

## NRT Quick Reference Guide: VX



GHS: Acute Toxicity, Category 1  
 H310 – Fatal in contact with skin  
 H330 – Fatal if inhaled

### 1. Agent Characteristics

#### Agent Characteristics

**Agent Classification:** Schedule 1 Chemical Warfare Nerve Agent; VX; CAS: 50782-69-9

**Description:** VX (O-ethyl S-(diisopropylaminoethyl) methyl phosphonothionate) is an odorless, oily, yellow/amber colored liquid when pure. VX, a “nerve agent,” is a lethal cholinesterase inhibitor with a mechanism of toxicity similar to organophosphate insecticides, though it is much more toxic. VX is a persistent agent with a very low vapor pressure. Aerosols or vapors of VX would be difficult to generate, maintain, and/or disperse unless sufficient heat or a dispersive device is used. VX may decompose when in contact with metals, producing highly flammable hydrogen gas. VX vapors can form explosive mixtures with air.

**Hydrolysis Breakdown Products:** VX is soluble in water, although relatively resistant to hydrolysis. Hydrolysis pathways and rates are dependent upon pH, with both somewhat acidic and alkaline conditions (pH < 6, pH > 10) predominantly forming the breakdown products methylphosphonic acid (MPA), ethyl methylphosphonic acid (EMPA), and diisopropyl ethyl mercaptoamine (DESH). MPA, EMPA, and DESH are considered to have low to moderate toxicity.

- At neutral to alkaline conditions, in pH range 7-10, the hydrolysis of VX predominately yields (S-(2-diisopropylaminoethyl) methylphosphonothioic acid), the breakdown product EA2192.
- EA2192 can also form in toxicologically significant amounts at pHs below 7 and above 10.
- EA2192 (CAS: 73207-98-4; Formula: C<sub>9</sub>H<sub>22</sub>NO<sub>2</sub>PS; Molecular Weight: 239.32 g/mol) is a white solid that is very soluble and stable in water.
- EA2192 is almost as toxic as VX itself by some exposure routes and is resistant to further hydrolysis by water. For most modeling, planning, and response needs, the physical and chemical properties of VX can be used where data for EA2192 is unavailable or missing.

**Persistence:** VX is considered a “persistent” chemical warfare agent. Vapor: hours to days; liquid: hours to months. Persistence will depend upon the amount and purity of the agent, method of release, environmental conditions, and the types of surfaces and materials impacted. Porous, permeable, organic, or polymeric materials such as carpets and vinyl tiles can accumulate VX, acting as “sinks,” thereby prolonging persistence. EA2192 will likely be formed following a VX release, and, out of an abundance of caution, it should be considered as persistent and toxic as VX, but more soluble in water and resistant to hydrolysis. These characteristics of EA2192 should be considered and could complicate the decontamination and recovery actions at a VX release site.

### 2. Physical Properties

#### Physical Properties

<b>Molecular Weight:</b> 267.37 g/mol	<b>Formula:</b> C <sub>11</sub> H <sub>26</sub> NO <sub>2</sub> PS
<b>Vapor Density:</b> 9.2 (air = 1)	<b>Flash Point:</b> 318.2°F/159°C
<b>Vapor Pressure:</b> 7.0–8.84 x 10 <sup>-4</sup> mm Hg (77°F/25°C)	<b>Liquid Density:</b> 1.008 g/mL (68°F/20°C)
<b>Volatility:</b> 10.5–12.7 mg/m <sup>3</sup> (77°F/25°C)	<b>Aqueous Solubility:</b> Slightly soluble (3 g/mL, 77°F/25°C)
<b>Boiling Point:</b> 557.6–568.4°F/292–298°C	<b>Non-aqueous Solubility:</b> Common organic solvents, alcohols, gasoline, oils, fats
<b>Melting/Freezing Point:</b> <-59.8°F/-51°C	<b>Hydrolysis (t<sub>1/2</sub>):</b> 60 hours (in water, 72°F/22°C)

**Note:** physical properties are listed at/near STP unless otherwise indicated

**Conversion Factors:** ppm = mg/m<sup>3</sup> x 0.09144; mg/m<sup>3</sup> = ppm x 10.936

### 3. Release Scenarios

#### Release Scenarios

**AIR RELEASE SCENARIOS ARE ASSUMED MOST PROBABLE; HOWEVER, OTHER RELEASE SCENARIOS AND EXPOSURE ROUTES SHOULD BE CONSIDERED.**

**Open Areas:** VX would be difficult to disperse in air as a gas due to its low volatility, but even small quantities can be lethal. It may be possible to disperse VX as a vapor/aerosol plume if sufficient heat or a dispersive device is employed; however, the size and extent of plume may be limited, posing localized hazards, which could persist from days to months depending upon environmental conditions. VX vapors, when present, are heavier than air, so vapors can accumulate in lower terrains. VX vapors can form explosive mixtures with air.

**Water/Water Systems:** If released into natural waters or water systems, VX will likely hydrolyze, with a half-life (t<sub>1/2</sub>) of 60 hours, but VX could potentially persist for weeks, depending on overall dilution and pH-dependent breakdown mechanisms. The hydrolysis breakdown product EA2192 is soluble and stable in water, posing an ingestion concern, and

may persist longer than VX itself. Therefore, both VX and EA2192 contamination may be present. If VX is released into water systems such as reservoirs, treatment plants, distribution systems, public fountains or pools, treatment processes can further break down VX. Certain water system components may act as sinks for VX, prolonging its persistence, so plumbing, surfaces, and equipment that have contacted contaminated water should be evaluated for decontamination, along with the bulk water.

**Indoor Facility:** Due to its low volatility, VX would be difficult to distribute effectively throughout a building or facility from a point source unless sufficient heat or a dispersive device is employed. Liquid VX will result in localized areas of surface contamination. VX vapors are heavier than air so vapors can accumulate in lower levels, basements, floor drains, floor HVAC air intakes, or utility corridors inside the buildings.

