



Reducing healthcare costs. Improving healthcare quality.

**MRI Safety Toolkit
September 2011**

Does your healthcare facility's safety culture include a comprehensive program on MRI Safety?

A partnership with Amerinet Quality Solutions can help ensure that your facility is educated on MRI Safety which could eliminate the risk of serious harm.

Value Proposition:

The Amerinet MRI Safety Toolkit provides a comprehensive, easily accessible resource containing best practice tools and information that will help facilities to maximize quality and safety outcomes while optimizing financial outcomes.

Table of Contents

- [Definition](#)
- [Extent of Problem](#)
- [Contributing Factors](#)
- [Incidence and Prevalence](#)
- [Risk Reduction Strategies](#)
- [Prevention Programs](#)
- [Educational Opportunities](#)
- [Educational Resources](#)
- [MRI Safety Informational Websites References](#)
- [Educational Tools](#)
- [Supplier Best Practice Information](#)
- [References](#)
- [Disclaimer](#)

Definition

- Magnetic resonance imaging, more commonly referred to as MRI, was introduced to healthcare in the late 1970s
- MRI provided, at the time, cutting edge technology which allowed two and three dimensional views of body tissue and structure
- Today, over 10 million MRI scans are performed in the United States each year
- While the capabilities of the MRI scanner are widely known, the inherent dangers are not

Source: The Joint Commission, SE Alert- Preventing accidents and injuries in the MRI suite, February 14, 2008

Extent of Problem

Types of injuries which can occur during an MRI Scanning Process:

- Projectile Effect
 - Magnetic pull or missile effect
 - Ferromagnetic and some non-ferromagnetic materials can be pulled into the magnet bore
 - Objects that have been reported include: oxygen cylinders, IV poles, transport stretchers, floor buffers, wheelchairs, drills, patient walkers, stethoscopes, IV pumps, scissors, hair barrettes, paper clips, and hairpins
 - Paper clips and hairpins near a 1.5T magnet can reach speeds of 40 mph
 - Projectile effect can be fatal, cause significant equipment damage, cause a quench of the magnet, and can result in large amounts of magnet downtime

Extent of Problem

Types of injuries which can occur during an MRI Scanning Process:

- Twisting or Torque Effect
 - Magnetic objects align parallel with the MRI magnetic field
 - Cochlear implants and cerebral aneurysm clips can twist in the body causing potentially fatal damage
 - Magnetic components can come loose from their base causing device failure or patient injury
- Acoustic
 - Injury caused by loud knocking noise produced by MRI scanner

Extent of Problem

Types of injuries which can occur during an MRI Scanning Process:

- Burns
 - External cable connections contacting the patient
 - Electrodes touching the patient
 - Conductive loops formed by poor patient positioning
 - Metallic tattoos or permanent eye liner containing iron oxide can heat to cause burns

- Image Artifacts
 - Presence of metals near the image site can product image artifacts; such as, signal voids or image distortion. Signal voids can mask pathology or be misinterpreted as pathology
 - RF emissions from equipment may be picked up by the MRI RF receiver resulting in image artifacts

Sources: VA National Center for Patient Safety, September 2008

The Joint Commission, SE Alert- Preventing accidents and injuries in the MRI suite, February 14, 2008

Extent of Problem

Types of injuries which can occur during an MRI Scanning Process:

- Device Malfunction
 - Devices with analog gauges, electric motors, transformers, relays, and switches can be adversely affected by the static magnetic field. For instance, battery-powered devices can suddenly fail ;some programmable infusion pumps may perform erratically; and pacemakers and implantable defibrillators may malfunction.
 - ECG waveforms can be distorted by interaction with the gradient magnetic field

Sources: VA National Center for Patient Safety, September 2008

The Joint Commission, SE Alert- Preventing accidents and injuries in the MRI suite, February 14, 2008

Contributing Factors



Images Provided By : www.simplyphysics.com

Contributing Factors

What Makes the MRI Environment Potentially Unsafe:

- MRI Magnets are powerful
 - Large invisible magnetic fields extend in three dimensions
 - MRI magnets are as much as three times as powerful as magnets used to lift cars
 - MRI magnets are always 'ON'
 - Shutting off the MRI magnet by quenching it have inherent risks of asphyxiation, frostbite, fire hazards, plus costs which range from \$20,000 – \$500,00
- Major incidents are severely under-reported
 - Near misses with no injury to staff or patients are rarely reported
 - Incidents with magnet damage are rarely reported
 - Media often discover and report near misses or major incidents

Contributing Factors

What Makes the MRI Environment Potentially Unsafe:

- Objects do not appear to be ferrous or contain ferromagnetic materials
 - Ferromagnetic materials can become projectiles or twist in the magnet bore
 - All metals are conductors which can become hot or cause image artifact

Examples of Ferromagnetic Objects		
Buffing Machines	Mops	Shrapnel
Chest Tube Stands (filings)	Nail Clippers and Files	Sandbags (metal)
Clipboards (patient charts)	Oxygen Cylinders	Steel Shoes
Gurneys	Pulse Oximeters	Stethoscopes
Hairpins	Pacemakers	Scissors
Hearing Aids	Pagers	Staples
Identification Badges	Paper Clips	Tools
Insulin Pumps	Pens and Pencils	Vacuum Cleaners
Keys	IV Poles	Watches
Medical Gas Cylinders	Prosthetic Limbs	Wheelchairs

Source: *The Joint Commission, SE Alert- Preventing accidents and injuries in the MRI suite, February 14, 2008*

Incidence and Prevalence

In 2005, an independent analysis of the FDA's MAUDE database was performed by Jason Lauenders, a medical physicist with the ECRI Institute.

