

REAC/TS Fact Sheet for Medical Providers

Fast Facts:

Exposed only: Individuals exposed to a discrete, intact radioactive source have radiation exposure, not contamination. They present no risk to treating medical personnel.

Contaminated: Radioactive contamination on bodies or clothing (external contamination) or have inhaled/ingested/absorbed (internal) radioactive contamination are at continued risk of worsening radiation injury and after life-saving treatment, should be decontaminated and / or treated for internal contamination. Risk may be mitigated to care-giver and patient.

Radiation Combined Injury: Trauma and / or burns in addition to radiation injury. These patients will increase 1-2 triage acuity categories and will have a worse prognosis.

0-1 Gy

≥1-2 Gy

5 - 6 Gy

10 Gy

6 Gy

Mitigating Risk: Rapid removal of casualties from blast site, getting out of the area around the blast site, treating in a solid shelter, when possible, and use of personal protective equipment as appropriate to the hazard or hazards.

Ionizing Radiation Injury:

Ionizing radiation induced biological effects are determined by:

- · Dose, Dose rate, Volume of body part exposed, Radiation type, Co-existing health conditions, Trauma, Burns
- · Main site of cellular injury: DNA (immature/rapidly dividing cells at high risk)

Acute Radiation Syndrome:

TREAT LIFE-THREATENING CONDITIONS FIRST!

The lethal dose (LD) that kills 50% of the exposed population (LD50) within 60 days after exposure (LD50/60) is: Dose leading to sub-syndromes:

Subclinical

Cutaneous

Hematopoietic

Gastrointestinal

Neurovascular

Healthy,	young	adults	without
therapy:	~3.5 -	4.0 Gy	/

Shifts to right with adults with antibiotics, supportive care and colony stimulating growth factors

Sub-syndromes:

Hematopoietic (bone marrow) syndrome: Loss of lymphocytes followed by neutrophils, and later, loss of red blood cells (RBCs) and platelets

Immune dysfunction, infections and sepsis, impaired wound healing, and hemorrhage may also occur

Treatment: Neutropenic precautions, consider prophylactic fluoroquinolones and/or other antimicrobials

U.S. FDA approved treatments:

G-CSF, PEGylated G-CSF, GM-CSF:

- Filgrastim: 10 mcg/kg, subcutaneous q day
- · PEGfilgrastim: 2 doses, 6 mg each given 1 week apart

Pediatric: < 10 kg: 0.1 mg/kg; 10-20 kg: 1.5 mg; 21-30 kg: 2.5 mg; 30-45 kg: 4 mg

- Sargramostim (GM-CSF):
 - Adults and Pediatric patients > 40 kg: 7mcg/kg
 - Pediatric patients 15 kg to 40 kg: 10 mcg/kg

Romiplostim: Thrombopoietin receptor agonist for thrombocytopenia: nplate_pi_hcp_english.pdf (amgen.com)





REAC/TS RADMED App

Search RADMED on Android or Apple



Gastrointestinal (GI) syndrome: May present with nausea, vomiting, diarrhea, bloody stool, and dehydration

GI bleed, bowel obstruction, acute renal failure, cardiovascular failure (8 - 14 days) may occur

Treatment: Antiemetics as indicated, enteral/parenteral nutrition, intensive care, consider bowel decontamination, stress ulcer prophylaxis

Neurovascular syndrome: Nausea and vomiting within 30 minutes, confusion and disorientation within minutes, severe hypotension, and fluid shifts, with possible cerebral edema, ataxia, seizures, coma

May be fatal within 24 - 48 hours

Treatment: Supportive and if resource adequate, intensive care

Triage/Dose Estimation: TREAT LIFE THREATENING CONDITIONS FIRST!