**Other:** VX will decompose when heated or when combusted in a fire to form highly toxic nitrogen oxides and sulfur oxides, and toxic gases, including phosphorous oxides. If VX is released into the air as a liquid spray (aerosol), it has the potential to contaminate agricultural products. If VX is released as a vapor, it is highly unlikely to contaminate agricultural products.

## 4. Health Effects

### Health Effects

**4.1. Onset:** Onset of symptoms is dose and route dependent. After exposure, symptoms may occur within seconds if VX is present in vapor form or within minutes to hours if in liquid form. Even a relatively low dose exposure to VX can be fatal and immediate administration of an antidote is critical (see First Aid below).

**4.2. Signs/Symptoms:** Initial symptoms will vary depending on dose and exposure route. The following is a general list of possible symptoms. The severity of effects depends upon the dosage. Initial effects from a sublethal amount of VX by vapor exposure are different from initial effects from a similar amount of liquid VX on skin.

**Mild/Moderate:**

- **Vapor, small dose:** Reduction in pupil size (miosis), dimness of vision, runny nose (rhinorrhea), tightness of chest (dyspnea), difficulty in breathing, headache, and salivation. *Time of onset:* seconds to minutes after exposure.
- **Liquid on skin, small to moderate dose:** Sweating or muscle twitching at site of exposure, nausea, vomiting, feeling of weakness. *Time of onset:* 10 minutes to 18 hours after exposure.

**Moderate/Severe:**

- **Vapor, large dose:** All of the above, plus sudden loss of consciousness, convulsions, temporary cessation of breathing (apnea), paralysis from reduced muscle tone (flaccid paralysis), copious nasal secretions, increased miosis (to level of pinpointing of pupils).
- **Liquid on skin, large dose:** All of the above, plus sudden loss of consciousness, convulsions, temporary cessation of breathing (apnea), paralysis from reduced muscle tone (flaccid paralysis), copious nasal secretions, diarrhea.

**Severe:** All of the above, plus severe breathing difficulty or cessation of breathing; generalized muscular twitching, weakness, or paralysis; convulsions; loss of consciousness, involuntary defecation and urination, coma, death. *Vapor, time of onset:* seconds to minutes after exposure; *Skin, time of onset:* minutes to an hour after exposure.

**4.3. Exposure Routes:**

**Inhalation:** A primary exposure route; inhalation of very small concentrations can produce health effects.

**Skin:** Direct contact with liquid agent is especially toxic. Moderate to severe signs/symptoms occur at, but are not limited to, the site of contact. Exposure can also result from dermal absorption of vapors.

**Eyes:** Eyes are the most sensitive target organs of nerve agent exposure. Miosis (reduction in pupil size) will typically be the first sign of exposure.

**Ingestion:** Contaminated drinking water and foods are the most likely route for ingestion of VX. Target organ from ingestion is the gastrointestinal tract.

## 5. Effect Levels

### Effect Levels

**5.1. Air (inhalation vapor hazard):** Acute Exposure Guideline Levels (AEGLs) for general population one-time exposure emergency scenarios for VX (complete definitions are available at: <https://www.epa.gov/aegl>).

AEGL Level in mg/m <sup>3</sup> , at various exposure durations	10 min.	30 min.	1 hr.	4 hr.	8 hr.
<b>AEGL 1:</b> Threshold mild effects	0.00057	0.00033	0.00017	0.00010	0.000071
<b>AEGL 2:</b> Potentially irreversible effects or impaired ability to escape	0.0072	0.0042	0.0029	0.0015	0.0010
<b>AEGL 3:</b> Threshold for severe effects/medical needs/increasing potential for lethality	0.029	0.015	0.010	0.0052	0.0038

American Industrial Hygiene Association (AIHA) Emergency Response Planning Guidelines (ERPG™) are not established/determined for VX.

**5.2. Dermal (liquid hazard):** Effects are likely to appear within minutes to hours of dermal exposure, with variation in latency and severity of reported health effects depending on dose, duration, and anatomical area of contact, as well as

temperature of surrounding area. Clinical manifestations range from sweating, muscle twitching, and gastrointestinal distress following limited exposure (one small drop on skin) to convulsions, paralysis, loss of consciousness, and death.

## 6. Exposure Guidelines

### Exposure Guidelines

**6.1. Airborne Exposure Limits (AELs):** CDC has issued recommendations for protecting human health from potential adverse effects of exposure to this agent.  
 (refer to: NIOSH ERSB-DB, [https://www.cdc.gov/niosh/ersbdb/emergencyresponsecard\\_29750005.html](https://www.cdc.gov/niosh/ersbdb/emergencyresponsecard_29750005.html))  
 [original source: CDC, National Center for Environmental Health (NCEH), Final Recommendations for Protecting Human Health from Potential Adverse Effects of Exposure to Agents GA (Tabun), GB (Sarin), and VX; 68 FR 58348-58351 (October 9, 2003). Cited in [USAPHC AR 40-5e, PHN No. 0711-02, July 2011](#)]

**CDC/NCEH IDLH** = 0.003 mg/m<sup>3</sup>; workers should remove themselves immediately from exposure if the concentration is reached at any point in time

**CDC/NCEH STEL** = 0.00001 mg/m<sup>3</sup> (1 × 10<sup>-5</sup>); exposure at the STEL should not exceed 15 minutes or occur more than once per day

**CDC/NCEH Worker Population Limit (WPL)** = 0.000001 mg/m<sup>3</sup> (1 × 10<sup>-6</sup>)

**CDC/NCEH General Population Limit (GPL)** = 0.0000006 mg/m<sup>3</sup> (6 × 10<sup>-7</sup>)

**6.2. Occupational:** (NA = not available)

**NIOSH IDLH** = NA

**OSHA PEL** = NA

**NIOSH REL-TWA** = NA

**ACGIH TLV-TWA** = NA

**NIOSH REL-STEL** = NA

**ACGIH TLV-STEL** = NA

**6.3. Population:**

**Soil: USAPHC Health Based Environmental Screening Levels (HBESL)** = 1.1 mg/kg over work life for Industrial Soil; 0.042 mg/kg over lifetime for Residential Soil.