Findings:

- Over a 10 year span, 389 reported MRI-related events, including nine deaths
- More than 70 percent of the 389 reports were burns
- 10 percent were projectile-related, another 10 percent were “other events”, including implant disturbances
- 4 percent were acoustic injuries, another 4 percent were fire-related
- 2 percent were internal heating-related

Risk Reduction Strategies

- With 99 percent sensitivity, the use of ferromagnetic detectors may help in screening patients for remaining objects
- In helping to prevent projectile cylinder accidents, implement protocols that permit maintenance and housekeeping personnel to enter MRI suite only after proper safety education and when no patient is in the suite
- Appoint a safety officer responsible for implementing and enforcing safety procedures

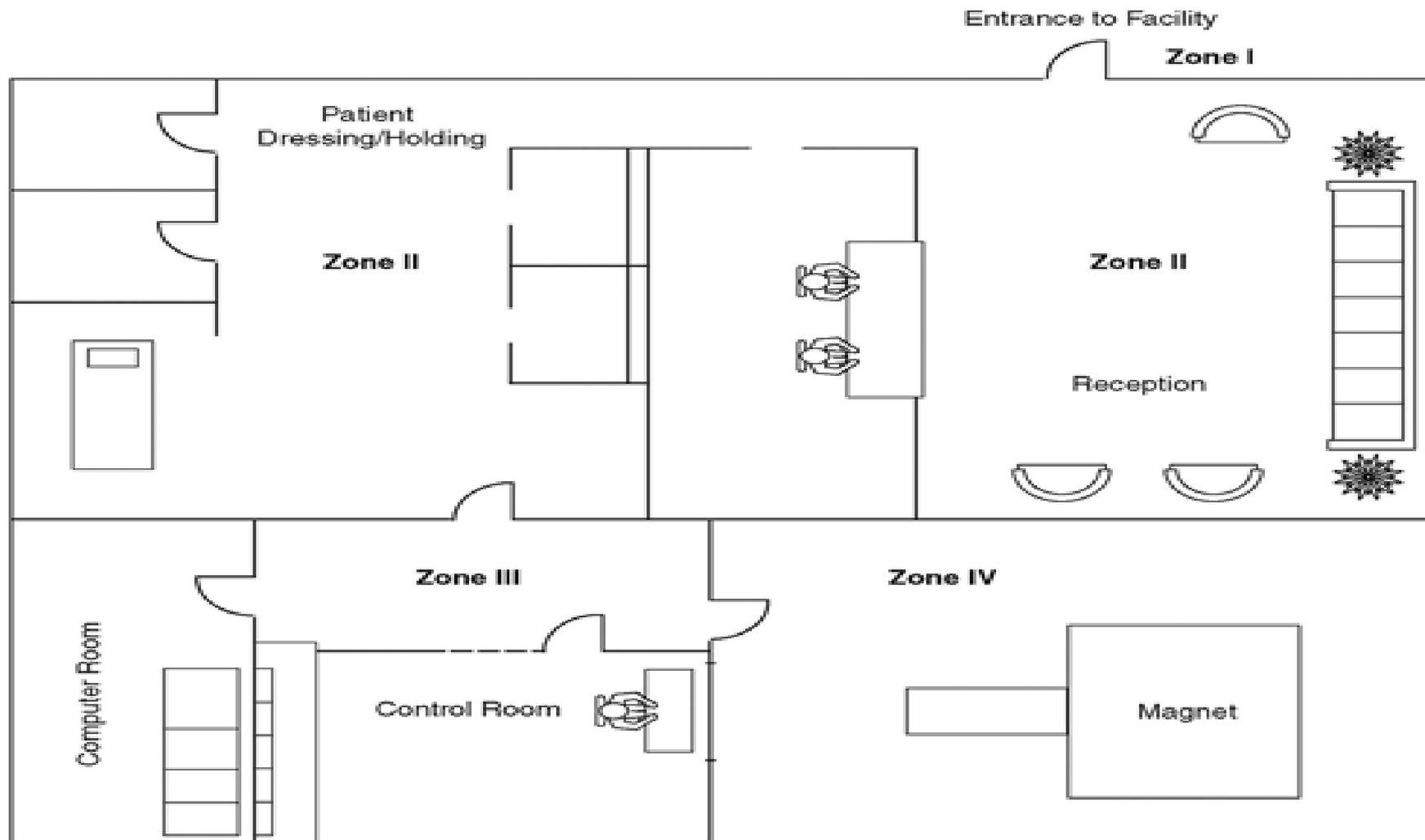
Risk Reduction Strategies

- Support safe MRI practices by establishing, implementing and maintaining written protocols and checklists
- Periodically review and assess your organizations MRI polices
- As general practice, only bring MR safe devices or equipment into the MRI environment
- Ensure that procedures are in place for reporting all adverse events, MR safety incidents, or “near incidents” that occur in a timely fashion (within 24 hours)

Risk Reduction Strategies

- Restrict access to all MRI sites by Implementing the Four Zone Concept
 - Zone I: General Public
 - Zone II: Unscreened MRI Patients
 - Zone III: Screened MRI Patients and Personnel
 - Zone IV : Screened MRI Patients under Constant Direct Supervision of Trained MR Personal

Model MR Facility Zone Configuration



Risk Reduction Strategies

The new MR safety marking standard will only be effective if it is well understood and properly implemented by device manufacturers and MR departments

