

# MRI Accreditation Program Requirements



## Overview

The MRI Accreditation Program evaluates the qualifications of personnel, the quality control program, MR Safety Policies and image quality specific to MRI. It involves the acquisition of clinical and phantom images and corresponding data for each unit. The acquisition of the phantom images involves the use of a designated MRI phantom appropriate to the unit or module listed on the application. Every unit must go through testing for the facility to be accredited.

Since the origin of the ACR MRI Accreditation Program, patterns of practice and use of units have evolved. In the 1990s, most units were used for general MRI so the mandate for brain, cervical spine, lumbar spine and knee examination from all units was reasonable. In recent years, units are frequently used for limited anatomic applications such as head or spine only. In addition, single application specialty units have been developed.

In response to this change in practice patterns, the ACR has modified the MRI Accreditation Program to offer modules based on these anatomic applications. The ACR offers the following modules:

- Head
- Spine
- Musculoskeletal (MSK)
- Body
- Magnetic Resonance Angiography (MRA)
- Cardiac

(See section on Accreditation Testing; page 8 for specific examinations per module.)

**Every unit must apply for all modules routinely performed on that unit for a facility to be accredited.**

## Mandatory Accreditation Time Requirements

Submission of all accreditation materials is subject to mandatory timelines. Detailed information about specific time requirements is located in the *Overview for the Diagnostic Modality Accreditation Program*. Please read and be familiar with these requirements.

<b>Materials Required</b>	<b>Due</b>
Renewal application	60 calendar days from date sent
Testing materials	45 calendar days from date sent
Repeat Option forms (after deficiency)	15 calendar days from date sent

## Withdrawn, Added, or Replacement Units

The MRI Accreditation Program is unit based. Consequently, facilities **must notify the ACR** if they have permanently **withdrawn** (i.e., removed) a unit from service, if they have **replaced** that unit with a new one or have **added** another unit. The type of accreditation options available for a new unit will depend on the amount of **time the facility has left on its current accreditation certificate**:

- **Over 13 months** – The facility needs to submit only unit information and additional testing materials. Once accreditation is approved, the new unit’s expiration date will be the same as the previous expiration date.
- **Less than 13 months** - The facility must renew accreditation for all units at the facility including the new one. Once approved, all of the units at the facility will have an expiration date that is three years from the old expiration date.

If the unit replaces an existing MRI unit after accreditation is granted, it will be treated as a new unit and follow the procedures above. If the unit begins performing examinations from a module that was not included on the original application, at least one examination from that module must be submitted. If less than thirteen months are left on the facility’s accreditation, it must renew the accreditation of all of its equipment at the same time.

## **Loaner unit**

Accredited facilities may use a “loaner” unit to temporarily replace an accredited unit that is out of service for repairs, etc. for up to six months without submitting clinical and phantom images for evaluation. The accredited facility must immediately notify the ACR of the installation date, manufacturer and model of the loaner. Any loaner unit that is in use for more than one month will be required to submit evidence of testing by a qualified medical physicist/MR scientist within 90 days of installation. If the loaner is in place for longer than six months, the facility must submit the unit for accreditation evaluation, including clinical and phantom image assessment and the corresponding fee.

## **Emergency Use of Units**

Facilities may use units that are not accredited in specific modules for other types of MR imaging in emergency cases without jeopardizing a facility’s accreditation status. An emergency situation would be one in which less than 10 examinations are performed outside a unit’s accreditation status in any 30 day period, or less than 50 examinations in any 12 month period. If the volume of examinations exceeds these limits, the facility must notify the ACR and submit testing for this module.

