

The American College of Radiology, with more than 30,000 members, is the principal organization of radiologists, radiation oncologists, and clinical medical physicists in the United States. The College is a nonprofit professional society whose primary purposes are to advance the science of radiology, improve radiologic services to the patient, study the socioeconomic aspects of the practice of radiology, and encourage continuing education for radiologists, radiation oncologists, medical physicists, and persons practicing in allied professional fields.

The American College of Radiology will periodically define new practice guidelines and technical standards for radiologic practice to help advance the science of radiology and to improve the quality of service to patients throughout the United States. Existing practice guidelines and technical standards will be reviewed for revision or renewal, as appropriate, on their fifth anniversary or sooner, if indicated.

Each practice guideline and technical standard, representing a policy statement by the College, has undergone a thorough consensus process in which it has been subjected to extensive review, requiring the approval of the Commission on Quality and Safety as well as the ACR Board of Chancellors, the ACR Council Steering Committee, and the ACR Council. The practice guidelines and technical standards recognize that the safe and effective use of diagnostic and therapeutic radiology requires specific training, skills, and techniques, as described in each document. Reproduction or modification of the published practice guideline and technical standard by those entities not providing these services is not authorized.

2008 (Resolution 22)*

ACR-SPR PRACTICE GUIDELINE FOR THE PERFORMANCE OF PEDIATRIC COMPUTED TOMOGRAPHY (CT)

PREAMBLE

These guidelines are an educational tool designed to assist practitioners in providing appropriate radiologic care for patients. They are not inflexible rules or requirements of practice and are not intended, nor should they be used, to establish a legal standard of care. For these reasons and those set forth below, the American College of Radiology cautions against the use of these guidelines in litigation in which the clinical decisions of a practitioner are called into question.

The ultimate judgment regarding the propriety of any specific procedure or course of action must be made by the physician or medical physicist in light of all the circumstances presented. Thus, an approach that differs from the guidelines, standing alone, does not necessarily imply that the approach was below the standard of care. To the contrary, a conscientious practitioner may responsibly adopt a course of action different from that set forth in the guidelines when, in the reasonable judgment of the practitioner, such course of action is indicated by the condition of the patient, limitations of available resources, or advances in knowledge or technology subsequent to publication of the guidelines. However, a practitioner who employs an approach substantially different from these guidelines is advised to document in the patient record information sufficient to explain the approach taken.

The practice of medicine involves not only the science, but also the art of dealing with the prevention, diagnosis, alleviation, and treatment of disease. The variety and complexity of human conditions make it impossible to always reach the most appropriate diagnosis or to predict with certainty a particular response to treatment.

Therefore, it should be recognized that adherence to these guidelines will not assure an accurate diagnosis or a successful outcome. All that should be expected is that the practitioner will follow a reasonable course of action based on current knowledge, available resources, and the needs of the patient to deliver effective and safe medical care. The sole purpose of these guidelines is to assist practitioners in achieving this objective.

I. INTRODUCTION

This guideline was developed collaboratively by the American College of Radiology (ACR) and the Society for Pediatric Radiology (SPR).

Computed tomography (CT) is a radiologic modality that provides clinical information in the detection, differentiation, and demarcation of disease. It is the primary diagnostic modality for a variety of presenting problems and is widely accepted as a supplement to other imaging techniques.

CT is a form of medical imaging that involves the exposure of patients to ionizing radiation. It should only be performed under the supervision of a physician with the necessary training in radiation protection to optimize examination safety. Medical physicists and trained technical staff must be available to evaluate the equipment and perform the examinations.

CT examinations should be performed only for a valid medical reason and with the minimum exposure that provides the image quality necessary for adequate diagnostic information.

Because children are more sensitive than adults to the effects of ionizing radiation, it is particularly important to tailor CT examinations to minimize exposure while providing diagnostic quality examinations. Protocols should include the sequence of studies, with and/or without contrast administration, and should specify CT radiation dose ranges based on the child's size, whether or not contrast is used, and the information required from the examination [1].

II. INDICATIONS

Primary indications for CT include, but are not limited to:

A. Chest

1. Evaluation of chest wall abnormalities [2-9], including:
 - a. Extent of chest wall developmental deformities, such as pectus excavatum, pectus carinatum, and thoracic insufficiency syndrome secondary to scoliosis or rib anomalies.
 - b. Chest wall injury, including penetrating trauma and injuries that are not adequately addressed by radiography such as sternal fractures, sternoclavicular dislocation, and occult rib fractures.
 - c. Chest wall mass and mass-like conditions including chest wall inflammatory/infectious processes. This also includes evaluation of post treatment complications and residual or recurrent mass.
2. Suspected extracardiac vascular disorder [10-16], including:
 - a. Congenital and syndromic vascular abnormalities such as vascular rings, pulmonary slings, pulmonary vein abnormalities (e.g., anomalous course), systemic-to-pulmonary collateral vessels, coarctation of the aorta, or other congenital processes including known or suspected bronchopulmonary sequestration.
 - b. Acquired disorders of the great vessels (e.g., medium or large vessels vasculitides, aneurysms, and stenoses, infectious or other inflammatory conditions) and posttraumatic evaluation. Assessment includes aortic dissection, transection, and pulmonary embolism.

For cardiac evaluation, see the [ACR-NASCI-SPR Practice Guideline for the Performance and Interpretation of Cardiac Computed Tomography](#).

3. Suspected tracheobronchial anomalies and stenoses, including tracheobronchial narrowing secondary to vascular anomaly, mass, inflammatory/infectious process, suspected foreign body, or congenital anomaly (such as tracheobronchus) [17-21].
4. Mediastinum – CT is used in the workup for a mediastinal mass to assess its anatomy as well as to indicate a specific diagnosis or differential diagnosis [22, 23].
 - a. Neoplastic conditions can be evaluated by CT. These include but are not restricted to germ cell tumors, lymphoma, or thymic tumors. Posterior mediastinal neurogenic tumors can also be imaged by CT, particularly with multidetector technology and reformats, but magnetic resonance imaging (MRI) is often more useful to depict neural foraminal, intraspinal, or vertebral/chest wall involvement.
 - b. CT is used in the evaluation of congenital abnormalities such as ectopic thymic tissue and bronchopulmonary foregut malformations that affect the mediastinum. The latter include bronchogenic cyst, esophageal duplication cyst, and neuroenteric cyst (although MRI is often necessary for any mass, cystic or solid, in a paraspinal location to assess for potential neural foraminal intraspinal or vertebral/chest wall component).
 - c. Infectious or inflammatory processes affecting the mediastinum can also be assessed, such as mediastinitis, abscess, or sternal osteomyelitis. The distribution of lymphadenopathy due to an infectious or inflammatory cause is evaluated by CT.
 - d. CT is used in the setting of trauma that is not adequately assessed by radiography.
5. Lung – CT is the primary cross-sectional imaging modality to evaluate the lung parenchyma [24-40].
 - a. These include imaging of infection/pneumonia complicated by involvement of the pleural space (such as parapneumonic effusion, empyema, or broncho-pleural fistula), the lung (such as cavitation/necrosis or abscess), or the pericardium (such as purulent pericarditis). In patients with persistent or recurrent pneumonias or whose plain film is atypical for pneumonia, CT is used to assess for possible underlying congenital lesion or mass. CT is also used to assess the sequelae of respiratory infections (such as bronchiectasis and bronchiolitis obliterans). In immunocompromised patients