ASTM (American Society for Testing and Materials) Terminology

- MRI Safe: an item that poses no known hazards in all MRI environments
- MRI Conditional: an item that has been demonstrated to pose no known hazards in a specified MRI environment with specified conditions for use; such as, field strength, spatial gradients, SAR, etc.
- MRI Unsafe: an item known to pose hazards in all MRI environments



Source: Pennsylvania Patient Safety Authority: Safety in the MRI Environment: Ferromagnetic Projectile Objects in the MRI Scanner Room, Vol. 6, No. 2, June 2009

Interventions/Prevention Strategies

11 Steps to Prevent Infections in the MRI Environment:

1. Develop an appropriate written infectious control policy which includes MRI cleaning procedures and schedule
2. Implement a mandatory hand washing / hand sanitizing procedure
3. Between each patient, clean the MRI system, tables, inside the bore of the MRI system and any other items that come into contact with the patient.
4. After each patient, clean all comfort pads and positioning sponges with an approved disinfectant.
5. Periodically inspect the pads and positioning pads with a magnifying glass, particularly at the seams, to identify fraying or tearing. If present, the pads should be replaced.
6. Regularly check all padding material with an ultraviolet (black) light and remove and biological material found.

Interventions/Prevention Strategies

11 Steps to Prevent Infections in the MRI Environment: (continued)

7. Replace damaged or contaminated pads with new pads incorporating permanent antimicrobial agents.
8. Use pillows with a waterproof covering that is designed to be surface wiped. Replace pillows when their barrier is compromised.
9. Promptly remove any body fluids and then disinfect surface and all other contaminated areas.
10. If a patient has an open wound or any history of MRISA or other infection:
 - Gloves and gowns should be worn by all staff coming in contact with the patient and removed before touching other areas not coming in contact with the patient.
 - Thoroughly clean the table and all the pads before the next patient is scanned. For patients with any known infections, add 10-15 minutes onto the scheduled scan time to assure there is enough time to thoroughly clean the room and all the pads.
11. All furniture should be cleaned periodically. Ideal surfaces are those that are waterproof and easily wiped. Infection control experts recommend this be done between each patient.

Interventions/Prevention Strategies

Redesign of the MRI environment:

- Identify and physically mark the 5 gauss line to stop MRI Unsafe items from entering the magnet room
- Identify and physically mark the 30 gauss line to prevent MRI Conditional items from being brought too close to the magnet
- Hang posters in multiple languages clearly explaining the dangers associated with the MRI environment
- Design MRI suites for safety and consider installing ferromagnetic detectors as a supplement to screening patients

Interventions/Prevention Strategies

Redesign of the MRI environment: (continued)

- Installation of piped in oxygen systems to reduce the risk that oxygen cylinders will be brought into the magnet room
- Control access to the MRI environment by instituting the AC Guidance Document for Safe MRI Practices – 2007 Zoning
- Provide a designated quarantine area so patients arriving in MRI Unsafe wheelchairs or walkers will have those items locked away during the MRI scan
- Finished surfaces should be designed to facilitate cleaning for infection control. All MRI environments should have a sink/hand-washing station

Interventions/Prevention Strategies

Training:

- Initial and refresher training on MRI safety for all MRI personnel and those who may need to enter the MRI environment. Included are MRI technologists, nurses, and physicians, plus staff from emergency, transport, maintenance, housekeeping, and security departments
- Outside agency training should include police and firefighters
- Conceptual and practical training for radiology residents and fellows
- Drills for all emergency conditions which may occur in the MRI environment should be conducted

Interventions/Prevention Strategies

Procedural:

- Assign MRI safety duties to appropriate staff to assure that MRI policies and procedures are in effect and personnel are trained
- Always assume the magnet is ON
- MRI Technologists should have control over access to the MRI environment
- All personnel entering the MRI environment should be screened for magnetic or conductive materials in, on, or attached to the body
- Check the patient's medical record for implants or other devices contraindicated for MRI scans

Interventions/Prevention Strategies

Procedural: (continued)

- Patients should don hospital-provided gowns acceptable to the MRI environment and lockers for their belongings should be provided
- Use a hand held magnet to screen objects on or accompanying patients and other items that need to be brought into the MRI environment
- Maintain a database of MRI Safe and MRI Conditional equipment and label the devices for their classification
- If the magnet is upgraded or a new magnet is purchased, the existing equipment should be tested for appropriate labeling

Interventions/Prevention Strategies

Procedural: (continued)

- Do not make any assumptions about implants, devices, or equipment being suitable for the MRI environment
 - Read the technical information about the product
 - Call the manufacturer of the product to obtain information about the suitability of the device in the MRI environment and obtain their written policy
 - Call the manufacturer of the MRI system
- Assume that blankets and clothing are hiding something missed during the screening process
- Place sensors and cables as far away as possible from RF coils and the magnet bore
- Remove all unneeded conductors from the magnet bore

Interventions/Prevention Strategies

Procedural: (continued)

- Do not let the patient contact the magnet bore and place insulation between the patient and bore if necessary
- Periodically check sensors for heating on sedated or unconscious patients
- Do not position the patient so their body tissue makes a loop
- All codes must be conducted outside of the MRI environment
- Report all incidents to the MRI Safety Officer and FDA
- Cryogen vents should be inspected annually

Interventions/Prevention Strategies

Purchasing:

- All equipment purchased for use in or near the MRI environment should be suitable for the specific environment
- Use manufacturer approved fiber optic, carbon fiber, or graphite leads rather than conductive leads
- Use manufacturer approved large surface area, low impedance ECG Electrodes
- Purchase sand bags containing sand, pillows without magnetic springs, and non-ferrous chairs for family in the magnet room
- Provide the patient with some type of alarm device suited for the MRI environment to alert the MRI staff