## Personnel Qualifications

As of July 1, 2007, the physician's and the medical physicist's/MR scientist's ongoing qualifications (experience and education) are required. All sites applying for the first time after July 1, 2007 will be required to meet the full requirements for CME and continuing experience at the time you renew your accreditation (as listed below for sites renewing after July 2009). Sites accredited prior to July 2007 will have the option to meet the following phase-in plan:

<b>Phase-In Plan for Continuing Education and Experience</b>	
<b>Continuing Education Requirement</b>	<b>Continuing Experience Requirement</b>
<b>Sites renewing after July 2007</b>	
Physicians and medical physicists/MR scientists must have earned at least 5 CME hours in the prior 12-month period. The 5 CME hours must be earned for each modality in which they are renewing (CT, MRI, nuclear medicine, PET and ultrasound).	Over the prior 12-month period, physicians reading: <ul style="list-style-type: none"> <li>• CT, MRI, nuclear medicine, PET and ultrasound examinations must have read an average of 9 exams per month.</li> </ul>
<b>Sites renewing in or after July 2008</b>	
Physicians and medical physicists/MR scientists must have earned at least 10 CME hours in the prior 24-month period. The 10 CME hours must be earned for each modality in which they are renewing (CT, MRI, nuclear medicine, PET and ultrasound).	Over the prior 24-month period, physicians reading: <ul style="list-style-type: none"> <li>• CT, MRI, nuclear medicine, PET and ultrasound examinations must have read an average of 9 exams per month.</li> </ul>
<b>Sites renewing in or after July 2009</b>	
Physicians and medical physicists/MR scientists must have earned at least 15 CME hours in the prior 36-month period. The 15 CME hours must be earned for each modality in which they are renewing (CT, MRI, nuclear medicine, PET and ultrasound).	Over the prior 36-month period, physicians reading: <ul style="list-style-type: none"> <li>• CT, MRI, nuclear medicine, PET and ultrasound examinations must have read an average of 9 exams per month.</li> </ul>

Note: The continuing education and continuing experience requirements are based on previous full calendar years. For example, if a site applies for accreditation in July 2009, the physicians and medical physicists/MR scientists at that site must have met the full requirement for continuing education from January 1, 2006 to December 31, 2008. Likewise, they must have met the full continuing experience requirements from January 1, 2007 to December 31, 2008. If they did not meet these requirements in the given timeframes, the ACR will accept continuing education credits or continuing experience obtained in 2009.

## Physician

The physician shall have the responsibility for all aspects of the study including, but not limited to, reviewing all indications for the examination, specifying the pulse sequences to be performed, specifying the use and dosage of contrast agents, interpreting images, generating written reports, and assuring the quality of both the images and interpretations.

All physicians who supervise and/or interpret MRI examinations must be a licensed medical practitioner who meets the following minimum criteria:

Requirements for Physicians Supervising and Interpreting MRI Examinations		
Qualifications	Radiologists	Other Physician
<b>Initial</b>	<ul style="list-style-type: none"> <li>• Board certification in radiology or diagnostic radiology by:               <ul style="list-style-type: none"> <li>○ ABR,</li> <li>○ American Osteopathic Board of Radiology,</li> <li>○ Royal College of Physicians and Surgeons of Canada, or</li> <li>○ Le College des Mediciens du Quebec, and</li> </ul> </li> <li>• Supervision and/or performance of, as well as interpretation and/or review and reporting of, 300 MRI examinations within the last 36 months.<sup>1</sup></li> </ul> <p style="text-align: center;">OR</p> <ul style="list-style-type: none"> <li>• Completion of an accredited diagnostic radiology residency program, and</li> <li>• Performance, interpretation, and reporting of 500 MRI examinations in the past 36 months.</li> </ul>	<p>(MR imaging limited to a specific anatomic area)</p> <ul style="list-style-type: none"> <li>• Completion of an accredited specialty residency, and</li> <li>• 200 hours of Category I Continuing Medical Education (CME) in MRI to include, but not limited to: MRI physics, recognition of MRI artifacts, safety, instrumentation, and clinical applications of MRI in the subspecialty area where MRI reading occurs, and</li> <li>• 500 MRI cases in that specialty area shall have been interpreted and reported in the past 36 months in a supervised situation. For neurologic MRI, at least 50 of the 500 cases shall have been MRA or the central nervous system.</li> </ul>
<b>Continuing Experience</b>	Upon renewal, physicians reading MRI examinations must have read an average of 9 exams per month over the prior 24-month period.	
<b>Continuing Education</b>	Upon renewal, physicians must have earned at least 15 CME in MRI (half of which must be category 1) hours in the prior 36-month period.	

