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 parameters 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 parameters and technical standards will be reviewed for revision or renewal, as appropriate, on their fifth anniversary or sooner, if indicated.

Each practice parameter 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 and approval. The practice parameters 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 parameter and technical standard by those entities not providing these services is not authorized.

Revised 2013 (Resolution 10)*

ACR–SCBT-MR–SPR PRACTICE PARAMETER FOR THE PERFORMANCE OF THORACIC COMPUTED TOMOGRAPHY (CT)

PREAMBLE

This document is an educational tool designed to assist practitioners in providing appropriate radiologic care for patients. Practice Parameters and Technical Standards 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 and our collaborating medical specialty societies caution against the use of these documents 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 practitioner in light of all the circumstances presented. Thus, an approach that differs from the guidance in this document, 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 this document 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 this document. However, a practitioner who employs an approach substantially different from the guidance in this document 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 the guidance in this document 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 this document is to assist practitioners in achieving this objective.

1 Iowa Medical Society and Iowa Society of Anesthesiologists v. Iowa Board of Nursing, ___ N.W.2d ___ (Iowa 2013) Iowa Supreme Court refuses to find that the ACR Technical Standard for Management of the Use of Radiation in Fluoroscopic Procedures (Revised 2008) sets a national standard for who may perform fluoroscopic procedures in light of the standard’s stated purpose that ACR standards are educational tools and not intended to establish a legal standard of care. See also, Stanley v. McCarver, 63 P.3d 1076 (Ariz. App. 2003) where in a concurring opinion the Court stated that “published standards or guidelines of specialty medical organizations are useful in determining the duty owed or the standard of care applicable in a given situation” even though ACR standards themselves do not establish the standard of care.

PRACTICE PARAMETER Thoracic CT / 1
I. INTRODUCTION

This practice parameter was revised collaboratively by the American College of Radiology (ACR), the Society of Computed Body Tomography and Magnetic Resonance (SCBT-MR), and the Society for Pediatric Radiology (SPR).

Computed tomography (CT) is a frequently used imaging modality for the diagnosis and evaluation of many thoracic diseases. Optimal performance of thoracic CT requires knowledge of normal anatomy, anatomic variants, pathophysiology, CT techniques, and the associated risks. This practice parameter outlines the principles for performing high-quality thoracic CT in adults and children.

The goal of thoracic CT is to demonstrate normal and pathologic anatomy and physiology within the chest.

II. INDICATIONS AND CONTRAINDICATIONS

A. Thoracic CT may be a complementary examination to other imaging studies such as chest radiography (see the ACR–SPR Practice Parameter for the Performance of Chest Radiography) or a stand-alone procedure. Indications for the use of thoracic CT include, but are not limited to:

1. Evaluation of abnormalities discovered on chest images [1].
2. Evaluation of clinically suspected cardiothoracic pathology.
3. Staging and follow-up of lung cancer and other primary thoracic malignancies, and detection and evaluation of metastatic disease [2-5].
4. Evaluation of cardiothoracic manifestations of known extrathoracic diseases [6-9].
5. Evaluation of known or suspected thoracic cardiovascular abnormalities (congenital or acquired), including aortic stenosis, aortic aneurysms, and dissection [10-12].
7. Evaluation of suspected pulmonary arterial hypertension [23].
8. Evaluation of known or suspected congenital cardiothoracic anomalies [24,25].
9. Evaluation and follow-up of pulmonary parenchymal and airway disease [26-33].
10. Evaluation of blunt and penetrating trauma [34,35].
11. Evaluation of postoperative patients and surgical complications [36,37].
12. Performance of CT-guided interventional procedures [38-41].
13. Evaluation of the chest wall [42-44].
15. Treatment planning for radiation therapy [47,48].
16. Evaluation of medical complications in the intensive care unit or other settings [49,50].

B. For more detailed evaluation of the use of CT in assessing a variety of pulmonary diseases see the ACR–SPR–STR Practice Parameter for the Performance of High-Resolution Computed Tomography (HRCT) of the Lungs in Adults.

C. There are no absolute contraindications to thoracic CT. As with all procedures, the relative benefits and risks of the procedure should be evaluated prior to the performance of thoracic CT, with or without the administration of intravenous iodinated contrast or oral contrast media. Appropriate precautions should be taken to minimize patient risks, including radiation exposure. (See the ACR–SPR Practice Parameter for the Use of Intravascular Contrast Media and the ACR Manual on Contrast Media.)

For the pregnant or potentially pregnant patient, see the ACR–SPR Practice Parameter for Imaging Pregnant or Potentially Pregnant Adolescents and Women with Ionizing Radiation.
III. QUALIFICATIONS AND RESPONSIBILITIES OF PERSONNEL

See the ACR Practice Parameter for Performing and Interpreting Diagnostic Computed Tomography (CT).

