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Revised 2013 (Resolution 8)*

A CR–ASNR PRACTICE PARAMETER FOR THE PERFORMANCE OF NON-BREAST MAGNETIC RESONANCE IMAGING (MRI) GUIDED PROCEDURES

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, 63 P.3d 1076 (Iowa 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.
I. INTRODUCTION

This practice parameter was revised collaboratively by the American College of Radiology (ACR) and the American Society of Neuroradiology (ASNR).

Recent hardware and software improvements in magnetic resonance imaging (MRI) have resulted in an expanding and evolving role for image guidance during interventional or surgical procedures in multiple organ systems. Pulse sequence improvements that have allowed the development of rapid imaging methods as well as advances in MRI compatible equipment.

The major benefits of using MRI for procedure guidance and monitoring are [1,2]:

1. The ability to continuously visualize vascular structures during the entire procedure. The high vascular conspicuity is due to flow-related enhancement effects inherent in the gradient echo sequences used for procedure guidance.
2. The multiplanar imaging capabilities that ensure precise targeted placement of the interventional device, e.g., biopsy needle along the axial as well as the craniocaudal dimensions in the anatomy of interest. In addition, imaging in any arbitrary plane allows the device trajectory to be tailored according to the individual case.
3. The ability to guide device navigation with continuous, near real-time imaging so the device can be redirected in a timely manner in order to avoid critical structures.
4. The ability to shift between T1-weighted and T2-weighted contrasts during the procedure to maximize the anatomic/pathologic conspicuity. For example, T2-weighted techniques allow sampling of the non-necrotic regions of complex masses, thus increasing the diagnostic tissue yield.
5. The ability to perform advanced physiologic imaging intraprocedurally, thus monitoring the effects of intervention. For example, diffusion weighted imaging can monitor tissue infarction, perfusion weighted imaging can assess tissue blood flow, and thermometry can assess tissue temperature.

II. DEFINITION

The term “procedural MRI” describes the use of MRI techniques for guidance and/or monitoring or control of minimally invasive diagnosis or therapy’ with the entire procedure performed in the procedural MRI suite. MRI-guided procedures encompass multiple approaches, including endoscopic, endovascular, percutaneous, focused ultrasound, and open techniques.

This practice parameter specifically excludes breast biopsy and localization procedures that can be safely and appropriately performed according to the ACR Practice Parameter for the Performance of Magnetic Resonance Imaging-Guided Breast Interventional Procedures.

III. INDICATIONS

A. Indications

Current indications for the use of MRI to guide/monitor procedures in near-real time can be classified under the following major categories:

1. Biopsy and aspiration
   The general indications for image-guided percutaneous tissue sampling can be reviewed in the ACR–SIR–SPR Practice Parameter for the Performance of Image-Guided Percutaneous Needle Biopsy (PNB) and the ACR–SIR–SPR Practice Parameter for Specifications and Performance of Image-Guided Percutaneous Drainage/Aspiration of Abscesses and Fluid Collections (PDAFC).

   Since MRI guidance is not intended to substitute for other less expensive modes of biopsy/aspiration guidance, MRI-guided procedures will assume a role when the patient would otherwise be subjected to
surgical exploration or open biopsy performed solely for the purpose of tissue diagnosis. Therefore, MRI-guided biopsy/aspiration will be most suited for patients having 1 or more of the following conditions [2-7]:

a. Lesions in areas of complex anatomy, e.g., suprahyoid neck, base of the neck adjacent to the brachial plexus or lung apex, and liver dome lesions.
b. Lesions close to vascular structures.
c. Transiently enhancing lesions.
d. Skeletal muscle or soft tissue lesions lacking sufficient tissue contrast on computed tomography (CT) and ultrasound.
e. Bone marrow infiltrative processes.
f. Tumors with heterogenous functional activity, necessitating the need for targeted MRI biopsies to the functionally active portions.
g. Lesions seen only by MRI and not on other imaging modalities under which biopsy could be performed.