- CT can be used in the absence of definite plain film abnormality to detect early manifestations of opportunistic infections.
- b. CT is used in the evaluation of interstitial lung disease, and “high-resolution” examinations are often utilized in this setting. This includes imaging of the various primary pulmonary interstitial lung diseases, as well as interstitial lung disease related to systemic processes such as collagen vascular, connective tissue, or autoimmune diseases. Some patients with cystic fibrosis may benefit from thoracic imaging by CT.
 - c. Congenital pulmonary abnormalities are also well assessed by CT. Bronchopulmonary foregut or congenital pulmonary airway malformations (CPAM) that affect the lung are well depicted by CT, such as congenital lobar emphysema, congenital cystic adenomatoid malformation, bronchogenic cyst, and pulmonary sequestration. CT can also be used to assess pulmonary agenesis, hypoplasia, and aplasia, and other hypogenetic lung complexes, bronchial atresia, horseshoe lung, and pulmonary arteriovenous malformation.
 - d. Neoplastic conditions (benign or malignant) affecting the lungs are imaged by CT. CT is the mainstay for surveillance in patients with a known underlying primary malignancy that metastasizes to lung. In immunocompromised patients, CT is used in the evaluation for lymphoproliferative disorder. Less commonly, due to the infrequency of these disorders, CT evaluates primary pulmonary neoplasms such as pulmonary blastoma.
 - e. CT is used in the setting of trauma that is not adequately assessed by radiography. Lung contusions and traumatic pneumatoceles are better seen on CT than on radiography.
 - f. CT is also used in assessing other nonspecific but clinically significant signs or symptoms in the respiratory system not adequately addressed by radiography.

B. Abdomen and Pelvis

1. Hollow viscera [41-58]
 - a. Suspected inflammatory or infectious processes affecting the GI tract, including the gastroesophageal junction, stomach, small intestine, colon, or appendix. These processes include, but are not limited to, appendicitis, infectious enteritis, inflammatory bowel disease, infectious, neutropenic colitis or radiation enteritis.
- b. Known or suspected congenital abnormalities, including gastrointestinal duplication cysts, and complications of omphalomesenteric duct remnants, such as Meckel diverticulitis.
- c. Benign and malignant neoplastic tumors, including but not limited to lymphoma (particularly Burkitt's lymphoma), gastrointestinal stromal tumor (GIST), lipoma, and postradiation enteritis.
- d. CT is frequently used in the setting of blunt or penetrating abdominal trauma. It can evaluate for primary or secondary signs of bowel injury in the setting of penetrating abdominal trauma or blunt trauma, or in nonaccidental injury to the duodenum or small bowel.
- e. CT can be used to further evaluate patients presenting with bowel obstruction.

2. Liver and gallbladder [59-69]

- a. Evaluation of patients with neoplastic processes associated with primary hepatic malignancies (benign or malignant), including but not limited to hemangiopericytoma, hepatoblastoma, and hepatocellular carcinoma, as well as liver metastases to evaluate for the presence and extent of tumor in the liver.
- b. Evaluation of blunt or penetrating abdominal trauma with suspected hepatic trauma to further assess the extent of parenchymal and hepatic vascular injury.
- c. Suspected hepatic infection, including pyogenic or amebic liver abscesses.
- d. Evaluation of disorders of the gallbladder and biliary tract to supplement ultrasound including gallbladder anomalies, thickening, bile duct dilation or stenosis, and bile leak.
- e. Evaluation of congenital abnormalities of the liver and biliary tree, including heterotaxy and associated anomalies, and all types of choledochal cyst.

3. Pancreas [70-75]

- a. CT can be used as a supplement to ultrasound or as the primary imaging tool to evaluate for complications of pancreatitis, including pancreatic hemorrhage or necrosis, peripancreatic vascular thrombosis, pseudocyst formation, secondary inflammation of hollow visceral structures within the anterior pararenal space, regional vessel thrombosis, or duct abnormalities, including stones or dilation.
- b. CT can be used in evaluating pancreatic tumors to further characterize the extent of

- lesion, staging, and involvement of adjacent structures.
- c. CT can be used in the setting of blunt or penetrating abdominal trauma with possible or suspected pancreatic trauma to evaluate the integrity of the gland, the extent of pancreatic injury including fracture and/or pancreatic ductal injury, and injury to adjacent solid or hollow visceral structures.
4. Kidneys [76-84]
- a. Diagnosis of suspected renal, ureteral, or bladder stones in the setting of hematuria. In children, CT should be used when ultrasound and radiographs do not provide enough information for optimal management.
 - b. Evaluation of renal and/or ureteral trauma. Delayed imaging may be useful if injury to the collecting system is suspected.
 - c. Detection and staging of renal tumors (benign and malignant), including vascular invasion, frequently as a supplement to ultrasonography.
 - d. Congenital anomalies of the genitourinary tract.
 - e. Obstruction of the genitourinary (GU) tract secondary to congenital anomalies, mass, infection/inflammation, or trauma.
 - f. Suspected or known infection of the GU tract, including focal or generalized pyelonephritis or abscess.
 - g. For renovascular evaluation in the setting of traumatic injury, hypertension, renal donor transplant evaluation, or regional masses, CT angiography is typically performed in the evaluation of hypertension and transplant donor assessment.
5. Adrenal gland [85-91]
- a. Evaluation of suspected adrenal hemorrhage as a potential supplement to ultrasound.
 - b. Evaluation of suspected adrenal trauma in the setting of blunt or penetrating trauma to the upper abdomen.
 - c. Primary evaluation of adrenal neoplasms, often as a supplement to ultrasound for suspected neuroblastoma or adrenocortical carcinoma.
6. Spleen [92-98]
- a. Primary evaluation of suspected splenic injury in the setting of blunt or penetrating trauma.
 - b. Further characterization of primary cystic or solid lesions of the spleen.
 - c. Other conditions such as infarction, sequestration (sickle cell disease), granulomatous disease, wandering spleen/torsions.
7. Abdomen and pelvis
- a. Mass or mass-like conditions of the pelvis organs including inflammatory/infectious processes and vascular malformations, and evaluation of lymph nodes.
 - b. Anomalies of the genital tract not adequately assessed by ultrasound where MRI is contraindicated or not available.
 - c. Evaluation of bladder rupture after trauma or bladder surgery.
8. Mesentery/omentum/peritoneum/abdominal wall/diaphragm [90-103]
- a. Suspected inflammatory or infectious processes affecting the mesentery, peritoneum, or omentum, including but not limited to abscess or generalized peritonitis.
 - b. Evaluation of peritoneal fluid in appropriate clinical circumstances.
 - c. Possible pneumoperitoneum.
 - d. Evaluation of cystic malformations, including mesenteric cyst and lymphangioma.
 - e. Benign or malignant neoplastic processes, including teratoma, sarcoma, and peritoneal spread of disease.
 - f. Focal omental infarction (cause of abdominal pain).
 - g. Post-traumatic processes.
 - h. Evaluation of congenital or post-traumatic abnormalities of the anterior abdominal wall.
 - i. Evaluation of congenital or post-traumatic abnormalities/defects of the diaphragm.
- C. Extremities/Musculoskeletal [104-135]
- CT should be used for the evaluation of bone lesions when the radiologist believes it is the proper modality or where MRI is contraindicated or not readily available. Some of the primary indications for CT in musculoskeletal imaging are further anatomic characterization of certain fractures, particularly those involving a joint or growth plate, and further evaluation of the internal characteristics of bone lesions seen by plain-film radiography.
1. Pelvis
- a. Inflammatory conditions such as osteomyelitis of pelvic bones, myositis, or complex inflammatory conditions where MRI is contraindicated or not available.
 - b. Evaluation of the extent of trauma to the pelvic bones or internal organs, and of hemorrhage.

