patient given medication to relieve anxiety
   b. Patient in given medication to relieve pain
   c. Pharmacology, sedation usually performed by radiologist/clinician with nurse
   d. Safer than deep sedation
3. Local anesthesia
   a. Most common
   b. Need patient cooperation
   c. Local skin anesthetized
   d. Least risk to patient
G. Sonographic Technique
1. Correlative image assessment
   a. CT, MRI, PET/CT, prior imaging
2. Patient preparation
   a. Consent
   b. Position
c. Location of lesion

3. Transducer selection

4. Sterile technique

5. Image optimization
   a. 2-D
   b. Doppler
      i) Avoid major blood vessels
      ii) Target areas of solid tumor with vascular blood flow; suggestive of viable tissue

6. Fusion technology
   a. US/CT
   b. US/MRI

7. Guidance Approach
   a. Transabdominal
   b. Endovaginal
   c. Transrectal
   d. Transperineal

8. Protocol
   a. Sonographic guidance
      i) Transducer guide attachment
      ii) Free hand technique
   b. Patient care techniques

H. Complications
1. Bleeding
2. Hematoma
3. Vasovagal Reaction
4. Arteriovenous Fistula
5. Pseudoaneurysm
6. Pneumothorax
7. Hemoptysis
8. Damage to Adjacent Organs
Section XI: Organ Transplant

I. List indications for organ transplant
II. Describe normal anatomy, organ function, sonographic technique and sonographic appearance related to organ transplant
III. Correlate clinical indications and laboratory values associated with organ transplant
IV. Describe associated pathologies and sequelae relative to organ transplants
V. Identify normal and abnormal flow characteristics and waveforms
VI. Correlate the method used to transplant the organ and expected sonographic findings

XI. Organ Transplant

A. Indications for Transplant
   1. Renal
      a. End-stage renal disease
         i) Adult polycystic kidney disease
         ii) Nephrotic syndrome
         iii) Renal failure
         iv) Diabetes
         v) Systemic lupus
         vi) Glomerulonephritis
         vii) Renovascular disease
         viii) Patients on renal dialysis
   2. Liver
      a. End-stage liver disease
         i) Cirrhosis
         ii) Fulminate liver failure
         iii) Cancer
         iv) Biliary atresia
         v) Cholestatic liver disease
         vi) Hepatocellular disease
         vii) Vascular disease

B. Transplant Anatomy
   1. Full organ
      a. Renal
         i) Single kidney
            • Donor
               o Cadaver
               o Living
ii) Double kidney
   - Donor
     - Cadaver
       - Pediatric en-bloc
       - Horseshoe kidney
   - Liver
     - Cadaver

2. Partial organ
   a. Liver
      i) Liver segment
         - Single hepatic artery
         - Single hepatic vein
         - Single portal vein
      ii) Donor
         - Cadaver
         - Living

C. Laboratory Values
   1. Organ and complication specific
      a. Kidney: Creatinine
      b. Liver: AST/ALT, Bilirubin, Alkaline Phosphatase
      c. Anti-rejection medication levels

D. Indications
   1. Baseline study
      a. Within 24 hours post-surgery
      b. Confirm patency of vessels, anastomoses
   2. Abnormal laboratory values
   3. Decreased urine output
   4. Follow progression of abnormal Doppler waveforms
   5. Monitor and follow complication
      a. Fluid collection
         i) Hematoma
         ii) Abscess
         iii) Lymphocele
         iv) Urinoma
      b. Post-transplant lymphoproliferative disorder; long-term complications
      c. Anastomotic leaks
      d. Anastomotic and vascular strictures
6. Vascular thrombosis

7. Rejection and other complications
   a. Organ specific
      i) Renal
         - Acute tubular necrosis
            - More common in cadaver transplants
            - Related to length of cold ischemic time of transplant prior to transplantation
         - Acute rejection
         - Chronic rejection
         - Vessel thrombosis
            - Arterial
            - Venous
         - Anti-rejection drug toxicity
      ii) Liver
         - Hepatic artery thrombosis
         - Hepatic artery stenosis
         - Portal vein stenosis
         - Portal vein thrombosis
         - Biliary complications
         - Reoccurrence of underlying disease

8. Obstruction

9. Biopsy guidance

10. Biopsy complications

E. Sonographic Technique

1. Imaging protocol for specific organ
   a. Gray scale images
      i) Anastomotic sites
      ii) Thrombus
      iii) Fluid collections
         - Hematoma
         - Abscess
         - Organ specific
            - Renal
               - Urinoma
               - Lymphocele
            - Liver

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iv) Evidence of an obstruction
   - Organ specific
     - Renal
       - Hydronephrosis
     - Liver
       - Dilated biliary ducts
b. Color Doppler images
   i) Anastomotic sites
   ii) Patency
   iii) Stenosis or kinking of vessel
   iv) Thrombus
v) Post biopsy
   - Pseudoaneurysm
   - Arteriovenous Fistula (AVF)
c. Spectral Doppler waveforms
   i) Anastomotic sites
   ii) Patency
   iii) Stenosis or kinking of vessel
   iv) Thrombus
       - Venous
         - Lack of venous signal
         - Bi-phasic or high resistive arterial signal
       - Arterial
         - Lack of arterial signal
         - Lack of venous signal
         - Evidence of collateral flow
v) Post biopsy
   - Pseudoaneurysm
   - Arteriovenous Fistula (AVF)
vi) Doppler indices
   - Resistive index (RI)
vii) Waveform analyses
   - Velocity
   - Turbulence
   - Flow direction
• Flow characteristics
  o Venous
    ~ Phasicity
    ~ Spontaneous
  o Arterial
    ~ Amount of diastolic flow
d. Correlative and/or prior imaging
Section XII: Breast

1. Describe the normal breast anatomy including the sonographic appearance
2. Describe the types of benign breast conditions, breast pathology and correlating sonographic appearances
3. Compare the roles of mammography, sonography, computerized tomography and magnetic resonance imaging (MRI)
4. Discuss various interventional procedures utilized in breast pathology diagnosis

XII. Breast

A. Embryology

B. Anatomy/Normal Variants

1. Surface of the breast
   a. Tail of Spence
   b. Nipple
      i) Areola
   c. Tissues
      i) Fat
      ii) Ligaments
      iii) Glandular tissue
      iv) Ductal system

2. Layers
   a. Subcutaneous layer
      i) Fat surrounded by connective tissue
      ii) Cooper’s ligaments
   b. Mammary layer
      i) Functional lobes
         • Acini
      ii) Connective tissue septa
         • Cooper’s ligaments
      iii) Lobules
      iv) Lobes
         • Lactiferous duct drains each lobe
         • Terminal Ductal Unit (TDLU)
            ~ Extralobular terminal duct
            ~ Intralobular terminal duct
            ~ Lobule (consisting of acini)
   c. Retromammary layer
      i) Muscles/chest wall
         • Pectoralis major
• Pectoralis minor
• Ribs
• Chest wall

3. Lymphatics
   a. Lymph vessels empty into lymph nodes
   b. Axillary lymph nodes
      i) Normal
      ii) Abnormal
   c. Pectoral nodes
   d. Internal mammary nodes
   e. Intramammary nodes

4. Variants
   a. Amastia
   b. Amazia
   c. Athelia
   d. Hypoplasia
   e. Polymastia
   f. Polythelia

C. Physiology and Function
   1. Terminal Ductal Unit (TDLU)
   2. Lobes (consisting of TDLUs) are drained by lactiferous ducts
   3. Lactiferous ducts converge to nipple and dilate to form lactiferous sinuses (ampullae)
   4. Lactiferous sinuses are milk reservoirs during lactation

D. Indications
   1. Palpable breast and/or axillary lump
   2. Breast pain
   3. Skin redness or nipple changes
   4. Correlation with mammography or MRI
   5. Pregnant or lactating patient
   6. Implant rupture after breast augmentation
   7. Post-surgical or post-irradiated breast
   8. Guidance for interventional procedures
   9. Screening in heterogeneous or dense breasts
   10. Detection and biopsy of abnormal axillary nodes

E. Sonographic Technique
   1. Whole breast versus targeted area
      a. Correlation with prior imaging
b. Correlation with palpable lump or pain

2. Transducer selection

3. Patient positioning

4. Image optimization

5. Annotation methods
   a. Laterality (right/left)
   b. Orientation
      i) Radial/antiradial
      ii) Sagittal/transverse
      iii) Oblique
   c. Location within breast
      i) Clock face
      ii) Distance from nipple
      iii) Labeled diagram of breast