MRI guidance may also be used for joint aspirations and/or injections for additional procedural safety compared to conventional fluoroscopically guided joint interventions [8].

2. Minimally invasive percutaneous procedures

Whenever feasible, minimally invasive approaches are preferred over other surgical procedures because of the following advantages:
- Decreased morbidity and mortality.
- Decreased length of hospitalization and expense.
- The potential for treatment or cure for patients who are not open surgical candidates.

These procedures may include, but are not limited to, the following:

a. Thermal tumor ablation
   The core contribution of MRI to interstitial thermotherapy is its ability to monitor the zone of thermal tissue destruction during the procedure, thereby providing real-time guidance for the deposition of thermal energy (MR thermometry). Through MRI monitoring, ablation zone size and configuration can be directly controlled by the operator and adjusted during the procedure in order to compensate for deviations from the preoperative predictions and to define the treatment endpoint without moving the patient from the operative or MRI suite. This is an attribute of MR imaging that cannot be reliably duplicated by any other currently used imaging modality [9,10]. It not only permits accurate tumor destruction, including the margins, but also extends the application of RF ablation to the safe destruction of tumor within the visceral organs and adjacent to vital neurovascular structures.

Interactive MRI can be used to guide and monitor tumor ablation with various sources of thermal energy such as:
- Radiofrequency [3,11-16].
- Laser [17-22].
- Cryotherapy [23-25].
- Focused ultrasound [26-32].
- Microwave [33,34].

Primary and metastatic liver tumors and primary renal malignancies are the tumors most commonly treated by MRI-guided percutaneous thermotherapy. Less common targets include, but are not limited to, breast tumors, uterine fibroids, and osteoid osteomas of bone.
b. Direct intralesional drug injection
Mixing the drug of interest with diluted gadolinium chelate enables interactive MRI guidance of the injection process and monitoring of drug distribution within the target tissue. Indications include, but are not limited to [35-39]:
- Chemical tumor ablation, e.g., via direct ethanol injection.
- Injection sclerotherapy of low-flow vascular malformations.
- Percutaneous injection of local anesthesia and steroids into a muscle or joint.

3. Open surgical MRI

The integration of MRI systems into the traditional surgical arena is currently implemented in some of the following procedures [40-50]:

a. Image guidance or monitoring
- Neurobiopsy or cyst aspiration.
- Deep brain stimulation/electrode placement.
- Thermal ablation of brain tumors.
- Trans-sphenoidal pituitary resection.

b. Procedure monitoring
- Craniotomy for brain and spine tumor resection, epileptogenic focus resection, or hematoma evacuation.

4. Catheter-based procedures

Using passive or active tracking techniques, catheter and/or guidewire navigation can be performed in the cardiovascular system in near-real-time to achieve better soft tissue visualization and to obtain more pertinent physiological information compared to conventional X-ray fluoroscopic techniques. Preliminary clinical data demonstrate the safety and feasibility of MRI guidance for diagnostic cardiac catheterization in patients with congenital heart disease and for interventional cardiac catheterization and radiofrequency ablation [51-53].

Vessel wall imaging using intravascular coils is an emerging application of MRI-guided procedures that has strong potential for clinical application in the near future due to its ability to map out vulnerable plaques that may not be identified with the “lumen-only” information obtained by conventional angiography [54,55].

Transcatheter intra-arterial perfusion (TRIP) MRI can be used as a first-pass intra-procedural perfusion technique to a) verify anticipated distribution of injected contrast media, b) monitor changes in tumor perfusion during transcatheter embolic therapies, and c) exclude potential unexpected collateral supply when used in conjunction with intravenous (IV) dynamic contrast enhancement (DCE) [56-58].

IV. QUALIFICATIONS AND RESPONSIBILITIES OF PERSONNEL

A. Physicians

MRI-guided procedures must be performed by, or under the supervision of, a physician who has the qualifications specified in sections 1 or 2 below. In addition to these qualifications, a physician must also have the training specified in section 3.

