# DEPARTMENT OF HEALTH AND HUMAN SERVICES Food and Drug Administration Center for Devices and Radiological Health

**ACCIDENTAL RADIATION OCCURRENCE REPORT** 

Form Approved: OMB Number 0910-0025 Expiration Date: February 28, 2026

See Burden Statement on page 5.

Note: Items with an asterisk (\*) require a response.

### SUBMITTER INFORMATION If you are not submitting this report representing the manufacturing establishment for the radiation-emitting product causing the problem, you may enter your own company name under Establishment Identification and Submitter Address, or enter N/A and provide your home or other address. **Contact Information** Contact Name (Title, first name, last name)\* Occupation Title Email Address\* Establishment Identification (Manufacturer of the radiation-emitting product being reported, if known) **Establishment Name Division Name** Submitter Address Address Telephone Number\* Street\* Fax Number City\* State\* Zip Code\* INFORMATION REGARDING PRODUCT MANUFACTURER Product Manufacturer Name (If known) Product Manufacturer Address (If known) Street (Line 2) Street (Line 1) City Zip or Postal Code Territory, Province, or State Country Product Model Designation (If known) Model Name or Number Model Family Designation **Brand Name** Please provide any other information known regarding the manufacturer of the product that was involved in the accidental radiation exposure incident. If you are aware that the manufacturer was informed about the incident, please provide the contact information below. Contact Information (Including whom you contacted and address) **Date Contacted**

#### PRODUCT INFORMATION

**Product Types** (Please select the best match (only one). Note that product types are grouped into radiation categories.)

Acoustic Radiation	Microwave EMF Radiation (Continued)
Therapeutic Ultrasonic Devices (Including diathermy and stimulators)	Microwave Identification, Safety, Security, and Surveillance Products
Ultrasonic Medical Devices (Miscellaneous) (Including	Industrial Dielectric Heaters
lithotriptors)	Microwave Medical Products
Diagnostic Ultrasound Devices	☐ Microwave Heating and Drying Products
Sonic Medical Products (Including hearing aids and vibrators)	Microwave Communication, Data Transmit, and Measurement Products (Including CB radios, cell phones,
Ultrasound Non-Medical Products (Including jewelry cleaners and intrusion security systems)	walkie-talkies, household remote controllers)  Nuclear Magnetic Resonance Devices
Sonic Non-Medical Products	Household ELF Products (Including electric blankets)
Utrasonic Products	Other Microwave Product
Utrasonic Products	
Other Sonic or Ultrasonic Product	Optical Radiation
Ionizing Radiation	Medical Laser Products (Including surgical devices and laser therapy)
Personnel Security Systems (Including backscatter and transmission x-ray systems)	Surveying, Leveling, Alignment Laser Products (Including laser pointers, laser levels)
Cargo Non-Intrusive Security Systems	Laser Light Show/Display Products
Cabinet X-Ray Systems, Non-Medical (Including baggage	Toy, Novelty, Play Laser Products
x-ray systems)	Safety, Security, Surveillance Laser Products (Including
Industrial X-Ray Systems (Excluding Cabinet)	night vision systems, traffic speed systems and intrusion detection systems)
Analytical X-Ray Systems, Non-Medical  High Voltage Vacuum Switches	Research, Scientific, Laboratory Laser Products
Industrial Particle Beam Systems	☐ Material Processing Laser Products (Including welders,
TVs and video monitors ( <i>Not</i> including flat-screen TVs)	cutters, engravers)
Medical Diagnostic X-Ray Equipment	Data Measurement, Transmit, Control Laser Products (Including fiber optic communication systems, laser vision
Dental Diagnostic X-Ray Equipment	systems and process control systems)
☐ Therapeutic X-Ray Systems	Utility/Peripheral Laser Products (Including laser printers,
Veterinary X-Ray Systems	bar code scanners, CD and DVD systems)
X-Ray Bone Densitometers	In Vitro and Other Medical Laser Products (Including Veterinary devices)
X-Ray Film and Film Processing Materials	Patient Positioning Medical Laser Products
Cabinet X-Ray Systems, Medical	Other Laser Products
Medical Accelerators	Sunlamp Products (Including sunlamps and tanning beds)
Non-Medical Accelerators	☐ Mercury Vapor Lamps
High Voltage Vacuum Tubes	Ultraviolet Medical Products
Cathode Ray Tube (Without Electronics Chassis)	Ultraviolet Commercial/Consumer Products
Cold-Cathode Gas Discharge Tubes	Ultraviolet Surveillance & Detection Products
Other X-Ray Product	Ultraviolet Hygiene Products (Including UV sanitizers)
Microwave EMF Radiation	General Optical Products, Medical (Including surgical lamps)
Microwave Ovens (Food Prep)	General Optical Products, Non-Medical (Including LEDs
── Microwave Hyperthermia Therapy Devices	and fluorescent lamps)
Microwave Diathermy Machines	