Interventions/Prevention Strategies

Anesthesia Considerations:

- Frequently have a large number of ferrous items on their person which are safe in the OR but not MRI
 - Stethoscopes
 - Cell phones, pagers, wallets (credit cards are erased by the magnet), shoes
 - Pens, clipboards to document the case
- Frequently use equipment which is MRI Conditional
 - Anesthesia machines
 - Laryngoscopes
 - Patient Monitors
 - Often use ferrous products purchased for joint use by MRI and CT, where they are safely used
 - Equipment must be kept outside the MRI with lines run through waveguides or equipment is tethered to the wall with chains and feet kept inside a specific area outlined on the floor

Interventions/Prevention Strategies

Anesthesia Considerations: (continued)

- Practice Advisory on Anesthetic Care for Magnetic Resonance Imaging, a report by the American Society of Anesthesiologists Task Force on Anesthetic Care for Magnetic Resonance Imaging, published in *Anesthesiology* 2009; 110:459-79.
 - Assist the anesthesiologist in decision making in patient care in the MRI
 - Highlights the dangers involved in anesthesia in MRI due to the use of ferrous equipment or anesthesia person carrying ferrous items into the magnet room
- The Child in MRI and CT, *Considerations and Techniques*, International Anesthesiology Clinics, Volume 47, Number 3, 15-23 by Cheryl K. Gooden, MD
 - Increasing use of MRI and CT on pediatric patients due to technological advances in diagnostic radiology
 - Anesthesiologist providing care to the pediatric patient in the MRI and CT settings often results in better and more accurate scans and patient outcome

Educational Opportunities

- MRI Safety Week was founded to celebrate and promote excellence in MRI safety. This is the anniversary of the tragic 2001 MRI death of young Michael Colombini, age 6, who died when a portable steel oxygen cylinder was brought into the MRI room during his exam.
- The 2011 observance marks the 10 year anniversary



[Colombini MRI Case: Root Cause Analysis - 10 Years Later](#)

Courtesy of : Reliability Center, Inc. Hopewell, VA. www.reliability.com

Educational Opportunities

- American College of Radiology
[Self-Assessment Module Online: Patient Safety](#)
- ECRI Institute e-Learn
[Screening Patients for MRI Procedures to Ensure Safety](#)
- Institute for Magnetic Resonance Safety, Education and Research
[MRI Safety Training Videos](#)
- Reliability Center, Inc
[MRI Coil Failures](#)

Educational Resources

- American College of Radiology
[ACR Guidance Document for Safe MR Practices: 2007](#)
- The PA Patient Safety Advisory
[Safety in the MR Environment: Ferromagnetic Projectile Objects in the MRI Scanner Room](#)

Informational Websites

- [American College of Radiology](#)
- [ECRI Institute's Patient Safety Blog](#)
- [FDA MAUDE Database](#)
- [FDA Medical Device Safety](#)
- [Institute for Magnetic Resonance Safety, Education, and Research](#)
- [National Quality Forum](#)

Educational Tools

- PA Patient Safety Authority [MRI Patient Screening Form](#)

Magnetic Resonance Imaging Patient Screening Form

Magnetic resonance (MR) facilities and units can refer to this sample MR screening form as a guide in developing a comprehensive MR screening form. All questions on the screening form should be answered completely to avoid confusion or misunderstanding as to the metal medical history and metal exposure history of the patient. The completed screening form should be reviewed with the patient (or patient's representative) by two separate MR personnel to verify completeness and accuracy.

Patient Identification No.: _____

Patient Name: _____ Age: _____ Date of Birth: _____ Today's Date: _____

Home Address: _____

Gender: _____ Height: _____ Weight: _____

Area of Body to Be Examined: _____

Reason for MRI: _____

Referring Physician: _____ Telephone No.: _____

1. Have you ever had a prior surgical procedure of any kind? Yes No

If yes, please indicate the date (approximate if unknown) and type of surgery:

Date _____	Type of Surgery _____
Date _____	Type of Surgery _____
Date _____	Type of Surgery _____

2. Have you ever had a prior diagnostic imaging study or examination (e.g., MRI, CT, X-ray)? Yes No

If yes, please list:

Body Part	Date	Facility
MRI _____	_____	_____
CT/CAT Scan _____	_____	_____
X-ray _____	_____	_____
Ultrasound _____	_____	_____
Nuclear Medicine _____	_____	_____
Other _____	_____	_____

3. Have you ever experienced any problem related to a previous MR procedure? Yes No

If yes, please explain: _____

4. Have you ever been a welder, grinder, or sheet metal worker? Yes No

If yes, please explain: _____

5. Have you had an eye injury involving a metallic object or fragment (e.g., metallic slivers, shavings)? Yes No

If yes, please explain: _____

Continued . . .