1. Certification in Radiology, Diagnostic Radiology or Interventional Radiology/Diagnostic Radiology (IR/DR) by the American Board of Radiology, the American Osteopathic Board of Radiology, the Royal College of Physicians and Surgeons of Canada or the Collège des Médecins du Québec. Also, the physician must have demonstrated competency as primary operator in MRI-guided procedures under the supervision of an on-site qualified physician, during which a minimum of 5 MRI-guided procedures were performed with acceptable success and completion rates documented by a case log.
b. Completion of a diagnostic radiology residency or fellowship program approved by the Accreditation Council for Graduate Medical Education (ACGME), the Royal College of Physicians and Surgeons of Canada (RCPSC), the Collège des Médecins du Québec, or the American Osteopathic Association (AOA) that included 6 months of training in cross-sectional imaging, including CT and MRI, and 3 months of training in image-guided interventional radiological techniques that included percutaneous biopsy and drainage procedures and vascular catheterization. This must include performance (under the supervision of a qualified physician) of at least 5 MRI-guided procedures with acceptable success and complication rates documented by a case log.

or

2. In the absence of the above requirements or other postgraduate certification and training that included comparable instruction and experience, physicians may meet the requirements for performing MRI-guided procedures by adhering to ALL of the following recommendations:
   a. Documentation of “hands-on” training in the performance of MRI-guided procedures.
   b. Performance and completion of at least 5 successful and uncomplicated MRI-guided procedures as primary operator under the supervision of an on-site qualified physician with acceptable success and complication rates.
   c. Substantiation in writing by the chair of the department of the institution in which the physician will be providing these services.

3. In addition, physicians involved in MRI-guided procedures are expected to demonstrate competence in the following fields of knowledge and skills:
   a. General medical background in order to be able to communicate appropriately with referring physicians from various specialties and to be able to discuss the appropriateness of the specific procedures in light of the patient’s integrated individual history, physical findings, comorbid conditions, and preoperative imaging findings.
   b. Basic anatomy (including congenital and developmental variants), physiology, and pathophysiology of the specific organ or tissue targeted for intervention.
   c. Fundamental MRI physics with particular emphasis on the technical parameters that influence device visualization and navigation. The ability to operate the scanner is strongly recommended for all physicians involved in MRI-guided procedures. A thorough understanding of MRI safety related to ferromagnetic attraction of instruments and equipment (e.g., oxygen tanks), instrument-related artifacts, and the generation of heat or current by magnetic induction of either permanently or temporarily implanted devices is mandatory.
   d. Initial management of clinical emergencies that may arise during surgical MRI, including, but not limited to, the administration of basic life support and the recognition and treatment of adverse reactions to administered sedatives and analgesics.

Maintenance of Competence

Physicians must perform a sufficient number of MRI-guided procedures to maintain their skills, with acceptable success and complication rates as laid out in this practice parameter. Continued competence should depend on participation in a quality improvement program that monitors these rates, as well as participation in postgraduate continuing medical education (CME) courses on diagnostic and technical advances in MRI-guided procedures.

Continuing Medical Education

The physician’s continuing education should be in accordance with the ACR Practice Parameter for Continuing Medical Education (CME).
B. Nonphysician Practitioners

Physician assistants and nurse practitioners can be valuable members of the interventional radiology team, but should not perform MRI-Guided Procedures independent of supervision by physicians with training, experience and privileges to perform the relevant procedures. See the ACR–SIR–SNIS–SPR Practice Parameter for Interventional Clinical Practice and Management.

C. Qualified Medical Physicist

See the ACR Practice Parameter for Performing and Interpreting Magnetic Resonance Imaging (MRI).

D. Radiologic Technologist

See the ACR Practice Parameter for Performing and Interpreting Magnetic Resonance Imaging (MRI).