2. Shoulder
 - a. Evaluation of glenoid morphology, glenoid dysplasia, asymmetry of rotator cuff musculature, and acquired glenohumeral deformity such as in patients with residual brachial plexus injury after birth.
 - b. Further characterization of fractures of the humerus, scapula and/or clavicle in or around the shoulder region where radiography is insufficient.
 - c. Diagnosis, further characterization and/or follow-up of benign and malignant bone tumors and infections if not adequately evaluated by MRI.
 - d. Assessment for radiodense foreign body.
3. Elbow
 - a. Further characterization of fractures of the humerus, ulna, and/or radius in or around the elbow joint.
 - b. Follow-up of fracture complications (such as premature growth plate fusion and intra-articular loose bodies).
 - c. Diagnosis and/or follow-up of osteochondritis dissecans.
 - d. Diagnosis, further characterization, and/or follow-up of benign and malignant bone tumors if not adequately evaluated by MRI, or supplemental to MRI.
 - e. Evaluation of suspected osteoid osteoma or treated osteoid osteoma around the elbow joint or long bones of the upper extremities.
 - f. Congenital malformations and infections not adequately assessed by conventional radiographs or ultrasound.
4. Hand and wrist
 - a. Further characterization of fractures in the hand or wrist related to acute trauma.
 - b. Evaluation of acute or chronic scaphoid fracture.
 - c. Evaluation of fracture complications (such as premature growth plate fusion).
 - d. Diagnosis and/or follow-up of osteochondritis dissecans.
 - e. Diagnosis and/or follow-up of osteoid osteoma.
 - f. Diagnosis and/or follow-up of benign and malignant bone tumors and infections if not adequately evaluated by MRI, or supplemental to MRI.
5. Hip and thigh
 - a. Further characterization of fractures in the bony pelvis and proximal femora.
 - b. Evaluation of patients with developmental dysplasia of the hip, including evaluation of relationship of femoral head and triradiate cartilage in patients in a SPICA cast.
6. Knee
 - a. Further characterization of fractures in the knee related to acute trauma.
 - b. Follow-up of fracture complications (such as premature growth plate fusion and intra-articular loose bodies).
 - c. Diagnosis and/or follow-up of osteochondritis dissecans.
 - d. Diagnosis and/or follow-up of osteoid osteoma.
 - e. Diagnosis and/or follow-up of benign and malignant bone tumors if not adequately evaluated by MRI, or supplemental to MRI.
 - f. Congenital malformations not adequately assessed by conventional radiographs.
- D. Foot and Ankle
 1. Further characterization of fractures in the foot or ankle related to acute trauma, including, but not limited to, triplane fracture of the ankle, and fracture involving the tibial plafond.
 2. Follow-up of fracture complications (such as premature growth plate fusion and intra-articular loose bodies).
 3. Diagnosis of tarsal coalition and follow-up of surgically corrected tarsal coalition.
 4. Diagnosis and/or follow-up of osteochondritis dissecans.
 5. Diagnosis and/or follow-up of benign and malignant bone tumors if not adequately evaluated by MRI, or supplemental to MRI.
 6. Congenital malformations not adequately assessed by conventional radiographs.

III. QUALIFICATIONS AND RESPONSIBILITIES OF PERSONNEL

See the [ACR Practice Guideline for Performing and Interpreting Diagnostic Computed Tomography \(CT\)](#).

IV. SPECIFICATIONS OF THE EXAMINATION

The written or electronic request for pediatric CT should provide sufficient information to demonstrate the medical necessity of the examination and allow for the proper performance and interpretation of the examination.

Documentation that satisfies medical necessity includes 1) signs and symptoms and/or 2) relevant history (including known diagnoses). The provision of additional information regarding the specific reason for the examination or a provisional diagnosis would be helpful and may at times be needed to allow for the proper performance and interpretation of the examination.

The request for the examination must be originated by a physician or other appropriately licensed health care provider. The accompanying clinical information should be provided by a physician or other appropriately licensed health care provider familiar with the patient's clinical problem or question and consistent with the state scope of practice requirements. (ACR Resolution 35, adopted in 2006)

Images should be labeled with the following: a) patient identification, b) facility identification, c) examination date, and d) the side (right or left) of the anatomic site imaged.

Additionally, an attempt should be made to obtain and review prior studies.

A. General Considerations

Pediatric CT may require different examination preparation and performance than in adults. Preparation includes ensuring appropriate NPO status if moderate sedation or general anesthesia is potentially necessary and adequate hydration and assessment of pediatric specific renal function prior to intravenous (IV) contrast media administration. For scan performance, single-phase scanning is the standard rather than the exception. Only the necessary scan coverage should be obtained, and changes in scan parameters — including beam collimation — tube current, gantry cycle time, and peak kilovoltage should be adjusted for the size of the child, the region scanned, and the clinical indications.

The physician responsible for the examination shall supervise patient selection and preparation, and be available in person or by phone for consultation.

Certain indications require administration of IV contrast media. IV contrast enhancement should be performed using appropriate injection protocols and in accordance with the institution's policy on IV contrast utilization. (See the [ACR-SPR Practice Guideline for the Use of Intravascular Contrast Media](#) and the [ACR Manual on Contrast Media](#).)

Appropriate emergency equipment and medications must be immediately available to treat adverse reactions associated with administered medications. The equipment and medications should be monitored for inventory and drug expiration dates on a regular basis. The equipment, medications, and other emergency support must also be appropriate for the range of ages and sizes in the patient population.