<sup>1</sup> Board certification and completion of an accredited radiology residency in the past 24 months will be presumed to be satisfactory experience for the reporting and interpreting requirement.

**Requirements for Physicians Supervising and Interpreting Cardiac MRI Examinations**

Qualifications	Radiologists	Other Physicians
<b>Initial</b>	<ul style="list-style-type: none"> <li>• Board certification in radiology or diagnostic radiology by:                             <ul style="list-style-type: none"> <li>○ ABR,</li> <li>○ American Osteopathic Board of Radiology,</li> <li>○ Royal College of Physicians and Surgeons of Canada, or</li> <li>○ Le College des Mediciens du Quebec, and</li> </ul> </li> <li>• Supervision and/or performance of, as well as interpretation and/or review and reporting of, 75 Cardiac MRI examinations within the last 36 months.<sup>1</sup></li> <li align="center">OR</li> <li>• Completion of an Accreditation Council for Graduate Medical Education (ACGME) Radiology Residency Program, AND</li> <li>• have supervised interpretation of 75 cardiac MRI cases in the past 36 months</li> <li align="center">AND</li> <li>• Completion of at least 40 hours of Category 1 Continuing Medical Education (CME) in cardiac imaging, including cardiac MRI, anatomy, physiology, and/or pathology or documented equivalent supervised experience in a center actively performing cardiac MRI.</li> </ul>	<p align="center"><b>CARDIOLOGISTS<sup>2</sup> (Cardiac Only)</b></p> <p align="center">Certification in cardiology by the American Board of Internal Medicine with completion of Level 2 training or higher</p> <p><b>Level 2 requirements</b></p> <ul style="list-style-type: none"> <li>• Board certification or eligibility, valid medical license, and completion of a 3-month (cumulative) specialty residency or fellowship in CMR</li> </ul> <p align="center">AND</p> <ul style="list-style-type: none"> <li>• 150 CMR examinations in which 50 where the candidate is physically present, involved in the acquisition and interpretation of the case,</li> </ul> <p align="center">AND</p> <p>Completion of 30 hours of courses related to MR in general and/or CMR in particular</p> <p><b>Level 3 requirements</b></p> <ul style="list-style-type: none"> <li>• Board certification or eligibility, valid medical license, and completion of a 12-month (cumulative) specialty residency or fellowship in CMR</li> </ul> <p align="center">AND</p> <ul style="list-style-type: none"> <li>• 300 CMR examination in which 100 where the candidate is physically present, involved in the acquisition and interpretation of the case,</li> </ul> <p align="center">AND</p> <ul style="list-style-type: none"> <li>• Completion of 60 hours of courses related to MR in general and/or CMR in particular</li> </ul> <p align="center"><b>NUCLEAR MEDICINE PHYSICIANS (Cardiac Only)</b></p> <ul style="list-style-type: none"> <li>• Completion of an ACGME approved training program in nuclear medicine</li> </ul> <p align="center">AND</p> <ul style="list-style-type: none"> <li>• Specific training in MRI within an ACGME accredited training program</li> </ul> <p align="center">OR</p> <ul style="list-style-type: none"> <li>• 160 hours of category 1 CME in MRI to include, but not limited to: MRI physics, recognition of artifacts, safety, instrumentation, and 40 hours specific to cardiovascular MRI.</li> </ul> <p align="center">AND</p> <ul style="list-style-type: none"> <li>• Interpretation and reporting under the supervision of a qualified physician of at least 75 cases of MRI of the cardiovascular system during the past 36 months.</li> </ul>
<b>Continuing Experience</b>	Upon renewal, radiologists reading Cardiac MRI examinations must have read 50 exams over the prior 24-month period. The cardiac examinations interpreted will count toward the overall continuing experience for other MR modules.	Upon renewal, cardiologists reading Cardiac MRI examinations must have continuing experience in accordance with level 2 requirements or higher – 50 examinations each year.
<b>Continuing Education</b>	Upon renewal, physicians must have earned at least 15 CME in MRI (half of which must be category 1) hours in the prior 36-month period and should include CME in Cardiac MRI as is appropriate to the physician's practice needs.	Upon renewal, cardiologists must have earned at least 30 hours of coursework in the prior 36 month period in accordance with level 2 requirements.