6. Tissue composition
   a. Homogenous - fat
   b. Homogeneous - fibroglandular tissue
   c. Heterogeneous

7. Mass evaluation
   a. Measurements
      i) Three measurements in two orthogonal planes
         • Largest measurement defines longest axis (plane) of the mass
         • Next measurement is perpendicular to first measurement
         • Final measurement is from a view orthogonal to the first plane
   b. Descriptors
      i) Shape
         • Oval
         • Round
         • Irregular
      ii) Orientation
         • Parallel
         • Not parallel
      iii) Margin
         • Circumscribed
         • Not circumscribed
            o Indistinct
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- Angular
- Mirolobulated
- Spiculated

iv) Echo pattern
   - Anechoic
   - Hyperechoic
   - Hypoechoic
   - Isoechoic
   - Heterogeneous
   - Complex
   - Cystic
   - Solid

v) Posterior features
   - No posterior features
   - Enhancement
   - Shadowing
   - Combined pattern

c. Calcifications
d. Associated features
   i) Architectural distortion
   ii) Duct changes
   iii) Skin changes
   iv) Edema
   v) Vascularity
e. Doppler
   i) Settings for low-flow states
   ii) Fremitus
f. Elastography
g. Special cases
   i) Simple cyst
   ii) Clustered microcysts
   iii) Complicated cyst
   iv) Mass in or on skin
   v) Foreign body including implants
   vi) Lymph node
   vii) Vascular abnormalities
• AVM
• Mondor disease
  viii) Postsurgical fluid collection
 ix) Fat necrosis

8. ACR BI-RADS Assessment Categories
  a. Category 0: Incomplete
  b. Category 1: Negative
  c. Category 2: Benign
  d. Category 4: Suspicious
     i) 4A = Low
     ii) 4B = Moderate
     iii) 4C = High
  e. Category 5: Highly suggestive of malignancy
  f. Category 6: Known Biopsy-proven malignancy

9. Pitfalls
  a. Pseudomass (from normal fat lobules)
  b. Large fatty breast
  c. Shadowing from fibroglandular tissue versus very large mass

F. Sonographic characteristics
1. Benign cyst
   a. Circumscribed, thin margins
   b. Oval shape
   c. Echogenicity: anechoic
   d. Posterior enhancement
   e. No internal vascularity (color Doppler)

2. Probably benign solid
   a. Circumscribed margins
   b. Oval shape (includes 2-3 large lobulations)
   c. Parallel orientation

3. Probably benign cystic
   a. Clustered microcysts
   b. Complicated cyst

4. Suspicious
   a. Irregular shape
   b. Not circumscribed margins
   c. Nonparallel orientation
   d. Complex cystic and solid
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e. Duct extension
f. Shadowing
g. Intraductal calcifications
h. Architectural distortion
   i) Straightening or thickening of Cooper’s ligaments
   ii) Echogenic rim
   iii) Obliteration of surrounding tissue planes
   iv) Distortion of ductal pattern
i. Skin overlying tumor is flattened or retracted or thickened
j. Echogenic extensions from mass or increased echogenicity anterior to mass (desmoplasia)

G. Pathology and Pathophysiology
   1. Cyst
      a. Clinical signs
      b. Sonographic findings
         i) Circumscribed
         ii) Anechoic
         iii) Posterior enhancement
   2. Complicated cyst
      a. Clinical signs
         i) May be related to inflammation or hemorrhage within cyst
      b. Sonographic findings
         i) Wall thickening or irregularities
         ii) Septations
         iii) Internal echoes
   3. Fibrocystic changes
      a. Clinical signs
      b. Sonographic findings
         i) Bilateral macrocysts; microcysts and cystic clusters possible
         ii) Increased fibrous stroma
         iii) Ductal ectasia
   4. Gynecomastia
      a. Causes
         i) Drugs
         ii) Endocrine active tumors
         iii) Reduced testosterone states
      b. Clinical findings
         i) Male subareolar breast lump
ii) Male breast tenderness

c. Sonographic findings
   i) Hypoechoic tissue with ducts radiating from the nipple
   ii) Color Doppler findings similar to female breast tissue

5. Solid mass – benign
   a. Fibroadenoma
      i) Most common breast tumor
      ii) Primarily in young women
      iii) Growth stimulated by estrogen
      iv) Clinical findings
         • Firm, rubbery, freely mobile
         • Slow growing
      v) Sonographic findings
         • Smooth, rounded margins
         • Low-level homogeneous internal echoes
         • Possible posterior acoustic enhancement
         • Typically hypoechoic
   b. Lipoma
      i) Fatty tissue
      ii) Clinical findings
         • May grow large before being clinically detected
         • Usually found in middle-aged/postmenopausal women
         • Large, soft, poorly demarcated, difficult to delineate from surrounding breast tissue
      iii) Sonographic findings
         • Typically smooth walls
         • Hypoechoic similar to breast fat
         • Posterior acoustic enhancement
         • Easily compressible
   c. Fat necrosis
      i) Causes
         • Trauma
         • Surgery
         • Radiation treatment
         • Mastitis
         • Other
      ii) Clinical findings
• Spherical nodule, superficial under layer of calcified necrosis

iii) Sonographic findings
• Irregular
• Complex low-level echoes
• May mimic malignant mass
• Acoustic shadowing may or may not be present

d. Abscess
i) Clinical findings
• Pain
• Swelling
• Redness
• Febrile
• Enlarged axillary nodes

ii) Sonographic findings
• Complex
• Diffuse increased echogenicity of the breast (if mastitis is present)
• Hyperemia with color and/or power Doppler

e. Papillomas
i) Arise from lining of the breast ducts

ii) Retroareolar area most common

iii) Clinical findings
• Bloody discharge from single duct

iv) Sonographic findings
• Tiny papilloma may not be detected
• May cause dilatation of single duct
• Intracystic papillomas
• Soft tissue mass growing into lumen of cystic lesion

f. Sebaceous cyst
i) Obstruction of sebaceous gland or hair follicle within dermal layer of skin

ii) Sonographic findings
• Rounded, well-defined mass with low to medium level echogenicity
• Wall calcification possible

g. Phyllodes
i) Borderline or transitional form between benign and malignant tumor

ii) Clinical findings
• Can grow quite large
6. Solid mass – malignant
   a. Ductal carcinoma in-situ (DCIS)
      i) Most common non-invasive carcinoma
      ii) Clinical findings
         • Asymptomatic
         • Palpable mass
         • Nipple discharge
      iii) Mammography findings
         • Microcalcifications
   iv) Sonographic findings
      • Intraductal mass
      • Microcalcifications
      • Irregular ductal dilatation
      • Architectural distortion
   b. Lobular carcinoma in-situ (LCIS)
      i) Generally affects premenopausal women
      ii) Distinct tumor not a feature
      iii) Sonographic findings
         • Bilateral
         • Multicentric
         • Mimics fibrocystic disease
   c. Invasive Ductal Carcinoma (IDC)
      i) Extension past duct and into stroma
      ii) Most common malignancy
      iii) Clinical findings
         • Hard and fixed mass
         • Skin dimpling or skin/nipple retraction
      iv) Mammography findings
         • Asymmetric
         • Radiopaque speculated mass
         • Microcalcifications
         • Thickened and retracted Cooper’s ligaments
      v) Sonographic findings
- Solid mass
- Hypoechoic
- Heterogeneous
- Taller-than-wide orientation
- Distal acoustic shadowing
- Possible
  - Microlobulation
  - Microcalcification
  - Ductal extension/branch pattern
  - Fascial plane disruption

d. Invasive lobular carcinoma (ILC)
  i) Bilateral
  ii) Multicentric
  iii) Multifocal
  iv) Mammography
     - Can underestimate extent
  v) Sonographic findings
     - Possible speculated mass mimicking invasive ductal carcinoma
     - Architectural distortion

e. Medullary
  i) Clinical findings
     - Circumscribed, non-tender, often large palpable mass
     - May be mildly compressible and movable
     - Often located in peripheral breast
     - Low incidence of lymph node involvement
     - Prevalence in younger females
  ii) Sonographic findings
     - Solid hypoechoic mass; round, oval or lobulated shape
     - Circumscribed margins; surface irregularities possible
     - Homogeneous/mildly heterogeneous
     - Acoustic enhancement
     - Hemorrhage or cystic degeneration may be noted

f. Colloid/mucinous
  i) Clinical findings
  ii) Sonographic findings
     - Well-defined margins
• Hypoechoic or isoechoic to fat
• Homogeneous
• Possible microlobulations

g. Papillary
   i) Clinical findings
      • Subareolar palpable mass
      • May protrude and effect skin
      • May cause skin dimpling and retraction
      • Possible bloody nipple discharge
   ii) Sonographic findings
      • Mimics non-invasive papillary carcinoma or papilloma

h. Phyllodes tumor (Phylloides)
   i. Paget’s disease

7. Post surgical findings
   a. Implant rupture
      i) Stepladder sign
      ii) Stepladder sign can be confused with normal double lumen implant
   b. Silicone rupture
      i) Snow storm appearance
      ii) Silicone lymph nodes