B. Examination Technique [136,137]

1. Chest

- a. Scanning parameters should be optimized to obtain diagnostic image quality while adhering to the as low as reasonably achievable (ALARA) principle. The scan area should be minimized according to the clinical indication. The scanning parameters, including kVp, tube current, and exposure time (mAs), should be changed according to body size, area of interest and clinical indication. This can be achieved by using weight-based tables or by using automatic exposure control (see the following Web site: www.imagegently.org). In addition, mAs can be further reduced if noncontrast scans are performed to evaluate calcifications only, or in cases in which only gross bony relationships are being evaluated, such as scans done for preoperative pectus excavatum evaluation. Consideration should be given to shielding superficial structures in the scan region such as breast and thyroid, depending on the specific equipment and protocols used [138].
- b. The examination may be conducted with or without IV contrast as clinically indicated. IV contrast may be necessary, based on the clinical indication for the imaging study. A dosage of 2 mL/kg (not to exceed the usual adult dose) is used routinely. Volume of contrast, rate of injection, scan delayed time, and hand/power injection should be determined according to the location, size, and type of the IV access, the child's body size, the underlying disease (such as congestive heart failure), and the clinical indication [139].
- c. High resolution scans may be useful if the primary indication is for the evaluation of

interstitial lung disease, as sharper algorithms are helpful in the evaluation of lung parenchyma in older children. The original data set can be reconstructed with both routine and high-resolution algorithms if both soft tissue and pulmonary parenchymal information is needed, without need to rescan the patient. It is important to remember that not all diagnostic chest CT studies in infants and children require imaging of the entire anatomy of the chest. In certain clinical situations, if only a sampling of the lung parenchyma is required to answer a specific clinical question (i.e., to rule out bronchiectasis or interstitial disease), a limited number (e.g., 4 to 6 slices) of 1 to 1.25 mm axial slices can be performed in a high resolution bone algorithm. The distance between the limited axial images increases incrementally as patient size increases. Sequential thin slices with primary lung algorithm scanning may be needed in some patients, and these additional images are typically acquired with nonsequential increments.

- d. Postprocessing 2D reformations and 3D reconstructions or 3D volume rendering may be useful adjuncts in displaying the anatomy. If this is anticipated, CT technique providing the thinnest original (usual axial) dataset should be used.

2. Abdomen

- a. Scanning parameters should be optimized to obtain diagnostic image quality while adhering to the ALARA principle. The scan area should be minimized according to the clinical indication. The scanning parameters, including kVp, tube current, and exposure time (mAs), should be changed according to body size, area of interest, and clinical indication. This can be achieved by using weight based tables or by using automatic exposure control (see www.imagegently.org). The testicles should not be included in the scanned area unless absolutely necessary for the clinical indication. Consideration should be given to shielding superficial structures in the scan region such as the testes. If precontrast images are needed solely to determine whether calcification is present, these can be done with additional decrease in mAs.
- b. IV contrast injection is usually used in the CT evaluation of the pediatric abdomen, since vascular structures and internal organs are much better visualized due to the paucity

of body fat in many pediatric patients. There are some exceptions, including renal stone evaluation. A routine dose of 2 mL/kg is generally used. Volume of contrast, rate of injection, scan delayed time, and hand/power injection should be determined according to the location, size, and type of the IV access, the child's body size, the underlying disease (such as congestive heart failure), and the clinical indication.

- c. Enteric contrast may be used in the CT evaluation of the pediatric abdomen. Exceptions would include, but are not limited to, renal stone protocol, CT angiography, and acute trauma.
- d. In the evaluation of the pediatric patient for suspected appendicitis, IV contrast is typically used, particularly to avoid repeat scans due to equivocal findings. Precontrast scans and delayed scans are not necessary, unless a renal anomaly requiring evaluation of the collecting system is incidentally identified. Some centers use oral or rectal enteric contrast material. If oral contrast is given, sufficient time should be allowed to elapse for the contrast to reach the right lower quadrant prior to scanning.
- e. Postprocessing 2D reformations and 3D reconstructions or 3D volume rendering may be useful adjuncts in displaying the anatomy.

3. Extremities

- a. Scanning parameters should be optimized to obtain diagnostic image quality while adhering to the ALARA principle. The scan area should be minimized according to the clinical indication. The scanning parameters, including kVp, tube current, and exposure time (mAs), should be changed according to body size, area of interest, patient age and size, and clinical indication.
- b. IV contrast may not be necessary if only evaluation of the bone structure is important. IV contrast should be used if indicated to assess vessels and soft tissues.
- c. Sharper reconstruction algorithms are needed for better spatial resolution and bone detail. Smoother (softer) algorithms are better for soft tissue evaluation and 3D postprocessing.

V. DOCUMENTATION

Reporting should be in accordance with the [ACR Practice Guideline for Communication of Diagnostic Imaging Findings](#).

VI. EQUIPMENT SPECIFICATIONS

In the interest of pediatric patient safety, it is necessary to have a general knowledge of the CT equipment including the use of weight adjusted mA and kVp, slice thickness, pitch, display field of view (DFOV) and dose modulation techniques. The equipment should be in good working order, meet manufacturer and regulatory standards, and be operated safely. The equipment needs to be tested for spatial and low-contrast resolution and be well-calibrated at all times. Technologists and radiologists should be aware of important artifacts and know how to avoid problems associated with them.

A. Performance Standards

To achieve acceptable clinical CT scans of body, the CT scanner should meet or exceed the following specifications:

1. Gantry rotation times: ≤ 2 seconds.
2. Slice thickness: ≤ 5 mm (≤ 2 mm is preferred).
3. Limiting spatial resolution: 8 lp/cm for ≥ 32 cm DFOV and ≥ 10 lp/cm for < 24 cm DFOV.
4. Table pitch: no greater than 2:1 for single-row-detector helical scanners.

B. Patient monitoring equipment and facilities for cardiopulmonary resuscitation, including vital signs monitoring equipment, support equipment, and an emergency crash cart, should be immediately available. Radiologists, technologists, and staff members should be able to assist with procedures, patient monitoring, and patient support. A written policy should be in place for dealing with emergencies such as cardiopulmonary arrest.

VII. RADIATION SAFETY IN IMAGING

Radiologists, medical physicists, radiologic technologists, and all supervising physicians have a responsibility to minimize radiation dose to individual patients, to staff, and to society as a whole, while maintaining the necessary diagnostic image quality. This concept is known as “as low as reasonably achievable (ALARA).”

Facilities, in consultation with the medical physicist, should have in place and should adhere to policies and procedures, in accordance with ALARA, to vary examination protocols to take into account patient body habitus, such as height and/or weight, body mass index or lateral width. The dose reduction devices that are available on imaging equipment should be active; if not, manual techniques should be used to moderate the exposure while maintaining the necessary diagnostic image quality. Periodically, radiation exposures should be measured and patient radiation doses estimated by a medical physicist in accordance with the appropriate ACR

Technical Standard. (ACR Resolution 17, adopted in 2006 – revised in 2009, Resolution 11)

A medical physicist and radiologist together should verify that any dose reduction devices or utilities maintain acceptable image quality while actually reducing radiation dose.

Dose estimates for typical examinations should be compared against reference levels described in the [ACR Practice Guideline for Diagnostic Reference Levels in Medical X-Ray Imaging](#).