**Drinking Water:** EPA Provisional Advisory Levels (PALs): see below for more information.

**EPA Provisional Advisory Level (PAL):** PALs represent chemical concentrations in air or water above which varying health effects (PAL1, PAL2, PAL3) are expected – they are developed for 24-hour, 30-day, and 90-day exposure durations. In the event of a nationally significant or large-scale chemical release, EPA can provide PALs to appropriate end-users and stakeholders as needed (PALs are not currently available to the public) to evaluate the severity of the situation, identify potential human health outcomes, and determine the most appropriate course of action; contact: [CESER@epa.gov](mailto:CESER@epa.gov), for information on and access to the PALs. (Note: PALs are not intended to define cleanup levels.)

## 7. Personnel Safety

### Personnel Safety

**Note:** Personal Protective Equipment (PPE) selection (levels A-D), medical surveillance requirements, First Aid options, and personnel decontamination may vary depending upon the amount and purity of agent, site conditions, and the release scenario. Additional information on personnel safety and PPE selection criteria can be found at: [www.cdc.gov/niosh/ersbdb](http://www.cdc.gov/niosh/ersbdb). We also recommend that responders check their own internal procedures (i.e., SOPs), if applicable.

**7.1. Medical:**

**Pre-incident:** A baseline cholinesterase activity determination and an annual physical and respiratory function exam.

**During Incident:** Conduct periodic on-site medical monitoring, observe for any signs and symptoms as per HEALTH EFFECTS section above and treat accordingly as per First Aid section below.

**Post-incident:** Post-incident medical surveillance is required.

**7.2. First Aid:** Immediately remove person from affected area and remove contaminated clothing and articles. Wash bare skin immediately with water, or warm, soapy water if available, at normal household pressures (~50-60 psi) for three minutes, ensure thorough soaking. Rinse eyes exposed to liquid agent with potable water for 15 minutes.

**ANTIDOTE: Atropine, 2-PAM Chloride injections (Duo Dote/Mark II kits)/ATNAA-Antidote Treatment Nerve Agent Autoinjector.** Antidote kit should only be administered as per pre-incident training. Recommended dosages of atropine and 2-PAM Chloride injections will vary depending on patient age and physical findings (mild/moderate or severe). Follow manufacturer's directions and adhere to all SOPs or recommendations developed by individual responder's agency.

**Other:** RSDL (Reactive Skin Decontamination Lotion), an FDA-cleared kit with a sponge impregnated with a lotion to remove or neutralize chemical warfare agents from contaminated skin. Apply RSDL immediately to area of skin with suspected exposure to a chemical warfare agent (do not wait for symptoms) and wipe affected area using a scrubbing action, rinse with water when time permits.

After administering first aid, send person for follow-up medical attention and evaluation. If cleared to resume work, continue to monitor for signs/symptoms and treat accordingly.

### 7.3. Personal Protective Equipment (PPE):

**GENERAL INFORMATION:** NIOSH-certified Chemical, Biological, Radiological, Nuclear (CBRN) Self Contained Breathing Apparatus (SCBA), NIOSH-approved Air Purifying Respirators (APR) or Powered Air Purifying Respirators (PAPR), full-face masks, and protective clothing should be used. Level A protection should be used until monitoring results confirm identity and concentration of contaminant. Pre-incident training and exercises on the proper use of PPE are recommended.

Per NIOSH guidance –

**LEVEL A:** Recommended for the initial response to a VX incident. NIOSH-certified CBRN full-face-piece SCBA operated in pressure-demand mode with Level A suit that provides protection against CBRN agents. Level A provides the greatest level of skin (totally-encapsulating chemical protective suit and chemical-resistant inner and outer gloves, along with chemical-resistant boots with steel toe and shank), respiratory (SCBA), and eye protection when the contaminant identity or concentration is unknown. Select Level A when the VX concentration is unknown or above the IDLH or AEGL-2, and when there is a potential of ocular or dermal exposure.

**LEVEL B:** Pressure-demand SCBA (NIOSH-certified CBRN full-face-piece SCBA) with Level B protective suit that provides protection against CBRN agents. Level B provides the highest level of respiratory protection (SCBA) when a lesser level of skin protection is required. Select Level B when the VX concentration is unknown or above the IDLH or AEGL-2, and when dermal exposure is less of a risk. Level B differs from Level A in that it typically incorporates a non-encapsulating, splash-protective, chemical-resistant outer suit that provides protection against most liquids but is not vapor tight (hooded chemical-resistant outer suit and chemical-resistant inner and outer gloves, along with chemical-resistant boots with steel toe and shank).

**LEVEL C:** May be selected when the contaminant identity and concentration are known and the respiratory protection criteria factors for the use of APR or PAPR (i.e., < IDLH, warning properties) are met. Level C may be appropriate when decontaminating personnel or equipment. Level C still incorporates hooded chemical-resistant outer suit that provides protection against CBRN agents, chemical-resistant inner and outer gloves, and chemical-resistant boots with steel toe and shank.

- For air levels greater than AEGL-2: NIOSH-certified CBRN tight-fitting PAPR with a filter or a combination organic vapor, acid gas, and particulate cartridge/filter combination or a continuous flow respirator.
- For air levels greater than AEGL-1: NIOSH-certified CBRN tight-fitting APR with a canister-type gas mask or CBRN PAPR.

**LEVEL D:** Select Level D when the contaminant is known and the concentration is below the appropriate occupational exposure limits or less than AEGL-1 for the stated duration times. PPE includes coveralls or other work clothes, boots, and gloves.

**Downgrading PPE levels can be considered only when the identity and concentration of the contaminant and the risks of dermal exposure are known, and must be accompanied by on-site monitoring.**

## 8. Personnel Decontamination

### Personnel Decontamination

#### 8.1. Personnel Decontamination Procedure:

Tents, berms, and collection vessels should be able to maintain copious amounts of wastewater in a contained and safe manner. Procedures should be in place to treat and replace contaminated materials used during the decontamination process as well as replace necessary chemicals and decontamination solutions.

