

Agent Characteristics

Agent Classification: Schedule 1 Chemical Warfare Nerve Agent; Sarin (GB); CAS: 107-44-8; Formula: C₄H₁₀FO₂P; Molecular Weight: 140.10 g/mol.

Description: GB (isopropyl methylphosphonofluoridate) is a colorless and odorless liquid when pure; brown liquid with a fruity odor in impure form. GB is a lethal cholinesterase inhibitor with a mechanism of toxicity similar to organophosphate insecticides, though it is much more potent. GB is more easily synthesized and more volatile than chemical warfare agents GA, GD, GF, HD, Lewisite, and VX. Environmental breakdown products of GB, including methylphosphonic acid (MPA) and isopropyl methylphosphonic acid (IMPA), are relatively non-toxic. Other breakdown products include fluoride ion, which may exist as hydrofluoric acid (HF) depending on the pH. GB can react violently with strong oxidizers and may decompose when in contact with metals, producing highly flammable hydrogen gas. GB vapors can form explosive mixtures with air.

Persistence: GB is considered a “very low persistent” chemical warfare agent. Vapor: minutes to hours; liquid: 2-24 hours. 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 GB vapors and liquids, acting as “sinks,” thereby prolonging persistence.

Physical Properties

Physical properties are listed at/near STP unless otherwise indicated. Conversion Factors: ppm = mg/m³ x 0.1745; mg/m³ = ppm x 5.730.

Vapor Density: 4.9 (air = 1)	Boiling Point: 297°F/147°C	Liquid Density: 1.09 g/mL (77°F/25°C)
Vapor Pressure: 2.9 mm Hg (77°F/25°C)	Melting/Freezing Point: -71°F/-57°C	Aqueous Solubility: miscible
Volatility: 22,000 mg/m ³ (77°F/25°C)	Flash Point: >280°F/138°C	Non-aqueous Solubility: Common solvents, alcohols, gasoline, oils, fats

Release Scenarios

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

Open Areas: GB has high volatility relative to other nerve agents but may still be present as a liquid or aerosol, and the primary release/attack scenario is an airborne release. GB is expected to degrade in the environment fairly rapidly; however, liquid GB on surfaces could persist for up to 24 hours or more.

Environmental conditions will affect the degradation and evaporation rates of GB with cooler conditions enhancing persistence. GB vapors are heavier than air, so vapors can accumulate in lower terrains. GB vapors can form explosive mixtures with air.

Water/Water Systems: If released into natural waters or water systems, GB will hydrolyze with a half-life of about 39 hours at pH 7, with persistence depending on released amount and environmental conditions. Hydrolysis byproducts have comparatively low toxicity so may only be of interest if large amounts of GB are released. Certain water system components may act as sinks for GB, prolonging its persistence long past the initial release.

Indoor Facility: Due to its volatility, GB could potentially be dispersed as a vapor or an aerosol inside a building or facility; HVAC systems could be impacted. GB vapors are heavier than air so vapors can accumulate in lower levels, basements, floor drains, or utility corridors inside the buildings.

Other: GB will decompose when heated or when combusted in a fire to form toxic gases including phosphorous oxides and fluoride. GB can contaminate food and agricultural produce but considered unlikely due to its volatility and lack of persistence.

Health Effects

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

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 GB by vapor exposure are different from initial effects from similar amount of liquid GB 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.

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 GB. Target organ from ingestion is the GI tract.

Effect Levels

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

AEGL Level in mg/m ³ , at various exposure durations	10 min.	30 min.	1 hr.	4 hr.	8 hr.
AEGL 1: Threshold mild effects	0.0069	0.0040	0.0028	0.0014	0.0010
AEGL 2: Potentially irreversible effects or impaired ability to escape	0.087	0.050	0.035	0.017	0.013
AEGL 3: Threshold for severe effects/medical needs/increasing potential for lethality	0.38	0.19	0.13	0.070	0.051

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

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.

Exposure Guidelines

Note: NA = not available.

Occupational

NIOSH IDLH = 0.1 mg/m ³	NIOSH REL-STEL = 0.0001 mg/m ³ (1 × 10 ⁻⁴)	ACGIH TLV-TWA = NA
NIOSH REL-TWA = NA	OSHA PEL = NA	ACGIH TLV-STEL = NA

Worker Population Limit (WPL) for DoD Worker [8-hr TWA] = 0.00003 mg/m³ (3 × 10⁻⁵)

Population

APHC Health Based Environmental Screening Levels (HBESL) = 32 mg/kg for Industrial Soil; 1.3 mg/kg for Residential Soil

Drinking Water: NA for civilians; Military Exposure Guidelines (MEG) at 5L/day, 15L/day = 28, 9.3 µg/L (not intended for civilian population); Military Field Water Standards (MFWS) at short term, 7-day exposure, 5L/day, 15L/day = 12, 4 µg/L (can address civilian applications); 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 (they 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, for information on and access to the PALs. (Note: PALs are not intended to define cleanup levels.)