VIII. QUALITY CONTROL AND IMPROVEMENT, SAFETY, INFECTION CONTROL, AND PATIENT EDUCATION

Policies and procedures related to quality, patient education, infection control, and safety should be developed and implemented in accordance with the ACR Policy on Quality Control and Improvement, Safety, Infection Control, and Patient Education appearing under the heading *Position Statement on QC & Improvement, Safety, Infection Control, and Patient Education* on the ACR web page (<http://www.acr.org/guidelines>).

Equipment monitoring and the continuous quality control program should be in accordance with the [ACR-AAPM Technical Standard for Diagnostic Medical Physics Performance Monitoring of Computed Tomography \(CT\) Equipment](#).

ACKNOWLEDGEMENT

This guideline was revised according to the process described under the heading *The Process for Developing ACR Practice Guidelines and Technical Standards* on the ACR web page (<http://www.acr.org/guidelines>) by the Guidelines and Standards Committees of the Commissions on Pediatric Radiology and General, Small, and Rural Practice in collaboration with the SPR.

Principal Drafters

Michael Callahan, MD
Sunny Pitt, MD
Boaz Karmazyn, MD
Ron Cohen, MD

ACR Pediatric Guidelines and Standards Committee

Marta Hernanz-Schulman, MD, Chair
Brian D. Coley, MD
Kristin L. Crisci, MD
Eric N. Faerber, MD
Lynn A. Fordham, MD
Laureen M. Sena, MD
Sudha P. Singh, MD, MBBS
Peter J. Strouse, MD
Donald P. Frush, MD, Chair, Commission

ACR GSR Guidelines and Standards Committee

Julie K. Timins, MD, Chair
William R. Allen, Jr., MD
Damon A. Black, MD
Richard A. Carlson, MD
James P. Cartland, MD
Laura E. Faix, MD
Mark F. Fisher, MD
Frank R. Graybeal, Jr., MD
Louis W. Lucas, MD
Matthew S. Pollack, MD
Diane C. Strollo, MD
Fred S. Vernacchia, MD
Susan L. Voci, MD
Geoffrey G. Smith, MD, Chair, Commission

Comments Reconciliation Committee

Kimberly E. Applegate, MD, MS, Chair
Michael J. Callahan, MD
Mervyn D. Cohen, MD, MB, ChB
Donald P. Frush, MD
Richard A. Geise, PhD
Marta Hernanz-Schulman, MD
James M. Hevezi, PhD
Alan D. Kaye, MD
David C. Kushner, MD
Paul A. Larson, MD
Lawrence A. Liebscher, MD
Lorenz T. Ramseyer, MD
Cynthia K. Rigsby, MD
William E. Sheehan, MD
Manrita K. Sidhu, MD
Geoffrey G. Smith, MD
Julie K. Timins, MD
David L. Vassy, Jr., MS
Ted S. Wen, MD

REFERENCES

1. Brenner DJ, Hall EJ. Computed tomography an increasing source of radiation exposure. *N Engl J Med* 2007;357:2277-2284.
2. Donnelly LF. Use of three-dimensional reconstructed helical CT images in recognition and communication of chest wall anomalies in children. *AJR* 2001;177:441-445.
3. Emans JB, Caubet JF, Ordonez CL, Lee EY, Ciarlo M. The treatment of spine and chest wall deformities with fused ribs by expansion thoracostomy and insertion of vertical expandable prosthetic titanium rib: growth of thoracic spine and improvement of lung volumes. *Spine* 2005;30:S58-68.
4. Haller JA, Jr., Kramer SS, Lietman SA. Use of CT scans in selection of patients for pectus excavatum surgery: a preliminary report. *J Pediatr Surg* 1987;22:904-906.
5. Miller LA. Chest wall, lung, and pleural space trauma. *Radiol Clin North Am* 2006;44:213-224, viii.
6. Morris BS, Maheshwari M, Chalwa A. Chest wall tuberculosis: a review of CT appearances. *Br J Radiol* 2004;77:449-457.
7. Schulman H, Newman-Heinman N, Kurtzbart E, Maor E, Zirkin H, Laufer L. Thoracoabdominal peripheral primitive neuroectodermal tumors in childhood: radiological features. *Eur Radiol* 2000;10:1649-1652.
8. Shamberger RC, Grier HE. Chest wall tumors in infants and children. *Semin Pediatr Surg* 1994;3:267-276.
9. Wong KS, Hung IJ, Wang CR, Lien R. Thoracic wall lesions in children. *Pediatr Pulmonol* 2004;37:257-263.
10. Bruckner BA, DiBardino DJ, Cumbie TC, et al. Critical evaluation of chest computed tomography scans for blunt descending thoracic aortic injury. *Ann Thorac Surg* 2006;81:1339-1346.
11. Katz M, Konen E, Rozenman J, Szeinberg A, Itzhak Y. Spiral CT and 3D image reconstruction of vascular rings and associated tracheobronchial anomalies. *J Comput Assist Tomogr* 1995;19:564-568.
12. Kim TH, Kim YM, Suh CH, et al. Helical CT angiography and three-dimensional reconstruction of total anomalous pulmonary venous connections in neonates and infants. *AJR* 2000;175:1381-1386.
13. Lee EY, Siegel MJ, Hildebolt CF, Gutierrez FR, Bhalla S, Fallah JH. MDCT evaluation of thoracic aortic anomalies in pediatric patients and young adults: comparison of axial, multiplanar, and 3D images. *AJR* 2004;182:777-784.
14. Melton SM, Kerby JD, McGiffin D, et al. The evolution of chest computed tomography for the definitive diagnosis of blunt aortic injury: a single-center experience. *J Trauma* 2004;56:243-250.
15. Qanadli SD, Hajjam ME, Mesurolle B, et al. Pulmonary embolism detection: prospective evaluation of dual-section helical CT versus selective pulmonary arteriography in 157 patients. *Radiology* 2000;217:447-455.
16. Stein PD, Fowler SE, Goodman LR, et al. Multidetector computed tomography for acute pulmonary embolism. *N Engl J Med* 2006;354:2317-2327.
17. Choo KS, Lee HD, Ban JE, et al. Evaluation of obstructive airway lesions in complex congenital heart disease using composite volume-rendered images from multislice CT. *Pediatr Radiol* 2006;36:219-223.
18. Heyer CM, Nuesslein TG, Jung D, et al. Tracheobronchial anomalies and stenoses: detection with low-dose multidetector CT with virtual tracheobronchoscopy--comparison with