8. Sonography-guided interventional procedures
   a. Cyst aspiration
   b. Fine needle aspiration cytology (FNAC)
   c. Drainage procedure
   d. Preoperative needle wire localization
   e. Core needle biopsy
   f. Vacuum assisted needle biopsy
   g. Sentinel node biopsy

9. Related imaging
   a. Mammography
   b. Computed tomography
   c. Magnetic resonance
   d. Computer Aided Detection (CAD)
   e. Elastography
   f. Positron Emission Mammography (PEM)
   g. Molecular Breast Imaging (MBI)
   h. Doppler examination
Section XIII: Neck

1. Describe the anatomy, physiology, pathophysiology, laboratory tests and pathology of the thyroid and parathyroid glands
2. Recite the protocol for sonographic assessment of the neck and relational anatomy
3. Discuss relational anatomy of the thyroid, parathyroid and other neck structures
4. Define other neck masses evaluated with sonography
5. Explain cervical lymph node mapping pre- and post-thyroidectomy

XIII. Neck

A. Embryology

B. Normal Thyroid Anatomy and Variants
   1. Description
      a. Right and left lobes located anterolateral to trachea and esophagus
   2. Structure
      a. Right and left lobe connected in midline by isthmus
         i) Normal variant is pyramidal lobe arising from isthmus
         ii) Four parathyroid glands
      b. Surrounding musculature
         i) Anterior
            • Strap muscles
               o Sternohyoid
               o Omohyoid
               o Sternothyroid
            • Sternocleidomastoid
         ii) Posterior
            • Longus colli muscle
      c. Vascular
         i) Arterial
            • Superior thyroid
               o Right and left
            • Inferior thyroid
               o Right and left
         ii) Venous
            • Superior and middle thyroid veins into jugular vein
            • Inferior thyroid vein into innominate vein

C. Function and Physiology
   1. Maintains body metabolism, growth, and development
a. Endocrine function  
   i) Secretes hormones  
      • Thyroxine (T4)  
      • Triiodothyronine (T3)  
      • Affect metabolic rate  
   ii) Calcitonin affects calcium metabolism  
   iii) Thyroid stimulating hormone (TSH)  
      • Controls thyroids hormone secretions  

b. Iodide ingestion  
   i) Necessary for production of thyroxine  

c. Laboratory tests  
   i) Serum  
      • Triiodothyronine (T3)  
      • Thyroxine (T4)  
      • Calcitonin  
      • Thyroid stimulating hormone (TSH)  
      • Thyroidglobulin  

D. Indications for Sonographic Examination  
   1. Palpable enlargement  
   2. Abnormal thyroid hormone level(s)  
   3. Palpable mass in neck/thyroid  
   4. Swelling of neck  
   5. Asymmetry of neck  
   6. Redness and/or tenderness  
   7. Hypercalcemia  
   8. Post thyroidectomy  

E. Sonographic Technique  
   1. Patient preparation  
   2. Transducer selection  
   3. Patient positioning  
   4. Image optimization  
   5. Examination protocol  

F. Pathology and Pathophysiology  
   1. Hypothyroidism  
      a. Under secretion of thyroid hormones  
         i) Clinical signs
• Myxedema
• Weight gain
• Hair loss
• Increased tissue around eyes
• Lethargy
• Intellectual and motor slowing
• Cold intolerance
• Constipation
• Deep, husky voice

ii) Causes
• Low intake of iodine (goiter)
• Thyroid hormone failure
• Diseases of the hypothalamus or pituitary

2. Hyperthyroidism
   a. Over secretion of thyroid hormones
      i) Clinical signs
         • Dramatic increase in metabolic rate
         • Weight loss
         • Increased appetite
         • Nervous energy
         • Tremor
         • Excessive sweating
         • Heat intolerance
         • Cardiac palpitations
         • Exophthalmos
      ii) Causes
         • Abnormal hormone secretion
         • Localized neoplasm cause overproduction of hormone(s)
         • Graves’ disease

3. Thyroid nodules
   a. Variable gland enlargement
   b. Causes
      i) Thyroiditis (ill-defined nodular areas)
      ii) Neoplasm (benign or malignant)
      iii) Nodular hyperplasia
      iv) Hashimoto’s thyroiditis (micronodules)
c. Sonographic findings
   i) Diffuse enlargement
   ii) Hypoechoic, heterogeneous without palpable nodules
   iii) Markedly increased vascularity
       • Increased peak systolic velocities in inferior thyroidal artery

4. Toxic multinodular goiter
   a. Grave’s disease
   b. Clinical signs
      i) Hypermetabolism
      ii) Diffuse toxic goiter
      iii) Exophthalmos
      iv) Cutaneous formations
         • Periorbital
         • Dorsum of feet
   c. Cause
      i) Autoimmune
   d. Sonographic findings
      i) Diffuse enlargement
      ii) Hypoechoic, heterogeneous without palpable nodules
      iii) Markedly increased vascularity
          • Thyroid inferno
          • Increased peak velocities in inferior thyroidal artery

5. Thyroiditis
   a. Inflammation of thyroid causing swelling and tenderness
   b. Types and clinical signs
      i) De Quervain’s (subacute thyroiditis)
         • Post viral inflammatory disorder
         • Focal or diffuse enlargement
         • Fever, neck pain, fatigue, increased sedimentation rate
         • Transient hyperthyroidism, hypothyroidism
         • Sonographic findings
            o Focal or diffuse enlargement
            o Ill defined, hypoechoic to heterogeneous regions with decreased to absent vascularity, adenopathy
      ii) Hashimoto’s
         • Most common form
         • Autoimmune - chronic inflammation
Abdomen and Superficial Structures Including Introductory Pediatric and Musculoskeletal

- Diffuse enlargement/possibly asymmetric
- Painless/may develop mild pain over time
- Eventual hypothyroidism
- Sonographic findings
  - Possibly hypoechoic/normal echo texture
  - Size of gland is variable
  - Microlobulations
  - Thick, fibrous strands or echogenic septae
  - Increased vascularity with color Doppler – acute phase
  - Decreased vascularity with color Doppler – chronic phase

6. Benign masses
   a. Cystic nodules
      i) Related to degeneration of an adenoma
      ii) Sonographic findings
         - Corresponds to degenerative changes
         - Mixed solid/cystic nodule
   b. Follicular neoplasm
      i) Benign or malignant
      ii) Sonographic findings
         - Isoechoic/hyperechoic
         - Thin hypoechoic peripheral halo
         - Color Doppler peripherally
         - Central cystic areas/degeneration

