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BIOENVIRONMENTAL ENGINEERING

Occupational and Environmental
Air/Gas Sampling

QUALIFICATION TRAINING PACKAGE

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STS Line Item 4.5.2.6.1: Detector Tubes or Chips (Draeger Civil Defense Simultest Kit)*

TRAINER GUIDANCE

Proficiency Code:	3c
PC Definition:	Can do all parts of the task. Needs only a spot check of completed work. Can identify why and when the task must be done and why each step is needed.
Prerequisites:	Draeger Civil Defense Simultest Kit (CDS), Web-Based Training.
Training References:	Equipment User's Manual. *The Occupational Environment: Its Evaluation, Control, and Management*
Additional Supporting References:	None
CDC Reference:	4B051
Training Support Material:	User's manual. CDS kit.
Specific Techniques:	Conducts hands-on training and evaluation of operation of equipment with verification of steps.
Criterion Objective:	Given a CDS kit, demonstrate how to prepare and perform the Set I (for S-Mustard, Phosgene, Hydrogen Cyanide, Lewisite, and N-Mustard) and Set V (for Nerve Agents, Phosgene, Cyanogen Chloride, Chlorine, and S-Mustard) Simultest sets successfully completing all checklist items with no trainer assistance.
Notes: *****The CDS kit has two different 5-tube Simultest sets (Sets I and V) that can identify various nerve, blood, lung, nose and throat irritating agents. Each test takes less than five minutes to complete and does not require warm-up time or calibration.	

TASK STEPS

PUMP LEAK TEST:
1. Insert an unopened detector tube into the pump socket.
2. Squeeze the pump completely and release.
3. After 15 minutes, look for the end-of-stroke indicator.[1]
4. If the pump is deemed leak-proof, remove the tube and reset the counter on the pump to zero.

PREPARE AND OPERATE ACCURO® PUMP WITH DETECTOR TUBES:
1. Determine which set tubes will be used for sampling.[2]
2. Open detector tubes only on pump side first (direction of arrow). [3]
3. Insert the open ends of the tube(s) into the tube adapter (flow arrows pointing in).
4. Score and break off tips on other side of set.
5. Activate the cyanogen chloride tube (Simultest Set V only).[4]
6. Connect the tube set to pump.[5]
7. Check stroke number.[6]
8. Operate the pump.[7]
9. Complete sampling.[8]
10. Remove the used tubes from the adapter and dispose of them correctly.[9]
11. Purge pump.[10]

RECORD DATA:
1. Utilize DOEHRS or equivalent as applicable.

LOCAL REQUIREMENTS: N/A

NOTES:
1. If the end-of-stroke indicator does not appear, the pump is leak-proof and you are ready to continue. If the end-of-stroke indicator appears, your pump is not leak-proof and therefore should not be used.

2. Some tubes contain ampoules that require additional steps, in these circumstances, refer to laminated card for specific instructions. The laminated instruction cards provide a quick reference for easy use and color change interpretations specific to each ampoule (detector tube). The laminated instruction cards also provide reference for groups of related contaminants.

3. Tube Opener Procedures:

   A. Simultest Tube Opener:
      - Carefully scrape the ceramic edge of the tube set opener against the ends of the tubes (at an angle) multiple times to score all five glass tips.
      - With the ceramic cutter UP, push opener completely over rubber tube block and apply pressure down

until all tube tips break off (if some tubes don't break off, re-scrape or use the singe-tube breaker, described below).

B. Deluxe Tube Opener (for individual tubes):
- Insert the tube into the cutter. Keep it pressed against the ceramic edge while turning it (scoring it).
- To open the tube, insert the tube into the opener and push on it at an angle until the tip breaks off (dispose of glass tip).

C. Pump Tube Opener (for individual tubes):
- To score and open an individual tube using the opener on the Accuro® pump; insert the tube into the opener. Press the tube against the ceramic edge while turning it to score the glass; then pull the tube at an angle until the tip breaks off (dispose of glass tip).
- When using the pump to open a tube, be sure to keep the pump facing downwards to prevent any glass splinters from entering the pump.

4. Before beginning the test using the Simultest Set V detector tubes, you will need to activate the cyanogen chloride tube. Remove the tub from the holder and bend the tube between the two black dots, allowing the reagent to flow onto the indicator until it is moistened. Place the tube back into the test holder, always keeping the arrows on the tube pointing towards the pump.

5. A special adapter is used to connect the Simultest Set to the pump. This adapter consists of a 5-slot manifold for the tube sets, 1 meter extension hose that connects to the Accuro® pump.

6. The number of strokes required is stated on the back of the Simultest Set holder and is also included in the Instructions for Use. Conveniently, both Simultest Set I and V require 50 strokes.

7. Grip and hold the Accuro® pump so that the end-of-stroke indicator and stroke counter are facing you. Squeeze the pump until it is fully compressed (the stroke indicator will advance) then release to allow the bellows to re-expand by itself until. A white dot reappears on the top of the pump and indicates re-expansion is complete (end of stroke).

8. Repeat STEP 6 until the number on the stroke counter corresponds to the number of required strokes.

9. All tubes, even those with negative results, cannot be reused and must be disposed of as hazardous waste.

10. The Accuro® pump must be purged after every use, regardless of test results. Flush out the pump by performing a few pump strokes (STEP 6) in clean air.

TRAINEE REVIEW QUESTIONS

STS Line Item 4.5.2.6.1: Detector Tubes or Chips (Draeger Civil Defense Simultest Kit)

1. When sampling using the Simultest Set V, what step must be performed prior to beginning the test?

PERFORMANCE CHECKLIST

STS Line Item 4.5.2.6.1: Detector Tubes or Chips (Draeger Civil Defense Simultest Kit)

Proficiency Code:	3c
PC Definition:	Can do all parts of the task. Needs only a spot check of completed work. Can identify why and when the task must be done and why each step is needed.

