

Cautions

- Orders must be customized for each event and patient!
- Specific drugs are suggested for function only, and patients may not need any/every category of drug listed. Consult the notes at the end of this document.
- This Adult Orders Prototype lists only FDA-approved medications as radioisotope countermeasures for internal contamination; see page 9-10. These drugs are currently in the Strategic National Stockpile. Prescribers should consult the FDA drug label for complete information.
- All dosages in this prototype are based on a 70 kg adult with normal renal and hepatic function. Appropriate dosage adjustments should be made based on age, weight, drug-drug interactions, nutritional status, renal and hepatic function. Pediatric doses are not referenced, except for Potassium Iodide.
- This Adult Orders Prototype does **not** address threshold levels of internal contamination that would trigger initiation, continuation, or discontinuation of decorporation treatment. See <u>REMM Countermeasures Caution and Comment</u> information that discusses this issue.
- After a mass casualty event, practitioners may encounter counterfeit drugs. This
 <u>FDA website</u> will provide information on avoiding and detecting counterfeit drugs
 and assist reporting of suspected counterfeit medications.
- See "Notes" at end of order list for additional information.



1. Administrative information	
Name: Unique Identifier:	
Address:Phone:	
Spoken language:	
Admit to:	
Hospital ward	Area
Team:	ICU
Physician:	Other
Next of kin contact information:	
Special needs:	
2. Diagnoses: Radiation, Other	
Radiation contamination: desc	cription
· See REMM Body Chart (page 14)) to record whole body radiation survey.
External contamination wi	th Isotope (Specify)
Internal contamination wit	th Isotope (Specify)
Contamination suspected,	Isotope uncertain
Radiation Exposure / Acute R	Radiation Syndrome (ARS)
Date of exposure Time of exposure Location of patient at time Estimated whole body/part Dose unknown	of exposure ial body dose, specify (dose)
· See REMM information on D	Oose Reconstruction.
Other potential complicating f	factors
Combined injuries? e.g. burn, Specify:	
Mass casualty incident	



Specific populations potentially requiring more customized management See REMM At-Risk/Special Needs Populations page __ Young age (e.g. children < 12-16 y) __ Older age (e.g., those > 65 y) __ Pregnant/Possibly pregnant __ Immunosuppressed __ History of prior significant chronic disease(s) or conditions. Specify each, including meds or special needs required for each: __ Precautions __ Contact __ Droplet __ Airborne __ External Radiation __ Internal Radiation ___ Reverse/Neutropenic **Urgent Consultations as indicted:** __ Hematopoietic Stem Cell Transplantation __ Radiation Oncology — Hematology / Oncology __ Transfusion Medicine ___ Mental Health / Psychiatry __ Endocrinology __ Ophthalmology ___ Pain Service __ Dermatology / Plastic Surgery __ Gastroenterology __ Radiation Safety ___ Burn Therapy 3. Condition: __ Good __ Fair __ Stable __ Guarded __ Critical 4. Vital Signs: __ q 2 hours X 4 __ q 4 hours X 4 __ Ward routine Notify physician for:



Temperature > 38.5 °C SBP > 180, <100 DBP > 100 < 50 HR >100 <50 RR >30 <8 O ₂ saturation < 92%
5. Special orders for patients with known or possible radiation contamination
Radiation precautions
 Universal precautions with gown, mask, cap, boots, and gloves Use medical facility procedures for discarding biological/physical/radioactive waste, including linens/towels/trash/personal protective equipment. Contact Radiation Safety Officer for additional instructions.
 Place radiation safety sign on door if patient has internal or external radioactive contamination. Notify pregnant staff that entry to room is prohibited if patient is/may be contaminated. Everyone entering room/area of contaminated patient must wear personal radiation dosimeter.
6. Allergies:
No Known Drug Allergies (NKDA)
Allergies (drugs, foods)
If yes, specify:
7. Activity:
Bed rest Bathroom privileges
Out of bed every hrs Ambulate as tolerated
8. Diet:
 Regular Diet Liquids (full, clear) NPO Advance as tolerated Neutropenic diet Other
Special dietary needs/requests
9. Height, weight, age:
Height: feet inches Weight: lbs oz cm kg