7. Malignant masses
   a. Papillary carcinoma
      i) Most common thyroid malignancy
         - Characterized by tiny calcifications called psammoma bodies
         - Major route of spread is through regional lymphatic system
         - May have macrocalcifications
         - Often hypoechoic with irregular margins or heterogeneous
      ii) Clinical signs
      iii) Asymptomatic
         - Painless lump in neck
      iv) Sonographic findings
         - Very hypoechoic
         - Predominately solid
• Microcalcifications, rim calcifications
• Lobulated, contour bulge, taller than wide
• Possible cervical lymph node metastasis

b. Follicular carcinoma
i) Cannot distinguish adenoma from carcinoma on fine needle aspiration
ii) Adenomas are more common than adenocarcinoma
iii) Sonographic findings
   • May be indistinguishable from adenoma
   • Tend to be hypoechoic, no halo, no cystic spaces
   • More common in older females

c. Medullary carcinoma
i) Less common thyroid malignancy
ii) Clinical signs
   • Palpable nodule
   • Abnormal serum calcitonin levels
iii) Sonographic findings
   • Overlap with papillary carcinoma, tend to be larger, homogeneous, greater cystic change and may contain calcifications.
   • Lymphadenopathy

d. Anaplastic carcinoma
i) Rare thyroid malignancy
ii) Clinical signs
   • Hard, fixed mass; rapid growth
   • Pain, pressure, tenderness
   • Locally invasive
iii) Sonographic findings
   • Large, hypoechoic, heterogeneous mass
   • Cystic areas due to necrosis
   • Amorphous calcifications
   • Vascular, lymphatic, extracapsular invasion
   • Hypervascularity
   • Macrocalcification
   • Refractory edge shadow
   • Possible cervical lymph node metastasis (often with microcalcifications or cystic)

e. Lymphoma
i) Clinical signs
• Rapidly growing mass in neck
• Pre-existing Hashimoto’s thyroiditis in more than 90% of cases

ii) Sonographic findings
• Hypoechoic, heterogeneous mass/nodules
• Echogenic septae

iii) May have an appearance similar to Hashimoto’s thyroiditis

8. Other neck masses
a. Thyroglossal duct cyst
   i) Congenital anomaly midline and anterior to trachea
   ii) Clinical signs
      • Palpable midline mass
      • Pain is associated with hemorrhage or infection
   iii) Sonographic findings
      • Cystic mass in midline anterior to the trachea
      • Internal echoes caused by hemorrhage or infection

b. Branchial cleft cyst
   i) Remnant of embryonic development usually lateral to thyroid
   ii) Sonographic findings
      • Primarily cystic
      • Possible low-level echogenicity, solid components

c. Cystic hygroma
   i) Congenital lymphatic malformation commonly posterolateral on neck
   ii) Webbed neck
   iii) Sonographic findings
      • Thin walled, cystic multiloculated mass

d. Fibromatosis colli
   i) Proliferation of the inferior sternocleidomastoid muscle
   ii) Often associated with neonatal torticollis

G. Correlative and/or Prior Imaging
1. Nuclear medicine
2. Computed tomography
3. Magnetic resonance
4. Radiography
5. Extracranial duplex Doppler examination
Section XIV: Prostate

1. Describe the normal anatomy and variants, physiology and sonographic appearance of the prostate
2. Describe the sonographic technique used to evaluate the prostate
3. Identify the clinical indications and laboratory values associated with prostate disease
4. Describe prostate pathology in terms of sonographic appearances, sequelae and associated pathologies
5. Explain sonographic technique for guided prostate biopsies

XIV. Prostate

A. Embryology

B. Anatomy and Normal Variants
   1. Oval-shaped, retroperitoneal structure
   2. Size
   3. Five lobes: anterior, middle, posterior, two lateral lobes
   4. Zonal anatomy
      a. Peripheral zone (PZ)
      b. Central zone (CZ)
      c. Transitional zone (TZ)
   5. Seminal vesicles
   6. Relational anatomy
      a. Prostate
         i) Inferior to seminal vesicles and urinary bladder
         ii) Anterior to rectum
         iii) Posterior to space of Retzius
      b. Seminal vesicles
         i) Superior to prostate
         ii) Inferior to urinary bladder
   7. Variants and congenital anomalies
      a. Agenesis of seminal vesicle
         i) Associated with ipsilateral renal agenesis
      b. Cysts
      c. Prostatic utricle cyst
         i) Associated with unilateral renal agenesis
      d. Ejaculatory duct cyst
      e. Müllerian duct cyst
      f. Agenesis of vas deferens

C. Function and Physiology
   1. Secretes alkaline fluid that provides a mode of fluid transport for sperm
2. Produces majority of fluid volume of ejaculate
3. Glandular tissue produces prostate specific antigen (PSA)
4. Testosterone and dihydrotestosterone regulate prostate growth and function
5. Most accurate method to calculate prostate function is by calculating prostate-specific antigen density (PSAD)

D. Laboratory Values
1. Prostate specific antigen (PSA)
2. PSA assays
   a. Monoclonal
   b. Polyclonal

E. Indications
1. Urinary symptoms
   a. Frequency
   b. Nocturia
   c. Dysuria
   d. Decreased force of urinary stream
2. Abnormal biochemistry
3. Abnormal digital examination
4. Pain
5. Hematospermia
6. Oligospermia
7. Infertility

F. Sonographic Technique
1. Transabdominal approach
   a. Patient preparation
   b. Transducer selection
   c. Patient positioning
   d. Image optimization
   e. Examination protocol
2. Transrectal (TRUS)
   a. Preferred technique
      i) Patient preparation
      ii) Transducer selection
      iii) Transducer preparation
      iv) Patient positioning
      v) Image optimization
      vi) Examination protocol
vii) Transducer cleaning and disinfectant

3. Measurements
   a. Maximum transverse width
   b. Antero posterior (AP)
   c. Length
   d. Prostatic volume

G. Pathology and Pathophysiology

1. Benign prostatic hypertrophy (BPH)
   a. Sonographic findings
      i) Enlarged gland
      ii) Hypoechoic inner region compared to peripheral region
      iii) May have echogenic nodules
      iv) Central and peripheral zones (PZ) may be compressed
      v) Calcification and cystic change may be seen
      vi) Thickened bladder wall
      vii) Distended bladder with thickened walls and trabeculations
      viii) Possible hydronephrosis, hydroureters
      ix) Asymmetric enlargement should be evaluated for malignant change

2. Prostatitis
   a. Acute inflammation
      i) Usually occurs in PZ
      ii) Clinical presentation
         • Pain
      iii) Complication
         • Abscess
      iv) Sonographic findings
         • Most commonly normal
         • Diffuse hyperechoic appearance of PZ
         • Hypervascularity with Doppler
         • Hypoechoic mass
         • Abscess
   b. Chronic inflammation
      i) Recurrent episodes of acute
      ii) Clinical presentation varies
      iii) Sonographic findings
         • Focal masses of varying echogenicity
         • Calcifications
3. Carcinoma (CA)
   a. The majority of prostatic carcinomas are located in PZ
   b. 25% are located in central gland
   c. Sonographic findings
      i) Small nodules are usually hypoechoic and located in PZ
      ii) May be present in the anterior and central gland - difficult to see with ultrasound
      iii) Isoechoic nodules (which are difficult to identify on sonography), hyperechoic or mixed
      iv) Enlarged prostate – usually asymmetric
      v) Distended thick bladder wall with trabeculations
      vi) Bilateral hydroureters and hydronephrosis
      vii) Loss of smooth contour in area of lesion indicating extracapsular spread
      viii) Increased vascularity on color or power Doppler

4. Cysts
   a. Usually asymptomatic
   b. Sonographic findings
      i) Smooth wall
      ii) Anechoic, posterior enhancement

5. US guided prostate biopsies
   a. Transrectal approach
   b. Transperienal approach
      i) Performed in patients with absent rectum
   c. Indications
      i) Diagnosis of prostate cancer in patients with elevated PSA
      ii) Diagnosis/drainage of prostate abscess
Section XV: Scrotum

1. Describe the normal anatomy, physiology, and sonographic appearance of the scrotum
2. Describe the sonographic technique used to evaluate the scrotum
3. Identify the clinical indications and laboratory values associated with scrotal disease
4. Describe scrotal pathology and pathophysiology including the sonographic appearance
5. Identify normal and abnormal flow characteristics and waveforms

XV. Scrotum

A. Embryology
B. Normal Anatomy and Variants
   1. Testicles
      a. Rete testis
      b. Mediastinum
         i) Thickened portion of tunica albuginea
      c. Seminiferous tubules
      d. Layers
         i) Tunica albuginea
         ii) Tunica vaginalis
   2. Epididymis
      a. Head, body, tail
   3. Vas deferens
   4. Seminal vesicles
   5. Variants
      a. Cryptorchidism
         i) Testicles or testis does not descend to normal position
      b. Polyorchidism or unilateral testis
C. Function and Physiology
   1. Testicles
      a. Produce sperm (maturing from basal layer of spermatic tubules to inner layer and then extruded)
   2. Epididymis
      a. Stores sperm
   3. Vas deferens
      a. Transports sperm from epididymis to prostatic urethra
   4. Seminal vesicles
      a. Produce fluid rich in fructose
D. Indications
   1. Pain
2. Enlargement
3. Palpable mass
4. Search for undescended testicle
5. Follow-up for patients with a previous orchiectomy for recurrent tumor
6. Trauma
7. Male infertility