DID THE TRAINEE…		YES	NO
PUMP LEAK TEST			
1. Insert an unopened detector tube into the pump socket?			
2. Squeeze the pump completely and release?			
3. After 15 minutes, look for the end-of-stroke indicator.?			
4. If the pump is deemed leak-proof, remove the tube and reset the counter on the pump to zero?			
PREPARE AND OPERATE ACCURO® PUMP WITH DETECTOR TUBES			
1. Determine which set tubes will be used for sampling?			
2. Open detector tubes only on pump side first (direction of arrow)?			
3. Insert the open ends of the tube(s) into the tube adapter (flow arrows pointing in)			
4. Score and break off tips on other side of set			
5. Activate the cyanogen chloride tube (Simultest Set V only)			
6. Connect the tube set to pump			
7. Check stroke number			
8. Operate the pump			
9. Complete sampling			
10. Remove the used tubes from the adapter and dispose of them correctly			

11. Purge pump			
RECORD DATA			
1. Utilize DOEHRS or equivalent as applicable?			
Did the trainee successfully complete the task?			

TRAINEE NAME (PRINT)

TRAINER NAME (PRINT)

ANSWERS

1. When sampling using the Simultest Set V, what step must be performed prior to beginning the test?

A: Before beginning the test using the Simultest Set V detector tubes, you will need to activate the cyanogen chloride tube. Remove the tub from the holder and bend the tube between the two black dots, allowing the reagent to flow onto the indicator until it is moistened. Place the tube back into the test holder, always keeping the arrows on the tube pointing towards the pump

(Source: Equipment User's Manual)

STS Line Item 4.5.2.6.2: PID/FID

TRAINER GUIDANCE

Proficiency Code:	3c
PC Definition:	Can do all parts of the task. Needs only a spot check of completed work. Can identify why and when the task must be done and why each step is needed.
Prerequisites:	None
Training References:	TVA-1000B Toxic Vapor Analyzer Instruction Manual TVA-1000B Operational Checklist (ESOH Service Center Website) Bioenvironmental Engineer's Guide to TVA-1000B (ESOH Service Center Website)
Additional Supporting References:	None
CDC Reference:	4B051
Training Support Material:	TVA-1000B Toxic Vapor Analyzer Instruction Manual TVA 1000B Calibration gases (isobutylene, methane, zero grade air) Tedlar bag
Specific Techniques:	Conduct hands-on training and evaluation of calibration and operation of equipment with verification of steps.
Criterion Objective:	Given a TVA 1000B, perform calibration and functional check on the analyzer, demonstrating how to operate the analyzer successfully completing all the checklist items with limited no trainer assistance.

Notes:

* The TVA-1000B Toxic Vapor Analyzer is a direct reading portable monitor that can display, monitor, and log data in either a flame ionization detector (FID), a photo ionization detector (PID), or both simultaneously to provide real-time measurements of organic, and some inorganic, vapor concentrations in air. It cannot confidently determine unknowns at low ppm (parts per million). Therefore, in these types of situations it is recommended that the analyzer be used more for approximations when exact concentrations are not required.

TASK STEPS

BEFORE STARTING THE UNIT, PERFORM THE FOLLOWING STEPS:*

1. Charge battery.
2. Connect sample probe.[1]
3. Fill/install hydrogen tank (FID versions).[2]
4. Open the hydrogen valve (FID versions).

TO USE THE UNIT, EXECUTE THE FOLLOWING START AND CALIBRATION PROCEDURE: [3]

1. Press ON.
2. Press CONTROL.
3. Press 3 to ignite.
4. Press 2 = setup.
5. Press 1 = calibrate
6. Press 2 = span concentration.
7. Enter span concentration for calibration gas being used.[4]
8. Press 3 = zero.
9. Press 1 = both.
10. Challenge analyzer with zero gas sample.
11. Press ENTER = start.
12. Wait to stabilize.
13. Press ENTER = start (write down zero counts)
14. Press 4 = span
15. Press 2 = PID.
16. Press ENTER = start.
17. Challenge analyzer with isobutylene span gas and wait for readings to stabilize.
18. Press ENTER to accept (wirite down span counts for PID)
19. Press 4 = span.
20. Press 3 = FID.
21. Press ENTER = start.
22. Challenge analyzer with methane span gas and wait for readings to stabilize.
23. Press ENTER = accept (write down span counts for FID)
24. Press 5 = response factor.
25. Confirm that response factor says "RF0:DEFAULT"
26. Verify zero count is within the correct range. (PID, 10.6eV 2000-8000/11.8eV 2000-20,000) (FID <5000)
27. Verify the detector sensitivity is within range. (PID, 10.6eV 3500-6000 ppm, 11.8eV 300-900 ppm) (FID 160-260 ppm)
28. Press EXIT two times to main menu.
29. Press 1 = run.
30. Take background reading.
31. Conduct monitoring.
32. Record data.
33. Power down unit. [5]
34. Remove the hydrogen gas tank from the side of the instrument (if using FID).

RECORD DATA

1. Utilize DOEHRS or equivalent as applicable.

LOCAL REQUIREMENTS:

NOTES:

1. The sample probe assembly is a hand-held device that enables you to take vapor samples at precise locations. It connects to the instrument by means of an umbilical. The umbilical has two quick-disconnect fasteners (one electrical, one sample line) at the instrument end.

2. The FID mode is used to detect most organic compounds and can be used to detect gaseous hydrocarbons in depressions or confined spaces. It is best suited to detect combustible compounds such as gasoline and methane. The PID mode is sensitive to compounds with double bonds, such as aromatic and chlorinated compounds. It can also measure some inorganic compounds that the FID does not detect, such as ammonia, carbon disulfide, carbon tetrachloride, chloroform, ethylamine, formaldehyde, and hydrogen sulfide. Calibration must be completed each day the TVA-1000B is used. The FID is calibrated with methane and the PID is calibrated with isobutylene.