<sup>21</sup> Board certification and completion of an accredited radiology residency in the past 24 months will be presumed to be satisfactory experience for the reporting and interpreting requirement.

<sup>2</sup> 2005 ACCF/AHA Clinical Competence Statement on Cardiac CT and MR

## Technologist and Medical Physicist/MR Scientist

Requirements for Technologists and Medical Physicists/MR Scientists		
Qualifications	Technologist	Medical Physicist/MR Scientist
<b>Initial</b>	<ul style="list-style-type: none"> <li>• ARRT, ARMRIT or CAMRT registered as an MR technologist</li> <li style="text-align: center;">OR</li> <li>• ARRT registered or unlimited state license, and</li> <li>• 6 months supervised MRI clinical scanning experience</li> <li style="text-align: center;">OR</li> <li>• Associate degree or bachelor degree in allied health field, and</li> <li>• Certification in another clinical imaging field (such as ARDMS or NMTCB), and</li> <li>• 6 months supervised MRI clinical scanning experience</li> <li style="text-align: center;">OR</li> <li>• Performing MRI prior to and continuously since October 1996, and</li> <li>• Evaluated by responsible physician to assure competence</li> </ul> <p style="text-align: center;"><b>For Cardiac Module</b></p> <ul style="list-style-type: none"> <li>• ARRT, ARMRIT or CAMRT registered as an MR technologist</li> <li style="text-align: center;">OR</li> <li>• ARRT registered and unlimited state license, and</li> <li>• 6 months supervised MRI clinical scanning experience</li> <li style="text-align: center;">AND (all technologists)</li> <li>• Cardiac MRI experience supervised by a qualified physician or a qualified technologist(required)</li> <li>• Experience in the intravenous administration of conventional MR contrast, supervised by a qualified physician or a qualified technologist. (required)</li> <li>• Maintain Basic Life Support (BLS) certification (recommended)</li> <li>• Be capable of using an automatic external defibrillator (AED).(recommended)</li> </ul> <p><b><i>Technologists practicing MRI scanning should be licensed in the jurisdiction in which he/she practices, if state licensure for MRI technologists exists.</i></b></p>	<ul style="list-style-type: none"> <li>• Board certification in diagnostic radiological physics or radiological physics (recommended)</li> <li style="text-align: center;">OR</li> <li>• A qualified MR scientist has obtained a graduate degree in a physical science involving nuclear MR (NMR) or MRI and should have three years of documented experience in a clinical MRI environment.</li> </ul>
<b>Continuing Education</b>	24 hours of Category A CME every two years (including MRI as is appropriate to the technologist's practice needs).(recommended)	Upon renewal Medical Physicists/MR Scientists must have earned at least 15 CME in MRI (half of which must be category 1) hours in the prior 36-month period.

In addition, the qualified medical physicist/MR scientist:

- Must be familiar with the principles of MRI safety for patients, personnel, and the public; the Food and Drug Administration's guidance for MR diagnostic devices; and other regulations pertaining to the performance of the equipment being monitored.
- Must be knowledgeable in the field of nuclear MR physics and familiar with MRI technology, including function, clinical uses, and performance specifications of MRI equipment, as well as calibration processes and limitations of the performance testing hardware, procedures, and algorithms.
- Must have a working understanding of clinical imaging protocols and methods of their optimization. This proficiency should be maintained by participation in continuing education programs of sufficient frequency to ensure familiarity with current concepts, equipment, and procedures.
- May be assisted in obtaining test data for performance monitoring by other individuals that must be properly trained and approved by the qualified medical physicist/MR scientist in the techniques of performing the tests, the function and limitations of the imaging equipment and test instruments, the reason for the tests, and the importance of the test results. The qualified medical physicist/MR scientist must review and approve all measurements and is expected to provide direct on-site supervision.