E. Sonographic Technique
1. Patient preparation
2. Transducer selection
3. Patient positioning
4. Image optimization
5. Examination protocol
   a. Measurements
   b. Color and/or power Doppler

F. Pathology and Pathophysiology
1. Benign conditions
   a. Hydrocele
      i) Abnormal accumulation of fluid in tunica vaginalis
      ii) Congenital or idiopathic
      iii) Often due to epididymitis
      iv) May be seen with orchitis, spermatic cord torsion and trauma
   b. Hema-tocele
      i) Blood in scrotal sac, usually after trauma or surgical intervention
   c. Pyocele
      i) Pus in scrotal sac, usually associated with epididymitis/orchitis
   d. Epididymitis
      i) Inflammation of epididymis
      ii) Most common cause of acute scrotal pain
      iii) Clinical findings
         • Fever
         • Pain
         • Dysuria
         • May have urethral discharge
      iv) Sonographic findings
         • Enlarged epididymis, usually of head
         • Thickened scrotal skin
      v) Decreased echogenicity with heterogeneous coarse echo pattern
vi) May be associated with hydrocele

vii) Increased Doppler flow in affected region when compared to contralateral side

e. Orchitis
    i) Inflammation of the testis
    ii) Associated finding in 20-30% of epididymitis
    iii) Chlamydia most common cause, mumps, viral, idiopathic

f. Spermatic cord torsion
    i) Due to “bell and clapper” deformity
    ii) Spermatic cord becomes rotated or twisted and cuts off blood supply
    iii) Acute
        - Scrotal pain and swelling
        - Nausea and vomiting
        - Sonographic findings
          - Varied and non-specific
          - Enlarged testicle and epididymal head with decreased echogenicity
          - Absent flow on color Doppler
          - Twisting or looping of arteries in the spermatic cord in inguinal canal
    iv) Chronic
        - Sonographic findings
          - Small heterogeneous testicle (infracted)
          - Scrotal wall thickening and possible hydrocele
          - No arterial flow on Doppler study
          - Increased flow in peritesticular soft tissues
    v) Intermittent torsion
        - Mobile testis with bell-clapper deformity with recurrent acute scrotal pain with rapid spontaneous resolution
        - Sonographic findings
          - Acute – similar to torsion
          - After acute presentation – nonspecific, with hydrocele is most common
    vi) Partial torsion
        - Sonographic findings
          - May show increased blood flow due to reactive hyperemia
          - Comparison must be made to contralateral side

g. Spermatocele
    i) Cystic lesion of epididymis containing sperm

h. Varicocele
    i) Enlargement of veins of spermatic cord
ii) Most common cause of infertility
iii) Majority occur on left side due to venous drainage into left renal vein
iv) Large, right-sided varicocele may be associated with renal or retroperitoneal tumor
v) Sonographic findings
   - Dilated veins
     - Valsalva maneuver or having patient stand will increase venous pressure
     - Reversal of flow occurs when intra-abdominal pressure increases

i. Cysts
   i) Intratesticular
   ii) Epidermoid

j. Abscess
   i) Most commonly caused from untreated epididymo-orchitis
   ii) Intratesticular or more commonly extra-testicular
   iii) Clinical findings
       - Fever
       - Scrotal pain
       - Swelling
   iv) Sonographic findings
       - Anechoic or complex mass
       - Increase blood flow around periphery of the mass
       - No blood flow in mass

k. Scrotal hernia

l. Hematoma
   i) Intratesticular more common than extratesticular
   ii) Associated with trauma

m. Extratesticular tumors
   i) Adenomatoid tumor
      - Most common extratesticular tumor
   ii) Non-germ cell neoplasms
      - Leydig cell tumor
      - Sertoli tumor
      - Cystadenoma
      - Dermoid cyst

G. Malignant Conditions
   1. Testicular cancer
      a. Clinical findings
i) Painless

ii) Unilateral enlargement

b. Pure tumors

i) Germ cell tumors
   - Seminoma
     o Most common germ cell tumor
     o Hypoechoic lesion
   - Embryonal carcinoma
     o Invasive producing ill-defined hypoechoic lesion with possible capsular distortion
   - Teratoma
     o Well-defined complex masses with possible calcification, often cystic changes on US
   - Choriocarcinoma

ii) Stromal cell tumors
   - Leydig cell
   - Sertoli
   - Granulose

c. Theca cell
d. Mixed germ cell tumors
   i) Second most common tumor after seminoma
e. Metastasis
f. Lymphoma

H. Correlative and/or prior imaging
  1. Nuclear medicine
  2. Computed tomography (CT)
  3. Magnetic resonance imaging (MRI)
Section XVI: Musculoskeletal Sonography

Rationale: Accurate assessment and performance of musculoskeletal sonograms requires sonographers to assemble a comprehensive knowledge of the anatomy, physiology, and pathophysiology of joints, tendons, muscles, bursae and ligaments that are sonographically accessible. An understanding of the sonographic appearances and dynamics of joints, tendons, muscles, bursae and ligaments are essential for the performance of high quality examinations.

1. Identify normal anatomic location and function of the tendons, ligaments, muscles, nerves and bursae
2. List indications for sonographic evaluation
3. Describe examination techniques and basic scanning protocols
4. Summarize basic protocols for musculoskeletal examinations
5. Describe applicable pathology

XVI. Musculoskeletal Sonography

A. Anatomy
   1. Muscles
      a. Pennate patterns
   2. Tendons
      a. Synovial sheath
   3. Ligaments
   4. Bursae
   5. Nerves
      a. Lower limb
         i) Sciatic
         ii) Popliteal
      b. Upper limb
         i) Suprascapular
         ii) Median
         iii) Radial
         iv) Ulnar
   6. Joints
   7. Bones
   8. Shoulder
      a. Biceps tendon/sheath
      b. Subscapularis tendon
      c. Supraspinatus tendon
      d. Infraspinatus tendon
      e. Acromioclavicular joint
      f. Subdeltoid bursa
      g. Supraspinatus, infraspinatus and teres minor muscles
9. Wrist
   a. Carpal tunnel
      i) Flexor retinaculum space
         • Flexor pollicis longus
         • Flexor digitorum tendons
      ii) Ulnar artery and veins
      iii) Scaphoid tubercle
      iv) Trapezium ridge
      v) Pisiform ridge
      vi) Pisiform bone
      vii) Hamate
      viii) Median nerve
   b. Wrist - Guyon’s canal
      i) Hamate and pisiform bones
      ii) Ulnar nerve

10. Ankle/leg/foot
    a. Achilles’ tendon
       i) Gastrocnemius muscle
       ii) Soleus muscle
       iii) Calcaneus
       iv) Paratendon
       v) Tibialis posterior
       vi) Peroneal tendons
       vii) Plantar fascia
    b. Popliteal fossa
       i) Femoral condyles and tibia
       ii) Popliteal bursa
          • Synovial fluid
       iii) Popliteal artery and vein

B. Indications
   1. Shoulder
      a. Shoulder pain or swelling
      b. Weakness with arm elevation
      c. Trauma
      d. Decreased range of motion
      e. Evaluation of soft tissue masses
   2. Wrist
a. Palpable lesion  
b. Loss or decrease of digital flexion/extension  
c. Pain  
d. Swelling  
e. Trauma  
f. Numbness of the middle and index fingers  
g. Weakness or clumsiness of the hand  
h. Tingling with nerve percussion  
   i) Tinel’s sign  
   ii) Phalen’s sign  

3. Achilles’ tendon  
a. Abnormal Thompson’s test  
b. Trauma  
c. Displacement of Kager’s fat pad on radiograph  
d. Knob or bulge over proximal tendon  
e. Audible pop or snap followed by sharp pain  
f. Inability to stand on toes  
g. Heel pain  
h. Decreased strength or mobility  
i. Postoperative  

4. Baker’s Cyst  
a. Swelling in the popliteal fossa  
b. Knee pain  

C. Sonographic Technique  
1. Shoulder-rotator cuff  
a. Patient position  
b. Arm positions  
   i) Palm up, elbow flexed straight external rotation  
   ii) Dynamic internal/external  
   iii) Place palm on back pocket  
c. Views  
   i) Longitudinal/transverse biceps/sheath  
   ii) Longitudinal/transverse subscapularis tendon  
   iii) Longitudinal/transverse supraspinatus and infraspinatus tendons  
   iv) Transverse posterior joint space/labrum  
   v) Acromioclavicular joint  
   • Longitudinal/transverse supraspinatus, infraspinatus and teres minor muscles
2. Carpal Tunnel  
   a. Patient position  
      i) Arm resting on a stand  
      ii) Palm supinated  
   b. Views  
      i) Longitudinal and transverse of area of interest  
      * Use gel or standoff pad