- flexible tracheobronchoscopy. *Radiology* 2007;242:542-549.
19. Honnep D, Wildberger JE, Das M, et al. Value of virtual tracheobronchoscopy and bronchography from 16-slice multidetector-row spiral computed tomography for assessment of suspected tracheobronchial stenosis in children. *Eur Radiol* 2006;16:1684-1691.
 20. Long FR. Imaging evolution of airway disorders in children. *Radiol Clin North Am* 2005;43:371-389.
 21. Pacharn P, Poe SA, Donnelly LF. Low-tube-current multidetector CT for children with suspected extrinsic airway compression. *AJR* 2002;179:1523-1527.
 22. Daltro P, Fricke BL, Kuroki I, Domingues R, Donnelly LF. CT of congenital lung lesions in pediatric patients. *AJR Am J Roentgenol* 2004;183:1497-1506.
 23. Franco A, Mody NS, Meza MP. Imaging evaluation of pediatric mediastinal masses. *Radiol Clin North Am* 2005;43:325-353.
 24. Allen GS, Cox CS, Jr. Pulmonary contusion in children: diagnosis and management. *South Med J* 1998;91:1099-1106.
 25. Brody AS. Imaging considerations: interstitial lung disease in children. *Radiol Clin North Am* 2005;43:391-403.
 26. Cooper P, MacLean J. High-resolution computed tomography (HRCT) should not be considered as a routine assessment method in cystic fibrosis lung disease. *Paediatr Respir Rev* 2006;7:197-201.
 27. Copley SJ. Application of computed tomography in childhood respiratory infections. *Br Med Bull* 2002;61:263-279.
 28. Donnelly LF. Maximizing the usefulness of imaging in children with community-acquired pneumonia. *AJR* 1999;172:505-512.
 29. Donnelly LF. Imaging in immunocompetent children who have pneumonia. *Radiol Clin North Am* 2005;43:253-265.
 30. Donnelly LF, Frush DP. Pediatric multidetector body CT. *Radiol Clin North Am* 2003;41:637-655.
 31. Fan LL, Deterding RR, Langston C. Pediatric interstitial lung disease revisited. *Pediatr Pulmonol* 2004;38:369-378.
 32. Hall A, Johnson K. The imaging of paediatric thoracic trauma. *Paediatr Respir Rev* 2002;3:241-247.
 33. Koh DM, Hansell DM. Computed tomography of diffuse interstitial lung disease in children. *Clin Radiol* 2000;55:659-667.
 34. Lim GY, Newman B, Kurland G, Webber SA. Post-transplantation lymphoproliferative disorder: manifestations in pediatric thoracic organ recipients. *Radiology* 2002;222:699-708.
 35. Newman B. Congenital bronchopulmonary foregut malformations: concepts and controversies. *Pediatr Radiol* 2006;36:773-791.
 36. Paterson A. Imaging evaluation of congenital lung abnormalities in infants and children. *Radiol Clin North Am* 2005;43:303-323.
 37. Paterson A, Frush DP. Dose reduction in paediatric MDCT: general principles. *Clin Radiol* 2007;62:507-517.
 38. Siegel MJ FD, Brody AS. Multidetector CT in pediatrics (RSP1504RC). *RSNA refresher course*. Oak Brook, IL; 2005.
 39. Thomas KE, Owens CM, Britto J, Nadel S, Habibi P, Nicholson R. Efficacy of chest CT in a pediatric ICU: a prospective study. *Chest* 2000;117:1697-1705.
 40. Tiddens HA. Chest computed tomography scans should be considered as a routine investigation in cystic fibrosis. *Paediatr Respir Rev* 2006;7:202-208.
 41. Balthazar EJ. CT of the gastrointestinal tract: principles and interpretation. *AJR* 1991;156:23-32.
 42. Balthazar EJ, Birnbaum BA, Yee J, Megibow AJ, Roshkow J, Gray C. Acute appendicitis: CT and US correlation in 100 patients. *Radiology* 1994;190:31-35.
 43. Blickman JG, Boland GW, Cleveland RH, Bramson RT, Lee MJ. Pseudomembranous colitis: CT findings in children. *Pediatr Radiol* 1995;25 Suppl 1:S157-159.
 44. Breen DJ, Janzen DL, Zwirewich CV, Nagy AG. Blunt bowel and mesenteric injury: diagnostic performance of CT signs. *J Comput Assist Tomogr* 1997;21:706-712.
 45. Bulas DI, Taylor GA, Eichelberger MR. The value of CT in detecting bowel perforation in children after blunt abdominal trauma. *AJR* 1989;153:561-564.
 46. Cox TD, Kuhn JP. CT scan of bowel trauma in the pediatric patient. *Radiol Clin North Am* 1996;34:807-818.
 47. Cox TD, Winters WD, Weinberger E. CT of intussusception in the pediatric patient: diagnosis and pitfalls. *Pediatr Radiol* 1996;26:26-32.
 48. Fitch SJ, Tonkin IL, Tonkin AK. Imaging of foregut duplication cysts. *Radiographics* 1986;6:189-201.
 49. Frager D, Medwid SW, Baer JW, Mollinelli B, Friedman M. CT of small-bowel obstruction: value in establishing the diagnosis and determining the degree and cause. *AJR* 1994;162:37-41.
 50. Gore RM, Balthazar EJ, Ghahremani GG, Miller FH. CT features of ulcerative colitis and Crohn's disease. *AJR* 1996;167:3-15.
 51. Jabra AA, Shalaby-Rana EI, Fishman EK. CT of appendicitis in children. *J Comput Assist Tomogr* 1997;21:661-666.
 52. Ruess L, Sivit CJ, Eichelberger MR, Gotschall CS, Taylor GA. Blunt abdominal trauma in children: impact of CT on operative and nonoperative management. *AJR* 1997;169:1011-1014.

