# UNIVERSITY OF TOLEDO

SUBJECT: Medical Event Involving Radiation Generating Equipment Procedure No: HM-08-048

## PROCEDURE STATEMENT

General administration requirements for a medical radiation-generating equipment Medical Event as defined in OAC 3701:1-66-02 (K-L)

### PURPOSE OF PROCEDURE

To define a medical event involving radiation generating equipment and reporting requirements.

#### **DEFINITION**

"Medical event" means one or more of the following criteria have occurred to a human patient:

- (a) Unintended skin dose to the same area in a single procedure greater than 2 sievert (200 rem);
- (b) Unintended dose other than skin dose in a single procedure greater than:
  - (i) 0.5 sievert (50 rem) to any organ; or
  - (ii) 0.05 sievert (5 rem) effective dose equivalent;
- (c) Wrong patient or wrong site for entire procedure when the resultant dose is:
  - (i) Greater than 0.5 sievert (50 rem) to any organ; or
  - (ii) Effective dose equivalent greater than or equal to 0.05 sievert (5 rem).

### **PROCEDURE**

Upon discovery of a medical event, the handler shall:

- 1. Contact the department regarding the medical event within one business day:
- 2. Provide a written report, including the analysis of the medical event, by a radiation expert to the department within fifteen business days of the medical event. The written report must include:
  - (a) The handler or registrant's name;
  - (b) The name of the prescribing physician;
  - (c) A brief description of the event including the body site, dose delivered and any critical structures involved;
  - (d) Why the event occurred;
  - (e) The effect, if any, on the individual who received the medical event;
  - (f) Actions, if any, that have been taken, or are planned, to prevent recurrence; and
  - (g) Certification that the handler notified the individual, or the individual's responsible relative or guardian, and if not, why not.
- 3. Provide a clinical summary to the prescribing physician and patient within fifteen business days; an
- 4. Maintain record of the medical event as part of the patient's permanent medical record.
- 5. The written report in (as defined in #2) shall not contain the individual's name or any other information that could lead to the identification of the individual.

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Joseph Agosti CNMT	
Radiation Safety Officer	
Individual Responsible for Radiation Protection	

Approved By: