

Approach to the <u>CBRN Event</u>: A Framework for Patient Care & Movement



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Inside the Mind of the Terrorist

• What characteristics of CWA would be most suitable for use in an attack?

oHigh volatility

Fast & effective absorption via skin and respiratory system

 \circ Rapid onset

 \circ Lethal / incapacitating effects



Chemical-Warfare Agents (CWAs)

Class	Representative Agents	Last Known Use or Attempted Use as a CWA*
Nerve agents (cholinesterase inhibitors)	G-series (sarin, soman, cyclosarin, tabun), V-series (VE, VG, VM, VX), organo- phosphates	Syria, 2017: sarin; Malaysia, 2017: VX
Asphyxiants (blood agents)	Hydrogen cyanide, cyanogen chloride	New York City subway, 2003: cyanide
Opioid agents	Fentanyl, carfentanil, remifentanil	Moscow theater, 2002: fentanyl or carfentanil (used to subdue terrorists)
Anesthetic agents	Chloroform, halothane, nitrous oxide	No known use as CWA
Anticholinergic (antimuscarinic) agents	3-Quinuclidinyl benzilate (BZ), Agent 15 (chemi- cally the same as or related to BZ), atropine	Syria, 2012: Agent 15
Vesicant agents	Mustards (nitrogen and sulfur), lewisite, phosgene oxime	Syria and Iraq, 2016: mustard gas
Caustic agents (acids)	Hydrochloric acid, hydrofluoric acid, sulfuric acid	London, 2017: sulfuric acid
Riot-control agents	Chloroacetophenone (CN), chlorobenzy- lidenemalononitrile (CS), bromobenzyl- cyanide (CA)	Falkland Islands, 1982: "tear gas" used on British troops
Trichothecene mycotoxins	T-2 toxin	Possible use in Vietnam War, 1970: T-2
Pulmonary agents	Chlorine, phosgene, diphosgene	Syria, 2017: chlorine
Botulinum toxin	Botulinum toxin	Tokyo, 1995: botulinum toxin used by Aum Shinrikyo

Ciottone GR. Toxidrome Recognition in Chemical-Weapons Attacks. N Engl J Med. 2018 Apr 26;378(17):1611-1620. doi: 10.1056/NEJMra1705224. PMID: 29694809.

Table 1. Classes of Chemical-Warfare Agents (CWAs) Likely to Be Used in a Civilian Attack.			
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The Objective!

• The classes of chemical-warfare agents that are most rapidly lethal (i.e., nerve agents, opioid agents and asphyxiants), should be quickly identified with the use of a toxidromebased system of rapid triage

DOES A FRAMEWORK ALREADY EXIST FOR MANAGEMENT OF CASUALTIES?





Hot Zone (DTC) Warm Zone Cold Zone

Primary Objectives:

- Complete the mission
- Treat the casualty
- Avoid additional casualties

Management Objectives:

- Massive hemorrhage control
- Casualty positioning
- Auto-injector antidotes

Hot Zone

Warm Zone (ITC) Cold Zone

Management Objectives:

- Airway Management
- Respirations
- Circulation
- Head Injury
- Hypothermia

Warm Zone Cold Zone (EVAC)

Hot Zone

Management Objectives:

- Evacuation to definitive therapy
- Frequent reassessment

HOW CAN WE ADAPT THAT FRAMEWORK FOR THE CBRN ENVIRONMENT?





MARCHE²

- M Mask up
- A Antidotes
- R Rapid Spot Decontamination
- C Countermeasures

 Antidotes are challenging in the HOT ZONE
 Medical management
- H Hypothermia
- E Extrication





*DeFeo DR, Givens ML. Integrating Chemical Biological, Radiologic, and Nuclear (CBRN) Protocols Into TCCC Introduction of a Conceptual Model - TCCC + CBRN = (MARCHE)2. J Spec Oper Med. 2018 Spring;18(1):118-123. PMID: 29533446.

Hot Zone (DTC) Warm Zone Cold Zone

What are the Primary Objectives?

- Don PPE
- Move casualty to safer position
- MARCHE²

What are the Management Objectives?

- Mask up
- Stop the poisoning (spot decon)
- Toxidrome-specific antidotes (limited)

Hot Zone (DTC)



• \mathbf{M} – Mask up



- A Antidotes (toxidrome-specific)

 ATNAA/CANA
 Naloxone IN/IM
 Amyl Nitrite
- **R** Rapid Spot Decon

 Life-threatening hemorrhage controlled
 - \circ Remove visible contamination
 - \circ Apply RSDL on affected areas
 - \circ Do NOT breach clothing further in HOT ZONE

What is the Purpose of Decon?



Medical countermeasure that mitigates the conversion of an exposure to a dose



Hot Zone

Warm Zone (ITC) Cold Zone

What are the Management Objectives?

- Gross decon will occur
- MAR<u>CHE</u>²
- If an intervention can be improved via vascular access, gain access AFTER decon
- Take note of interventions performed in the HOT ZONE / WARM ZONE before decon





 TQ assessment, downgrade/conversion, decontamination

 TQ exchange is situation dependent

• Effective and necessary vs. source of ongoing exposure for the casualty



- **C** Countermeasures
 - Antidotes

 Atropine
 CANA (additional)





Airway Management

 Secure airway
 Replace interventions prior to decon



C - Countermeasures

- Respiration

 Nerve Agents
 Atropine
 - Opioids







- Naloxone (10 mg)
- Cyanide (Asphyxiants)
 - Hydroxocobalamin
- Hydrofluoric Acid (Caustics)
 - Calcium gluconate
 - Bronchodilators
- o Blister Agents
 - Bronchodilators





- **H** Hypothermia • Minimize heat loss
 - o Address "wet and naked"

 \circ Cover burns







• **E** – Extrication





Hot Zone Warm Zone Cold Zone (EVAC) What are the Management Objectives?

- Reassessment of interventions
- Enhanced monitoring
- Considerations of delayed onset presentations
- Pain management

Cold Zone (EVAC)





- Definitive airway control
- Use of oxygen
- Cardio-resp monitoring





Cold Zone (EVAC)



- Cardio-resp monitoring
- Delayed Respiratory failure



1.5 mL of a 10% calcium gluconate injection to 6 mL of sterile water in nebulizer



Cold Zone (EVAC)







- Delayed burn presentation
- Pain management



Let's Summarize

- Decon <u>Medical countermeasure</u> that mitigates the conversation of an exposure to a dose
- Toxidrome-based approach for recognition and rapid management of suspected chemical-warfare agents
- MARCHE² is dynamic
- Hot Zone $\underline{M}ask$, $\underline{A}ntidotes$, $\underline{R}apid spot decon$
- Hot Zone Antidotes ATNAA/CANA, Naloxone IN/IM, Amyl Nitrite
- Warm Zone <u>C</u>ountermeasures based on suspected toxidrome, preventing <u>Hypothermia</u>, initiating <u>E</u>xtraction
- Cold Zone Reassessment and advanced monitoring