3. Achilles’ tendon  
   a. Patient position  
      i) Prone with foot hanging over edge of cart/table  
   b. Views  
      i) Longitudinal and transverse of area of interest  
      ii) Dorsiflexion/plantar flexion  
      iii) Thompson’s test

4. Popliteal fossa  
   a. Patient position  
      i) Prone with towel roll under ankle  
   b. Views  
      i) Sagittal and transverse of area of interest

5. Normal sonographic findings  
   a. Tendons  
   b. Ligaments  
   c. Muscles  
   d. Bursae

6. Artifacts  
   a. Anistrophy

D. Pathology/Pathophysiology

1. Shoulder  
   a. Biceps tendon  
   b. Subluxation/dislocation  
   c. Rotator cuff tear  
   d. Tendinopathy  
   e. Acromiocravicular joint arthritis  
   f. Subdeltoid effusion/bursitis  
   g. Fatty degeneration of cuff muscles

2. Wrist  
   a. Carpal tunnel syndrome
b. Ganglion and synovial cysts  
c. Tears of triangular fibrocartilage  
d. Tenosynovitis  
e. Tumor  
f. Foreign body  
g. Tendon tear  
h. Inflammatory arthritis  
i. Tendon entrapment  
3. Baker’s cyst  
   a. Enlargement of gastrocnemius-semimembranosus bursa caused by meniscal tear or osteoarthritis, rheumatoid arthritis, joint effusion  
      i) May dissect inferiorly into calf muscles or superiorly into thigh  
   b. Sonographic findings  
      i) Anechoic fluid collection, may be complex (loose bodies, septations, internal echoes), may dissect into calf muscles  
E. Correlative and/or prior imaging  
   1. Radiography  
   2. Arthrography  
   3. Magnetic resonance  
   4. Computed tomography
Section XVII: Pediatric Hip

Rationale: Accurate assessment and performance of pediatric hip sonograms requires sonographers to assemble a comprehensive knowledge of the anatomy, physiology, and pathophysiology of the pediatric hip. An understanding of the sonographic appearance and dynamics of the pediatric hip are essential for the performance of high quality examinations.

1. Describe the anatomical structures of hip joint and relational anatomy
2. Describe the indications and sonographic techniques for assessment of the pediatric hip
3. Discuss pathological conditions of the hip

XVII. Pediatric Hip

A. Embryology
   1. Mesoderm
   2. Osteogenesis
   3. Myoblasts and mesenchyme
   4. Medial rotation of lower limbs

B. Anatomy
   1. Bones and joints
      a. Pelvic girdle
         i) Ilium
         ii) Ischium
         iii) Pubis
      b. Femur
         i) Femoral head
      c. Fascia lata
      d. Muscles
         i) Anterior/extensor
         ii) Medial/adductor
         iii) Posterior/hamstring and gluteal
   2. Femoral triangle
      a. Anterior of upper thigh
         i) Inguinal ligament
         ii) Adductor longus muscle
         iii) Sartorius muscle
   3. Gluteal region
   4. Hip joint
      a. Acetabulum
         i) Acetabular labrum
      b. Femoral head
c. Ileofemoral ligament

C. Functions
1. Flexion
2. Extension
3. Abduction
4. Adduction
5. Medial rotation
6. Lateral rotation

D. Indications for Sonographic Examination
1. Developmental displacement in neonate
   a. Risk factors
      i) First born child
      ii) Female
      iii) Breech birth
      iv) Family history
      v) Intrauterine conditions
   b. Causes
      i) Genetic
      ii) Mechanical
      iii) Physiologic
2. Abnormal hip examination
3. Response to treatment

E. Sonographic Technique
1. Patient preparation
2. Transducer selection
3. Patient position
4. Image optimization
5. Examination protocol
   a. Coronal with neutral hip position
      i) Push/pull movement of hip
      ii) Document signs of subluxation/dislocation
   b. Coronal with hip flexion
      i) Push/pull movement of hip
      ii) Barlow maneuver
         • Adduction with gentle pushing
      iii) Document signs of subluxation
   c. Transverse with hip flexion
i) Push/pull movement of hip  
ii) Barlow maneuver  
iii) Ortolani test  
d) Transverse with neutral position of hip  
e) Measurements using Graf’s technique  

6. Assessment of hip during treatment  
a) May be performed while patient is in Pavlik harness  
b) Contraindications  

F. Pathology/Pathophysiology  
1. Dislocation of the hip  
a) Acquired (traumatic or non-traumatic)  
b) Teratogenic  
c) Developmental displacement of the hip (DDH)  
   i) Common congenital anomaly especially in females  
   ii) Dysplastic, subluxed, dislocatable or dislocated hips  
d) Sonographic findings  
   i) Femoral head  
      • Normal, subluxed, dislocated  
   ii) Stability  
      • Normal, dislocatable, reducible, irreducible  
   iii) Acetabulum  
      • Normal, immature, dysplastic  

2. Hip effusion  
a) Widening of joint space with displacement of femur laterally  
   i) Toxic synovitis  
   ii) Septic arthritis  
b) Scan plane is anterior and parallel to femoral neck, patient supine, hip in neutral position  
   i) Hips should be symmetrical in sonographic appearance  
c) Sonographic findings  
   i) Anterior recess of joint capsule distends anteriorly  
   ii) Echogenicity of fluid may indicate hemorrhage, transudate, or exudate  

3. Proximal femoral focal deficiency  
a) Congenital underdevelopment of proximal femur with varying degrees of severity  

G. Correlative and/or Prior Imaging  
1. Radiography  
2. Computed tomography  
3. Magnetic resonance
Section XVIII: Neonatal Brain

Rationale: Accurate assessment and performance of neonatal brain sonograms requires the sonographer to assemble a comprehensive knowledge of the anatomy, physiology and pathophysiology of the infant brain. An understanding of the embryologic development and gestational age-adjusted sonographic appearances of the premature and term infant brain are essential for the performance of high quality examinations.

1. Describe the embryology, normal anatomy, and sonographic appearances of the premature and term infant brain
2. Describe the sonographic technique, procedure and protocol used to evaluate the infant brain
3. Describe clinical indications for sonographic examination of the infant brain in both premature and term infants
4. Describe the major pathologies diagnosed with sonographic imaging of the neonatal brain

XVIII. Neonatal Brain

A. Embryology
   1. Forebrain
      a. Prosencephalon
      b. Diencephalons
      c. Telencephalon
   2. Midbrain
      a. Mesencephalon
   3. Hindbrain
      a. Myelencephalon
      b. Metencephalon

B. Anatomy
   1. Fontanelles and sutures
   2. Meninges
      a. Dura mater
      b. Arachnoid
      c. Pia mater
   3. Falx cerebri
   4. Tentorium cerebelli
   5. Ventricular system
      a. Lateral ventricles
         i) Anterior horn
         ii) Body
         iii) Posterior horn
         iv) Temporal horn
   6. Foramen of Monro
   7. Trigone
   8. Corpus callosum
9. Cavum septum pellucidum
10. Thalamus
11. Aqueduct of Sylvius
12. Foramen of Luschka
13. Foramen of Megendie
14. Choroid plexus
15. Cisterns
   a. Subarachnoid
      i) Cisterna magna
16. Cerebrum
   a. Two hemispheres
      i) Connected by corpus callosum
   b. Gray and white matter
17. Lobes of the brain
   a. Frontal
   b. Parietal
   c. Occipital
   d. Temporal
18. Gyri and sulci
19. Fissures
   a. Interhemispheric
   b. Sylvian fissure
   c. Quadrigeminal fissure
20. Basal ganglia
   a. Caudate nucleus
   b. Lentiform nucleus
   c. Claustrum
   d. Thalamus
21. Brain stem
   a. Midbrain
   b. Pons
   c. Medulla oblongata
22. Cerebellum
   a. Hemispheres
      i) Vermis
   b. Nerve tracts
23. Cerebrovascular system
C. Function and Physiology

1. Cerebellum
   a. Skeletal muscle movement

2. Cerebral hemispheres
   a. Frontal lobe
      i) Speech
      ii) Emotions
      iii) Personality
      iv) Morality
      v) Intellect
   b. Parietal lobe
      i) Somesthetic
      ii) Pain
      iii) Temperature
      iv) Spatial ability
   c. Occipital lobe
      i) Vision
   d. Temporal lobe
      i) Auditory
      ii) Olfactory
   e. Other
      i) Judgement
      ii) Memory
      iii) Reasoning