History and physical; time to vomiting; geographic location and time in area of blast; clinical prodrome from above; Complete Blood Count (CBC) with differential every 6-12 hours; and dicentric chromosome analysis (will take minimum 4 days)

Biodosimetry Based on Acute Photon-Equivalent Exposures

	Onset of vomiting		Lymphocyte count (x10º/liter) by day*			Lymphocyte depletion rate	Number of	dicentrics			
Dose [Gy]	ማං	Time [hr]	0.5	1	2	4	6	8	Rate constant	Per 50 cells	Per 1000 cells
0			2.45	2.45	2.45	2.45	2.45	2.45		0.05 - 0.1	1-2
1	19		2.30	2.16	1.90	1.48	1.15	0.89	0.126	4	88
2	35	4.63	2.16	1.90	1.48	0.89	0.54	0.33	0.252	12	234
3	54	2.62	2.03	1.68	1.15	0.54	0.25	0.12	0.378	22	439
4	72	1.74	1.90	1.48	0.89	0.33	0.12	.044	0.504	35	703
5	86	1.27	1.79	1.31	0.69	0.20	0.06	.020	00.63	51	1024
6	94	0.99	1.68	1.15	0.54	0.12	0.03	.006	0.756		
7	98	0.79	1.58	1.01	0.42	.072	.012	.002	0.881		
8	99	0.66	1.48	0.89	0.33	.044	.006	<.001	1.01		
9	100	0.56	1.39	0.79	0.25	.030	.003	<.001	1.13		
10	100	0.48	1.31	0.70	0.20	.020	.001	<.001	1.26		

* The normal range for lymphocytes in human blood is between 1.4 and 3.5 x 10° per liter.

Lymphocyte depletion rate is based on the model $L_{r} = 2.45 \times 10^{\circ}$ /liter x e^{4c0x} where L_{r} equals the lymphocyte count (x10°/liter), 2.45 x 10°/liter equals the a constant representing the consensus mean lymphocyte count in the general population, k equals the lymphocyte depletion rate constant for a specific acute photon dose, and t equals the time after exposure (days)

Cutaneous Radiation Injury/Syndrome:

- Acute effects (days to weeks post exposure): Redness, swelling, blisters, ulceration, tissue necrosis
- Long-term issues (month to years post exposure): Fibrosis, atrophy (sclerosis), and telangiectasia formation

Treatment: Topical Class II/III steroids, antihistamines, antibiotics, and moisturizers (Aquaphor[®]), Pentoxifylline with α-tocopherol; growth factors; artificial skin/bioengineered constructs; debridement; and other surgical techniques

Internal Contamination/Countermeasures:

- Enters body through airways/ingestion/wound contamination and incorporation into body tissues
- Bioassay of Urine/Feces to assess internal contamination

Treatment:

Potassium lodide: blocks I¹³¹ (nuclear detonation/reactor failure)

- Treat before exposure or within 6-12 hours of exposure
- · Maintain until no longer being exposed
- I¹³¹ Risk greatest to children, infants, and young adults

Prussian Blue (Radiogardase®): for Cesium – drives excretion via feces – U.S. FDA approved

• Dose: adults/children: 3 grams orally, 3 times a day/1 gram orally, 3 times a day - assess via bioassay

DTPA: U.S. FDA approved for Plutonium, Americium

- · Every 24 hour Dosing
- Dose: Initially 1 gm Ca-DTPA IV and then Zn form 1 gm IV until decision to cease by bioassay
- · For inhalation intake use nebulizer (1:1 dilution with water/saline)

Dose [Gy]	Sign	Timing
3	Epilation	Begins around day 14 - 17
6	Erythema Distinguish from thermal burn	Minutes to weeks, depending on dose
10 - 15	Dry desquamation	2 - 3 weeks post-exposure, depending upon dose
15 - 20	Moist desquamation	2 - 3 weeks post-exposure, depending upon dose
25	Deep ulceration Radionecrosis	21 days

Wasalenko JK et al. Ann Intern Med. 2004

U.S. FDA Potassium Iodide Guidelines

Age Category	Predicted Absorbed Dose to the Thyroid cGy ^b	Kl Dose (mg)°	Number of 130 mg Tablets
Adults 40 y	500	130	1
Adults 18-40 y	10	130	1
Pregnant or lactating women	5	130	1
Adolescents 12-18 y°	5	65	0.5
Children 3-12 y	5	65	0.5
1 month - 3 y	5	32	0.25
Birth - 1 month	5	16	0.125