### **Product Description** Description of product and its intended use **ACCIDENTAL RADIATION OCCURRENCE INFORMATION** Location of Occurrence Please provide the physical location where the Accidental Radiation Occurrence took place (e.g., at a residence, a factory, a tanning salon, school, restaurant, airport, etc.). If you do not know the exact address, provide responses to the best of your ability, or enter "Unknown." Location or Establishment Name Specific Section of Location or Establishment (If applicable) Address Telephone Number Street Fax Number City State Zip Code Date of Event\* From To Web Address Persons Involved Number of people exposed in the Number of people Number of unexposed Number of potentially exposed people Accidental Radiation Occurrence\* adversely affected\* people who were involved\* who have not exhibited any adverse reactions\* Type of reportable event Death Serious Injury Malfunction Other Description of the nature and magnitude of exposure and/or injuries

PRODUCT INFORMATION (Continued)

ACCIDENTAL RADIATION OCCURRENCE INFORMATION (Continued)
Description of the Radiation Occurrence
Is this a new Accidental Radiation Occurrence (ARO) report or a supplement to a previous ARO report filed by you or your organization? (Please select one.)*
New ARO report Supplement to previous ARO report (Enter date of previous report below.)
Date of previous ARO report, if applicable (mm/dd/yyyy) (Required entry* only if "Supplement to previous ARO report" is selected.)
Description of circumstances surrounding the accidental radiation occurrence (Please include a description of the activities leading up to the event and actions that occurred during the event, as well as any suspected causes of the occurrence.)*
Actions Taken
The actions described below are those taken to control, correct, or eliminate the causes and to prevent reoccurrence. If unknown, you may state "Unknown" below.
Description of specific actions, to date, taken by the manufacturer in response to the Accidental Radiation Occurrence*

# Actions Taken (Continued) Description of future actions to be taken by the manufacturer, if known, in response to the Accidental Radiation Occurrence (If this is a preliminary ARO report from the manufacturer, please indicate that further investigation is ongoing.)\* If this involved a medical device, has a Medical Device Report (MDR) been submitted to FDA?\* Unknown Yes No N/A Other Important Information (Please enter below) Feel free to send in medical documentation regarding the incident and injuries. Please mail this completed FORM FDA 3649 U.S. Food and Drug Administration Center for Devices and Radiological Health to the address to the right: Document Mail Center - WO66-G609 10903 New Hampshire Avenue Silver Spring, MD 20993-0002 This section applies only to requirements of the Paperwork Reduction Act of 1995.

**ACCIDENTAL RADIATION OCCURRENCE INFORMATION (Continued)** 

#### \*DO NOT SEND YOUR COMPLETED FORM TO THE PRA STAFF EMAIL ADDRESS BELOW.\*

The burden time for this collection of information is estimated to average 2 hours per response, including the time to review instructions, search existing data sources, gather and maintain the data needed and complete and review the collection of information. Send comments regarding this burden estimate or any other aspect of this information collection, including suggestions for reducing this burden, to:

Department of Health and Human Services Food and Drug Administration Office of Chief Information Officer Paperwork Reduction Act (PRA) Staff PRAStaff@fda.hhs.gov "An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB number."