IV. SPECIFICATIONS OF THE EXAMINATION

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

Documentation that satisfies medical necessity includes 1) signs and symptoms and/or 2) relevant history (including known diagnoses). 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)

B. A typical CT of the thorax should include axial images from the lung apices to the posterior costophrenic sulci usually reconstructed at ≤5 mm with both soft tissue and a high spatial frequency (lung or bone) reconstruction algorithm. Images using 1-2 mm slice thickness with a high spatial frequency reconstruction algorithm may be helpful. Sagittal and coronal reconstructions are recommended.

C. During most examinations, scans should be obtained at suspended full inspiration. Cine imaging may be appropriate for evaluating certain disease states such as tracheomalacia. Expiratory scans may be included to evaluate for abnormalities such as air-trapping. Respiratory-gated CT may be helpful in certain applications such as radiation therapy planning. Imaging should be obtained through the entire area of interest [51]. The field of view should be optimized for each patient.

D. The examination may be conducted with or without intravenous iodinated contrast media as clinically indicated.

E. Anatomically appropriate window and level settings should be used to view the lung parenchyma, mediastinal, chest wall, and skeletal structures, and any visible portions of the lower neck and upper abdomen. Review on a PACS (picture archiving and communication system) workstation facilitates evaluation of many studies, particularly those with large data sets. Multiplanar viewing is encouraged to facilitate the display of anatomy and pathology, and sliding-slab maximum-intensity projections (MIPs) are encouraged for detecting lung nodules [52,53].

F. Although many of the operations of a CT scanner are automated, a number of technical parameters remain operator-dependent. Because these parameters can significantly affect the diagnostic quality of a CT examination, the supervising physician must be familiar with the following:

1. Radiation exposure factors (mAs, kVp).
2. Collimation.
3. Display section thickness for multidetector systems.
4. Table increment or pitch.
5. Acquisition and reconstruction field of view.
6. Window settings (width and center).
7. Reconstruction algorithm, filter, or kernel.
8. Image reconstruction interval or increment.
9. Detector configuration for multidetector systems.
10. Automatic exposure control (angular and longitudinal tube current modulation) and image quality reference parameter.
12. Reformatted images (MPR, curvilinear, Max IP, MinIP, and 3D surface or volume rendered).
13. Reconstruction techniques such as filtered backprojection or iterative reconstruction.
14. ECG-gating, this may be helpful to reduce cardiac or aortic motion artifacts.

G. Optimizing CT examination technique requires the supervising physician to select an appropriate CT protocol based on careful review of the patient history (including risk factors that might increase the likelihood of adverse reactions to contrast media) and clinical indications, as well as relevant prior imaging studies when available. This optimization process includes determining if the thoracic CT examination is clinically appropriate.

H. Protocols may be prepared according to region of interest and clinical indication. Techniques should be selected that provide image quality consistent with the diagnostic needs of the examination at optimal radiation dose levels to answer the clinical questions posed. For each area of interest or indication, the protocol should indicate the following:

1. If gastrointestinal contrast material is used, the volume, type, route of administration (oral or via nasogastric or other tube), and the time intervals during which it should be delivered.
2. If intravenous contrast material is used, the type, volume, rate of administration, and time delay between administration and scan initiation. Bolus tracking should be used whenever indicated to optimize results.
3. Detector configuration.
4. Pitch (table increment and feed).
5. Slice thickness.
6. Reconstruction interval or increment.
7. Reconstruction kernel, algorithm, or filter.
8. kVp and effective mAs per section as appropriate for adult or pediatric patients. With tube current modulation, a prescribed image quality with maximum and minimum mAs as appropriate for adult or pediatric patients.
10. Superior and inferior extent of the region of interest to be imaged.
11. Field of view.
12. Protocols for sending images to PACS (e.g., section thickness and plane of reformations such as coronal, sagittal, and other oblique projections), and the Medical Image Processing System as needed.
13. 3D reconstructions where appropriate to further delineate known or suspected abnormalities.
14. For every CT examination, the information in the radiation dose report (CTDI and dose length product) should be retained in the radiological record (such as the PACS, in a radiation dose monitoring software tool, and/or the radiology report) for reference.

These protocols should be reviewed and updated annually, and dated copies should be available to appropriate physicians, radiologic technologists, medical physicists, and administrative personnel at the facility.

I. For all patients, particularly pediatric patients and small adults, efforts should be directed to:

1. Minimize radiation dose when diagnostically feasible increasing pitch, using low mA or kVp, and tightly restricting the scan range to the body region of clinical concern. While bismuth shields have been shown to reduce radiation doses, there are several disadvantages associated with their use, especially when used with automatic exposure control (tube current modulation). Other techniques exist that can provide the same level of anterior dose reduction at equivalent or superior image quality without these disadvantages. The AAPM recommends that these alternatives to bismuth shielding be carefully considered and implemented when possible (see the AAPM Position Statement on the Use of Bismuth Shielding for the Purpose of Dose Reduction in CT Scanning at http://www.aapm.org/publicgeneral/BismuthShielding.pdf).
2. Minimize motion artifact with short scan times (balanced against any changes in mA in order to maintain appropriate mAs), tightly restricting the scan range to the body region of clinical concern, and using appropriate sedation.