53. Siegel MJ, Evans SJ, Balfe DM. Small bowel disease in children: diagnosis with CT. *Radiology* 1988;169:127-130.
54. Sivit CJ, Taylor GA, Bulas DI, Bowman LM, Eichelberger MR. Blunt trauma in children: significance of peritoneal fluid. *Radiology* 1991;178:185-188.
55. Sivit CJ, Taylor GA, Bulas DI, Kushner DC, Potter BM, Eichelberger MR. Posttraumatic shock in children: CT findings associated with hemodynamic instability. *Radiology* 1992;182:723-726.
56. Taylor GA, Eichelberger MR, O'Donnell R, Bowman L. Indications for computed tomography in children with blunt abdominal trauma. *Ann Surg* 1991;213:212-218.
57. Taylor GA, Fallat ME, Eichelberger MR. Hypovolemic shock in children: abdominal CT manifestations. *Radiology* 1987;164:479-481.
58. Taylor GA, Fallat ME, Potter BM, Eichelberger MR. The role of computed tomography in blunt abdominal trauma in children. *J Trauma* 1988;28:1660-1664.
59. Baron RL. Computed tomography of the biliary tree. *Radiol Clin North Am* 1991;29:1235-1250.
60. Bonaldi VM, Bret PM, Reinhold C, Atri M. Helical CT of the liver: value of an early hepatic arterial phase. *Radiology* 1995;197:357-363.
61. Brick SH, Taylor GA, Potter BM, Eichelberger MR. Hepatic and splenic injury in children: role of CT in the decision for laparotomy. *Radiology* 1987;165:643-646.
62. Dachman AH, Pakter RL, Ros PR, Fishman EK, Goodman ZD, Lichtenstein JE. Hepatoblastoma: radiologic-pathologic correlation in 50 cases. *Radiology* 1987;164:15-19.
63. Dalen K, Day DL, Ascher NL, et al. Imaging of vascular complications after hepatic transplantation. *AJR* 1988;150:1285-1290.
64. Finn JP, Hall-Craggs MA, Dicks-Mireaux C, et al. Primary malignant liver tumors in childhood: assessment of resectability with high-field MR and comparison with CT. *Pediatr Radiol* 1990;21:34-38.
65. Holbert BL, Baron RL, Dodd GD, 3rd. Hepatic infarction caused by arterial insufficiency: spectrum and evolution of CT findings. *AJR* 1996;166:815-820.
66. Jeffrey RB, Jr., Federle MP, Laing FC, Wing VW. Computed tomography of blunt trauma to the gallbladder. *J Comput Assist Tomogr* 1986;10:756-758.
67. Noda T, Todani T, Watanabe Y, Yamamoto S. Liver volume in children measured by computed tomography. *Pediatr Radiol* 1997;27:250-252.
68. Pobiel RS, Bisset GS, 3rd. Pictorial essay: imaging of liver tumors in the infant and child. *Pediatr Radiol* 1995;25:495-506.
69. Shirkhoda A. CT findings in hepatosplenic and renal candidiasis. *J Comput Assist Tomogr* 1987;11:795-798.
70. Balthazar EJ, Robinson DL, Megibow AJ, Ranson JH. Acute pancreatitis: value of CT in establishing prognosis. *Radiology* 1990;174:331-336.
71. Daneman A, Gaskin K, Martin DJ, Cutz E. Pancreatic changes in cystic fibrosis: CT and sonographic appearances. *AJR* 1983;141:653-655.
72. Herman TE, Siegel MJ. CT of the pancreas in children. *AJR Am J Roentgenol* 1991;157:375-379.
73. Hernanz-Schulman M, Teele RL, Perez-Atayde A, et al. Pancreatic cystosis in cystic fibrosis. *Radiology* 1986;158:629-631.
74. King LR, Siegel MJ, Balfe DM. Acute pancreatitis in children: CT findings of intra- and extrapancreatic fluid collections. *Radiology* 1995;195:196-200.
75. Siegel MJ, Sivit CJ. Pancreatic emergencies. *Radiol Clin North Am* 1997;35:815-830, 814.
76. Agrons GA, Wagner BJ, Davidson AJ, Suarez ES. Multilocular cystic renal tumor in children: radiologic-pathologic correlation. *Radiographics* 1995;15:653-669.
77. Dacher JN, Boillot B, Eurin D, Marguet C, Mitrofanoff P, Le Dosseur P. Rational use of CT in acute pyelonephritis: findings and relationships with reflux. *Pediatr Radiol* 1993;23:281-285.
78. Fanney DR, Casillas J, Murphy BJ. CT in the diagnosis of renal trauma. *Radiographics* 1990;10:29-40.
79. Fernbach SK, Feinstein KA, Donaldson JS, Baum ES. Nephroblastomatosis: comparison of CT with US and urography. *Radiology* 1988;166:153-156.
80. Kawashima A, Sandler CM, Goldman SM, Raval BK, Fishman EK. CT of renal inflammatory disease. *Radiographics* 1997;17:851-866; discussion 867-858.
81. Montgomery P, Kuhn JP, Afshani E. CT evaluation of severe renal inflammatory disease in children. *Pediatr Radiol* 1987;17:216-222.
82. Smith RC, Rosenfield AT, Choe KA, et al. Acute flank pain: comparison of non-contrast-enhanced CT and intravenous urography. *Radiology* 1995;194:789-794.
83. Smith RC, Verga M, McCarthy S, Rosenfield AT. Diagnosis of acute flank pain: value of unenhanced helical CT. *AJR* 1996;166:97-101.
84. Weinberger E, Rosenbaum DM, Pendergrass TW. Renal involvement in children with lymphoma: comparison of CT with sonography. *AJR* 1990;155:347-349.
85. Abramson SJ. Adrenal neoplasms in children. *Radiol Clin North Am* 1997;35:1415-1453.

86. Berdon WE, Ruzal-Shapiro C, Abramson SJ, Garvin J. The diagnosis of abdominal neuroblastoma: relative roles of ultrasonography, CT, and MRI. *Urol Radiol* 1992;14:252-262.
87. Burks DW, Mirvis SE, Shanmuganathan K. Acute adrenal injury after blunt abdominal trauma: CT findings. *AJR* 1992;158:503-507.
88. Cirillo RL, Jr., Bennett WF, Vitellas KM, Poulos AG, Bova JG. Pathology of the adrenal gland: imaging features. *AJR* 1998;170:429-435.
89. Kenney PJ, Robbins GL, Ellis DA, Spirt BA. Adrenal glands in patients with congenital renal anomalies: CT appearance. *Radiology* 1985;155:181-182.
90. Sivit CJ, Ingram JD, Taylor GA, Bulas DI, Kushner DC, Eichelberger MR. Posttraumatic adrenal hemorrhage in children: CT findings in 34 patients. *AJR* 1992;158:1299-1302.
91. Westra SJ, Zaninovic AC, Hall TR, Kangarloo H, Boechat MI. Imaging of the adrenal gland in children. *Radiographics* 1994;14:1323-1340.
92. Caslowitz PL, Labs JD, Fishman EK, Siegelman SS. The changing spectrum of splenic abscess. *Clin Imaging* 1989;13:201-207.
93. Dodds WJ, Taylor AJ, Erickson SJ, Stewart ET, Lawson TL. Radiologic imaging of splenic anomalies. *AJR* 1990;155:805-810.
94. Donnelly LF. CT imaging of immunocompromised children with acute abdominal symptoms. *AJR* 1996;167:909-913.
95. Donnelly LF, Foss JN, Frush DP, Bisset GS, 3rd. Heterogeneous splenic enhancement patterns on spiral CT images in children: minimizing misinterpretation. *Radiology* 1999;210:493-497.
96. Freeman JL, Jafri SZ, Roberts JL, Mezwa DG, Shirkhoda A. CT of congenital and acquired abnormalities of the spleen. *Radiographics* 1993;13:597-610.
97. Urban BA, Fishman EK. Helical CT of the spleen. *AJR* 1998;170:997-1003.
98. Wadsworth DT, Newman B, Abramson SJ, Carpenter BL, Lorenzo RL. Splenic lymphangiomatosis in children. *Radiology* 1997;202:173-176.
99. Balthazar EJ, Lefkowitz RA. Left-sided omental infarction with associated omental abscess: CT diagnosis. *J Comput Assist Tomogr* 1993;17:379-381.
100. Hamrick-Turner JE, Chiechi MV, Abbott PL, Ros PR. Neoplastic and inflammatory processes of the peritoneum, omentum, and mesentery: diagnosis with CT. *Radiographics* 1992;12:1051-1068.
101. Miller PA, Mezwa DG, Feczkó PJ, Jafri ZH, Madrazo BL. Imaging of abdominal hernias. *Radiographics* 1995;15:333-347.
102. Pickhardt PJ, Siegel MJ. Abdominal manifestations of posttransplantation lymphoproliferative disorder. *AJR Am J Roentgenol* 1998;171:1007-1013.
103. Rao PM, Rhea JT, Novelline RA. CT diagnosis of mesenteric adenitis. *Radiology* 1997;202:145-149.
104. Aisen AM, Martel W, Braunstein EM, McMillin KI, Phillips WA, Kling TF. MRI and CT evaluation of primary bone and soft-tissue tumors. *AJR* 1986;146:749-756.
105. Burk DL, Jr., Mears DC, Kennedy WH, Cooperstein LA, Herbert DL. Three-dimensional computed tomography of acetabular fractures. *Radiology* 1985;155:183-186.
106. Chan DP, Abujudeh HH, Cushing GL, Jr., Novelline RA. CT cystography with multiplanar reformation for suspected bladder rupture: experience in 234 cases. *AJR* 2006;187:1296-1302.
107. Chandnani VP, Beltran J, Morris CS, et al. Acute experimental osteomyelitis and abscesses: detection with MR imaging versus CT. *Radiology* 1990;174:233-236.
108. Cone RO, 3rd, Nguyen V, Flournoy JG, Guerra J, Jr. Triplane fracture of the distal tibial epiphysis: radiographic and CT studies. *Radiology* 1984;153:763-767.
109. Eggli KD, King SH, Boal DK, Quiogue T. Low-dose CT of developmental dysplasia of the hip after reduction: diagnostic accuracy and dosimetry. *AJR* 1994;163:1441-1443.
110. Einstein DM, Singer AA, Chilcote WA, Desai RK. Abdominal lymphadenopathy: spectrum of CT findings. *Radiographics* 1991;11:457-472.
111. Emery KH. Splenic emergencies. *Radiol Clin North Am* 1997;35:831-843.
112. Ferrozza F, Bova D, Draghi F, Garlaschi G. CT findings in primary vascular tumors of the spleen. *AJR* 1996;166:1097-1101.
113. Fishman EK, Wyatt SH, Bluemke DA, Urban BA. Spiral CT of musculoskeletal pathology: preliminary observations. *Skeletal Radiol* 1993;22:253-256.
114. Frush DP, Siegel MJ, Bisset GS, 3rd. From the RSNA refresher courses. Challenges of pediatric spiral CT. *Radiographics* 1997;17:939-959.
115. Glass RB, Rushton HG. Delayed spontaneous rupture of augmented bladder in children: diagnosis with sonography and CT. *AJR* 1992;158:833-835.
116. Godfroid N, Stalens JP. Thigh pain due to obturator internus phlegmon: a diagnostic challenge. *Eur J Pediatr* 1995;154:273-274.
117. Herman TE, Siegel MJ. CT of acute splenic torsion in children with wandering spleen. *AJR* 1991;156:151-153.
118. Hernandez RJ, Tachdjian MO, Poznanski AK, Dias LS. CT determination of femoral torsion. *AJR* 1981;137:97-101.