## Equipment

The MR equipment specifications and performance shall meet all state and federal requirements. The requirements include, but are not limited to, specifications of maximum static magnetic field strength, maximum rate of change of magnetic field strength (dB/dt), maximum radiofrequency power deposition (specific absorption rate), and maximum auditory noise levels.

## Quality Control

### Acceptance Testing

Acceptance testing is intended to measure quantifiable system parameters, which may then be compared to the manufacturer's specifications. A complete evaluation of the system performance should be performed by a qualified medical physicist/MR scientist after completion of installation and prior to patient imaging.<sup>3</sup>

### Quality Control Testing

All facilities applying for accreditation must maintain a documented quality control (QC) program and must comply with the minimum frequencies of testing outlined below. Detailed instructions for each of the QC tests listed below are contained in the *2004 ACR MRI Quality Control Manual*. Upon acceptance of a facility's initial application, the manual will be sent to the MRI supervising physician at the practice site address under separate cover.

The ongoing QC program assesses relative changes in system performance as determined by the technologist, service engineer, qualified medical physicist/MR scientist, or supervising physician. A

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<sup>3</sup> A suggested protocol for acceptance testing is contained in "Acceptance Testing of Magnetic Resonance Imaging Systems: Report of American Association of Physicists in Medicine (AAPM) Nuclear Magnetic Resonance Task Group No. 6, Medical Physics. 1992; 19:217-219. This document is meant only to serve as a reference. The substance of this document is not intended to be incorporated by reference into the ACR Practice Guideline for Performing and Interpreting Magnetic Resonance Imaging (MRI).

qualified medical physicist/MR scientist **must** have the responsibility for overseeing the equipment quality control program and for monitoring performance upon installation and routinely thereafter. All facilities applying for accreditation or renewal must demonstrate compliance with the ACR requirements for quality control (QC) by including a copy of the facility's most recent **Annual MRI System Performance Evaluation** (must be dated within 1 year of the date of ACR MRI submission for accreditation and **must be performed** by a medical physicist/MR scientist) and copies of the facility's weekly on-site QC data (forms on pages 64, 65, and 66 of the *2004 ACR MRI Quality Control Manual*) for the most recent quarter. If the facility has been conducting QC for less than one quarter, the facility will submit whatever they have on these forms. Additionally, if the **Annual MRI System Performance Evaluation** and/or QC files show performance deficits (e.g. problems with the system and/or data outside of the action limits), the facility must state what steps it has taken to correct the problems. All QC testing must be carried out in accordance with the written procedures and methods outlined in the *ACR 2004 MRI Quality Control Manual*.

## Continuous Quality Control

The following is a list of QC tests and frequencies that must be performed by technologists and medical physicists/MR scientists:

Technologist's Weekly QC Tests		Physicist/MR Scientist's Annual QC Tests
<ul style="list-style-type: none"> <li>• Center Frequency</li> <li>• Table Positioning</li> <li>• Setup and Scanning</li> <li>• Geometric Accuracy</li> <li>• High-Contrast Resolution</li> </ul>	<ul style="list-style-type: none"> <li>• Low-Contrast Resolution</li> <li>• Artifact Analysis</li> <li>• Film Quality Control</li> <li>• Visual Checklist</li> </ul>	<ul style="list-style-type: none"> <li>• Magnetic Field Homogeneity</li> <li>• Slice Position Accuracy</li> <li>• Slice Thickness Accuracy</li> <li>• Radiofrequency Coil Checks</li> <li>• Soft-Copy Displays (Monitors)</li> </ul>

## Preventive Maintenance

Preventive maintenance shall be scheduled, performed and documented by a qualified service engineer on a regular basis. Service performed to correct system deficiencies shall also be documented and service records maintained by the MR site.