Age: ____ years Repeat body weight: q ____ hours q ____ days 10. Peripheral IV management: ___ IV Fluids: _____ @ ____ cc/hr, with additive _____ ___ IV Fluids: _____ @ ____ cc/hr, with additive _____ 11. __ Foley catheter management Use radiation precautions for urine and feces for patients with internal radiation contamination. 12. __ Monitor I / O Frequency _ Use radiation precautions for urine and feces for patients with internal radiation contamination. 13. Deep Venous Thrombosis (DVT) prophylaxis¹: __ TED hose to Bilateral Lower-Extremities Seguential Compression Devices (SCD) ___ Anticoagulation regimen _____ __ Other The potential benefit of anticoagulation (e.g. heparin^{1,2}) should be balanced against the risk of excessive bleeding in patients with severe thrombocytopenia or significant gastrointestinal toxicity. 14. Respiratory Therapy: (Radiation precautions needed if patient is contaminated.) Room air Chest tube care (Specify) ___ Titrate oxygen supplementation for Oxygen saturation > ____% ___ Nebulizer treatment (Specify) ______ 15. Wound care¹: (see also item 22: burn therapy) ___ Decontaminate external wounds if there is external contamination. See REMM contaminated wound care recommendations.

__ Sterile dressing to wounds daily



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Monitor waste: Use medical facility procedures for discarding biological/radioactive/physical waste and linens/towels/trash/personal protective equipment. Radiation precautions needed if patient is contaminated.
Silvadene (Silver Sulfadiazine) 2 cream topically to burns
Other wound management per Burn team/Dermatology/Surgery: Pager Phone
16. Orthopedic care:
Splint/brace/cast
Other orthopedic management procedure per orthopedics: Pager Phone
17. Admission labs / imaging studies / other:
CBC w/differential
Comprehensive Metabolic Panel (CMP) / Chem 14
Cardiac enzymes
PT / PTT
Urinalysis
Urine culture
Blood culture x 2
Urine HCG
Serum HCG
Thyroid Function Tests (Specify)
Serologies: Herpes Simplex Virus type 1 (HSV-1) Herpes Simplex Virus type 2 (HSV-2) Cytomegalovirus (CMV) Varicella-zoster virus (VZV)
Electrocardiogram
Chest x-ray PA/Lat Portable
Other imaging studies Specify:

18. Standing labs / studies:



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CBC w/diff q hours, x days, Followed by q until further orders		
Comprehensive Metabolic Panel (CMP) / Chem 14 Followed by q hours, x days Followed by q until further orders		
19. Electrocardiogram		
Electrocardiogram STAT Electrocardiogram for chest pain, notify physician		
20. Biodosimetry/Bioassay tests: See REMM's Explaining Biodosimetry.		
For biodosimetry: See REMM for more on the <u>Dicentric chromosome assay</u> .		
Dicentric Chromosome Assay: Draw extra green top tube on: date time		
Send this tube ON ICE for outside lab study		
To the Attention of:		
Name of Lab:Address of Lab:		
See REMM for location of <u>laboratories that perform this test</u> .		
For Bioassay: tests evaluating/managing internal decontamination:		
Spot urine for name of radioactive isotope		
24-hour urine for name of radioactive isotope		
Spot fecal specimen for name of radioactive isotope		
24-hour fecal specimen for name of radioactive isotope		
Send specimen to:		
Special requirements for containment, labeling, and shipping of specimen:		
Note: Consult senior radiation event medical managers for name and location of specialized laboratories if your facility cannot perform these assays.		
21 Type and cross match		
Type and screen		
For units of packed red blood cells For units of platelets		

 $\boldsymbol{\cdot}$ Use only leukoreduced AND irradiated products, if available, unless



it is known with certainty that the patient was exposed to a low dose of radiation, e.g. less than 100 cGy.

- · If dose is not known with certainty, leukoreduced AND irradiated products are preferred, if available.
- · See <u>REMM blood use page</u> for additional information.

