

### Confirmation and Sources of Assistance and Support (cont.)

- For help in projecting clinical effects, contact
  - nuclear medicine physician
  - Medical Radiological Advisory Team (MRAT) at Armed Forces Radiobiology Research Institute (AFRRI) 301-295-0530
- Obtain complete blood count
  - absolute lymphocyte count  $<1000 \text{ mm}^3$  suggests moderate exposure
  - absolute lymphocyte count  $<500 \text{ mm}^3$  suggests severe exposure
  - Acute, short-term rise in neutrophil count
- Swab both nostrils
- Collect 24 hour stool if GI contamination is possible
- Collect 24 hour urine if internal contamination with radionuclides is possible
- CDC ATSDR Hotline 770-488-7100

### Decontamination Considerations

- Exposure to a beam of radiation generally does not contaminate a patient. Patient contamination generally results from contact with radioactive particles.
- Treating contaminated patients before decontamination may contaminate the facility: plan for decontamination before arrival
- Exposure without contamination requires no decontamination (RSO measurement)
- Exposure with contamination requires Standard Precautions, removal of patient clothing, and decontamination with soap and water
- For internal contamination, contact the RSO and/or Nuclear Medicine Physician
- Patient with life-threatening condition: treat, then decontaminate  
Patient with non-life-threatening condition: decontaminate, then treat

### Treatment Considerations

- If life-threatening conditions are present, treat them first
- If external radioactive contaminants are present, decontaminate
- If radioiodine (reactor accident) is present, consider protecting the thyroid gland with prophylactic potassium iodide if within first few hours only (ineffective later). (Table 3)
- Review <http://www.afri.usuhs.mil/www/outreach/pdf/2edmmrhandbook.pdf> or <http://vaww.oqp.med.va.gov/cpg>

### Institutional Reporting

- If reasonable suspicion of a radiation event, contact hospital leadership (Chief of Staff, Hospital Director, etc)
- Immediately discuss hospital emergency planning implications

### Public Health Reporting

- Contact local public health office (city, county or state)
- If needed, contact the FBI (for location of nearest office, see <http://www.fbi.gov/contact/fo/fo.htm>)

## TERRORISM WITH IONIZING RADIATION GENERAL GUIDANCE Pocket Guide

### Diagnosis: Be Alert to the Following

- Acute radiation syndrome (table 1) follows a predictable pattern after substantial exposure or catastrophic events
- Victims may also present individually, as described in table 2, over a longer period of time after exposure to contaminated sources hidden in the community
- Specific syndromes of concern, especially with a 2-3 week prior history of nausea and vomiting, are
  - thermal burn-like skin lesions without documented heat exposure
  - immunological dysfunction with secondary infections
  - a tendency to bleed (epistaxis, gingival bleeding, petechiae)
  - marrow suppression (neutropenia, lymphopenia, and thrombocytopenia)
  - hair loss

### Understanding Exposure

- Exposure may be known and recognized or clandestine as
  - large radiation exposures, such as a nuclear bomb or catastrophic damage to a nuclear power station
  - small radiation source emitting continuous gamma radiation producing chronic intermittent exposures (such as radiological sources from medical treatment or industrial devices.)
  - skin contamination with radioactive material (“external contamination”)
  - internal radiation from absorbed, inhaled, or ingested radioactive material (“internal contamination”)

### Confirmation and Sources of Assistance and Support

- Contact radiation safety officer (RSO) for help



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VA access card: <http://www.oqp.med.va.gov/cpg/cpg.htm>

DoD access card: <http://www.qmo.amedd.army.mil>

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**TABLE 1: ACUTE RADIATION SYNDROME**

1 Gray (Gy) = 100 rads 1 centiGray (cGy) = 1 rad

Whole body radiation from external radiation or internal absorption							
Phase of Syndrome	Feature	Subclinical range		Sublethal range		Lethal range	
		0 – 100 rad or cGy	100-200 rad 1-2 Gy	200-600 rad 2-6 Gy	600-800 rad 6-8 Gy	800-3000 rad 8-30 Gy	>3000 rad >30 Gy
<b>Prodromal Phase</b>	Nausea, vomiting	none	5-50%	50 - 100%	75-100%	90-100%	100%
	Time of onset		3-6 hrs	2-4 hrs	1-2 hrs	<1 hr	Minutes
	Duration		<24 hrs	<24 hrs	<48 hrs	<48 hrs	N/A
	Lymphocyte count	Unaffected	Minimally decreased	< 1000 at 24 hr	< 500 at 24hr	Decreases within hours	Decreases within hours
	CNS function	No impairment	No impairment	Cognitive impairment for 6-20 hrs	Cognitive impairment for >24 hrs	Rapid incapacitation, often after a lucid period of up to several hours	
<b>Latent Phase (subclinical)</b>	Absence of Symptoms	> 2 wks	7-15 days	0-7 days	0-2 days	None	
<b>Acute Radiation Illness or "Manifest illness" phase</b>	Signs and symptoms	none	Moderate leukopenia	Severe leukopenia, purpura, hemorrhage Pneumonia Hair loss after 300 rad/3 Gy		Diarrhea Fever Electrolyte disturbance	Convulsions, Ataxia, Tremor, Lethargy
	Time of onset		> 2 wks	2 days - 2 wks		1-3 days	
	Critical period		none	4-6 wks - Most potential for effective medical intervention		2-14 days	1-48 hrs
	Organ system	none		Hematopoietic and respiratory (mucosal) systems		GI tract Mucosal systems	CNS
<b>Hospitalization</b>	%	0	<5%	90%	100%	100%	100%
	Duration		45-60 days	60-90 days	90+ days	weeks to months	days to weeks
<b>Mortality</b>		None	Minimal	Low with aggressive therapy	High	Very high, significant neurological symptoms indicate lethal dose	

**TABLE 2: SYMPTOM CLUSTERS AS DELAYED EFFECTS AFTER RADIATION EXPOSURES**

Headache	Partial and full thickness skin damage
Fatigue	Hair loss
Weakness	Ulceration
Anorexia	Lymphopenia
Nausea	Neutropenia
Vomiting	Thrombopenia
Diarrhea	Purpura
	Opportunistic infections

**TABLE 3: POTASSIUM IODIDE DOSAGES:**

The dose of potassium should be taken once a day until a risk of significant exposure to radioiodines no longer exists \*

Age group	Dosage
Infants < 1 month	16 mg
Children 1 months-3 yrs	32 mg
Children 3-18 yrs	65 mg
Adults	130 mg

\* For information regarding preparation of potassium iodine solution:  
<http://www.fda.gov/cder/drugprepare/kiprep.htm>