

DIAGNOSTIC MISADMINISTRATION REPORT

Licensee name				Licensee number					
Address (number, street)				Event date			Report date		
				Month	Day	Year	Month	Day	Year
City	State	Zip code							

PATIENT DATA

Name of Authorized User	Patient Identifier
Name of Allied Health Personnel	Name of Patient's Referring Physician

Type of Misadministration <input type="checkbox"/> Wrong radiopharmaceutical <input type="checkbox"/> Wrong patient <input type="checkbox"/> Dosage differing from prescribed by 50% <input type="checkbox"/> Wrong route	Did the misadministration involve an isotope of iodine <input type="checkbox"/> Yes <input type="checkbox"/> No				Additional Description of Event:						
	Intended <input type="checkbox"/> No clinical procedure <input type="checkbox"/> Nuclear medicine study (Complete "Intended" and "given" sections) <input type="checkbox"/> X-ray study <input type="checkbox"/> Ultrasound <input type="checkbox"/> CT study <input type="checkbox"/> NMR study <input type="checkbox"/> Other:				Intended mCi Isotope Chemical Form Study				Given mCi Isotope Chemical Form Study		

Precipitator <input type="checkbox"/> Referring physician <input type="checkbox"/> Ward nurse <input type="checkbox"/> Ward clerk <input type="checkbox"/> Nuclear pharmacy			<input type="checkbox"/> Authorized user			<input type="checkbox"/> Hot Lab technologist <input type="checkbox"/> Imaging technologist <input type="checkbox"/> Clinical receptionist <input type="checkbox"/> Scheduling technologist <input type="checkbox"/> Patient <input type="checkbox"/> Other:		
Name of nuclear pharmacy			City	State				

Error					
Hot Lab		Referral	Administration	Other	
<input type="checkbox"/> Mislabeled a syringe <input type="checkbox"/> Mislabeled a vial or vial shield <input type="checkbox"/> Reconstituted wrong reagent kit <input type="checkbox"/> Placed reconstituted vial in wrong shield		<input type="checkbox"/> Selected wrong vial when drawing dosage <input type="checkbox"/> Set dose calibrator improperly <input type="checkbox"/> Misread dose calibrator <input type="checkbox"/> Misunderstood radiopharmaceutical or dosage order	<input type="checkbox"/> Misunderstood referring physician's request <input type="checkbox"/> Requested wrong study <input type="checkbox"/> Requested study for wrong patient	<input type="checkbox"/> Selected wrong patient <input type="checkbox"/> Answered waiting room page intended for other patient <input type="checkbox"/> Brought wrong patient to clinic <input type="checkbox"/> Selected wrong syringe from dosage cart	<input type="checkbox"/> Specify

Contributing Factors		Action Taken to Prevent Recurrence	
<input type="checkbox"/> Student technologist <input type="checkbox"/> New employee <input type="checkbox"/> Foreign language <input type="checkbox"/> Patient incoherent or unconscious <input type="checkbox"/> ID bracelet not checked		<input type="checkbox"/> Requisition not checked <input type="checkbox"/> Patient chart not checked <input type="checkbox"/> New procedure <input type="checkbox"/> Heavy workload <input type="checkbox"/> Other:	
		<input type="checkbox"/> Implement new procedures for: <input type="checkbox"/> Verification of request <input type="checkbox"/> Radiopharmaceutical labeling and handling <input type="checkbox"/> Verification of patient identification <input type="checkbox"/> Re instruct personnel <input type="checkbox"/> Reprimand personnel	
		<input type="checkbox"/> Improve supervision of personnel <input type="checkbox"/> No action <input type="checkbox"/> Other	

Effect on Patient(s) None apparent Dose Estimate Below Patient Informed See abstract

TARGET ORGAN	ESTIMATED RADIATION DOSE	Abstract (If more space is required, attach additional sheets.)

Prepared by	Radiation Officer (printed name)	Signature	Telephone number	Dated
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IMPORTANT: Retain this record for 5 years from the report date.