General Population Limit (GPL) for DoD chemical demilitarization [24-hr TWA] = 0.000001 mg/m³ (1 × 10⁻⁶); inhalation exposure

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/ershdb. We also recommend that responders check their own internal procedures (i.e., SOPs), if applicable.

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.

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. Send person for follow-up medical attention and evaluation. If cleared to resume work, continue to monitor for signs/symptoms and treat accordingly.

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

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 GB 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 GB 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 GB 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.

Personnel Decontamination

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 (continued)

Personnel Decontamination Procedure (continued):

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: https://www.cdc.gov/NIOSH/ershdb/EmergencyResponseCard_29750001.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.

Step / Zone		Personnel Decontamination Procedures (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 GB 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 GB 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 GB.

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.

Note: NA = not applicable.

Minimum Screening Ranges/ Levels for Air Vapor	JCAD M4A1 (pre-concentrator)	AP2C/ AP4C	MX-908 Vapor Mode	Dräger (CDS Kit)	MINICAMS (Near real-time)	Minimum Screening Ranges/ Levels for Liquid/ Vapor	M256/ M256A1	M8	Screening Levels for Water	M272 (water)
ppm	0.02 (0.0002)	0.0015-0.003	0.1	0.025	0.000005	ppm	0.0008-0.0009	0.02 ml		0.02 mg/L
mg/m ³	0.10 (0.001)	0.01-0.03	0.57	0.14	0.00003	mg/m ³	0.005	NA		NA

Environmental Sampling

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

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, (www2.epa.gov/emergency-response/environmental-response-laboratory-network). For questions on environmental sampling for GB call EPA/HQ-EOC at 202-564-3850.

Environmental Sampling (continued)

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 GB has very low persistence on surfaces, ambient air sampling to help to “clear areas” should be included in the sampling plan.

Note: GB breaks down in most environmental conditions to numerous breakdown products, especially fluoride ion, methylphosphonic acid (MPA), and isopropyl methylphosphonic acid (IMPA), which may be used as markers to determine the extent of contamination of the parent GB. See ANALYSIS section below to ensure sampling procedures are compatible with all analytes. Some preparation techniques for both GB and 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.

Types of Samples:

Air (Vapors are heavier than air): Samples are collected using appropriate solid phase absorbent media (tubes) or air sampler (e.g., SUMMA canister) at breathing zone level (~5 ft.) to assess inhalation exposure and at ground levels (~6 in.) to assess off gassing at surfaces.

Water: Water should be collected in appropriate containers with addition of appropriate de-chlorinating agents and preservatives to minimize GB degradation and hydrolysis prior to analysis.

Soil: For localized “hot spot” areas where soil deposition may occur, surface soil samples should be taken from a non-vegetated area to a depth of less than one inch. Sub-surface soil samples may not be necessary unless a large amount of liquid was poured on the ground, or if an underlying aquifer is endangered.

Surface Wipes: Wipe samples are often desired to indicate absence of GB on non-porous surfaces. Concurrent air monitoring is recommended.

Bulk: For hot spot areas where liquid GB deposition may occur on porous surfaces (e.g., concrete, asphalt), actual pieces 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 GB into these materials. Concurrent air monitoring is recommended.

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

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 GB-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 GB 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 GB 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/m³ 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 GB, 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.

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 GB, particularly for large numbers of samples and for the various types of environmental media. EPA’s ERLN labs (www2.epa.gov/emergency-response/environmental-response-laboratory-network) that are specially trained and equipped for the analysis of GB, 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 GB analysis, and methods provided in EPA’s ESAM, contact EPA/HQ-EOC at 202-564-3850. The ERLN also maintains EPA’s Laboratory Compendium, 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 CWA, including GB and its breakdown products, can be obtained by querying the database of laboratories listed in EPA’s Laboratory Compendium (<https://cfext.epa.gov/cetl/lblogin.cfm?action=None>); prior registration is necessary.

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, that 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 monitoring, 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.

Environmental Decontamination/Cleanup

Decontamination/Cleanup Planning:

Once site controls are in place, develop a site-specific decontamination/cleanup plan. 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 facility, wastes generated, as well as the time the facility or item will be out of service and any socio-economic, psychological, 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).

Disposal Option: The urgency to restore a facility as quickly as possible may result in the outright and timely removal and disposal of contaminated materials. Certain materials may be resistant to decontamination formulations, or may be cheaper to discard and replace than to decontaminate and restore.