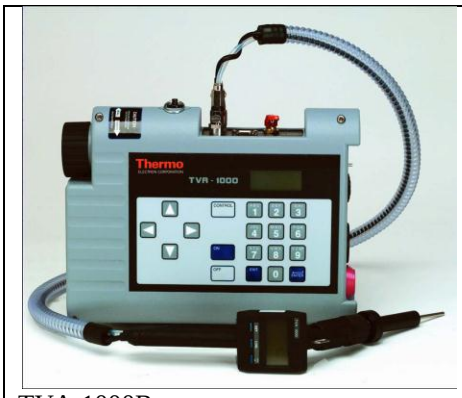
   To refill the hydrogen tank, follow these steps:

   1. Turn the supply tank valve to OFF.
   2. Attach the tank filler adapter to the supply cylinder with valve and manifold valves in OFF position.
   3. Attach the TVA-1000B hydrogen tank to the tank filler adapter (left hand thread-do not over tighten).
   4. Open the supply tank valve.
   5. Move the fill adapter valve to FILL position.
   6. Wait for TVA-1000B tank to fill (may take two to three minutes because of flow restrictors in the tank and fill adapter). Do not fill past 2200psi.
   7. Once full, close the fill adapter valve.
   8. Remove the TVA-1000B tank.
   9. Close the supply cylinder valve.
   10. Reopen the adapter valve to release gas remaining in tank fill adapter (removes pressure and makes adapter removal easier).
   11. Remove the tank fill adapter. (Always remove tank from instrument for storage.)

3. Calibration must be completed each day the TVA-1000B is used. Prior to performing calibration, the instrument must be on and warmed up for approximately 30 minutes. The pump must be ON, the PID lamp must be ON, and the FID must be ignited throughout the warm-up period.

4. If PID only, enter concentration of isobutylene. If FID only, enter concentration of methane. If dual, enter concentration of both gases.

5. To power down this instrument, press and hold the OFF key. With FID versions, you must also shut off the gas valve to avoid depleting the tank supply.



TVA-1000B

***Key Functions:**

ON: The ON key enables power from the battery to the instrument.

OFF: The OFF key disables power from the battery to the instrument.

CONTROL: The CONTROL key is multi-function and is used to turn the pump, PID, and

FID: on or off, and to ignite the FID.

EXIT: The EXIT key clears any entry made in error or bypasses information that you do not want to change, and clears error or warning screens.

ENTER: The ENTER key has three functions:

- Press ENTER if you have typed one or more characters and wish to
- Keep that information.
- Press ENTER to respond to a menu question.
- Press ENTER instead of the LOG key on the standard probe to initiate logging.

Left/Right Arrows: The left and right arrow keys move character entry positions.

Up/Down Arrows: The up and down arrow keys make page selections or scroll through options in SETUP entry screens.

Alphanumeric: The alphanumeric keys enable you to type letters or numbers into various menus. If a display asks for a number only, simply press the desired key. Two steps are required to type an alphanumeric character. First, press the key with the desired letter or number. The screen then displays a selection prompt at the bottom in which 1 = first letter, 2 = second letter, 3 = third letter, and 0 = number. Press the appropriate key to execute the selection. Three uses:

- Select menu options
- Enter numbers, 0-9, using single keystroke
- Enter alphanumeric data, A-Z, 0-9, SPACE, using 2 keystrokes per character

TRAINEE REVIEW QUESTIONS

STS Line Item 4.5.2.6.2: PID/FID

1. Which detector(s) within the TVA-1000B is/are able to distinguish inorganic material?
2. How should the TVA-1000B be decontaminated?
3. How does the ion potential (IP) directly relate to the detection capabilities of the TVA-1000B?
4. How is the response factor calculated?

PERFORMANCE CHECKLIST

STS Line Item 4.5.2.6.2: PID/FID

Proficiency Code:	3c
PC Definition:	Can do all parts of the task. Needs only a spot check of completed work. Can identify why and when the task must be done and why each step is needed.

DID THE TRAINEE…		YES	NO
BEFORE STARTING THE UNIT			
1. Charge battery?			
2. Connect sample probe?			
3. Fill/install hydrogen tank (FID versions)?			
4. Open the hydrogen valve (FID versions)?			
TO USE THE UNIT, EXECUTE THE FOLLOWING START AND CONFIGURE PROCEDURE			
1. Press ON?			
2. Press CONTROL?			
3. Press 3 to ignite?			
4. Press 2 = setup?			
5. Press 1 = calibrate?			
6. Press 2 = span concentration?			
7. Enter Span Concentration for calibration gas being used?			
8. Press 3 = zero?			
9. Press 1 = both?			
10. Challenge analyzer with zero gas sample?			
11. Press ENTER = start?			

12. Wait to stabilize?			
13. Press ENTER = start?			
14. Press 4 = span?			
15. Press 2 = PID?			
16. Press ENTER = start?			
17. Challenge analyzer with isobutylene span gas and wait for readings to stabilize?			
18. Press ENTER to accept?			
19. Press 4 = Span?			
20. Press 3 = FID?			
21. Press ENTER = start?			
22. Challenge analyzer with methane span gas and wait for readings to stabilize?			
23. Press ENTER = accept?			
24. Press 5 = response factor?			
25. Confirm that response factor says "RF0:DEFAULT"?			
26. Verify zero count is within the correct range. (PID, 10.6eV 2000-8000/11.8eV 2000-20,000) (FID <5000)			
27. Verify the detector sensitivity is within range. (PID, 10.6eV 3500-6000 ppm, 11.8eV 300-900 ppm) (FID 160-260 ppm)			
28. Press EXIT two times to main menu?			
29. Press 1 = run?			
30. Take background reading?			
31. Conduct monitoring?			
32. Record data?			
33. Power down unit?			
34. Remove the hydrogen tank from the side of the instrument (if using FID)?			