E. Nursing Services

Nursing services are an integral part of the team for preprocedural and postprocedural patient management and education and may assist the physician in monitoring the patient during MRI-guided procedures.

V. SAFETY GUIDELINES AND POSSIBLE CONTRAINDICATIONS

For contraindications to MRI scanning, see the ACR Practice Parameter for Performing and Interpreting Magnetic Resonance Imaging (MRI), the ACR Manual on Contrast Media, and the ACR Guidance Document on MR Safe Practices [59,60].

For relative contraindications to percutaneous needle procedures, see the ACR–SIR–SPR Practice Parameter for the Performance of Image-Guided Percutaneous Needle Biopsy (PNB) and the ACR–SIR–SPR Practice Parameter for Specifications and Performance of Image-Guided Percutaneous Drainage/Aspiration of Abscesses and Fluid Collections (PDAFC).

Relative contraindications to percutaneous thermal tumor ablation include, but are not limited to:

1. Large size of the targeted tumor. There is no universal cut-off diameter for eligibility for percutaneous thermal ablation. However, the larger the tumor the greater the likelihood of incomplete necrosis and development of postablation syndrome. Tumors smaller than 3 to 4 cm in diameter are generally suitable for percutaneous ablation.
2. Proximity to vital structures, such as the gallbladder, renal pelvis, or bowel wall, if protective measures are not possible.
3. Proximity to large blood vessels. The presence of flowing blood in the vicinity of the tumor being ablated compromises the efficacy of treatment due to the heat sink effect.
4. Amenability to surgical resection with respect to both technical feasibility and patient fitness.

VI. SPECIFICATIONS OF THE EXAMINATION

A. Request for the Examination or Interventional Consultation

The written or electronic request for MRI-guided procedures 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)

In practice settings in which the interventional radiologist or interventional neuroradiologist performs procedures on the basis of physician referrals or direct patient referrals, an appropriate radiology clinic or consultation note should document preprocedure patient evaluation, procedure indication, and procedure medical necessity.

B. Facility Requirements

The basic requirements for setting up a procedural MRI suite are [61-67]:

1. Hardware requirements

   a. An appropriate MRI system should be available to facilitate the patient access necessary for performing the procedures. Many different MRI system designs have been used to interactively guide procedures. Each of them has advantages and disadvantages, with a constant tradeoff between signal-to-noise, patient access, useable field of view, and expense.

   b. For scans performed with low-field MRI systems, low-noise receiver chains should be available to provide relatively high signal-to-noise ratios and sufficient image quality to guide interactive procedures. Image acquisition times should be tailored to the indication to provide appropriate temporal resolution.

   c. Facilities should be the ability to operate the scanner and view images in the operative suite within the magnetic field through an in-room, high-resolution, RF-shielded monitor. Combined with the patient access allowed by an appropriate MR imaging system, this capability allows the entire procedure to be performed with the operator maintaining constant device or instrument control while simultaneously viewing the images.

   d. Facilities should have the availability of MRI-compatible needles, probes, catheters, guidewires, etc. that are undeflected by the magnetic field and that create little or no field distortions or image degradation except as desired for passive device localization. The potential for specific devices, particularly catheters and guidewires, to be heated by imaging gradients should be understood prior to clinical use [68].

   e. A surgical or MRI table capable of safely transferring a patient from the operating position into the imager in case of intraoperative MR imaging should be present.

   f. The 5-gauss line should be clearly defined.

   g. MRI-compatible patient monitoring equipment should be available and used as appropriate to the procedure [59].