6. Have you ever been injured by a metallic object or foreign body (e.g., BB, bullet, shrapnel)? Yes No
If yes, please explain: _____
7. Have you recently had a small bowel endoscopy study with ingestion of a small camera capsule? Yes No
If yes, how recent: _____
8. Are you currently taking or have you recently taken any medication or drug? Yes No
If yes, please list medications or drugs: _____
9. Are you allergic to any medication or drug? Yes No
If yes, please list medications or drugs: _____
10. Do you have anemia or any disease(s) that affects your blood, a history of renal (kidney) disease, renal (kidney) failure, renal (kidney) transplant, high blood pressure (hypertension), liver (hepatic) disease, or seizures? Yes No
If yes, please explain: _____
11. Do you have a history of asthma, allergic reaction, respiratory disease, or reaction to contrast medium or dye used for MRI, CT, or X-ray procedures? Yes No
If yes, please explain: _____
12. Have you ever had a reaction to or have been told that you should not have contrast medium injections for imaging studies? Yes No
If yes, please explain: _____

For Female Patients:

13. Date of last menstrual period: _____ Are you postmenopausal? Yes No
14. Are you or could you be pregnant or experiencing a late menstrual period? Yes No
15. Are you taking oral contraceptives or receiving hormonal treatment? Yes No
16. Are you taking any type of fertility medication or having fertility treatments? Yes No
If yes, please explain: _____
17. Are you currently breastfeeding? Yes No

Continued . . .

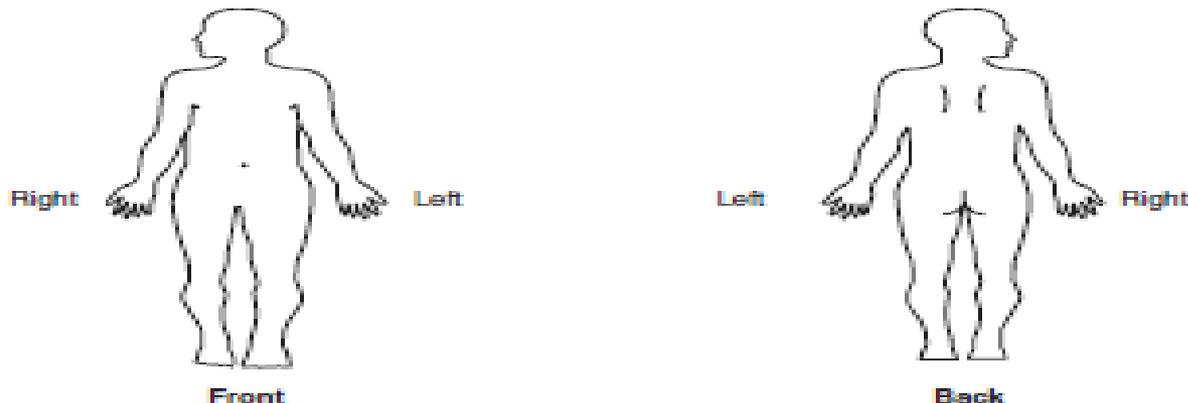
CAUTION

Magnetic resonance imaging (MRI) systems use strong magnetic fields and radio-frequency energy for imaging soft tissue in the body. Certain implants, devices, or objects may pose a hazard to individuals in close proximity to the magnet of the MRI system and/or may interfere with the MRI procedure.

Please indicate if you currently have or ever had any of the following:

- | | |
|---|---|
| <p>Aneurysm clip(s) <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Cardiac pacemaker <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Implanted cardioverter defibrillator (ICD) <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Electronic Implant or device <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Magnetically activated implant or device <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Neurostimulator <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Spinal cord stimulator <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Bone growth/bone fusion stimulator <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Internal electrodes or wires <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Cochlear, otologic, or other ear implant (including hearing aid) <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Insulin or other infusion pump <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Implanted drug infusion device <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Any type of prosthesis (e.g., eye, penile) <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Heart valve prosthesis <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Blood clot filter <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Eyelid spring or wire <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Artificial or prosthetic limb <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Metallic stent, filter, or coil <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Shunt (spinal or intraventricular) <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Vascular access port and/or catheter (e.g., Broviac, Port-A-Cath, Hickman) <input type="checkbox"/> Yes <input type="checkbox"/> No</p> | <p>Radiation seeds or implants <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Swan-Ganz or thermodilution catheter <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Medical patch (transdermal) (e.g., Nicotine, Nitroglycerine) <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Any metallic fragment or foreign body <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Wire mesh implant <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Tissue expander (e.g., breast) <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Surgical staples, clips, or metallic sutures <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Joint replacement (hip, knee, etc.) <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Bone/joint pin, screw, nail, wire, plate, etc. <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Intrauterine device (IUD), diaphragm, or pessary <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Braces, dentures, or partial plates <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Tattoo or permanent makeup <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Body piercing jewelry <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Wig or hair implants <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Hair accessories (e.g., hairpins) <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Other Implant <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Breathing problem or motion disorder <input type="checkbox"/> Yes <input type="checkbox"/> No</p> <p>Claustrophobia <input type="checkbox"/> Yes <input type="checkbox"/> No</p> |
|---|---|

Please mark on the figure(s) below the location of any implants, metal, tattoos, or permanent makeup inside or on your body.





Patient Instructions

Please use the hospital-supplied hearing protection (e.g., earplugs, headphones) during the MRI scan because the MRI scanner produces significant acoustic noise that may affect your hearing or that you may find uncomfortable.

Please remove all metallic objects before entering the MRI scan room, including the following:

Jewelry (e.g., earrings, rings, body piercings), hairpins, hair clips, dentures, false teeth, partial dental plates, hearing aids, eyeglasses, watch, pager, cell phone, keys, safety pins, paper clips, money clip, any magnetic strip cards (e.g., bank, credit), coins, pens, pocketknife, nail clipper, tools, and clothing with metal fasteners or containing metal thread.

It may be necessary for you to remain still for up to one hour while lying on your back during the MRI procedure. If you do not believe you can remain still for that long, please discuss this with the MRI technologist or radiologist before entering the MRI scan room.