J. When sedation is used, it should be administered in accordance with the ACR–SIR Practice Parameter for Sedation/Analgesia.

V. DOCUMENTATION

Reporting should be in accordance with the ACR Practice Parameter for Communication of Diagnostic Imaging Findings.

VI. EQUIPMENT SPECIFICATIONS

A. Performance Guidelines

To achieve acceptable clinical CT scans of the thorax, a CT scanner should meet or exceed the following capabilities:

1. Multirow detector acquisition.
2. Scan rotation time: ≤1 sec.
3. Acquired slice thickness: ≤2 mm.
4. Limiting spatial resolution: ≥8 lp/cm for ≥32-cm display field of view (DFOV) and ≥10 lp/cm for <24 cm DFOV.

B. 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.

C. Images should be available on a PACS workstation for review by the radiologist. Remote viewing of images should also be available to authorized health care providers. Equipment should be capable of providing a digital means of conveying the dataset.

VII. EQUIPMENT QUALITY CONTROL

The quality control program for CT equipment should be designed to minimize patient, personnel, and public radiation risks and to maximize the quality of the diagnostic information. The program should be supervised by a Qualified Medical Physicist. Each imaging facility should have documented policies and procedures that include:

1. A list of tests to be performed and the frequency of performance.
2. A list identifying which individual or group will perform the tests.
3. A written description of the procedure that will be used for each test, including the technique factors to be employed, the equipment to be used for testing, the acceptability limits of each test, and sample records from each test.
4. Periodic tests for CT technologists to assure that they understand CT principles and are complying with dose reduction protocols for multidetector CT imaging.

VIII. RADIATION SAFETY IN IMAGING

Radiologists, medical physicists, registered radiologist assistants, radiologic technologists, and all supervising physicians have a responsibility for safety in the workplace by keeping radiation exposure to staff, and to society...
as a whole, “as low as reasonably achievable” (ALARA) and to assure that radiation doses to individual patients are appropriate, taking into account the possible risk from radiation exposure and the diagnostic image quality necessary to achieve the clinical objective. All personnel that work with ionizing radiation must understand the key principles of occupational and public radiation protection (justification, optimization of protection and application of dose limits) and the principles of proper management of radiation dose to patients (justification, optimization and the use of dose reference levels).

Nationally developed guidelines, such as the ACR’s Appropriateness Criteria®, should be used to help choose the most appropriate imaging procedures to prevent unwarranted radiation exposure.

Facilities should have and adhere to policies and procedures that require varying ionizing radiation examination protocols (plain radiography, fluoroscopy, interventional radiology, CT) to take into account patient body habitus (such as patient dimensions, weight, or body mass index) to optimize the relationship between minimal radiation dose and adequate image quality. Automated dose reduction technologies available on imaging equipment should be used whenever appropriate. If such technology is not available, appropriate manual techniques should be used.

Additional information regarding patient radiation safety in imaging is available at the Image Gently® for children (www.imagegently.org) and Image Wisely® for adults (www.imagewisely.org) websites. These advocacy and awareness campaigns provide free educational materials for all stakeholders involved in imaging (patients, technologists, referring providers, medical physicists, and radiologists).

Radiation exposures or other dose indices should be measured and patient radiation dose estimated for representative examinations and types of patients by a Qualified Medical Physicist in accordance with the applicable ACR technical standards. Regular auditing of patient dose indices should be performed by comparing the facility’s dose information with national benchmarks, such as the ACR Dose Index Registry, the NCRP Report No. 172, Reference Levels and Achievable Doses in Medical and Dental Imaging: Recommendations for the United States or the Conference of Radiation Control Program Director’s National Evaluation of X-ray Trends. (ACR Resolution 17 adopted in 2006 – revised in 2009, 2013, Resolution 52).

A Qualified 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 to reference levels described in the ACR–AAPM Practice Parameter for Diagnostic Reference Levels in Medical X-Ray Imaging.

IX. 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 website (http://www.acr.org/guidelines).

For specific issues regarding CT quality control, see the ACR Practice Parameter for Performing and Interpreting Computed Tomography (CT).

Equipment performance monitoring should be in accordance with the ACR–AAPM Technical Standard for Diagnostic Medical Physics Performance Monitoring of Computed Tomography (CT) Equipment.

ACKNOWLEDGEMENTS

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 website (http://www.acr.org/guidelines) by the
Committee on Thoracic Radiology of the ACR Commission on Body Imaging and the Guidelines and Technical Standards Committee of the ACR Commission on Pediatric Radiology in collaboration with the SCBT-MR and the SPR.

**Collaborative Committee** – members represent their societies in the initial and final revision of this guideline

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REFERENCES


*Practice parameters and technical 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 practice parameters and technical standards published before 1999, the effective date was January 1 following the year in which the practice parameter or technical standard was amended, revised, or approved by the ACR Council.

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