Instructions on use of RADIATION CASUALTY ASSESSMENT TOOL

METER Course, v2.4 (5/09)

This information packet ('tool') is designed to help with the assessment and management of casualties of an incident involving radiation. Use one packet per casualty, labelling each page. It should become part of the permanent record for that casualty. You do not have to use those parts of the tool that do not apply to that casualty.

1. Triage Guide Question 1: Is patient I □ "NO" then □ "YES" then go to Question]	 filled out by triage MD or RN used to establish initial priority (i.e. immediate treatment vs. immediate decontamination vs delayed treatment and/or decontamination) designed to look and function like the SARS screening tool
2. History and Physical form (2 pages) NameAgeM/F DateTime of Arrival Physician:Time seenh Mode of arrival: self □ EMS □ other □ HISTORY AND F Vitals: HR BP/ Temp°C RR sats% onRA/Lpm	 filled out by treating MD used to record findings on history and physical prompts physician to obtain specifics relevant to treatment and disposition decisions unique to radiation exposure and/or contamination includes biodosimetry estimates using three clinical measures
3. Body Mapping form for Skin Contamination and Injury Name Age M/F Date Time of Arrival M/F Physician: Time seen h BODY MAPPING Injuries, burns, skin changes seconda Oracle for details be	 filled out by treating MD or RN used to facilitate recording location of skin contamination contaminated areas are recorded (with initial count and description) as they are discovered by person performing survey. All contaminated areas must be decontaminated, with final counts recorded as well also used to record location of injuries
 4. Standing Orders ALLERGY ALERT No known drug allergy Known allergies: DATE TIME i.v.: DNS vs D 	 filled out by treating MD prompts physician to order specific labs, specimens, and medications relevant to treatment of radiation exposure and/or contamination
5. Severity Scoring form (2 pages) SEVERITY Time of Exposure Time of Symptom Onset Time of Assessment 1. NEUROLOGICAL (Circle mos Acute Symptom 2 1 (mild) 2 (mode	 reference material for treating MD allows physician to estimate severity of injury due to radiation exposure when the exposure dose has not been determined. This may help with disposition decision lists some decorporating agents for internal contamination, table of 'time of onset of vomiting' as biodosimetry marker

 Name_____Age___M/F

 Date_____Time of Arrival_____h
 ___Age____M/F Triaged by: _____ Time seen _____h Mode of arrival: self
EMS
ambulatory
stretcher
METER Course, v2.4 (5/09)

PLACE ID **STICKER HERE**

TRIAGE

YT	Question 1: Is p	patient <u>medically</u> <u>stable?</u>
LABILI	□ "NO" then →	 Cover with sheet, assume contaminated Move immediately to Contaminated Treatment Area
S	□ "YES" then go to	Question 2

Question 2: Does patient have measurable skin
<u>contamination</u> during 2 minute survey with Geiger
Counter in triage?

CONTAMINAT	□ "YES" then →	 Identify as contaminated (<i>i.e.</i> red bracelet) Record sites/activity of contamination (p 5) Prioritise for decon, move patient to decon site, then integrate into cohorted stream of uncontaminated ED patients
0		4. Further assess for Exposure ASAP
0	\square "NO" then \longrightarrow	1. Identify patient as uncontaminated (<i>i.e.</i> green bracelet)
		2. go to Question 3

RE	Question 3: Does patient have history, signs and symptoms of possible <u>exposure</u> to radiation?	
Inso	□ "YES"	 New onset of nausea, vomiting, diarrhea or skin changes? New onset of weakness, confusion, unexplained low BP?
EXP	□ "NO"	 Prioritise for treatment integrate into cohorted stream of uncontaminated ED patients

NO

RADIATION CASUALTY ASSESSMENT TOOL	
Nama M/E	PLACE ID
NameAgeM/FDateTime of Arrivalh	STICKER HERE
Physician: Time seen h	METER Course, v2.4 (5/09)
HISTORY AND	PHYSICAL Form
Vitals: HR BP/ Temp°C RR sats% onRA/Lpm	Details of radiation contamination/exposure:
Chief complaint:	
HPI:	Isotope known: unknown Type of particle: α □ β □ γ □ X-rays □ neutrons □ State: solid/powder □ liquid □ gas/steam □
	Contaminationsee diagramExternal contamination: yesnounknownExtent of contamination (see diagram):localised (skin/hair)WoundGeneralisedInternal contamination: yesnounknown
	$\frac{\text{Decontamination}}{\text{Location: in field } \square \text{ at ED } \square \text{ , done by}}$
	Exposure yes □ no □ unknown □ Time of exposure:h, Duration:hmin Whole body □ Parts of Body □
	Past Medical History Immunosuppression
	Cancer (radiation chemo , when?)
Review of Systems (selected)	Previous fluoroscopy/Nuc Med testing/occupational
Neuro: Confusion D Fatigue D	exposure?
Changes in: speech vision dizzy headache	Other:
Vomiting: yes 🛛 or no 🗤 # of times: (began ath, =h after exposure)	
Motor/sensory deficits? Cognitive deficits?	Medications (include dose & freq if known):
Blood: Active bleeding? Bruising Petechiae Petechiae	
Derm : Redness or Rash □ (Time of onset:h) Swelling □ Blisters □ Ulcers □ Desquamation □ Hair loss □ Onycholysis □ Dysaesthesia/pruritis □	Allergies to meds: NKDA/
GI: Nausea - (severity:/10) Anorexia - Abdominal pain - Blood -/mucus - in stool Diarrhea - (began ath; # of times:)	Social history:
if female: LMP, Pregnant: yes/no/?	

Name	Ane M/F	
Date	AgeM/F Time of Arrivalh	STICKER HERE
	Time seen h	
·		METER Course, v2.4 (5/09
Physical exam:		Labs & Investigations:
		Blood samples
		$\Box CBC: WBC x10^3.$
		Abs Lymphoocytes Abs Neutrophils
		Hgb mg/dL, Plt $x10^3$
		HgbRg/dL, PltK_10 ³ □ Chem 7: NaClKCO2 BUNCreatGlc
		Pregnancy test (all females): neg/pos
		□ Thyroid : TSH, T3, free T4
		 Cytogenetics (green-top tube; keep at room temp; send ASAP) if exposure potentially > 0.5 Gray
		□ HLA typing (green-top tube; hold if potential for
		requiring bone marrow transplant))
		Specimens
		(scan with Geiger Counter, then label & save)
		Nasal swabs (labeled L&R): activity: yes/no
		Mouth Swab: activity yes/no
		 Urine sample: activity yes/no Stool sample: activity yes/no
		 Emesis sample: activity yes/no
		ECG:
		imaging studies:
	ng different methods of estimating	
	; use REMM Tool or tables p7-8 to	
calculate estimated d		Course in ED:
1 Time of exact of y		
	romiting (see Table on page 8) xposure & onset vomiting: h	
- Estimated dose:		
2 Ab b - b - b		
2. Absolute Lymphoc	tyte depletion rate (use REMM)	
- serial ALC's:2 nd	10 ³ ,hrs post-exposure 10 ³ ,hrs post-exposure	
- Estimated dose:	Gray	Reassessed: Time h:
3 Response Categor	y: Neurological <i>:</i> 1 2 3 4	
	Hematologic: 1 2 3 4	
	Dermatological: 1 2 3 4	Diagnosis: 1)
	Gastrointestinal:1234ISE CATEGORY:1234	2)3)
	e from 4 individual categories above)	Decorporating agent considered: Yes No No
Consistent biodosii	metry estimate using all 3	Disposition: home , transfer (to:), admit
methods is sugges	tive of radiation exposure at	Follow-up: RTED if:
the indicated dose		FP/ED indays (pt aware □) outpt labs □
(source: REMM□, ot	ther:)	Prescriptions
	24/7 throughout Canada):	
🗆 Health Canada: (6		
🛛 Radiation Trauma	Unit (UHN in Toronto):	see RADIATION STANDING ORDERS
	(416) 603-5800 ext 5098	Signature:timet
🗆 REAC/TS: (865) 57	76-3131, □ www.remm.nlm.gov	see continuation sheet 🗆

PLACE ID

Name	Age	M/F
Date	Time of Arrival	<u>h</u>
Physician:	Time seen	<u>h</u>

PLACE ID STICKER HERE

METER Course, v2.4 (5/09)

BODY MAPPING Form



Name	Age	M/F
Date	Time of Arrival	h
Physician:	Time seen	h

PLACE ID STICKER HERE

METER Course, v2.4 (5/09)

PHYSICIANS ORDERS

ALLERGY ALERT

No known drug allergy
 Known allergies: _____

DATE	TIME	PHYSICIAN'S SIGNED ORDERS	Initial
		□ i.v.: □NS vs □other, initial boluscc, thencc/hr	
		□ O ₂ @L/min by □NP □non-rebreather	
		Monitor: Dardiac Do2 sats	
		Labs: □CBC & manual diff q6hx4	
		Lytes, BUN, creatinine, glucose	
		Qualitative HCG (ICON)	
		□ TSH, T3, free T4	
		Tube for chromosomal analysis	
		🗆 other:	
		Specimens (note: label specimen, test with Geiger Counter, then save)	
		□ Nasal swab (L&R)	
		Skin wipe	
		🗆 Urine sample	
		Stool sample	
		Vomit sample	
		🗆 other:	
		Medications	
		🗆 Pain:	
		Nausea/vomiting:	
		Anti-diarrheal agent:	
		 Home Meds (itemize home meds including dose/route/schedule on separate page) 	
		Decorporating agent ¹ :	
		other:	
		□ see additional order sheet Signed: MD	

¹ See 'Severity Scoring Form', pages 7-8 PAGE 6

Name	Age	M/F
Date	Time of Arrival	h
Physician:	Time seen	h

PLACE ID STICKER HERE

METER Course, v2.4 (5/09)

SEVERITY SCORING Form

Time of Exposure	Based on Waselenko JK et al. Ann Internal Med 2004;140(12):1037-1051,
Time of Symptom Onset	also Fliedner TM <i>et al</i> . Oxford: British Institute of Radiology; 2001: 64pp
Time of Assessment	also refer to REMM website (<u>www.remm.nhs.gov</u>)

1. NEUROLOGICAL (Circle most appropriate description for each symptom)

Acute Symptom ²	1 (mild)	2 (moderate)	3 (severe)	4 (most severe)
Nausea	Mild	Moderate	Severe	Unbearable
Vomiting	~ 1 per day	~ 2-5 per day	~ 6-10 per day	> 10 per day
Anorexia	Mildly decreased appetite	Moderately decreased appetite	Severely decreased appetite	Unable to eat
Fatigue Syndrome	No functional impairment	Moderate functional impairment	Severe functional impairment	Unable to function
Fever	37.5-38 °C	38.1 - 40 °C	>40 °C for <24h	>40 °C for >24 h
Headache	Mild	Moderate	Severe	Unbearable
Hypotension	HR>100, BP>100/70	BP<100/70	BP <90/60 (transient)	BP <80/60 (persistent)
Neurological deficits	Minor deficit; no functional impairment	Moderate deficit; moderate functional impairment	Marked deficit; marked functional impairment	Severe deficit; loss of consciousness
Cognitive deficits	Mild cognitive impairment	Moderate cognitive impairment	Severe cognitive impairment	Profound cognitive impairment

2. HEMATOLOGIC (Circle most appropriate description for each symptom)

Acute Symptom ²	1 (mild)	2 (moderate)	3 (severe)	4 (most severe)
Abs Lymphocyte	≥1.5 x 10 ⁹ /I	1.0-1.5 x 10 ⁹ /l	0.5-1.0 x 10 ⁹ /l	<0.5 x 10 ⁹ /l
Abs Granulocyte	≥2.0 x 10 ⁹ /I	1.0-2.0 x 10 ⁹ /l	0.5 – 1.0 x 10 ⁹ /l	<0.5 x 10 ⁹ /l
Abs Platelet count	≥100 x 10 ⁹ /l	50-100 X 10 ⁹ /I	20-50 x 10 ⁹ /l	<20 x 10 ⁹ /l
Infection ³	Local; no antibiotics required	Local; topical or oral antibiotics	Systemic; oral antibiotics	Sepsis; i.v. antibiotics
Bleeding ³	Petechiae; easy bruising; normal Hgb	Mild blood loss; <10% decrease in Hgb	Gross blood loss; 10- 20% decrease in Hgb	Spontaneous bleeding; >20% decrease in Hgb