Discuss any questions or concerns that you may have or if you are unsure if an item should be removed with the MRI technologist or radiologist prior to entering the MRI scan room.

I have reviewed the above information and attest that the information is accurate to the best of my knowledge. I have read and understand the entire contents of this form and had the opportunity to ask questions regarding this information and the MRI procedure.

Patient Name (please print): _____

Patient Signature: _____

Date: _____

Patient Guardian or Authorized Representative Name (please print): _____

Patient Guardian or Authorized Representative Signature: _____

Date: _____

Relationship to Patient: _____

Form Information Reviewed by: _____

Date: _____

MRI Technologist Registered Nurse Radiologist Other: _____

For more information, go to <http://www.patientsafetyauthority.org>.

This form accompanies the following:

Safety in the MR environment: MR safety screening practices.

Pt Patient Saf Alerts

2009 Mar;6(1):20-6.

Adapted with permission from:

Frank G. Shellock, PhD, Adjunct Clinical Professor of Radiology and Medicine, Keck School of Medicine; Adjunct Professor of Clinical Physical Therapy, Division of Biokinesiology and Physical Therapy, School of Dentistry, University of Southern California; Director for MRI Studies of Biomimetic MicroElectronic Systems, National Science Foundation, Engineering Research Center, University of Southern California; Institute for Magnetic Resonance Safety, Education, and Research; President, Shellock R & D Services, Inc.

References

Appendix 3: safety screening form, MR hazard checklist, and patient instructions. In: Kanal E, Barkovich A, Bell C, et al. ACR guidance document for safe MR practices: 2007. *Am J Roentgenol* 2007 Jun;188(6):1447-74.

MRI patient checklist [online]. [cited 2009 Feb 23]. Available from Internet: <http://www.walacradiology.com/LinkClick.aspx?link=MRIChecklist.pdf&tabid=1&mid=930>.

MRI safety checklist and patient consent form [online]. 2005 Jan [cited 2009 Feb 23]. Available from Internet: <http://www.nedodl.com/documents/mriinfoconsent.pdf>.

MRI safety screening [form online]. [cited 2009 Feb 23.] Available from Internet: http://www.urmcch.edu/~fmri/safety_screening.pdf.

Shellock FG. Magnetic resonance (MR) procedure screening form for patients [online]. [cited 2009 Feb 23]. Available from Internet: http://www.mrisafety.com/screening_form.asp.

Educational Tools

MRI Safety Poster



MRI Safety: "Sandbags" May Not Be What You Think...

Pennsylvania Patient Safety Reporting System (PA-PSRS)

Janet Johnston, RN, MSN, JD; William Marella, MBA; Edward Finley, BS



Background
More than 400 Pennsylvania hospitals, ambulatory surgery centers, birthing centers, and selected abortion facilities report to PA-PSRS patient-related adverse events and near misses. PA-PSRS collects such reports, analyzes the data, and provides feedback to reporting facilities about lessons learned, including evidence-based safety strategies. Since its inception in June 2004, more than 300,000 reports have been submitted to PA-PSRS.

Problem
The following PA-PSRS report was submitted:
A post cardiac catheterization patient with a sandbag placed on the left groin went to MRI. When the technician moved the patient into the MRI, the magnet pulled the sandbag from the patient's groin to the outer housing of the MRI unit.

Application to Patient Safety
Healthcare workers may be unaware of the risk of ferromagnetic "sandbags" — manufacturer labels may not indicate that a sandbag contains metal pellets, or whether it is MRI compatible. The sandbag may be concealed with a sheet or blanket. Order forms/catalogs, invoices, or vendor packing slips may fail to indicate that the product contains metal rather than sand. Disseminating such information reduces the potential of patient injury in the MRI setting.

Other MRI Incompatible Items Reported to PA-PSRS

- Ferromagnetic sandbags
- Dentures
- Rings/wrist/watches/hairstrips
- Knives
- Cell phones
- Metal implants/sutures/clips
- Permanent makeup/tattoos
- Crutches/walkers
- Laryngoscopes
- Capsule endoscopes
- Medication patches
- Neuraxial realizers
- Ovidewires/probes/stents
- Implants
- Tracheostomy tubes: metal/metal fibers

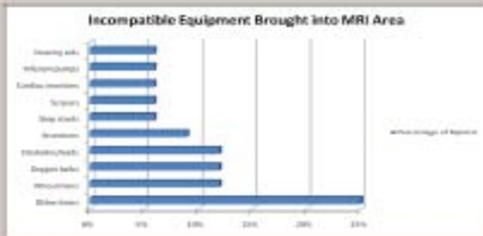
Patient Safety Strategies

- If feasible, do not allow MRI-incompatible sandbags in your facility.
- Assume that items are MRI incompatible until proven otherwise.
- Use a powerful hand magnet (>1000G) — NOT the MRI magnet — to evaluate sandbags and confirm they do not contain metal.
- Have only sandbags in the MRI environment labeled as MRI compatible.
- Do not allow sandbags from other departments into the MRI area unless confirmed that they are non-ferromagnetic.
- Add ferromagnetic sandbags to lists of MRI-incompatible equipment posted in the MRI area.
- Revise MRI screening checklists to include evaluating patients for ferromagnetic sandbags.
- If the facility must use ferromagnetic sandbags, clearly label them as containing iron and that they are not for use in the MRI area.
- Ensure that potential magnetic objects are not covered/concealed/stored on transport equipment or the patient. Look under patient gowns, blankets, sheets, and towels.
- Transfer patients to MRI-compatible equipment before they enter the MRI area.
- Prior to MRI, check patients' medical records to determine whether a recent procedure/complication may have required the use of a sandbag (such as cardiac catheterization).
- Assign trained healthcare personnel the responsibility for physically evaluating the patient and securing the MRI area.
- Heighten awareness of healthcare providers of this potential patient safety problem.