22. General Medications ¹ :
For gastric acid suppression:
Lansoprazole (Prevacid) ² 15-30 mg PO daily
For radiation-induced nausea & vomiting:
 Ondansetron (Zofran)² 4 mg IV q 8h PRN nausea/emesis Lorazepam (Ativan)² 0.5 mg – 1 mg PO q 6-8h PRN anxiety/insomnia/breakthrough nausea Prochlorperazine (Compazine)² 10 mg PO/IM/IV q 6-8h PRN anxiety/insomnia/breakthrough nausea
 See <u>American Society of Clinical Oncology 2006 Anti-emetic Guidelines</u>³ See NEJM June 5, 2008 article: <u>chemotherapy induced nausea and vomiting</u>³
For Fever:
Acetaminophen (Tylenol) ² 650 mg PO q 6 – 8h PRN temperature > 38 °C
For diarrhea:
 Loperamide hydrochloride (Imodium)²: Recommended initial dose is 4 mg (2 capsules) followed by 2 mg (1 capsule) after each unformed stool. Daily dose should not exceed 16 mg (8 capsules)
For constipation:
Senna (Senokot) ² 2 tabs PO BID, hold for loose stools
Docusate (Colace) ² 100 mg PO BID, hold for loose stools. Per FDA monograph: 50 to 360 mg QD or divided BID for adults
For rash:
Topical sterile dressing
Diphenhydramine hydrochloride (Benadryl) ² 25-50 mg PO q 4-6 hours for pruritis, not to exceed 300 mg/24 hours

For pain:



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Morphine sulphate ² mg route frequency				
For skin burns: (see also item 15: wound care)				
Burn topical regimen				
Replace body fluid				
Other burn therapy				
For oral mucositis:				
Mouth care regimen				

23. For radioisotope decorporation or blocking:

- Note: Only FDA approved radiation countermeasures are listed in table below.
- See <u>REMM Radiation Countermeasures for Treatment of Internal Contamination</u> table for longer list of countermeasures which have been recommended by some experts but are not FDA approved as radiation countermeasures.

Medical	Administered	Route of	Dosage	Duration
	for	Administration	Dosage	Daration
Countermeasure	101	Administration		
Ca-DTPA ^{2,4}	Americium	IV ² :	IV:	Ca-DTPA for
Zn-DTPA ^{2,4}	(Am-241) ²	Give once daily as a	1 g in 5 cc 5%	the first dose
		bolus or as a single	dextrose in	 Give Zn-DTPA
See REMM's DTPA		infusion, i.e., do not	water (D5W) or	for any follow-
<u>information.</u>	(Cf—252) ³	fractionate the dose.	0.9% sodium	up doses (i.e.,
			chloride (normal	maintenance
See FDA's Zn-	Cobalt	DTPA is FDA-approved	saline, NS) slow	as indicated)
DTPA drug label.	(Co-60) ³	for intravenous Rx of	IV push over 3-	 Duration of
		known or suspected	4 minutes	therapy
		internal contamination		depends on
DTPA drug label.	(Cm-244) ²	with Am, Cm, and Pu	OR	total body
		only.		burden and
	Plutonium		1 g in 100-250	response to
	(Pu-238 and		cc D5W or NS as	treatment
	Pu-239) ²		an infusion over	
		Nebulized	30 minutes	
		inhalation ² :		
	(Y-90) ³	DTPA is FDA-approved	Nebulized	
		for nebulized inhalation	inhalation:	
		in adults only, and if	1 g in 1:1	
		the route of	dilution with	
		contamination is	sterile water or	
		through inhalation.	NS over 15-20	
			min	



Medical	Administered	Route of	Dosage	Duration
iviculcai	for	Administration	Dosage	Duration
Countermeasure	101	Administration		
Potassium iodide ² See REMM's KI summary information. See FDA's KI information.	Iodine (I-131)	PO	years: 130 mg/day (for projected	dose of KI. Incident managers may recommend additional doses if ongoing radioactive iodine ingestion or inhalation represents a continuing threat.
Prussian blue, insoluble ² See REMM's Prussian Blue summary information. See FDA's Prussian Blue drug label.	(Cs-137) Thallium (TI-201)	PO	FDA package insert) OR 1 - 3 g PO tid with 100-200 mL water, up to 10-12 g/day	 Minimum 30 days course per FDA Obtain bioassay and whole body counting to assess treatment of efficacy Duration of therapy depends on total body burden and response to treatment



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24. Neutropenia therapy, if indicated^{1, 5}:

- Although the 3 drugs listed below are FDA-approved for the treatment of chemotherapy induced neutropenia, none is approved either for radiation-induced neutropenia or as prophylactic treatment prior to the onset of neutropenia.
- · See additional REMM information on white cell growth factors/cytokines.
- In a mass casualty radiation event, use of these drugs would be off-label or require a formal Emergency Use Authorization.