## Quality Assurance

### Physician Peer-Review Requirements

Examinations should be systematically reviewed and evaluated as part of the overall quality improvement program at the facility<sup>4</sup>. Monitoring should include evaluation of the accuracy of interpretation as well as the appropriateness of the examination. Complications and adverse events or activities that may have the potential for sentinel events must be monitored, analyzed and reported as required, and periodically reviewed in order to identify opportunities to improve patient care. These data should be collected in a manner that complies with statutory and regulatory peer-review procedures in order to ensure the confidentiality of the peer-review process<sup>3</sup>.

All sites initially applying for ACR accreditation and all sites renewing their accreditation must actively participate in a physician peer review program that performs the following functions:

<sup>4</sup> 2005 ACR Guidelines and Technical Standards. ACR Position Statement on Quality Control and Improvement, Safety, Infection Control, and Patient Education Concerns. Page IV.

- Includes a double reading (2 MDs interpreting the same study) assessment.
- Allows for random selection of studies to be reviewed on a regularly scheduled basis.
- Exams and procedures representative of the actual clinical practice of each physician.
- Reviewer assessment of the agreement of the original report with subsequent review (or with surgical or pathological findings).
- A classification of peer review findings with regard to level of quality concerns (One example is a 4-point scoring scale).
- Policies and procedures for action to be taken on significant discrepant peer review findings for the purpose of achieving quality outcomes improvement.
- Summary statistics and comparisons generated for each physician by imaging modality.
- Summary data for each facility/practice by modality.

There are several options available to meet this requirement. Sites may develop their own peer review program, use a vendor product or RADPEER, a peer review process developed by the ACR. For information about RADPEER or eRADPEER please visit the ACR web site at: [http://www.acr.org/SecondaryMainMenuCategories/quality\\_safety/radpeer.aspx](http://www.acr.org/SecondaryMainMenuCategories/quality_safety/radpeer.aspx).

## MR Safe Practice Guidelines

Safety guidelines, practices, and policies shall be written, enforced, reviewed, and documented at least annually by the MR supervising physician. For complete information regarding MR safety, see the ACR Guidance Document for Safe MR Practices. In: Kanal E, Barkovich AJ, Bell C, et al. *ACR Guidance Document for Safe MR Practices; 2007*. Leesburg, VA: American Journal of Roentgenology; 2007. The ACR Guidance Document for Safe MR Practices may also be downloaded from the American College of Radiology website at [www.acr.org](http://www.acr.org).

## Accreditation Testing

### Clinical Images

Between four and six examinations per unit are required for accreditation. The exact number of examinations depends on the number of modules for which the unit is used.

Specific examinations are required for each module (see table below). At least one of the examinations chosen for each module must be a specialty examination. *Asterisks denote the specialty examinations.*

Examination choices for MR Accreditation by module (specialty examinations denoted by asterisk*)		
Head/Neck	Spine	MSK
<ul style="list-style-type: none"> <li>• Brain for transient ischemic attack (TIA)</li> <li>• Internal auditory canal (IAC/temporal bone) for hearing loss</li> <li>• Brain for suspected demyelinating disease*</li> <li>• Pituitary with dynamic contrast enhancement*</li> <li>• Orbits for vision loss*</li> </ul>	<ul style="list-style-type: none"> <li>• Lumbar Spine</li> <li>• Thoracic Spine</li> <li>• Cervical Spine*</li> <li>• Cervical Spine with contrast for intramedullary disease*</li> </ul>	<ul style="list-style-type: none"> <li>• Knee such as for internal derangement</li> <li>• Shoulder such as for internal derangement</li> <li>• Wrist such as for internal derangement*</li> <li>• Elbow such as for internal derangement*</li> <li>• Forefoot for Morton's neuroma*</li> </ul>
Body	MRA	Cardiac
<ul style="list-style-type: none"> <li>• Male pelvis such as for prostate cancer</li> <li>• Renal</li> <li>• Hepatobiliary to Include MRCP*</li> <li>• Female pelvis such as for uterine or adnexal disease*</li> </ul>	<ul style="list-style-type: none"> <li>• Brain</li> <li>• Carotid</li> <li>• Thoracic aorta</li> <li>• Distal peripheral runoff</li> <li>• High resolution arch and carotid*</li> <li>• Abdomen for renal artery stenosis *</li> </ul>	<ul style="list-style-type: none"> <li>• Black blood</li> <li>• Basic</li> <li>• Delayed enhanced cine 1</li> <li>• Delayed enhanced cine 2</li> <li>• Delayed enhanced cine + black blood*</li> </ul>