Prior to entering the hot zone, all personnel are required to familiarize themselves with the site-specific personnel decontamination procedures.

Personnel decontamination should take place in a decontamination area comprised of two decontamination corridors (one for entering and one for exiting). Position corridors upwind and uphill of release area; exit should be upwind and uphill of entrance. Detergent and water solution (pH>8, but <10.5), soft brushes, and durable 6-mil polyethylene bags should be provided.

Personnel decontamination area workers need to wear appropriate PPE as indicated below. Be aware that absorbed agent can be released from clothing and skin as a gas.

Conduct personnel decontamination per NIOSH ERSB-DB:

[https://www.cdc.gov/niosh/ersbdb/emergencyresponsecard\\_29750005.html](https://www.cdc.gov/niosh/ersbdb/emergencyresponsecard_29750005.html)

- **Emergency Responders:** Use soft brush to wash PPE with soap and detergent solution in a downward motion, getting into all folds. Repeat washing and rinsing until thoroughly clean. Remove PPE by rolling downward from head; avoid pulling PPE over the head. Remove SCBA last, and place all PPE in polyethylene bags.
- **Patient/victim:** Remove all clothing down to at least undergarments, and place in polyethylene bags. Thoroughly wash and rinse skin with soap and water solution, taking care not to break the skin and covering all open wounds. Cover patient/victim (e.g., blanket, towels, Tyvek) and move to treatment area. If available in decontamination kit, apply RSDL immediately to area of skin with suspected exposure to a chemical warfare agent (do not wait for symptoms) and wipe affected area using a scrubbing action, rinse with water when time permits.

**8.2. Personnel Decontamination Procedures by Zone/Step:** (attendants will verbally direct personnel through each step)

**Conducted in Hot Zone (exclusion zone)**

1	Equipment Drop	Place equipment taken into the Hot Zone on a plastic covered table or container provided prior to entering the contamination reduction corridor. Equipment will either be reused if more than one entry is planned or will be decontaminated later.
---	----------------	-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------

**Conducted in Warm Zone (contamination reduction zone)**

2	Sample Drop	Place samples in a container provided for sample decontamination. Care needs to be taken to ensure that workers maintain chain-of-custody of samples. It is recommended that samples are decontaminated in a separate decontamination line.
3	Outer Boot and Glove Wash	The purpose of this step is to enable physical removal of gross contamination if contamination is visible. If gross contamination is not visible, this step may be skipped. Wash outer boots and then outer gloves using designated decontaminating agents as specified in HASP (e.g., soap and water, trisodium phosphate substitute, or diluted bleach).
4	Glove, Boot, and Suit Wash	Wash all outer surfaces in a contained area (e.g., kiddie pool) using a pressurized spray with designated decontamination solution. Start with decontaminating boots and gloves, then work on suit from the top down, including SCBA/PAPR casing. Decontamination personnel should conduct this step. Care should be taken to ensure that all areas are decontaminated, including around zipper, arms, front torso, and any other area that could have come in contact with contamination. The solution used for decontamination should be contained, collected, and disposed of properly from the decontamination line.
5	Outer Glove, Boot, and Suit Removal	While sitting on a stool, remove outer boots and outer gloves. Undo the SCBA/PAPR belt and hold in hand. While touching only the inside of suit, carefully roll suit in an outward motion from shoulders down to feet. Dispose of boots, gloves, and suit in a designated container. This step may require decontamination personnel to assist either by holding SCBA/PAPR unit or assisting in suit removal.
6	Mask Removal	With inner gloves, remove the mask. Remove cartridge filters and place into designated container. Put mask into mask wash. Decontamination personnel will clean each mask and SCBA/PAPR assembly prior to return to service.
7	Inner Glove Removal	Remove inner gloves by only touching outside of first glove and then only inside of second glove. Place gloves into designated container.

**Conducted in Cold Zone (support zone)**

8	Quadrant Monitoring	Using appropriate VX air monitoring equipment, screen personnel for residual contamination by dividing body into 4 sections: upper right and left sides of the body, and lower left and right sides. If positive, perform spot decontamination immediately and direct person to showers.
9	Personal Shower	Personnel should shower using copious quantities of soap and water for a minimum of 5 minutes and change into clean clothes. If a personal shower is not immediately available then, at the minimum, hands and face should be washed thoroughly.
10	Medical Monitoring	Report to medical monitoring station for post-entry monitoring and report to appropriate personnel for debriefing. Observe for any obvious sign of VX exposure. Using criteria listed above in PERSONNEL SAFETY section of this QRG, administer Duo Dote antidote and notify site Health and Safety officer.

**Emergency Egress Corridor:** Establish an emergency egress line to use for quickly decontaminating personnel who have medical emergencies while in the hot zone. Personnel must be decontaminated prior to receiving treatment from emergency medical technicians or transported to a hospital.

**Hand-Wash Station:** A hand-wash station should be available for personnel to clean up following entry. However, this may not be available initially at the scene or weather conditions may prohibit their use. If a hand-wash station is not available, personnel should wash their hands and face as soon as possible.

**Caution:** Avoid waterless hand cleaners, which contain solvents (alcohols) that could increase risk of dermal exposure to VX.

**9. Field Detection**

**Field Detection**

**Real-time field screening tools (results not confirmatory or quantitative):** Caution should be given to equipment that has not been properly evaluated. False positive and false negatives may occur in the presence of interferents common in the environment. The following is a summary of minimum screening concentration ranges or levels for equipment procured by many EPA and HAZMAT response teams. Other screening tools may be used by these teams and other agencies and responders, some with similar capabilities and limitations.