RECORD DATA			
1. Utilize DOEHRS or equivalent as applicable?			
Did the trainee successfully complete the task?			

TRAINEE NAME (PRINT)

TRAINER NAME (PRINT)

ANSWERS

1. Which detector(s) within the TVA-1000B is/are able to distinguish inorganic material?

   A: Photoionization Detector

   (Source: Bioenvironmental Engineer's Guide to TVA-1000B)

2. How should the TVA-1000B be decontaminated?

   A: The TVA-1000B can be decontaminated by wiping the exterior with a moist towel. Do NOT decontaminate the TVA-1000B by submerging it in water. If the TVA-1000B is accidentally contaminated by drawing a liquid sample into the probe head, it is recommended to call the manufacturer, Thermo Fisher Scientific.

(Source: Bioenvironmental Engineer's Guide to TVA-1000B)

3. How does the ion potential (IP) directly relate to the detection capabilities of the TVA-1000B?

   A: Dependent upon what lamp (10.6 eV or 11.8 eV) is installed for the PID will correlate to the detection of the chemical in question. For example, acetaldehyde has an IP of 10.21 therefore it can be detected by either a 10.6 eV or an 11.8 eV PID lamp but chlorine has an IP of 11.47 so it cannot be detected by the 10.6 eV. An 11.8eV lamp would have to be used.

(Source: Bioenvironmental Engineer's Guide to TVA-1000B)

4. How is the response factor calculated?

   A: Response Factor = Actual Concentration / Measured Concentration

(Source: Bioenvironmental Engineer's Guide to TVA-1000B)

STS Line Item 4.5.2.6.3: Portable GC/MS (HAPSITE®)*

TRAINER GUIDANCE

Proficiency Code:	3c
PC Definition:	Can do all parts of the task. Needs only a spot check of completed work. Can identify why and when the task must be done and why each step is needed.
Prerequisites:	Complete Computer Based Training.
Training References:	Inficon Equipment User's Manual. *HAPSITE® Smart Plus Chemical Identification System Operating Manual (ESOH Service Center Website)*
Additional Supporting References:	*Fundamentals of Industrial Hygiene,* 5th Edition, Chapter 17. *Technical Report on BE HAPSITE® Preventive Maintenance and KD Analytical Support Guidance,* July 21, 2010 *HAPSITE® GC/MS Training Guide – United States Training Version,* 2002
CDC Reference:	4B051
Training Support Material:	HAPSITE® (GC/MS) VOC test sample
Specific Techniques:	Conduct hands-on training and evaluation.
Criterion Objective:	Given a HAPSITE® (GC/MS), perform pre-operational check and operate instrument successfully completing all checklist items with NO trainer assistance.

Notes:

*****The HAPSITE® is a gas chromatograph/mass spectrometer (GC/MS) proven to provide verifiable data for critical health-risk decisions. The HAPSITE® systems deliver fast, dependable on-site analysis of volatile organic compounds (VOCs) in air, water, and soil for emergency response, environmental, hazardous waste, industrial hygiene, process monitoring, and medical applications. The HAPSITE® Headspace sampling system supports the HAPSITE® Smart Chemical Identification System in detecting and identifying VOCs in water or soil on-site or from another location.

INFICON recommends storing the HAPSITE® Smart in extended standby mode. This keeps the NEG (pump) operating at 400°C and the ion pump ON to maintain proper vacuum conditions. Extended standby ensures the battery is charged and ready for deployment/response. While extended standby is recommended, it is not a substitute for system use and it is not a feature to extend the time period between system operations. Using the system or running a weekly Blank Run is the best method to ensure overall operational readiness.

The Guidance Document (HAPSITE® Field Guide) referenced above is designed to provide user's the capability to maximize the use of deployment technology at both garrison and deployed environments in both routine and emergency response situations.

Completion of this Craftsman QTP Training Module also satisfies Craftsman QTP Training 4.15.3.5.1.

TASK STEPS

START UP FROM STANDBY MODE:

(These steps are ONLY for resuming use when the HAPSITE® has been placed in STANDBY MODE.)

1. Using your thumbs, Open front panel of HAPSITE®.[1]
2. Insert purple-banded Nitrogen gas canister into the opening with the purple stripe[2].
3. Insert yellow-banded Internal Standard gas canister into bottom canister opening marked with yellow stripe.[2]
4. Insert a fully charged battery into the rectangular opening to the left of the canister openings.[3]
5. Ensure the sample loop is installed.[4]
6. Naviigate to main menu.
7. Allow the HAPSITE® to boot up and run auto tune check (self-calibration).[5]