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You must submit localizer or scout sequences with each clinical examination with cross-reference locations for each sequence. Sites cannot submit examinations performed on models or volunteers. All clinical images **must** be from actual patients. **Use of volunteers or models may result in withholding, denial or revocation of accreditation.** The images submitted for each individual exam must be from the same patient (i.e., all brain images must be from the same brain study). The only exception to this is in the event that the facility is only submitting one examination for the cardiac module, in which case it is acceptable to have the black blood sequence from a different patient than the delayed enhanced cine examination.

Facilities are strongly encouraged to submit images on CD. Images submitted electronically **must be** in DICOM format with an embedded viewer that meets the minimum requirements (see *MR Accreditation Clinical Image Quality Guide*, available on the ACR website. Click on “Accreditation”). **However, all submissions for the Cardiac module must submit on CD, in DICOM format.** Please refer to *ACR MRI Clinical Image Quality Guide* (Available on the ACR website. Click on “Accreditation”) for list of requirements for an embedded viewer.

See the table below for the number of examinations required based on the number of modules your site selected on the application:

Number of modules on application	Number of examinations per module	Total exams
One module	Four examinations are required, at least one specialty examination	Four
Two modules	Select one specialty examination and one basic examination from each module	Four
Three modules	Select one specialty examination from <i>each</i> module, <i>and</i> select one basic examination from <i>any</i> of the three modules	Four
Four modules	One specialty examination from each module	Four
Five modules	One specialty examination from each module	Five
Six modules	One specialty examination from each module	Six

Each set of clinical images will be evaluated for:

- A. Pulse sequences and image contrast
- B. Filming technique (If submitting hard copy images)
- C. Anatomic coverage and imaging planes
- D. Spatial resolution
- E. Artifacts
- F. Exam ID (All patient information annotated on clinical exams will be kept confidential by the ACR.)

Please refer to the *MRI Accreditation Clinical Image Quality Guide* for specific recommendations and requirements for each examination.

## Phantom Testing and Image Quality

Clinical image review and phantom review are intended to complement each other for a comprehensive evaluation of the quality of MRI services. The criteria for evaluation are independent of field strength and can be applied uniformly so that all magnets are measured against a single standard. Depending on the type of scanner, one of two phantoms will be required for the facility to scan and submit images on. The ACR will provide this information in the testing materials for each unit once the application is processed.

Each facility is required to submit phantom images for each unit using the ACR protocols for T1 and T2 weighting *and* phantom images using its own routine T1 and T2 weighted scan protocol.

The images and testing data will be used to assess:

- Limiting high-contrast spatial resolution
- Slice thickness accuracy
- Distance measurement and accuracy
- Signal uniformity
- Image ghosting ratio
- Low-contrast detectability
- Slice positioning accuracy
- Image artifacts

Phantom data must be submitted in the form of uncompressed DICOM images, on a CD-ROM without a viewer. There may be additional costs associated with phantom data translation. If the facility is unable to convert their phantom data to DICOM CD-ROM, the facility must pay for a phantom on-site data analysis, and an ACR medical physicist will come to the site to review the phantom data.