**9.1. Minimum Screening Ranges/Levels for Air/(Vapor):**

Field Equipment:	ppm	mg/m <sup>3</sup>
JCAD M4A1, at 30 secs [30 mins/pre-concentrator] (detected as Nerve agent)	0.01 [0.0001]	0.1 [0.001]
AP2C / AP4C (detected as V-agent)	0.001 / 0.001	0.01 / 0.01
MX-908 Vapor Mode	0.10	1.1
Dräger (CDS Kit)	0.025	0.27
MINICAMS™ (Near real-time; at 15 minutes)	0.000001	0.00001

**9.2. Minimum Screening Ranges/Levels for Vapor/Liquid:**

Field Equipment:	ppm (vapor)	mg/m <sup>3</sup> (vapor)	mL (liquid)
M256 / M256A1 (13-15 mins) (detected as Nerve agent)	0.01 / 0.002	0.1 / 0.02	NA
M8 (absence/presence as Nerve V-agent)	NA	NA	0.020

Note: M256 is combined 2 kits with 12 disposable sampler/detectors and a booklet of M8 paper.  
 NA = not applicable

**9.3. Minimum Screening Ranges/Levels for Water:**

Field Equipment:	mg/L
M272 (detected as V-agent)	0.02

**10. Environmental Sampling**

**Environmental Sampling**

*Note: This section on sampling contains general guidelines and does not replace the need for a site-specific sampling plan*

**10.1. Sampling Concerns:** Detection, sampling equipment and procedures, and analytical techniques will be site-specific and depend on: 1) physical state of the agent; 2) type of surfaces contaminated (e.g., porous vs. non-porous); 3) the purpose of sampling (e.g., characterization, decontamination efficacy, and clearance); and 4) specific laboratory requirements. The U.S. Environmental Protection Agency (EPA) has set up mobile and fixed labs and analytical assets for chemical agent analysis of environmental samples under their Environmental Response Laboratory Network (ERLN), see ANALYSIS section below, (<https://www.epa.gov/emergency-response/environmental-response-laboratory-network>). For questions on environmental sampling for VX and its toxic breakdown products, call EPA/HQ-EOC at 202-564-3850.

**10.2. Sample Locations and Planning:** Initially consider air monitoring to ensure worker safety and to determine if there is a vapor plume that could impact other areas. Characterization sampling is initiated by targeted or judgmental sampling to identify “hot spots,” potential agent flow paths, and media or objects potentially acting as sinks. Additional biased or random sampling can be used to determine the extent of potential contamination or to verify the efficacy of decontamination. More thorough probabilistic sampling (e.g., grid, statistical approach) may be required for the clearance phase or if there are large uncertainties about the area impacted or the amount released. Because VX is a persistent agent, sample priorities should include surfaces that are potentially contaminated with liquid, or precipitated fumes or aerosols (e.g., release site, low lying areas, HVAC, utility corridors) and areas that people are likely to contact or where food or agricultural products are present.

**Note:** VX breaks down in most environmental conditions to numerous breakdown products, especially EA2192, methylphosphonic acid (MPA), and ethyl methylphosphonic acid (EMPA), which may be used as markers to determine the extent of contamination of the parent VX. See ANALYSIS section below to ensure sampling procedures are compatible with all analytes. Some preparation techniques for both VX or its breakdown product EA2192 and other breakdown products are available in EPA’s Sample Information Collection Documents (<https://www.epa.gov/esam/sample-collection-information-documents-scids>). These provide general information regarding sampling procedures for different media, sampling supplies, sample size, container, holding time, preservation, packaging, and shipping, supporting collection of samples. Environmental sampling and analytical methods for EA2192 are limited, as such, methods used for VX can be used as substitutes when methods for EA2192 are missing. **Caution:** Pre-incident training on administration of antidote kit is recommended (see First Aid under PERSONNEL SAFETY section above).

**10.3. Types of Samples:**

**Air** (Vapors are heavier than air): Samples are collected using appropriate solid phase absorbent media (tubes) at breathing zone level (~5 ft.) to assess inhalation exposure. To assess off gassing from surfaces and at ground levels, collect air samples at ~6 in. Concurrent air monitoring for VX is recommended.

**Water:** Water should be collected in appropriate containers with addition of appropriate de-chlorinating agents and preservatives to minimize VX degradation and hydrolysis prior to analysis. VX is expected to dissipate via breakdown, hydrolysis, and dilution. To rule out EA2192 contamination concerns, analyses should include EA2192 if possible. Concurrent air monitoring for VX is recommended.

**Soil:** For localized “hot spot” areas where soil deposition may occur (i.e., neat liquid, aerosol or liquid droplets), surface soil samples should be taken from a non-vegetated area to a depth of less than one inch. Sub-surface soil samples are typically not necessary unless a large amount of liquid was poured on the ground, or if an underlying aquifer is endangered. Since most soils contain water, VX may undergo hydrolysis to form EA2192, as such, soil sampling analyses should include testing for the presence of EA2192. Concurrent air monitoring for VX is recommended.

**Surface Wipes:** Wipe samples are often desired to indicate absence of VX and its toxic breakdown products (EA2192) on non-porous surfaces. Concurrent air monitoring for VX is recommended.

**Bulk:** For hot spot areas where liquid VX deposition may occur on porous surfaces (e.g., concrete, asphalt), actual pieces (chips) or cores of contaminated surface may be obtained using appropriate tools (scabbling, coring, or drills) for subsequent laboratory extraction analysis. Bulk samples of suspected sink materials may be recommended to rule out secondary vapor phase disposition or absorption of VX into these materials. Concurrent air monitoring for VX is recommended.

**Other Sample Matrices:** Contact EPA/HQ-EOC at 202-564-3850 for sampling instructions.

## 11. Packaging/Shipping: CWA Environmental Samples for Site Characterization

### Packaging and Shipping: CWA Environmental Samples For Site Characterization

The packaging and shipping of environmental samples potentially contaminated with a chemical warfare agent (CWA) would be subject to complex and restrictive regulations established primarily by DOT for ground transportation (49 CFR Parts 171-180), and by DOT, ICAO, and IATA for air transportation (in addition to other regulations by CDC, USPS, OSHA). Transportation of VX-contaminated waste for treatment and disposal is covered under the WASTE MANAGEMENT section below.