D. Indications

1. Cranial abnormality found on prenatal sonogram
2. Increasing head circumference with or without increasing intracranial pressure
3. Acquired or congenital inflammatory disease
4. Prematurity
5. Diagnosis of hypoxia, hypertension, hypercapnia, hypernatremia, acidosis, pneumothorax, asphyxia, apnea, seizures, coagulation defects, patent ductus arteriosus
6. History of birth trauma or surgery
7. Genetic syndromes and malformations

E. Sonographic Technique
1. Patient preparation and care
   a. Special considerations for premature and neonatal care

2. Transducer selection

3. Patient positioning

4. Image optimization

5. Examination protocol
   a. Coronal scan planes
      i) Anterior
         - Orbits, anterior horns, and lateral ventricles
      ii) Mid
         - Lateral ventricles
         - Cavum septum pellucidum
         - Third ventricle-foramen of Monro
         - Corpus callosum
      iii) Posterior
         - Ambient wings and cisterna magna
         - Choroids
         - Glomus of choroids
         - Occipital lobe
   b. Sagittal and bilateral parasagittal scan planes/imaging
      i) Midline
         - Cavum septum pellucidum
         - Corpus callosum
         - Third ventricle
         - Brain stem
         - Cerebellum (tentorium)
         - Cisterna magna
      ii) Thalamus and caudate
      iii) Lateral ventricle
         - Anterior horn, body, posterior horn
         - Temporal horn may be seen in cases of ventriculomegaly or hydrocephalus
         - Caudalthamic groove
      iv) Lateral
         - Lateral ventricle to show white matter and Sylvian fissure (middle cerebral artery)
   c. Special views
      i) Axial
ii) Posterior fontanelle
iii) Submastoid

6. Doppler
a. Color and spectral Doppler may be used to demonstrate and/or evaluate intracranial blood flow
   i) Decreased blood flow/ischemia/infarction
   ii) Vascular abnormalities
   iii) Cerebral edema
   iv) Hydrocephalus
   v) Intracranial tumors
   vi) Near-field structures

F. Pathology/Pathophysiology
1. Chiari malformations
   a. Caudal displacement of the cerebellar hemispheres due to presence of spina bifida
   b. Types
      i) Chiari I: caudal displacement of the cerebellum without displacement of fourth ventricle or medulla
      ii) Chiari II: most common; associated with meningomyelocele
      iii) Chiari III: cervical encephalomeningocele containing cerebellum, fourth ventricle and medulla
      iv) Chiari IV: severe hypoplasia of cerebellum without displacement
   c. Sonographic findings Chiari Type II
      i) Small posterior fossa
      ii) Possible myelomeningocele formation
         • Decompression of ventricles
      iii) Small, displaced cerebellum
      iv) Widening of third ventricle
      v) Cerebellar tonsils and vermis herniated into spinal canal through enlarged foramen magnum
      vi) Pons and medulla inferiorly displaced
      vii) Fourth ventricle becomes elongated
      viii) Possible enlargement of the massa intermedia
         • Third ventricle may be slightly enlarged
      ix) Frontal horns of the ventricles are small
         • Bat wing sign
         • Posterior horns enlarged
         • Cavum septum pellucidum may be partially or completely absent
      x) Interhemispheric fissure may be widened
      xi) Tentorium low and hypoplastic
2. Holoprosencephaly
   a. Grossly abnormal brain with common large central ventricle due to failure of cleavage of the prosencephalon into separate cerebral hemispheres
      i) Varying degrees of severity and associated midline facial defects
         • Alobar holoprosencephaly (most severe)
            o Fused thalami anterior to a fused choroid plexus
         • Semilobar holoprosencephaly
            o Single ventricle; brain parenchyma present with portions of falx and interhemispheric fissure in the posterior segment
         • Lobar holoprosencephaly (least severe)
            o Near complete separation of hemispheres; development of falx and interhemispheric fissure
               ~ Only frontal lobes may be fused
               ~ Septum pellucidum is absent
               ~ Anterior horns of ventricles fused

3. Dandy-Walker Complex
   a. Congenital anomaly of the roof of the fourth ventricle with concurrent occlusion of aqueduct of Sylvius and foramina of Magendie and Luschka
   b. Variant: enlarged cisterna magna that communicates with fourth ventricle; normal or hypoplastic cerebellar vermis
   c. Sonographic findings
      i) Enlarged 4th ventricle cyst occupies the area of cerebellum
      ii) Secondary dilatation of the 3rd and lateral ventricles
      iii) Possible absent vermis
      iv) Hydrocephalus

4. Callosal Dysgenesis
   a. Complete or partial absence of the connection tissue between cerebral hemispheres
      i) May be an isolated finding or associated with other anomalies
   b. Sonographic findings
      i) ‘Longhorn’ or ‘moosehead’ configuration of anterior horns and third ventricle
      ii) Lateral ventricles with parallel configuration
      iii) Marked separation of lateral ventricles
      iv) Widening of occipital horns and 3rd ventricle
      v) Gyri in a straightened or ‘sunray’ appearance

5. Ventriculomegaly
   a. Enlargement (unilateral or bilateral) of the ventricles without increased head circumference
   b. Types
      i) Communicating
ii) Non-communicating
iii) Result of cerebral atrophy
c. Sonographic findings
   i) Ventricle(s) demonstrate(s) greater than normal size first noted in trigone and occipital horn areas
   ii) Visualization of the third and possibly, fourth ventricle
   iii) Choroid plexus appears to “dangle” within ventricular atrium
   iv) Thinned brain mantle in cases of cerebral atrophy

6. Hydrocephalus
   a. Enlargement (unilateral or bilateral) of the ventricle(s) with increased head circumference
   b. Types
      i) Communicating
      ii) Non-communicating
c. Sonographic findings
   i) Blunted lateral angles of enlarged lateral ventricles
   ii) Possible rupture of interhemispheric fissure
   iii) Thinned brain mantle
   iv) Head circumference is enlarged
   v) In cases of non-communicating hydrocephalus, point of obstruction may be seen

7. Aqueductal stenosis
   a. Aqueduct of Sylvius is narrowed or is a small channel with blind ends
   b. Sonographic findings
      i) Widening of lateral and 3rd ventricles and normal 4th ventricle

8. Hydranencephaly
   a. Prenatal occlusion of internal carotid arteries resulting in necrosis of cerebral hemispheres
      i) Absence of both cerebral hemispheres with presence of falx, thalamus, cerebellum, brain stem, and portions of occipital and/or temporal lobes
   b. Sonographic findings
      i) Large fluid-filled cranial vault with identifiable falx
      ii) Intact cerebellum and midbrain
      iii) May mimic holoprosencephaly

9. Cephalocele
   a. Herniation of portion of the neural tube through a defect in skull
      i) Occipital defect most common
      ii) May contain variable amounts of brain tissue
         • If cephalocele is large, head circumference may be decreased
   b. Sonographic findings
10. Hemorrhagic pathology
   a. Subependymal - intraventricular hemorrhage (SEH – IVH)
      i) Pre-term infants are at risk
      ii) Grades
         • Grade I SEH without ventricular enlargement
         • Grade II IVH without ventricular enlargement
         • Grade III IVH with ventricular enlargement
         • Grade IV parenchymal hemorrhage
   b. Sonographic findings
      i) Appearance/location depends on grade
      ii) Echogenicity depends on the age of hemorrhage/clot

11. Intraparenchymal hemorrhage
   a. Sonographic findings
      i) Zones of increased echogenicity in white matter

12. Intracerebellar hemorrhage
   a. Types
      i) Primary
      ii) Venous infarction
      iii) Traumatic laceration
      iv) Extension from SEH-IVH
   b. Sonographic findings
      i) Areas of increased echogenicity within echogenic cerebellar parenchyma