2. Software requirements

   The application of fast gradient-echo pulse sequences that allow a wide range of tissue contrast in a time frame sufficient for device tracking even at the low field strengths of open magnets and with the suboptimal coil position sometimes required to access the puncture site [69-74].
C. Performance Guidelines

1. Minimally invasive procedures
   a. Biopsy and aspiration
      The MRI-compatible biopsy needle is advanced into the targeted lesion under MRI guidance using short TR/short TE gradient-echo sequences. The process uses a continuous imaging mode that allows automated sequential acquisition, reconstruction, and in-room display of multiple sets of images to guide needle insertion with respect to the 3-dimensional geometry of the lesion. This may be performed through acquisition of a set of contiguous, parallel thin (e.g., 5 mm) slices centered on the needle position, or through the acquisition of image sets in multiple scan planes oriented along the shaft of the needle. Depending on the operator’s preference, the procedure can be performed using free-hand methods, needle holders, or real-time guidance from optically linked systems [75-80].
   
b. Thermal ablation
      The needle, thermal electrode, or laser fiber is introduced into the targeted area of pathology using the basic technique of MRI guidance described above for biopsy and aspiration.

      For thermal tumor ablation, real-time monitoring of the evolving ablation zone can be achieved using a rapid gradient-echo sequence. In cases of MRI-guided radiofrequency tumor ablation, imaging interference caused by the RF source can be overcome by software and hardware modifications that allow RF energy to be deposited during imaging [81]. Alternatively, TSE T2-weighted and/or TSE STIR images can be simply acquired intermittently between the RF deposition cycles. Regardless of the source of thermal energy used, the size and shape of the developing ablation zone are directly observed as an enlarging low-signal area surrounded by high-signal tissue reaction. Electrode repositioning into persistent foci of high-signal tumor as detected on the T2 and/or STIR images is performed in the scanner under continuous MRI guidance in an interactive manner similar to that used for initial electrode placement. The “guide-ablate-monitor” sequence of events is repeated until the induced ablation zone is noted to encompass the entire tumor and a small cuff of normal adjacent tissue or the developing ablation zone approaches adjacent vital structures [82].

c. Lesion injection and chemical ablation
   If MRI is used to guide intrallesional drug injection rather than thermal ablation, the injected drug (alcohol, sclerosing agent, or anesthetic/steroid) may be mixed with an MRI contrast agent to facilitate interactive monitoring of drug distribution during injection [83].

2. Open surgical MRI
   The performance of open surgical procedures that require interactive image guidance of devices - such as biopsy needles, aspiration devices, thermal electrodes, laser fibers, or curettes with continuous or intermittent monitoring of the result - should follow the previously described guidelines for “minimally invasive procedures,” as appropriate.

   The status of tumor resection, hematoma evacuation, or other invasive procedures may be intermittently imaged so that a satisfactory result can be documented. This requires only relatively minor modification to standard imaging systems because patient access is not necessary during the monitoring process. These types of procedures may not require an open operating room MRI scanner but may be performed on conventional cylindrical superconducting systems provided that patient transfer between the surgical position and the imager is secured as discussed above in “hardware requirements.” Integration with a frameless stereotactic localization system may be added in these procedures to facilitate guidance to areas of residual pathology [84-92].

3. Catheter-based procedures
   After vascular access is performed, a coronal 2-dimensional steady-state free-precession sequence is used to localize the abdominal aorta and to interactively guide the advancement of catheters and guidewires in real time from the introducer sheath to the heart or the vascular anatomy of interest. The imaging plane
can subsequently be modified to best fit the route of catheter manipulation within the vascular system [93].

Visualization of catheters and guidewires is usually achieved via “passive tracking” techniques by incorporating a metal ring within the dilator tip [94] and by using catheters and guidewires modified with a ferrite mixture to induce susceptibility artifacts [95]. Another simple technique that may be used to render a catheter MRI-visible using the same theory of passive tracking is to fill the balloon of a standard angiographic catheter with a small amount of carbon dioxide gas [51].

Other methods to track catheters within the vascular system are achieved by creating “controlled locally induced field inhomogeneities.” One way is to generate current in a thin copper wire wrapped around the catheter shaft. Another is to use “active tracking” techniques achieved by mounting a single or multiple micro receive coils connected to the receive channel of the MRI system to the tip of the catheter [96-106]. Augmented steering of catheters can currently be achieved by mechanical tip-deflecting catheters [106-109].