Samples can be collected from environmental media that include surface and subsurface soil, groundwater, surface water, drinking water, dust, air, and solids other than soil (e.g., building materials). Given the wide range of potential environmental media and complex regulatory requirements, the approach would likely be situationally dependent.

**CAUTION:** Environmental samples potentially contaminated with CWA should not be introduced into commercial transportation as an undeclared hazardous material. Hazard classification, packaging, and hazard communication are the shipper's responsibility under DOT's Hazardous Materials Regulations (49 CFR Parts 171-180).

A summary of key packaging and shipping considerations for environmental samples with unknown concentrations of a potential unknown CWA is:

- Transport of pure VX is forbidden other than via military (Technical Escort Unit) transport in accordance with 49 CFR §173.7.
- If the collected sample contains or is suspected to contain hazardous materials, as defined in 49 CFR §171.8, the shipper must determine the appropriate UN ID Number, the Proper Shipping Name (PSN), and the Packing Group (PG) from the Hazardous Materials Table in 49 CFR §107.101. The table will then direct the shipper to the type of hazard and handling labels needed, the appropriate packaging (inner and outer packaging), and any special provisions.
- The designated shipper (EPA personnel or contractors) must be trained and certified according to the requirements found in 49 CFR §172.704 (a)(2) and/or by IATA Dangerous Goods (DG) 1.5 requirements for shipments by air.
- Contact the sample-receiving laboratory to determine if they have additional packaging, shipping, or labeling requirements.

Note that there is no UN ID for VX listed in the Hazardous Materials Table (49 CFR §172.101). Therefore, the most likely classification would be UN3381, Toxic by inhalation liquid, n.o.s. with an LC50 lower than or equal to 200 mL/m<sup>3</sup> and saturated vapor concentration greater than or equal to 500 LC50, PG I. In the US, non-bulk packaging would then be in accordance with 49 CFR §173.226.

**Use of Mobile labs:** Another consideration would be use of an on-site mobile laboratory for CWA analysis. This could eliminate the shipper's responsibility for transporting the collected samples containing a substance that might be considered forbidden for transport by air or a hazardous material or DG by ground or air transport to an off-site laboratory. In addition, there may be public concern about shipping samples off-site, or reluctance of commercial shipping companies to accept and transport samples from a known CWA-contaminated site. EPA maintains mobile laboratory assets (PHILIS mobile laboratories: <https://www.epa.gov/emergency-response/philis-portable-high-throughput-integrated-laboratory-identification-system>) in NJ and CO that are capable of analyzing CWAs, including VX, in environmental matrices, down to health-based risk clearance levels. Access to the PHILIS mobile labs for a CWA incident can be obtained from EPA HQ/EOC at 202-564-3850. EPA also has access to the US Army CBRNE assets, including shipping and analysis, through inter-agency agreements as described in the COORDINATION WITH OTHER AGENCIES section below.

## 12. Analysis

### Analysis

**CAUTION: Many labs may not be able to perform analysis on all matrices (e.g., wipes and soil).** Few laboratories currently have the capability to determine VX or its most toxic breakdown product EA2192, particularly for large numbers of samples and for the various types of environmental media. Environmental analytical standards and methods for EA2192 are limited, as such, methods used for VX can be used as substitutes when methods for EA2192 are missing. EPA's ERLN labs (<https://www.epa.gov/emergency-response/environmental-response-laboratory-network>) that are specially trained and equipped for the analysis of VX, will use sample prep and analytical methods from EPA's Environmental Sampling and Analytical Methods (ESAM) Programs (<https://www.epa.gov/esam>). For access to the nearest ERLN laboratory specially trained and equipped for VX analysis, and methods provided in EPA's ESAM, contact EPA/HQ-EOC at 202-564-3850. The ERLN also maintains EPA's Compendium of Environmental Testing Laboratories (CETL), a database of commercial, federal, state, and academic laboratories, which can be queried for specific analyses and matrices. Analysis on environmental matrices for toxic organics, metals, biological and radiological agents, as well as several of the CWAs, including VX and its breakdown products, can be obtained by querying the database of laboratories listed in EPA's CETL (<https://cfext.epa.gov/cetl/lblogin.cfm?action=None>); prior registration for access to CETL website is necessary.

## 13. Coordination with Other Agencies: CWA Field Activities

### Coordination With Other Agencies: CWA Field Activities

Numerous agencies other than EPA may be involved in a chemical agent response incident. Every attempt should be made to integrate assets and design a uniform approach to sampling procedures, quality assurance, and data sharing. Every attempt should be made to coordinate activities, share data, and maintain chain-of-custody integrity throughout all phases of the response, amongst all agencies involved.

**Civilian:** The National Guard Civil Support Team (CST) and the U.S. Coast Guard “Strike Teams” deploy survey teams, response vehicles, and mobile labs to hazardous chemical incidents throughout the United States. Many CSTs and Strike Teams have the capabilities to sample, prepare, and analyze certain types of environmental samples for CWA analysis. CSTs have analytical equipment that can provide screening or presumptive data results for CWAs. The OSC should discuss site-specific types of samples, data quality, and chain-of-custody requirements with Strike Teams and CSTs before integrating their capabilities into the overall CWA response. Other agencies, such as the FBI, may be present on-site performing tasks, such as evidence retrieval, which are specific for their agency.

**Military:** EPA’s Special Teams (ERT and CMAD) have access to Department of Defense (DOD) assets through Inter-Agency Agreements (IAA) with the US Army’s Combat Capabilities Development Command, Chemical Biological Center (CBC) at Aberdeen Proving Ground, MD. The CBC has expertise and deployable assets for CWA air monitoring (i.e., MINICAMSTM), sampling, analysis, decontamination, and waste transport and disposal through their ongoing demilitarization activities at former chemical agent munitions facilities in the US and abroad.