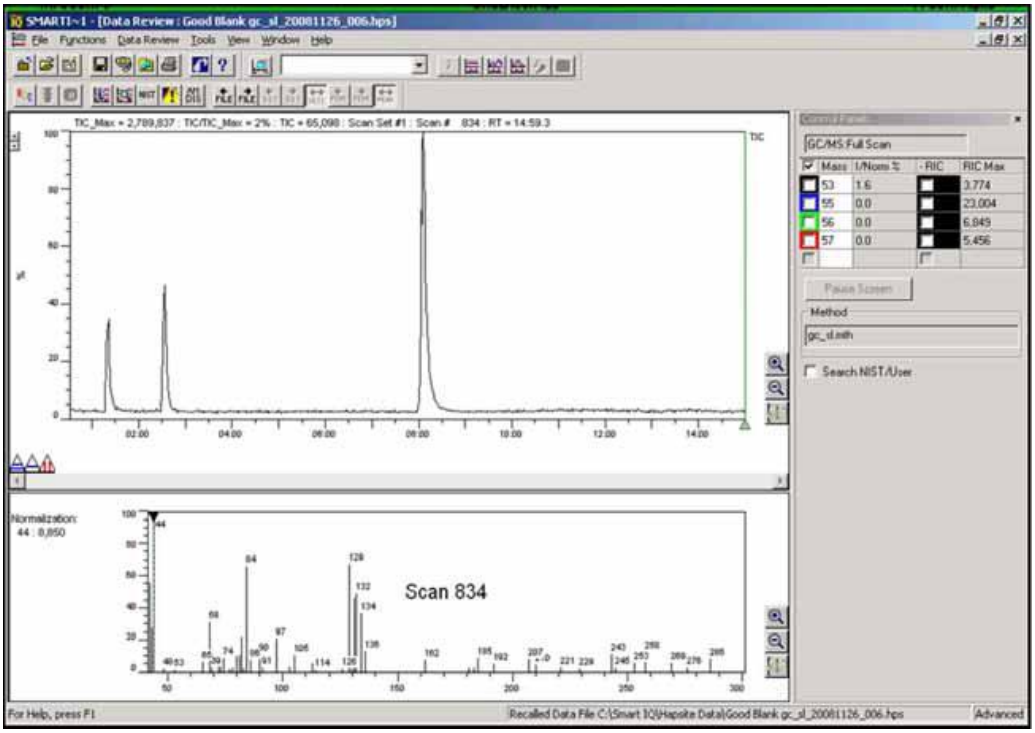
HAPSITE® SEQUENCE OF OPERATION (SURVEY MODE):

1. Ensure HASPITE is turned on and warmed up.
2. Navigate to main screen.
3. Choose "Return to Main Menu"
4. Choose "Run Method"
5. Choose JPMESG Rev 2 Methods
6. Choose JPMESG Survey
7. Ensure Tune parameters are OK,
8. Press **Run** and sample background in ambient air surrounding for about a minute to allow the background to drop and stabilize.[6]
9. Get a volatile organic compound (VOC) sample to test (e.g., toluene, acetone, gasoline).
10. Hold probe over sample for up to one minute while monitoring the TIC count. Look for a response (spike) and pull probe away. (Remember: TIC count over 60 million is indicative of oversaturation.)
11. Keep running the HAPSITE® for at least one minute away from the sample and allow background to drop again.
12. After the clean background has been obtained leave the HAPSITE® running in the clean area for a minimum of a minute prior to entering a suspected contaminated area.
13. When entering an area ensure the TIC count is being observed at all times, if the TIC count reaches 60,000,000 back away from the area.
14. Return to the clean area and let HAPSITE® run for 1 minute
15. Select **Escape** to end the method and return to main menu.
16. Review findings

HAPSITE® SEQUENCE OF OPERATION (SAMPLE LOOP BLANK):

1. Ensure HASPITE is turned on and warmed up.
2. Navigate to main screen.
3. Choose "Return to Main Menu"
4. Choose "Run Method"
5. Choose JPMESG Rev 2 Methods
6. Choose JPMESG GCMS
7. Ensure the Sample Loop is installed with the correct cover. Sample Loop cover will have **Sample Loop** written on it.
8. Select **JPMESG Loop Method**
9. **Select gc_sl**.
10. Press **Run**.
11. HAPSITE® will start sampling as soon as the user selects the run button.
12. Sample collection time is 60 seconds, collection of sample is indicated on the bottom of the screen as "loop fill"
13. When complete, review the blank run. It should show the following:[8]
    - Air Peak at 1:20 +/- 10 seconds
    - Internal Standard #1 at 2:30 minutes +/- 10 seconds (TRIS)
    - Internal Standard #2 at: 8:00 minutes +/- 10 seconds (BPFB)
    - No additional peaks and low background

    All four criteria constitute a satisfactory blank run. See the figure below for an example of a good Sample Loop blank run with all three peaks identified and no additional peaks.



Sample Loop Blank Chromatogram

HAPSITE® SEQUIENCE OF OPERATION (TRI-BED CONCENTRATOR BLANK):

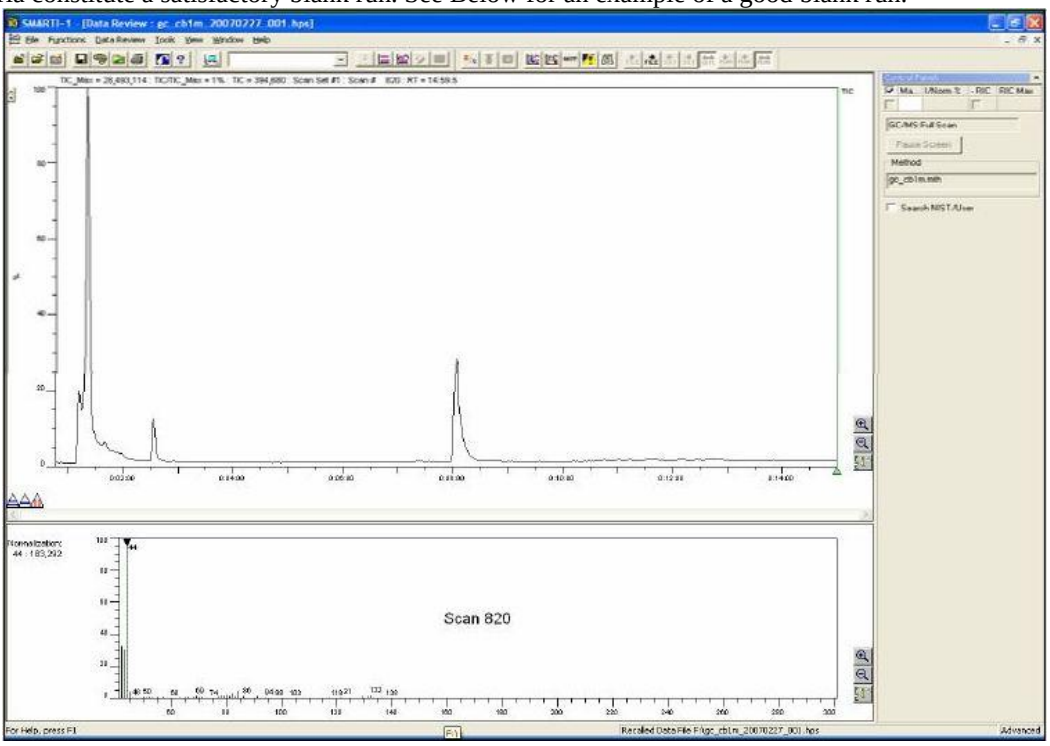
1. Ensure HASPITE is turned on and warmed up.
2. From Main screen, choose press the ESC button
3. Ensure that the Tri-bed concentrator is installed with groove facing up and the appropriate cover is attached. If a problem should occur when running a concentrator, the following message may appear:

   No Concentrator Installed/Incorrect Concentrator:

   (1) Indicates the concentrator cover may not be on.

   (2) Indicates the concentrator is not being recognized; may be due to a chipped end at the base of tube. Chipped concentrator will show Low Column Pressure Warning.[9]
4. Choose "Run Method"
5. Choose JPMESG Rev 2 Methods
6. Choose JPMESG GCMS
7. Choose JPMESG Concentrator
8. Select **JPMESG Concentrator Clean-out (gc cbcl)**. Press **Run** and observe the maximum TIC during this three-minute run. If the TIC is greater than 500,000 at the end of run, repeat clean-out.
9. Note the number of clean-outs required to get TIC below 500,000, and note the actual TIC in comments.[10]
10. Choose "Return to Main Menu"
11. Choose "Run Method"
12. Choose JPMESG Rev 2 Methods
13. Choose JPMESG GCMS
14. Choose JPMESG Concentrator
15. Select **JPMESG Tri-bed concentrator method (gc_cb1m)**, and press **Run**. (Check the Tune Report if you have not already done so.)
16. Sample collection time is 60 seconds, collection of sample is indicated on the bottom of the screen as "conc fill"
17. When complete, review the blank run. It should show the following:[11]
    - Air Peak at 1:20 +/- 10 seconds
    - Internal Standard #1 at 2:30 minutes +/- 10 seconds (TRIS)
    - Internal Standard #2 at: 8:00 minutes +/- 10 seconds (BPFB)
    - No additional peaks and low background.

All four criteria constitute a satisfactory blank run. See Below for an example of a good blank run.



Tri-bed Concentrator Blank Chromatogram

HAPSITE® SEQUENCE OF OPERATION (HEADSPACE SAMPLING SYSTEM (HSS) SET-UP METHOD):[12]

1. Ensure HAPSITE® is in Extended Standby Mode.
2. Attach "Y"-Cable Power Splitter. Connect the single connector end of the cable to the Convertec, power supply. Connect one of the split ends of the cable to the left side of the HAPSITE® and the other to the back of the Headspace unit.

Back of HAPSITE® Analytical Module

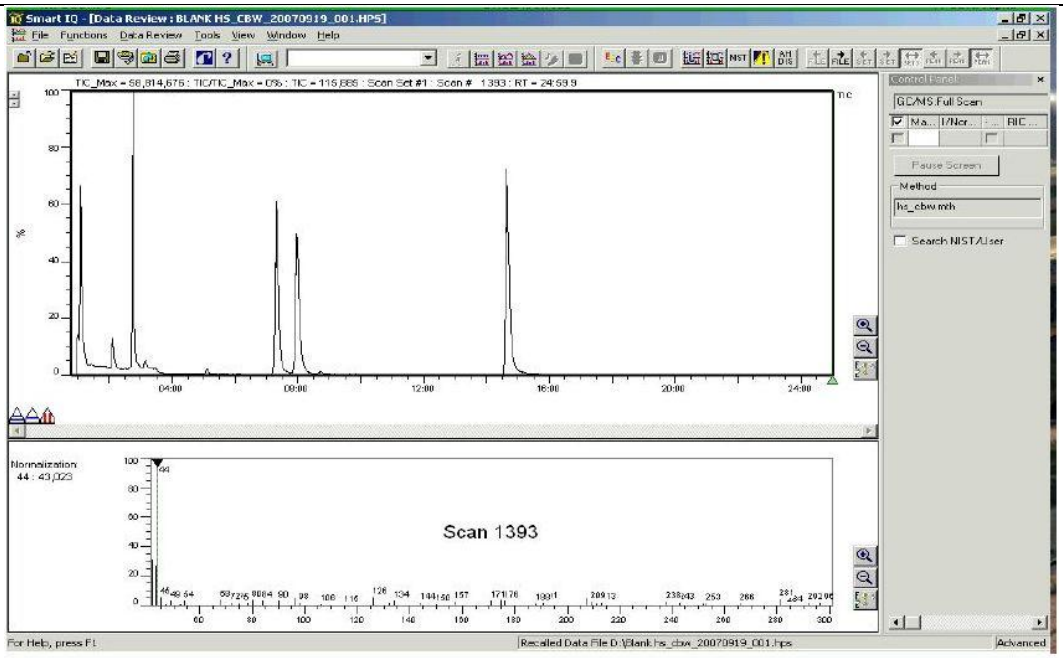
3. Remove probe from the HAPSITE®. Connect the HSS transfer line and ensure that the end of the HSS transfer line with the yellow label marked "This End to HAPSITE®" is connected to HAPSITE®. Connect the end of transfer line with the white label marked "This End to Headspace" to the back of the HSS.

