

Pre-MRI X-ray – Medical Directive

<sup>e</sup> Diagnostic Imaging Quality Council Approved: 18MAR2022

Harmonized

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## Authorizing Prescriber(s)

Radiologists working at Lakeridge Health (LH)

#### Authorized to Whom

Medical Radiation Technologists-Magnetic Resonance (MRT(MR)) working at any of the LH sites who have the knowledge, skill and judgement to order a relevant x-ray prior to a patient's Magnetic Resonance Imaging (MRI) Scan. As the initiator of the MRI scan request, the referring physician will be indicated as the ordering provider.

#### **Patient Description/Population**

Any patient scheduled for an MRI Scan

#### **Order and/or Procedure**

X-ray of the orbit to exclude the possibility of a metallic foreign body in the eye

X-ray of any other body part to exclude the possibility of a known or suspected electronic or metallic foreign body

#### Indications to the Implementation of the Directive

• A request for an MRI scan has been received

AND the patient has one or more of the following:

- A known injury to the eye with metal
- Previous eye surgery e.g. retinal repair, where there is a question of foreign body
- Has a known or suspected metallic or electronic foreign body that requires confirmation prior to MRI to ensure patient safety

#### **Contraindications to the Implementation of the Directive**

The directive must not be implemented in any of the following circumstances:

- Patient does not consent to the x-ray or MRI Scan
- Prior imaging of the required body part has already been obtained
- Patient or Substitute Decision Maker (SDM) withdraws or refuses diagnostic investigation

#### Consent

Verbal consent will be obtained by the MRT(MR)

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Lakeridge Health

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#### **Documentation Requirements**

In addition to standard documentation practices, the MRT(MR) implementing this medical directive must ensure the following is included in the electronic health record:

- The name of this medical directive
- The procedure implemented
- The name of the implementer
- The date and time
- Signature of implementer including credentials/electronic signature

#### **Review/Evaluation Process**

The medical directive will be reviewed every two years as directed by the Diagnostic Imaging Program



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\*\*\*This table must **not** be used independently apart from the Medical Directive\*\*\*

### **Order Table Form**

| Order                     | Indication                                                                                                                             | Contraindication                                                                               | Notes<br>(Optional)                                                                            |
|---------------------------|----------------------------------------------------------------------------------------------------------------------------------------|------------------------------------------------------------------------------------------------|------------------------------------------------------------------------------------------------|
| X-ray for<br>foreign body | A request for an MRI scan has been received                                                                                            | Patient does not consent to the X-ray or MRI Scan                                              | Special consideration-<br>pregnancy                                                            |
|                           | AND the patient has one or more of the following:                                                                                      | Prior imaging of the required body part has already been obtained                              | For additional information,<br>refer to MRI Safety - Implants<br>and Foreign Bodies Policy and |
|                           | A known injury to the eye with<br>metal previous eye surgery e.g.<br>retinal repair, where there is a<br>question of foreign body      | Patient or Substitute Decision Maker<br>(SDM) withdraws or refuses<br>diagnostic investigation | Procedures                                                                                     |
|                           | Has a known or suspected metallic<br>or electronic foreign body that<br>requires confirmation prior to MRI<br>to ensure patient safety | Any additional, unrelated contraindication to the MRI procedure                                |                                                                                                |