## Accreditation Fees

Checks should be made payable to the American College of Radiology (include modality accreditation ID#, if available). American Express, MasterCard, and Visa are accepted. The charge for the phantom is paid directly to the manufacturer.

Accreditation Fees	
Cycle	Fees
<b>Accreditation</b> (Initial cycle and renewal)	\$2400 for the first unit up to four modules, \$2600 for five modules, \$2800 for six modules  \$2300 each additional unit at one site location applying for four modules, \$2500 for five modules, \$2700 for six modules
<b>Repeat</b>	\$800 per unit for clinical or phantom images.  \$1600 per unit if repeating both.
<b>Reinstate/Corrective Action Plan</b>	\$2400 for the first unit up to four modules, \$2600 for five modules, \$2800 for six modules  \$2300 each additional unit at one site location applying for four modules, \$2500 for five modules, \$2700 for six modules
<b>Add units (mid cycle)</b>	\$1600 per unit
<b>Add module (mid cycle)</b>	\$1600 per unit
<b>Replacement Certificate</b>	\$65 per certificate
<b>Large Phantom</b>	\$1050 (includes shipping and handling)
<b>Small Phantom</b>	\$780 (includes shipping and handling)

*Note: Fees subject to change without notice.*

## For Additional Information

For further information log on to the ACR Web site at [www.acr.org](http://www.acr.org), click on “Accreditation” and click on “MRI”. A link to “Frequently Asked Questions” is available in the MRI menu, along with other useful information about accreditation and many of the program’s forms. To contact the ACR MRI Accreditation Program office by phone, dial (800) 770-0145.

## ACR Practice Guidelines and Technical Standards

The following ACR Practice Guidelines and Technical Standards are applicable to accredited facilities performing MRI examinations.

[ACR Practice Guidelines for Performing and Interpreting Magnetic Resonance Imaging \(MRI\)](#)

[ACR Practice Guidelines for the Performance and Interpretation of Magnetic Resonance Imaging \(MRI\) of the Brain](#)

[ACR Practice Guidelines for the Performance and Interpretation of Pediatric and Adult Cerebrovascular Magnetic Resonance Angiography \(MRA\)](#)

[ACR Practice Guidelines for the Performance and Interpretation of Magnetic Resonance Imaging \(MRI\) of the Head and Neck](#)

[ACR Practice Guidelines for the Performance and Interpretation of Magnetic Resonance Imaging \(MRI\) of the Adult Spine](#)

[ACR Practice Guidelines for the Performance and Interpretation of Magnetic Resonance Imaging \(MRI\) of the Wrist](#)

[ACR Practice Guidelines for the Performance and Interpretation of Magnetic Resonance Imaging \(MRI\) of the Shoulder](#)

[ACR Practice Guidelines for the Performance and Interpretation of Magnetic Resonance Imaging \(MRI\) of the Elbow](#)

[ACR Practice Guidelines for the Performance and Interpretation of Magnetic Resonance Imaging \(MRI\) of the Hip and Pelvis for Musculoskeletal Disorders](#)

[ACR Practice Guidelines for the Performance and Interpretation of Magnetic Resonance Imaging \(MRI\) of the Knee](#)

[ACR Practice Guidelines for the Performance and Interpretation of Pediatric and Adult Body Magnetic Resonance Angiography \(MRA\)](#)

[ACR Practice Guidelines for the Performance and Interpretation of Cardiac Magnetic Resonance Imaging \(MRI\)](#)

[ACR Practice Guidelines for the Performance and Interpretation of Magnetic Resonance Imaging \(MRI\) of the Liver](#)

[ACR Practice Guidelines for the Performance and Interpretation of Magnetic Resonance Imaging \(MRI\) of the Abdomen \(excluding the liver\)](#)

[ACR Technical Standard for Diagnostic Medical Physics Performance Monitoring of Magnetic Resonance Imaging \(MRI\) Equipment](#)