HAPSITE® Headspace Transfer Line

4. Insert a nitrogen canister and a charged battery into the HSS and turn on the power.

HAPSITE® SEQUENCE OF OPERATION (HEADSPACE TRI-BED PERFORMANCE STANDARD):

1. Select **Run** Method from Main Menu.
2. Select JPMESG GCMS Methods
3. JPMESG HeadSpace
4. Select Headspace. Select **hs slwqc** method.
5. HAPSITE® warm-up heaters window will appear. This process takes approximately 15–20 minutes.
6. Automatic tune will initiate (see HAPSITE® LCD screen). The message **Instrument is Tuned** should appear.
7. Measure 20 mL of deionized or sterile water into a 40 mL vial. Inject 1 µL of the Headspace Performance Standard into the 20 mL of de-ionized water through the septum. Gently mix, and then place in the Headspace. Place a clean empty vial in the Headspace next to the vial with Performance Standard.
8. Close yellow cover and press **Run** to start method. Observe on HAPSITE® or laptop screen. Data file name will automatically be generated on HAPSITE® LCD screen.
9. After the run is complete, follow screen directions. Put needle in clean vial and press SEL to purge. Purging takes approximately two minutes.
10. Press **SEL** to view results from front panel LCD.



Headspace Tri-bed Performance Test Method

RECORD DATA:

1. Utilize DOEHRS or equivalent as applicable.

LOCAL REQUIREMENTS:

NOTES:

1. Place hands on top of front panel, using thumbs, pull panel down and outward to open. Care should be taken not to tear the seal.

2. Insert instructions for canisters are located on the inside of the front panel and require the operator to press and hold the PUSH button located to the right of the containment area while inserting canister. With canister pushed in, release the button and this should engage the canister to stay in the containment area. If you can pull it out then it was not inserted properly.

3. The battery is loaded in the opening to the left of the canisters. The INFICON name will be in the upper left corner of the battery and the TEST button in the upper right corner when the battery has been inserted correctly. When the HAPSITE® is in extended standby a battery should be in the machine. The battery will be recharged while in extended standby.

4. Sample loop is located to the right of the canisters. When installing the Sample Loop do **not** over tighten.

5. When tune check is complete, *PRESS ANY BUTTON TO CONTINUE* will appear at the top of the display screen. Any button you press on the HAPSITE® will cause the display window to show the MAIN MENU.

6. TIC generally should be less than 200,000. If not, check area for inteferents such as chemicals that may be in the area.

7. Instrument will continuously run until you stop it while in Survey Mode.

8. Monitoring what a normal blank looks like is one step in verifying the operation of the HAPSITE® and determining if there is a problem. If there are additional peaks in the blank spectrum, and they cannot be removed with additional blank runs, review your blank chromatogram, note the additional analytes, and contact the ESOH Service Center.

9. It is important to blow out the ferrule chamber to ensure broken pieces of the chipped tube are not imbedded.

10. Clean-outs required.

11. If there are additional peaks in the blank spectrum, and they cannot be removed with additional blank runs, AND they are not getting in the way of other analytes, note the additional analytes, and adjust your sample spectrum accordingly. Remember that in future samples, if the chemical that showed up in the blank run is sampled, there will be an increase in peak heights.

12. The Headspace Performance Standard is a test of the HAPSITE® and Headspace connections using the Tri-bed concentrator.

PERFORMANCE CHECKLIST

STS Line Item 4.5.2.6.3: Portable GC/MS (HAPSITE®)

Proficiency Code:	3c
PC Definition:	Can do all parts of the task. Needs only a spot check of completed work. Can identify why and when the task must be done and why each step is needed.

DID THE TRAINEE…		YES	NO
START UP FROM STANDBY MODE			
1. Use thumbs to open front panel of HAPSITE®?			
2. Insert purple-banded Nitrogen gas canister into the opening with the purple stripe?			
3. Insert yellow-banded Internal Standard gas canister into bottom canister opening marked with yellow stripe?			
4. Insert a fully charged battery into the rectangular opening to the left of the canister openings?			
5. Ensure the sample loop is installed?			
6. Press and hold the power button located on the outside of the HAPSITE®s face panel?			
7. Allow the HAPSITE® to boot up and run auto tune check (self-calibration)?			
HAPSITE® SEQUENCE OF OPERATION (SURVEY MODE)			
1. Ensure HASPITE is turned on and warmed up?			
2. From Main screen, choose P**ress the ESC button**?			
3. Choose **Return to Main Menu**?			
4. Choose **Run Method**?			
5. Choose **JPMESG Rev 2 Methods**?			
6. Choose **JPMESG Survey**?			

7. Ensure Tune parameters are OK			
8. Press **Run** and sample background in ambient air surrounding for about a minute to allow the background to drop and stabilize?			
9. Get a volatile organic compound (VOC) sample to test?			
10. Hold probe over sample for up to one minute while monitoring the TIC count?			
11. Keep running the HAPSITE® for at least one minute away from the sample and allow background to drop again repeating steps 4 and 5 two or three times?			
12. After the clean background has been obtained leave the HAPSITE® running in the clean area for a minimum of a minute prior to entering a suspected contaminated area?			
13. When entering an area ensure the TIC count is being observed at all times?			
14. Return to the clean area and let HAPSITE® run for one minute?			
15. Select **ESC** to end the method and return to main menu?			
16. Review findings?			
HAPSITE® SEQUENCE OF OPERATION (SAMPLE LOOP BLANK)			
1. Ensure HASPITE is turned on and warmed up?			
2. From Main screen, choose press the ESC button?			
3. Choose **Return to Main Menu**?			
4. Choose **Run Method**?			
5. Choose **JPMESG Rev 2 Methods**?			
6. Choose **JPMESG GCMS**?			
7. Ensure the Sample Loop is installed with the correct cover?			
8. Select **JPMESG Loop Method?**			
9. Select **gc_sl?**			
10. Press **Run?**			
11. Allow sample collection time of 60 seconds?			
12. When complete, review the blank run?			