13. Ischemic - Hypoxic lesions
   a. Types
      i) Selective neuronal necrosis
      ii) Status marmoratus
      iii) Parasagittal cerebral injury
      iv) Periventricular leukomalacia (PVL) or white matter necrosis (WMN)
      v) Focal brain lesions
   b. Sonographic findings
      i) Variable
         • Areas of increased echogenicity in subcortical and deep white matter in basal ganglia
            o Necrosis
14. **Ventriculitis**
   a. **Sonographic findings**
   
   i) Thin septations extending from walls of lateral ventricles

15. **Ependymitis**
   a. **Sonographic findings**
   
   i) Thickened, hyperechoic ependyma

16. **Cerebral edema**
   a. Swelling of the brain associated with hypoxic events or metabolic abnormalities in full-term infants
   
   i) May be related to metabolic disorders, birth asphyxia, infection, and congenital malformation
   
   ii) May lead to infarction and subsequent porencephaly or encephalomalacia
   
   b. **Sonographic findings (initial)**
   
   i) Diffusely echogenic brain with loss of visualization of normal landmarks
   
   ii) Slit-like ventricles
   
   iii) Speckling of parenchyma and poorly defined sulci
   
   c. **Sonographic findings (resolved)**
   
   i) Patchy increased periventricular echogenicity
   
   ii) Development of cystic areas
   
   • Also known as periventricular leukomalacia (PVL), cystic encephalomalacia or white matter necrosis (WMN)

G. **Miscellaneous**

1. **Lissencephaly**

2. **Lipoma of the corpus callosum**

3. **Papilloma of the corpus callosum**

4. **Schizencephaly**

5. **Septooptic dysplasia**

6. **Sub-arachnoid cyst**

7. **Sub-dural hemorrhage**

8. **Vein of Galen aneurysm**

9. **Calcifications**

H. **Correlative and/or prior imaging**

1. **Computed tomography**

2. **Magnetic resonance**

3. **Angiography**

4. **Radiography**
Section XIX: Neonatal Spine

Rationale: Accurate assessment and performance of neonatal spine sonograms requires the sonographer to assemble a comprehensive knowledge of the anatomy, physiology and pathophysiology of the infant spine. An understanding of the embryologic development and sonographic appearances are essential for the performance of high quality examinations.

1. Describe the anatomy of the neonatal spine
2. List the common pathologic conditions of the neonatal spine that relate to sonographic diagnosis
3. Describe the indication and sonographic techniques for neonatal spine imaging
4. Describe the major pathologies diagnosed with sonographic imaging of the neonatal spine

XIX. Neonatal Spine

A. Embryology
   1. Neural plate
   2. Neurulation
   3. Caudal cell mass development
   4. Neural tube and canal
   5. Spinal meninges
   6. Paraxial mesoderm
      a. Vertebrae

B. Anatomy
   1. Vertebrae
   2. Sacrum
   3. Intervertebral discs
   4. Ligaments and nerves
   5. Spinal cord
      a. Conus medullaris
      b. Filum terminale
      c. Anterior median fissure
      d. Posterior media fissure
      e. Roots of spinal nerves
         i) Cauda equina
   6. Meninges of spinal cord
      a. Dura mater
      b. Arachnoid mater
      c. Pia mater

C. Function and Physiology
   1. Spinal cord carries nerve impulses to and from brain and to rest of body
   2. Bony spine encloses and protects the spinal cord
D. Indications
   1. Midline cutaneous abnormalities
   2. Sacral dimple(s)
   3. Hemangioma
   4. Raised midline
   5. Hairy patch
   6. Tail-like projection of lower spine
   7. Diagnosis of myelomeningocele or myeloschisis
   8. Lower extremity deformity

E. Sonographic Technique
   1. Patient positions
      a. Prone
      b. Lateral decubitus
      c. Upright
      d. Spine flexed
      e. Head should be more elevated than caudal end of spinal canal, gravity may pull fluid into sacs or cystic spaces that may not be seen otherwise
   2. Transducer selection
   3. Sonographic technique
      a. Stand-off pad may be utilized
      b. Determine level of the conus medullaris
         i) Begin at sacrum - possible to have 6 lumbar vertebrae
         ii) Counting from lowest rib bearing vertebra
   4. Sonographic findings
      a. Vertebral bodies
      b. Spinous processes
      c. Dura mater
      d. Coccyx
      e. Spinal cord
         i) Central echo complex
      f. Nerve roots
      g. Filum terminale

F. Pathology/Pathophysiology
   1. Tethered cord
      a. Fixation of cord at a more caudal location
      b. Diminished cord movement with stretching and ischemia of cord
      c. Sonographic findings
i) Visualization of cord caudal to normal termination
ii) Diminished cord pulsations may be noted
iii) Eccentric cord location within the canal

2. Lipoma
   a. Mass of filum terminale
   b. Frequently associated with tethered cord
   c. Sonographic findings
      i) Echogenic mass

3. Hydromyelia
   a. Dilatation of the central canal which may be diffuse or focal
   b. Associated with myelomeningocele and diastematomyelia; may mimic or co-exist with syringomyelia
   c. Sonographic findings
      i) Separation of echogenic linear structures of central canal

4. Diastematomyelia
   a. Cord is split at one or more sites by septum
   b. Associated with meningocele or myelomeningocele
   c. Sonographic findings
      i) Split segments best seen on transverse views

5. Cysts of spinal cord
   a. May be noted in cauda equine or filum terminale; possible arachnoid cyst; thought to be normal variant
   b. Related to tethered cord

6. Myelomeningocele
   a. Spina bifida with low termination of spinal cord and a sac/pouch containing cerebral spinal fluid and nerves
   b. Sonographic findings
      i) Pre-operative sonographic examination can differentiate between myelomeningocele and a meningocele (sac/pouch containing only cerebral spinal fluid)
      ii) Detection of associated anomalies
         • Hydromelia, diastematomyelia, lipoma or thickened filum terminale

G. Correlative and/or prior imaging
   1. Radiography
   2. Computed tomography
   3. Magnetic resonance
### Abbreviations

<table>
<thead>
<tr>
<th>Abbreviation</th>
<th>Description</th>
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<tbody>
<tr>
<td>AAA</td>
<td>Abdominal Aortic Aneurysm</td>
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<td>ACTH</td>
<td>Adrenocorticotropic Hormone</td>
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<td>ADPKD</td>
<td>Autosomal Dominant Polycystic Kidney Disease</td>
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<td>AFP</td>
<td>Alpha-Fetoprotein</td>
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<tr>
<td>AG</td>
<td>Adrenogenital Syndrome</td>
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<tr>
<td>AI</td>
<td>Acceleration-Index</td>
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<tr>
<td>AIDS</td>
<td>Acquired Immune Deficiency Syndrome</td>
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<tr>
<td>ALP</td>
<td>Alkaline Phosphatase</td>
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<tr>
<td>ALT</td>
<td>Alanine Aminotransferase</td>
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<tr>
<td>AP</td>
<td>Anterior-Posterior</td>
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<tr>
<td>APCKD</td>
<td>Adult Polycystic Kidney Disease</td>
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<tr>
<td>APN</td>
<td>Acute Pyelonephritis</td>
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<tr>
<td>ARDS</td>
<td>Acute Respiratory Distress Syndrome</td>
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<td>ARPKD</td>
<td>Autosomal Recessive Polycystic Kidney Disease</td>
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<td>Arteriovenous Fistula</td>
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<td>Blood Urea Nitrogen</td>
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<td>Carcinoma</td>
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<td>Common Bile Duct</td>
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<td>CHA</td>
<td>Common Hepatic Artery</td>
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<td>Common Hepatic Duct</td>
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<td>CHF</td>
<td>Congestive Heart Failure</td>
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<tr>
<td>CMJ</td>
<td>Corticomedullary Junction</td>
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<tr>
<td>C/RL</td>
<td>Caudate to Right Lobe Ratio Technique</td>
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<tr>
<td>CSF</td>
<td>Cerebral Spinal Fluid</td>
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<tr>
<td>CT</td>
<td>Computed Tomography</td>
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<td>Central Zone</td>
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<tr>
<td>DCIS</td>
<td>Ductal Carcinoma In-Situ/Intraductal Carcinoma</td>
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<td>DDH</td>
<td>Developmental Dysplasia of the Hip</td>
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<td>DSR</td>
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<td>FFP</td>
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<td>FNH</td>
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<td>Abbreviation</td>
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<tr>
<td>HIV</td>
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<td>HPS</td>
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<tr>
<td>IBD</td>
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<tr>
<td>IDC</td>
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<tr>
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<td>IMV</td>
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<td>WMN</td>
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Utilized References

10. Sonography Canada: Canadian Association of Registered Diagnostic Ultrasound Professionals (CARDUP) National Competency Profiles. http://www.sonographycanada.ca/Apps/Pages/home-csdms