D. Patient Care

1. Preprocedural care

a. The clinical history and relevant physical findings, including the vital signs, must be reviewed and recorded in a patient’s medical record by the physician performing the procedure. Specific inquiry should be made with respect to relevant medications, prior allergic reactions, and bleeding/clotting status.

b. The indication(s) for the procedure, including (if applicable) documentation of prior therapy, must be recorded. Any relevant prior studies should be reviewed.

c. Explanation of the procedure along with its benefits, risks, and alternatives should be discussed with the patient by an appropriate member of the procedural team in accordance with the ACR–SIR Practice Parameter on Informed Consent for Image-Guided Procedures.

d. The patient should be screened for contraindications for MRI and for MRI contrast media.

2. Procedural care

a. Adherence to the Joint Commission’s current Universal Protocol for Preventing Wrong Site, Wrong Procedure, Wrong Person Surgery™ is required for procedures in non-operating room settings including bedside procedures. The organization should have processes and systems in place for reconciling differences in staff responses during the “time out.”

Anesthesia equipment, surgical scalpels, and electrocautery systems should be made of MRI-compatible materials. Physiologic monitors must be either nonferromagnetic or kept outside the fringe field of the magnet.

b. Patients undergoing MRI-guided procedures must have IV access in place for the administration of any required fluids and medications.

c. Vital signs should be obtained at regular intervals during the procedure, and a record of these measurements should be maintained.
d. If the patient is to receive conscious sedation, pulse oximetry must be used. Administration of sedation for MRI-guided procedures should be in accordance with the ACR–SIR Practice Parameter for Sedation/Analgesia as appropriate. A registered nurse or other appropriately trained personnel should be present and have primary responsibility for monitoring the patient. A record of medication doses and times of administration should be maintained.

e. 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 Parameter for the Use of Intravascular Contrast Media.)

f. 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 size in the patient population.

3. Postprocedural care

a. A postprocedural MRI scan should be obtained prior to transferring the patient from the procedural MRI suite in order to document the procedural result and the presence and extent of any complications, and to provide a baseline for future comparison imaging.

b. A procedural or operative note should be documented in the patient’s medical record detailing the procedure, any immediate complications, and the patient’s status at the conclusion of the procedure.

c. During the initial postprocedural period, patients will usually require bed rest, and appropriately trained personnel should monitor the patient’s vital signs and observe for bleeding.

d. Postprocedural pain should be treated as deemed appropriate by the physicians performing the procedure, nonphysician practitioners or, in certain practice settings, by the patient’s referring physician or surgeon.

e. Patient discharge

   Group I - Patients undergoing uneventful MRI-guided biopsy or aspiration, sclerotherapy, drug injection, catheter-based procedures, and other minor procedures may be discharged after an observation period of 1 to 6 hours.

   Group II - Patients subjected to percutaneous thermal or chemical tumor ablation may be admitted to the hospital for overnight observation and to be examined by the treating physician prior to discharge. In selected cases, when the percutaneous approach to the tumor is straightforward and the ablation is uneventful, the patient may be discharged after careful physical examination following an initial observation period of 6 hours.

   Group III - The disposition (transfer or discharge) of patients undergoing open surgical MRI will be managed by the primary surgical team.

In all cases, the physician, surgeon, or health care surrogate must be available for continuing care both during hospitalization and after discharge. The patient’s status at discharge must be documented in the patient’s medical record. The patient should be educated about possible delayed complications.
E. Surgical and Emergency Support

When MRI-guided procedures are performed in a freestanding diagnostic center, detailed protocols for prompt transport or admission of the patient to an emergency department with a surgical facility should be formalized in writing.