Approximate **equivalent exposure doses** corresponding to different overall Response Categories: $1 \sim 1-2$ Gy, $2 \sim 3-4$ Gy, $3 \sim 6-7$ Gy, and $4 \sim >8-10$ Gy (note: high individual variability)

² Acute symptoms are those that began after the radiation exposure, and not thought to be attributable to another acute cause ³ Only present subacutely

Name	Age	M/F
Date	Time of Arrival	h
Physician: _	Time seen	h

PLACE ID STICKER HERE

METER Course, v2.4 (5/09)

			suproving cach sympt	
Acute Symptom ²	1 (mild)	2 (moderate)	3 (severe)	4 (most severe)
Erythema	Minimal, transient	Moderate; isolated patches <10cm2; <10% of body surface area (BSA)	Marked; isolated patches or confluent; 10 -40% BSA	Severe; isolated patches or confluent; erythroderma; >40% BSA
Sensation/ itching	Occasional pruritis	Slight, intermittent pain	Moderate; persistent pain	Severe; persistent pain
Swelling / Edema	Mild; asymptomatic	Moderate; symptomatic	Severe; symptomatic	Compartment syndrome
Blistering	Vesicles, with sterile fluid	Vesicles, with haemorrhage	Bullae, with sterile fluid	Bullae, with haemorrhage
Desquamation	Mild	Patchy, dry	Patchy, moist	Confluent, moist
Ulcer/ necrosis	Epidermal only	Dermal	Subcutaneous	Muscle / bone involvement
Hair loss ³	Thinning, not striking	Patchy , visible	Extensive	Complete and most likely irreversible
Onycholysis ³	Minimal	Moderate	Severe	Complete

3. CUTANEOUS (*Circle most appropriate description for each symptom*)

4. GASTROINTESTINAL (*Circle most appropriate description for each symptom*)

Acute Symptom ²	1 (mild)	2 (moderate)	3 (severe)	4 (most severe)
Stool frequency	2 - 3 stools per day	4 - 6 stools per day	7 - 9 stools per day	>10 stools per day; intractable diarrhea
Mucosal loss with diarrhea	Rare	Intermittent, with moderate patches	Persistent, with larger patches	Continuous, with large patches
Bleeding with diarrhea	Occult	Intermittent	Persistent	Gross hemorrhage
Abdominal cramping & pain	Minimal	Tolerable	Intense	Excruciating

Decorporating agents (for use with internal contamination) ⁴ :
Cesium \rightarrow Prussian Blue (1g in 200mL of water tid x 2-3 days)
Iodine \rightarrow KI (note: dose of KI is age dependent; 50-130mg
given po)
Plutonium, Americium \rightarrow DTPA (given as Ca-DTPA initially, then
Zn-DTPA)
Uranium \rightarrow Sodium bicarbonate (250mL of 1.4% NaHCO ₃)
Tritium \rightarrow water (>6 litres/day)
Radium \rightarrow Ca-gluconate (10mL of 20% solution bid)
Strontium \rightarrow Barium sulphate (300g po single dose), Ca-

gluconate Other decorporating agents: Deferoxamine, Dimercaprol (BAL), and Penicillamine

Onset of vomiting Dose (hours after exposure) (Grays) duration 0.5-2.0 >6, or absent <24 hours 2.0-3.5 2-6 12-24 3.5-5.5 1-2 24 >5.5 Minutes 48

Time interval prior to onset of vomiting for initial biodosimetry

² Acute symptoms are those that began after the radiation exposure, and not thought to be attributable to another acute cause

³ Only present subacutely

⁴ For prescribing information and other decorporating agents, refer to REMM; for local availability refer to Disaster Plan