119. Holland DG, Quint LE. Traumatic rupture of the diaphragm without visceral herniation: CT diagnosis. *AJR* 1991;157:17-18.
120. Janzen DL, Connell DG, Munk PL, Buckley RE, Meek RN, Schechter MT. Intraarticular fractures of the calcaneus: value of CT findings in determining prognosis. *AJR* 1992;158:1271-1274.
121. Kuhn JP, Berger PE. Computed tomographic diagnosis of osteomyelitis. *Radiology* 1979;130:503-506.
122. Laasonen EM, Jokio P, Lindholm TS. Tibial torsion measured by computed tomography. *Acta Radiol Diagn (Stockh)* 1984;25:325-329.
123. Lee MS, Harcke HT, Kumar SJ, Bassett GS. Subtalar joint coalition in children: new observations. *Radiology* 1989;172:635-639.
124. Mahboubi S. CT appearance of nidus in osteoid osteoma versus sequestration in osteomyelitis. *J Comput Assist Tomogr* 1986;10:457-459.
125. Mahboubi S, Horstmann H. Femoral torsion: CT measurement. *Radiology* 1986;160:843-844.
126. Murphy SB, Simon SR, Kijewski PK, Wilkinson RH, Griscom NT. Femoral anteversion. *J Bone Joint Surg Am* 1987;69:1169-1176.
127. Pereira SJ, O'Brien DP, Luchette FA, et al. Dynamic helical computed tomography scan accurately detects hemorrhage in patients with pelvic fracture. *Surgery* 2000;128:678-685.
128. Siegel MJ. Pelvic tumors in childhood. *Radiol Clin North Am* 1997;35:1455-1475.
129. Silber JS, Flynn JM, Katz MA, Ganley TJ, Koffler KM, Drummond DS. Role of computed tomography in the classification and management of pediatric pelvic fractures. *J Pediatr Orthop* 2001;21:148-151.
130. Skiadas VT, Koutoulidis V, Eleytheriades M, et al. Ovarian masses in young adolescents: imaging findings with surgical confirmation. *Eur J Gynaecol Oncol* 2004;25:201-206.
131. Stewart NR, Gilula LA. CT of the wrist: a tailored approach. *Radiology* 1992;183:13-20.
132. Stiletto RJ, Baacke M, Gotzen L. Comminuted pelvic ring disruption in toddlers: management of a rare injury. *J Trauma* 2000;48:161-164.
133. Viani RM, Bromberg K, Bradley JS. Obturator internus muscle abscess in children: report of seven cases and review. *Clin Infect Dis* 1999;28:117-122.
134. Vo NJ, Gash J, Browning J, Hutson RK. Pelvic imaging in the stable trauma patient: is the AP pelvic radiograph necessary when abdominopelvic CT shows no acute injury? *Emerg Radiol* 2004;10:246-249.
135. Wechsler RJ, Schweitzer ME, Deely DM, Horn BD, Pizzutillo PD. Tarsal coalition: depiction and characterization with CT and MR imaging. *Radiology* 1994;193:447-452.
136. Fricke BL, Donnelly LF, Frush DP, et al. In-plane bismuth breast shields for pediatric CT: effects on radiation dose and image quality using experimental and clinical data. *AJR* 2003;180:407-411.
137. Greess H, Lutze J, Nomayr A, et al. Dose reduction in subsecond multislice spiral CT examination of children by online tube current modulation. *Eur Radiol* 2004;14:995-999.
138. Coursey C, Frush D, Yoshizumi T, et al. Pediatric chest MDCT and tube current modulation: effect on radiation dose with breast shielding. *AJR* 2008;190:W54-W61.
139. Siegel MJ. *Pediatric Body CT*. 2nd ed. Philadelphia, Pa: Lippincott, Williams and Wilkins; 2007.

*Guidelines and standards are published annually with an effective date of October 1 in the year in which amended, revised, or approved by the ACR Council. For guidelines and standards published before 1999, the effective date was January 1 following the year in which the guideline or standard was amended, revised, or approved by the ACR Council.

Development Chronology for this Guideline

2008 (Resolution 22)

Amended 2009 (Resolution 11)