VII. DOCUMENTATION

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

VIII. SUCCESS AND COMPLICATION RATES AND THRESHOLDS

Indicator thresholds may be used to assess the efficacy of ongoing quality improvement programs. For the purpose of these practice parameters, a threshold is a specific level of an indicator, e.g., a complication rate that should prompt a review. MRI techniques are used to guide and monitor an increasing array of diverse percutaneous and open surgical procedures. The success and complication rates and thresholds are therefore expected to demonstrate significant variations depending on the nature of the procedure and on the targeted organ. As the primary aim of using MRI to guide and monitor procedures is to increase patient safety by improving the visualization of vascular and other vital structures and by enhancing the intraprocedural or intraoperative detectability of residual pathology, complication thresholds should never exceed those used for equivalent procedures performed under the guidance of other imaging modalities or for open surgery performed without MRI monitoring.

Routine periodic review is strongly encouraged for all cases, with particular attention to unexpected or unsatisfactory outcomes. When complication rates exceed a threshold value, a review should be performed to determine the causes and to implement any necessary changes.

Similarly, success rates and thresholds should be viewed in light of each specific procedure, and institutional results should be compared to the published data for standard equivalent procedures and surgeries.

The only exception would involve the technical success rate. The ability to use appropriate MRI techniques to adequately visualize the target lesion and procedural device, to navigate it safely into the tissue of interest, and to unequivocally discriminate pathological tissue from adjacent structures is essential, so the technical success rate should be 100%.

At the time these practice parameters were written, there were no published reports of comprehensive success or complication rates of institutional procedural MRI programs.

IX. EQUIPMENT SPECIFICATIONS

The MRI equipment specifications and performance must meet all state and federal requirements. These 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 acoustic noise levels.

X. 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).
Specific policies and procedures related to MRI safety should be in place with documentation that is updated annually and compiled under the supervision and direction of the supervising MRI physician. Guidelines should be provided that deal with potential hazards associated with the MRI examination of the patient as well as to others in the immediate area. Screening forms must also be provided to detect those patients who may be at risk for adverse events associated with the MRI examination.

Equipment monitoring should be in accordance with the ACR Technical Standard for Diagnostic Medical Physics Performance Monitoring of MRI Equipment.

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tumors: initial clinical experiences using a 0.5 T open MR system. *J Magn Reson Imaging* 2002;16:576-
35. 583.
35. Lewin JS, Merkle EM, Duerk JL, Tarr RW. Low-flow vascular malformations in the head and neck: safety
and feasibility of MR imaging-guided percutaneous sclerotherapy—preliminary experience with 14
36. Shinmoto H, Mulkern RV, Oshio K, Silverman SG, Colucci VM, Jolesz FA. MR appearance and spectral
features of injected ethanol in the liver: implication for fast MR-guided percutaneous ethanol injection
40. Dort JC, Sutherland GR. Intraoperative magnetic resonance imaging for skull base surgery. *Laryngoscope*
2001;111:1570-1575.
42. Jolesz FA, Talos IF, Schwartz RB, et al. Intraoperative magnetic resonance imaging and magnetic
43. Lewin JS, Metzger A, Selman WR. Intraoperative magnetic resonance image guidance in neurosurgery. *J
44. Patel NK, Plaha P, O'Sullivan K, McCarter R, Heywood P, Gill SS. MRI directed bilateral stimulation of
the subthalamic nucleus in patients with Parkinson's disease. *J Neurol Neurosurg Psychiatry* 2003;74:1631-
1637.
46. Schulder M, Carmel PW. Intraoperative magnetic resonance imaging: impact on brain tumor surgery.
47. Tan TC, Mc LBP. Image-guided craniotomy for cerebral metastases: techniques and outcomes.
*Neurosurgery* 2003;53:82-89; discussion 89-90.
48. Vitaz TW, Hushek SG, Shields CB, Moriarty TM. Interventional MRI-guided frameless stereotaxy in
54. Ladd ME, Quick HH, Debatin JF. Interventional MRA and intravascular imaging. *J Magn Reson Imaging*
58. Wang D, Bangash AK, Rhee TK, et al. Liver tumors: monitoring embolization in rabbits with VX2 tumors-


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