Outcomes

- An article was published in the *PA-PSRS Patient Safety Advisory* (quarterly publication) regarding this hazard that included several patient safety strategies to reduce the potential for serious patient injury from sandbags in the MRI environment.
- In a survey of 186 Pennsylvania Patient Safety Officers conducted in December 2006, 37% indicated that this article helped drive change in their facilities.

Incompatible Equipment Brought into MRI Area



Item	Percentage of Reports
Hearing aids	~10%
Implants/links	~10%
Cardiac monitors	~10%
Sponges	~10%
Step stools	~10%
Staircases	~10%
Wheelchairs/beds	~10%
Sponges/beds	~10%
Wheelchairs	~10%
Other items	~10%

© 2007 Pennsylvania Patient Safety Authority

For more information visit:
www.psa.state.pa.us

This poster was adapted from
"Sandbags" May Not Be What You Think."
PA-PSRS Patient Safety Advisory,
September 2006, Vol. 3, No. 3.

MRI Safety Supplier Best Practice Information

- Kopp Development



Kopp Development, Inc. Company Overview

- Founded in January, 2004
- Product lines patented in U.S. and Europe
- Company website: www.koppdevelopment.com
- Worldwide network of manufacturer's representatives, distributors, and OEM partners



Kopp Development, Inc. Company Overview

- Kopp Development FerrAlert™ Halo Prescreen and FerrAlert™ Halo Entry are the only Ferromagnetic Detectors Made in America
- Product Lines
 - FerrAlert™ Halo Prescreen Ferromagnetic Detector
 - FerrAlert™ Halo Entry Ferromagnetic Detector
 - GaussAlert™ Magnetic Field Strength Alarm System
 - FerrAlert™ SOLO Patient and Staff Screener (New Product – not on contract at this time)
- FerrAlert Halo Devices are installed in over 500 facilities on 4 continents



Development Inc.



Product Descriptions

- **FerrAlert™ Halo Prescreen** is intended for use with outpatients, family members, and hospital staff to detect small ferrous objects on the person down to the size of a typical hairpin. The recommended placement for this device is near the patient lockers or changing area. One method of installation is parallel to a wall, about six (6) inches away to allow a patient or family member to walk into the FerrAlert Prescreen, turn around, and walk out, giving you the security that the front, back, and sides of the person have been screened. Additional information is available at: www.koppdevelopment.com/prescreen.html
- **FerrAlert™ Halo Entry** is installed unobtrusively onto the frame of the magnet room door, outside or inside the magnet room, depending on the swing of the door. FerrAlert Entry has the capability to detect ferromagnetic mass of moderate to large size, similar to a bandage scissors or larger, and can significantly help to protect your investment in the magnet and to avoid a catastrophic accident, usually caused by large ferrous projectiles, which can severely injure those in the magnet room; the technologist, patient, family member, or other hospital staff; and, could damage the magnet itself. Additional information is available at: www.koppdevelopment.com/entry.html

Product Descriptions

- Both **FerrAlert** devices have 24 sensors, 12 on each side, which change from green to yellow when a ferrous object has been detected, alerting you to the presence and location of the object; in addition, both **FerrAlert** Products have an adjustable width ultrasonic sensor built into the top crossbar. In the most open position, a person who has a ferrous object will be detected and the alarm will sound and LED's change to yellow at the location of the ferrous object approximately 15 inches before the person enters the **FerrAlert** device.
- **FerrAlert Entry** has received Compatibility Certification by Siemens Medical Solutions on their 1.5T and 3T magnets with no noise, interference, or image artifact with the power ON or OFF. Testing was done using a FerrAlert Entry mounted inside the magnet room. The Test Certificate may be downloaded at:
www.koppdevelopment.com/pdfs/Testcertificate_Kopp_FerrAlertHaloEntry.pdf



Development Inc.



Product Descriptions

- **GaussAlert Magnetic Field Strength Alarm System** is available in three standard product alarm threshold point configurations:

GaussAlert Model 501-10 (10 gauss or 0.1 mT) is intended to be mounted on MRI Unsafe equipment which should never be brought into the magnet room as it may become a projectile or pose other risks of injury to a patient or MRI personnel, or cause damage the magnet. Products such as emergency crash carts, defibrillators, oxygen cylinders, medical gas cylinders, IV poles, and micro infusion pumps fit into this category.

GaussAlert Model 501-30 (30 gauss or 0.3 mT) is intended to be mounted on equipment identified as MRI Conditional or the older MRI Compatible rating to prevent a user from bringing the equipment inside the 30 gauss field where it may be attracted to the magnet and pulled toward the bore or even become a projectile, potentially injuring staff, a patient, or damaging the magnet. If there is any question about the equipment, use the equipment manufacturer's documents to determine if it is appropriate to use any model **GaussAlert** with the device.

GaussAlert Model 501-100 (100 gauss or 1.0 mT) is intended to prevent monitored equipment from inadvertently being moved into the exclusion zone by alarming when the equipment is inside the 100 gauss line. If the monitored equipment's manufacturer recommended field strength limit is exceeded, image degradation as well as projectile effect, or torque effect may occur, resulting in the injury of a patient or staff, damage to the magnet, or the monitored equipment.

The **GaussAlert APPLICATION NOTES** is available at: <http://www.koppdevelopment.com>



Development Inc.