Monitored Natural Attenuation: GB degrades 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.

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 separate and immobilize the agent contamination from coming into contact with the environment or the public. This can be a temporary or permanent solution.

Decontamination Strategy: A decontamination strategy can be developed by designating contaminated areas into four broad categories: 1) surfaces or hot spots, 2) large volumetric spaces, 3) sensitive equipment or items, and 4) 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.

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 GB liquid should be absorbed using vermiculite or dry sand and transferred into a sealed container and disposed according to WASTE MANAGEMENT section below. Decontamination of GB occurs mainly through hydrolysis, which may be catalyzed (sped up), by various approaches:

- 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 GB with efficacy achieved with contact time of 15-60 minutes depending on surface material. Calcium hypochlorite, present in commercial products, such as HTH (10% hypochlorite solution), is better for surfaces with high concentrations of liquids in localized areas.
- 2) Hydroxide (e.g., sodium, potassium – 10% solution) reacts rapidly with GB, but solutions are very damaging to many surfaces and should be rinsed thoroughly after use.
- 3) Other high pH solutions, such as sodium carbonate (10% solution) have been shown to decontaminate surfaces but slower than decontamination with sodium or potassium hydroxide.
- 4) Proprietary decontamination technologies such as DF-200®, Dahlgren Decon®, CASCAD®, Decon Green®, or L-Gel® have been shown to be effective against GB on the order of minutes to hours, but not all have been thoroughly tested. Availability, cost, and the need for specialized equipment may limit their use early in the response.

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

- 1) Monitored Natural Attenuation is more passive than other decontamination options and is non-destructive to materials. This option may be preferable given the scope and severity of contamination, especially given GB’s relative high volatility.
- 2) Forced or Hot Air ventilation methods are recommended for vapor plume contamination or low concentration of GB in large volumetric spaces or open areas; efficacy typically can be achieved in hours to days with less waste and adverse impacts to materials.

Sensitive Equipment and Items: Forced or Hot Air ventilation may be used for GB and can be used either in-situ or ex-situ to decontaminate these items.

Aqueous Solutions: GB degrades via hydrolysis but may persist in aqueous solutions, depending on initial concentration and environmental conditions. 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: Water systems, plumbing, surfaces, and equipment frequently consist of polymers that may act as sinks for GB, so should be evaluated for decontamination. While GB has high water solubility, it may persist in water systems through various mechanisms, including entrapment in hydraulic dead ends and sorption to certain component materials, leading to recontamination after contaminated water is removed from the system. It may be necessary to isolate affected portions of the system while implementing system decontamination. A contaminated water system may transfer contamination to building and premise plumbing, which then may also require decontamination.

Environmental Decontamination/Cleanup (continued)

CAUTION: Decontamination products may have unique safety/PPE requirements due to their own toxicity or that of breakdown products during use (e.g., bleach results in chlorine vapors). Dirt, grime, and other coatings 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 spread.

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

Waste Management

Transportation:

Federal requirements for the commercial transport of hazardous materials and procedures for exemptions are specified in www.fmcsa.dot.gov/safety-security/hazmat/complyhmregs.htm#hmp. GB 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 to discuss specific cases. Additional resources on packaging, labeling, and shipping are available at <https://www.phmsa.dot.gov/about-phmsa/offices/office-hazardous-materials-safety>. Detailed state regulations can be found at www.envcap.org/.

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. GB is not listed under 40 CFR §261.31-33, but GB-contaminated waste can be reactive hazardous waste, chemical code D003, if it produces toxic fumes when mixed with water (40 CFR §261.23). Listed or characteristic waste may be land disposed only if the waste meets the applicable treatment standards (40 CFR §268.40). Sarin (GB) is also regulated as an Extremely Hazardous Substance under the Emergency Planning and Community Right-to-Know Act (EPCRA), with a reportable quantity threshold of 10 lbs (40 CFR §355 Appendices A and B).

The states (except for Alaska and Iowa) have the primary responsibility to implement the hazardous waste regulations and can impose more stringent requirements than the federal program, so it is critical to open a dialogue with state regulators as early as possible. 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 contaminated residues, soils, and debris.

Management of toxic decomposition products, associated residual decontamination solutions, local waste acceptance criteria, and transportation and handling requirements should be considered. EPA has developed **I-WASTE**, a web-based tool that contains links to waste transportation guidance, treatment and disposal facilities, state regulatory offices, packaging guidance, and guidance to minimize the potential for contamination. Access to EPA's web-based disposal tool requires pre-registration (<http://www2.ergweb.com/bdrtool/login.asp>).

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/>).