Cytokine ³	Adult dose	Pregnant Women ⁶
G-CSF or filgrastim ³ (<u>Neupogen</u>)	 Subcutaneous administration 5 ug/kg/day via single daily injection Continue until absolute neutrophil count > 1.0 x 10⁹ cells/L 	Class C ⁶ (Same as adults)
Pegylated G-CSF or pegfilgrastim ³ (Neulasta)	 1 subcutaneous dose, 6 mg Consider second 6 mg dose 7 or more days after initial dose, if significant neutropenia persists 	Class C ⁶ (Same as adults)
GM-CSF or sargramostim ³ (Leukine)	 Subcutaneous administration 250 ug/m²/day Continue until absolute neutrophil count > 1.0 x 10° cells/L 	Class C ⁶ (Same as adults)

See Practice Guidelines for myeloid growth factors

- National Comprehensive Cancer Network
- · American Society of Clinical Oncology

Antimicrobial prophylaxis¹:

- · Use as appropriate for each patient.
- · Drugs listed are examples only.

Anti-bacterial prophylaxis

__ Levofloxacin (Levaquin)² 500 mg PO/IV qd

Anti-viral prophylaxis

__ Acyclovir (Zovirax)² 400 mg PO q12h, or __ Acyclovir (Zovirax)² 250 mg/m² IV q12h

Anti-fungal prophylaxis

- __ Fluconazole (Diflucan)² 400 mg PO/IV daily beginning when absolute neutrophil Count (ANC) becomes < 1000, or</p>
- __ Posaconazole (Noxafil)² 200 mg PO tid with food beginning when absolute Neutrophil Count (ANC) becomes < 1000</p>

25. Fever and Neutropenia¹



Blood cultures x 2 sets	Urinalysis w/culture			
Sputum culture + sensitivity	Chest x-ray			
Cefepime (Maxipime) ² 2 gm IV q 8	3h			
Vancomycin (Vancocin) ³ 1gm IV o	12h, consider trough level before 4th dose			
See current Fever and Neutropenia Guidelines from • IDSA Infectious Diseases Society of America • ASCO American Society of Clinical Oncology				
Antifungal therapy (consider one of tl	ne following ¹):			
Liposomal amphotericin B (Amb	isome) ² 3mg/kg/day IV over 1-4h			
Amphotericin B lipid complex (A	belcet) ² 3mg/kg/day IV over 1-4h			
Voriconazole (Vfend) ³ 6mg/kg IV	q 12h for two doses, then 4mg/kg IV q 12h			
Caspofungin (Cancidas) ² 70mg I	V once then 50mg IV q 24h			

NOTES

- 1. Suggested drugs are listed as representatives of a functional class, and no specific medication endorsement is implied. Dosages are based on a 70 kg adult with normal baseline renal and hepatic function. Appropriate dosage adjustments should be made based on age, weight, drug-drug interactions, nutritional status, renal and hepatic function, and any other patient-specific characteristics that may apply.
- 2. FDA approved for this indication
- 3. This drug is **not** approved by the FDA for this indication. If used, this would be an "off label use", and physician discretion is strongly advised.
- 4. Ca-DTPA and Zn-DTPA have not been approved by FDA for treating internal contamination with californium, thorium, and yttrium. For initial treatment, Ca-DTPA is recommended, if available, within the first 24 hours after internal contamination. Zn-DTPA is preferred for maintenance after the first 24 hours, if available, due to safety concerns associated with prolonged use of Ca-DTPA.
- 5. When to initiate treatment with cytokines
 - Initiation of treatment should be strongly considered for victims who develop an absolute neutrophil count of < 0.500 x 10⁹ cells/L and are not already receiving colony-stimulating factor.
 - Evidence from **animal studies** indicates that outcomes may be improved if colony stimulating factors are administered as soon as possible after radiation exposure, and prior to the onset of neutropenia.
 - Although most therapy for ARS is directed at actual clinical signs and symptoms, some clinical effects of ARS can be anticipated and potentially



mitigated, as with the use of prophylactic <u>white cell cytokines</u>. This prophylactic use is also off label.

- <u>Emergency Use Authorization</u> will be required for use of cytokines for radiation induced neutropenia in a mass casualty setting.
- See published guidelines links in section 24.

6. For pregnant women:

- Experts in biodosimetry must be consulted.
- Any pregnant patient with exposure to radiation should be evaluated by a health physicist and maternal-fetal specialist for an assessment of risk to the fetus
- Class C refers to U.S. Food and Drug Administration Pregnancy Category C, which indicates that studies have shown animal, teratogenic, or embryocidal effects, but there are no adequate controlled studies in women; or no studies are available in animals or pregnant women.



Body Chart for Recording Results of Radiation Survey