Access to EPA’s IAA for support to Federal OSCs at a CBRN response or incident can be arranged through EPA/HQ-EOC at 202-564-3850. Consultation or training for EPA personnel and partners that do not need to go through EPA/HQ-EOC can be arranged via the IAA with EPA’s ERT-Special Team at 732-321-6660. Additional CBRNE support can be obtained via the IAA with EPA’s CMAD-Special Team, including support for chemical, biological, and radiological agent response through EPA/HQ-EOC at 202-564-3850.

## 14. Environmental Decontamination/Cleanup

### Environmental Decontamination/Cleanup

#### 14.1. Decontamination/Cleanup Planning:

Once site controls are in place, develop a site-specific decontamination/cleanup plan. Environmental decontamination may require a “tiered approach” using a variety of techniques and products. Call EPA/HQ-EOC at 202-564-3850 for more information.

**General Considerations:** A cost vs. benefit evaluation should be undertaken for each decontamination strategy and approach that considers public safety, total cost, impact on the area, wastes generated, time the area or item will be inaccessible and/or out of service, as well as any socio-economic, public health, and/or security impacts that may result. Large volumes of decontamination wastes may be generated that will need to be collected, treated, and disposed of properly. Waste handling and disposal must be addressed as early in the decontamination and cleanup process as possible (see WASTE MANAGEMENT section below). Decontamination/cleanup wastes may contain EA2192, which may require further treatment, **so it is important to thoughtfully apply decontamination strategies that limit EA2192 formation to avoid unintended consequences of inappropriate decontamination. Any site with a suspected VX release that has been sprayed with water may contain EA2192.**

**Disposal Option:** The urgency to restore an area or item as quickly as possible may result in the outright and timely removal and disposal of contaminated materials. Certain materials may be impacted by the decontamination products, and/or may be cheaper to discard and replace than to decontaminate and restore.

**Monitored Natural Attenuation:** VX degrades slowly (see AGENT CHARACTERISTICS section above) via natural processes. Environmental monitoring must be maintained during decontamination and recovery phases. Monitored natural attenuation may require institutional controls (e.g., access restriction and contaminant containment measures). The time to achieve clearance must be considered in the overall cost/benefit evaluation. This option is more passive than other options but is non-destructive to materials. Depending on the initial level of contamination, VX amounts remaining on surfaces may not reach expected cleanup goals for weeks or months. The presence of EA2192 should also be considered and addressed, as its environmental fate and transport may differ from VX.

**Fix-in-Place Option:** The contaminated area may be resistant to decontamination products or may be unable or impractical to be treated. Physical barriers can be used to immobilize the contamination and prevent it from coming into contact with the environment or the public. This can be a temporary or permanent solution.

#### 14.2. Decontamination Strategy:

A decontamination strategy can be developed by designating contaminated areas into five broad categories: 1) surfaces or hot spots, 2) large volumetric spaces, 3) sensitive equipment or items, 4) aqueous solutions, and 5) water systems. Areas in each category may be treated using one or more unique decontamination processes in a tiered approach to the overall site-specific decontamination strategy.

#### Cautions:

- VX may hydrolyze and form the toxic breakdown product EA2192, with greatest yields between pH levels 7 and 10 but toxicologically significant amounts may be formed outside this range, especially at higher pH. Presence of



EA2192 may pose significant challenges in verifying site decontamination and in waste management. While it is possible to destroy EA2192 with high concentrations of oxidants, this can be a slow process and may require verification, so it is advisable to limit the formation of EA2192. Decontamination solutions containing activated peroxide, which decrease the relative formation rates of EA2192 compared to non-toxic byproducts, may help limit EA2192 formation.

- Decontamination products may have unique safety/PPE requirements due to their own toxicity or that of breakdown products during use (e.g., use of bleach results in release of chlorine vapors). Strong oxidizers, such as hypochlorite, may react violently with organics. Decontamination products should be chosen that minimize the formation of toxic EA2192.
- Dirt, grime, and other coatings (organic load) can reduce the efficacy of decontamination; pre-cleaning surfaces with soap and water may be needed before the application of decontamination formulations but resulting pre-cleaning rinsates require containment to avoid agent and toxic breakdown products (EA2192) spread.

For additional information, contact the EPA/HQ-EOC at 202-564-3850.

**Surfaces/Hot Spots:** This category is for areas smaller in size but with higher levels of agent contamination. They may require more rigorous decontamination products and methods. Excess VX liquid should be absorbed using, e.g., vermiculite or dry sand, and transferred into a sealed container and disposed according to WASTE MANAGEMENT section below. Application of the following decontamination solutions and formulations may be efficacious by following applicable manufacturers' directions.

- 1) Hypochlorite-containing solutions: Hypochlorite can be corrosive to certain surfaces and materials and should be rinsed thoroughly afterwards. Household bleach solutions ( $\geq 5\%$  sodium hypochlorite) are effective for VX with efficacy achieved with contact time of 15-60 minutes depending on surface material. Diluted bleach (10x) has been shown to yield lower efficacy against VX and also shown to increase formation of EA2192 and should be avoided. Calcium hypochlorite, present in commercial products, such as HTH (10% hypochlorite solution), is better for surfaces with high concentrations of liquids in localized areas. Note that lowering the pH of hypochlorite solutions is not required and may be counterproductive. Site-specific conditions may alter the pH into a range where EA2192 formation is a concern.
- 2) Proprietary decontamination technologies such as EasyDecon DF-200®, Decon7 (D7), Dahlgren Decon®, CASCAD®, Decon Green®, or L-Gel® have been shown to be effective against VX on the order of minutes to hours, but not all have been thoroughly tested. Availability, cost, and the need for specialized equipment to apply the decontaminant may limit their use early in the response. Most of these products contain peroxide or peroxyacetic acid as active ingredients, which helps in VX destruction versus EA2192 formation, although site-specific conditions may require investigation of EA2192 formation.

**Large Volumetric Spaces:** This category is for areas larger in size but with lower levels of agent contamination. These areas may require less aggressive but more broadly applied decontamination products and methods.