Kopp Website Resources

- **Economic Justification for FerrAlert**

To analyze the potential financial implications of the acquisition of the FerrAlert ferromagnetic detection system in terms of capital outlay compared with the expected accrual of potential operational savings through the device's use.

The Economic Justification worksheet or a spreadsheet to insert your site's information is available at: www.koppdevelopment.com

- **Who recommends Ferromagnetic Detectors?**

Summary listing of agencies recommending Ferromagnetic Detectors. Document is available at: www.koppdevelopment.com

- **Customer references, Veterans Administration Inspector General's Review, Editorials, and several MRI Safety papers** are available at: www.koppdevelopment.com

- **Technical Drawings and Wiring Diagrams for Architects, Designers and Builders**

To better assist you in the architectural and design phase of your MRI facility, we will be happy to guide you to download our products' Technical Drawings and Wiring Diagrams. Sitting Ferromagnetic Detectors at MRI facilities will help you to comply with Joint Commission (JCAHO) and Facility Guidelines Institute (FGI) standards. A link to request this information is available at:

www.koppdevelopment.com



Development Inc.



Kopp Website Resources

- **2010 Edition of the Guidelines for Design and Construction of Health Care Facilities in U.S. published by the Facility Guidelines Institute (FGI)**
Suites for MRI equipment shall be planned to conform to the four-zone screening and access control protocols identified in the American College of Radiology's "Guidance Document for Safe MRI Practices." A number of states have adopted the FGI 2010 Edition as their code for construction of Health Care Facilities.

The Section for MRI suites extracted from the Guidelines is:

2.2-3.4.4.2 DESIGN CONFIGURATION OF THE MRI SUITE:

- 1. Suites for MRI equipment shall be planned to conform to the four-zone screening and access control protocols identified in the American College of Radiology's "Guidance Document for Safe MRI Practices."*
- 2. The layout shall include provisions for the following functions:*
 - a. Patient interviews and clinical screening*
 - b. Physical screening and changing areas (as indicated)*
 - c. Siting of ferromagnetic detection systems*
 - d. Access control*
 - e. Accommodation of site-specific clinical and operational requirements*
- 3. An anteroom visible from the control room shall be located outside the MRI scanner room so that patients, health care personnel, and other employees must pass through it before entering the scanning area and control room. This room shall be outside the restricted areas of the MRI's magnetic field.*
- 4. Any area in which the magnetic field strength is equal to or greater than 5 gauss (0.5 millitesla) shall be physically restricted by the use of key locks or pass-key locking systems.*

The FGI website is available at: fgiguidelines.org



Development Inc.



Additional Information on Kopp Website

- Willis HRH STRATEGIC OUTCOMES PRACTICE Technical Advisory Bulletin is available at:
www.koppdevelopment.com/articles
- The New MRI Design Guide from Department of Veterans Affairs Recommends Use of Ferromagnetic Detectors. Document is available at:
[www.koppdevelopment.com/pdfs/VA MRI Design Guide-08](http://www.koppdevelopment.com/pdfs/VA_MRI_Design_Guide-08)
- A number of MRI Safety related papers are available at:
www.koppdevelopment.com



Kopp Development Inc. Contact Information

- Contract # VQ10189
- **Kopp Development Inc.**
785 N.E. Dixie Hwy
Jensen Beach, FL. 34957
Toll Free Number: (888) 838-5677
Telephone: (772) 225-6932
Fax: (772) 225 – 6291
Contact: James S. Schmidt, Sales Manager
E-mail: jschmidt@koppdevelopment.com
- **Kopp Development Inc. Europe**
Travessera de Gracia 56, 3^o 1^a
Barcelona 08006
Telephone: (+34) 670 – 740 – 451
Fax: (+34) 932 – 405 – 040
Contact: Gio Kopp Juluhaze
E-mail: gja@koppdevelopment.com

References

- American College of Radiology www.acr.org
- ECRI Institute www.ecri.org
- Institute of Magnetic Resonance Safety, Education, and Research www.MRIsafety.com
- PA Patient Safety Authority www.patientsafetyauthority.org
- Reliability Center, Inc. www.reliability.com
- The Joint Commission www.jointcommission.org

Disclaimer

TERMS OF USE

Copyright © Amerinet 2011

All Rights Reserved

The contents of all material available on this internet site are copyrighted by Amerinet unless otherwise indicated.

Copyright is not claimed as to any part of an original work prepared by a U.S. or state government officer or employee as part of that person's official duties. All rights are reserved by Amerinet, and content may not be reproduced, downloaded, disseminated, published, or transferred in any form or by any means, except with the prior written permission of Amerinet.

Affiliates and members of Amerinet may download pages or other content for their own use, consistent with the mission and purpose of Amerinet and the contractual relationship between the two parties. However, no part of such content may be otherwise or subsequently reproduced, downloaded, disseminated, published, or transferred, in any form or by any means, except with the prior written permission of, and with express attribution to Amerinet.

Copyright infringement is a violation of federal law subject to criminal and civil penalties. Amerinet and such other designated names appearing on this Internet site are registered service marks of Amerinet.

The following trademarks and service marks of Amerinet appear on this web site. This list does not represent all such marks that Amerinet uses and/or holds:

Amerinet Inc.

Amerinet Choice

Amerinet Clinical Advantage

Amerinet QualityTouch

Amerinet Options

DealerBuy

Amerinet Diagnostix

TargetBuy

Disclaimer Although Amerinet includes links providing direct access to other internet sites, Amerinet takes no responsibility for the information contained on these external sites, and does not exert any editorial or other control over them.