- 1) Forced or Hot Air ventilation methods are recommended for vapor plume contamination or low surface concentration of VX in large volumetric spaces or open areas; efficacy may be typically achieved in days to weeks with less waste and adverse impacts to materials. HVAC systems in large indoor spaces may require a separate decontamination strategy that could include the use of Hot Air ventilation or fumigation. The low volatility of VX may necessitate high operating temperatures. Capture technologies, such as activated carbon containing air filters, would be required to prevent transfer of the VX vapor to the outside environment or prevent recirculation into other surrounding spaces.
- 2) Fumigation with modified vaporous hydrogen peroxide (mVHP®; a combination of ammonia and hydrogen peroxide vapor) has been reported to be effective against VX. In this case, the peroxide directly oxidizes VX, and unless water is present, EA2192 formation by hydrolysis is unlikely. Prolonged hydrogen peroxide fumigation (up to six days) at lower vapor concentrations ( $< 25$  ppm) was found to be non-efficacious.
- 3) Fumigation with chlorine dioxide ( $\text{ClO}_2$ ) has been reported to be effective against VX on surfaces. However, this research did not address the formation of EA2192, which may occur in the presence of expected high relative humidity ( $\geq 70\%$ ) conditions during chlorine dioxide fumigation.
- 4) Steam application has been reported to be effective against VX on surfaces. Steam also transfers VX into the condensate. The high-water vapor content during steam application at neutral pH may favor VX hydrolysis into EA2192, which was not assessed in this research.

**Sensitive Equipment or Items:**

- 1) Forced or Hot Air ventilation may be used for VX and can be used either in-situ or ex-situ to decontaminate these items. The low volatility of VX may necessitate high operating temperatures. Collective protection using activated carbon air filters would be required to prevent transfer of the VX vapor to the outside environment or prevent recirculation into other surrounding spaces.
- 2) Modified VHP® fumigation can be used on these items with less corrosion to electronics than hypochlorite solutions. As noted above, the peroxide directly oxidizes VX.

**Aqueous Solutions:** VX degrades via hydrolysis but may persist in aqueous solutions or form EA2192, depending on initial concentration and environmental conditions, such as pH and temperature. Aqueous solutions that result from decontamination operations should be adjusted for pH to minimize the formation of EA2192. Avoid any additional release to water systems, drains, or sewers (e.g., through inappropriate disposal). Contain liquid or transfer liquid to appropriate containers and dispose according to WASTE MANAGEMENT section below.

**Water Systems:** Hydrolysis and removal of contaminated water will lessen VX and EA2192 contamination in water systems, but VX may persist in hydraulic dead ends and via sorption to system components (e.g., plastics) that act as sinks for VX. It may be necessary to isolate potentially affected portions of the system to evaluate them and implement decontamination. A contaminated water system may transfer VX to building and premise plumbing, which then may also require decontamination. EA2192 is much more soluble than VX so may have different mobility in water systems.

**Verification of Decontamination:** Site and situation specific. Please contact EPA/HQ-EOC at 202-564-3850 for further assistance.

## 15. Waste Management

### Waste Management

#### 15.1. Transportation:

Federal requirements for the commercial transport of hazardous materials and procedures for exemptions are specified in How to Comply with Federal Hazardous Materials Regulations, available at:

<https://www.fmcsa.dot.gov/regulations/hazardous-materials/how-comply-federal-hazardous-materials-regulations>.

VX should not be offered for commercial transportation without being rendered safe by neutralization. Contact the PHMSA Hazardous Materials Information Center at 1-800-467-4922 or [infocntr@dot.gov](mailto:infocntr@dot.gov) to discuss specific cases. Additional resources on packaging, labeling, and shipping are available at: <https://www.phmsa.dot.gov/standards-rulemaking/hazmat/hazardous-materials-regulations>. Detailed state regulations can be found at [www.envcap.org/](http://www.envcap.org/).

This QRG is intended to apply to Federal OSCs in the first 24-48 hours of a response. Once determined, the concentrations of VX in individual waste streams should be used to determine which transportation requirements apply. For instance, certain requirements may apply to waste streams with concentrated agent, but may not apply to waste streams such as soil containing dilute concentrations of agent.

#### 15.2. Waste Management:

Under the Resource Conservation and Recovery Act (RCRA), waste is classified as hazardous waste (subtitle C) or solid waste (subtitle D). The RCRA regulations generally define a waste to be hazardous if it is: (1) a listed waste (40 CFR §261.31-§261.32); (2) exhibits specific characteristics (40 CFR §261.21-§261.24); or (3) is a discarded commercial chemical product, off specification species, container residue, or spill residue listed in 40 CFR §261.33. VX is not listed under 40 CFR §261.31-33 and VX-contaminated waste is generally not considered a RCRA characteristic hazardous waste (as defined in 40 CFR Part 261 subpart C) under 40 CFR §261.21-24. It is the responsibility of the waste generator to make a hazardous waste determination (40 CFR §262.11).

The states (except for Alaska and Iowa) have the primary responsibility to implement the hazardous waste regulations and can impose more stringent requirements or requirements broader in scope than the federal program. Several states, including CO, IN, KY, MD, OR, and UT, have their own waste designations for chemical agents, which may be applicable for the cleanup of VX-contaminated residues, decomposition products, soils, and debris. It is critical to open a dialogue with state regulators as early as possible.

Management of toxic decomposition products, associated residual decontamination solutions, local waste acceptance criteria, and transportation and handling requirements should be considered. High pH aqueous decontamination solution waste may be considered corrosive hazardous waste, chemical code D002, if it has a pH greater than or equal to 12.5 (40 CFR §261.22).

EPA/CMAD can provide Federal OSCs with information and support to address knowledge gaps for dealing with wastes contaminated with dilute concentrations of CWA; contact EPA/HQ-EOC at 202-564-3850.

EPA also recommends the creation of pre-incident waste management plans as a preparedness measure for chemical agent releases, and has created an **All-Hazards Waste Management Planning Tool** to help state, local, territorial, and tribal waste management officials coordinate and prepare these plans. Access to the All-Hazards Waste Management Planning Tool requires pre-registration (<https://wasteplan.epa.gov/>).