HAPSITE® SEQUIENCE OF OPERATION (TRI-BED CONCENTRATOR BLANK)			
1. Ensure HASPITE® is turned on and warmed up?			
2. From Main screen, choose the **ESC** button?			
3. Ensure that the Tri-bed concentrator is installed with groove facing up and the appropriate cover is attached?			
4. Choose **Return to Main Menu?**			
5. Choose **Run Method?**			
6. Choose **JPMESG Rev 2 Methods?**			
7. Choose **JPMESG GCMS?**			
8. Choose **JPMESG Concentrator?**			
9. Select **JPMESG Concentrator Clean-out (gc_cbcl)?**			
10. Press **Run** and observe the maximum TIC during this three-minute run?			
11. Note the number of clean-outs required to get TIC below 500,000, and note the actual TIC in comments?			
12. Choose **Return to Main Menu?**			
13. Choose **Run Method?**			
14. Choose **JPMESG Rev 2 Methods?**			
15. Choose **JPMESG GCMS?**			
16. Choose **JPMESG Concentrator?**			
17. Select **JPMESG Tri-bed Concentrator Method (gc_cb1m)?**			
18. Press **Run?**			
19. Allow a sample collection time of 60 seconds?			
20. When complete, review the blank run?			
HAPSITE® SEQUENCE OF OPERATION (HEADSPACE SAMPLING SYSTEM (HSS) SET-UP METHOD)			
1. Ensure HAPSITE® is in Extended Standby Mode?			

2. Attach "Y"-Cable Power Splitter?			
3. Remove probe from the HAPSITE®?			
4. Connect the HSS transfer line?			
5. Ensure that the end of the HSS transfer line with the yellow label marked "This End to HAPSITE®" is connected to HAPSITE®?			
6. Connect the end of transfer line with the white label marked "This End to Headspace" to the back of the HSS?			
7. Insert a nitrogen canister and a charged battery into the HSS and turn on the power?			
HAPSITE® SEQUENCE OF OPERATION (HEADSPACE TRI-BED PERFORMANCE STANDARD)			
1. Select **Run Method** from Main Menu?			
2. Select **JPMESG GCMS Methods?**			
3. Select Headspace?			
4. Select **hs_slwqc** method?			
5. Wait until the message **Instrument Is Tuned** appeared?			
6. Measure 20 mL of deionized or sterile water into a 40 mL vial?			
7. Inject 1 µL of the Headspace Performance Standard into the 20 mL of de-ionized water through the septum?			
8. Gently mix, and then place in the Headspace?			
9. Place a clean empty vial in the Headspace next to the vial with Performance Standard?			
10. Close yellow cover?			
11. Press **Run** to start method?			
12. Observe on HAPSITE® or laptop screen?			
13. After the run completed, follow screen directions?			
14. Put needle in clean vial and press **SEL** to purge?			
15. Press **SEL** to view results from front panel LCD?			
RECORD DATA			

Utilize DOEHRS or equivalent as applicable			
Did the trainee successfully complete the task?			

TRAINEE NAME (PRINT)

TRAINER NAME (PRINT)

STS Line Item 4.5.2.6.4: Combustible Gas Meters MSA Passport® Personal Alarm*

TRAINER GUIDANCE

Proficiency Code:	3c
PC Definition:	Can do all parts of the task. Needs only a spot check of completed work. Can identify why and when the task must be done and why each step is needed.
Prerequisites:	None
Training References:	MSA Passport® Personal Alarm Instruction Manual
Additional Supporting References:	MSA Passport® Personal Alarm Technical Manual http://media.msanet.com/NA/USA/PortableInstruments/CombinationInstrumentsandCombustibleGasIndicators/PassportPersonalAlarm/803919.pdf
CDC Reference:	4B051
Training Support Material:	MSA Passport® Personal Alarm with fully charged battery pack/power source Calibration gas
Specific Techniques:	Conduct hands-on training and evaluation of calibration and operation of equipment with verification of steps.
Criterion Objective:	Given a Passport® Personal Alarm, demonstrate how to operate and calibrate it successfully completing all the checklist items with no trainer assistance.

Notes:
***** The Passport® Personal Alarm detects oxygen (O_2), carbon monoxide (CO), hydrogen sulfide (H_2S), and sulfur dioxide (SO_2). The Passport® Personal Alarm detects gases and vapors in air only. It cannot measure combustible or toxic gases in reducing atmospheres, furnace stacks, or environments with inert gas backgrounds. The Passport Alarm measures combustible gases and vapors; however, it cannot measure the presence of combustible airborne mists such as lubricating oils.

TASK STEPS

1. Turn on Passport in clean, fresh air environment.[1]
2. Observe readings to verify no gas present.
3. Check battery condition.
4. Perform calibration check.[2]
5. Attach sampling lines and related equipment, if available and collecting a sample from a remote or inaccessible location.[3]
6. Expose instrument to environment.
7. Record meter readings.
8. Utilize DOEHRS or equivalent.

LOCAL REQUIREMENTS:

NOTES:

1. When the unit is turned on it responds with the following:
   - backlight flashes
   - screen flashes
   - alarm sounds
   - alarm lights flash
   - major electronic components are tested automatically

2. Calibration checks must be made frequently if materials such as silicone, silicates, or lead-containing compounds such as leaded gasoline are suspected to be present in the tested atmosphere. If you do not recalibrate, the instrument may give false readings and endanger life and health. To perform a calibration check, do the following steps:

   a. Attach the pump module or calibration cap to the Passport Alarm, orienting the inlet fitting to point toward the battery pack
   b. Attach the calibration adapter to the calibration cap or pump module
   c. Attach the regulator to the cylinder
   d. Connect the black tubing to the regulator
   e. Open the valve on the regulator and connect the other end of the tubing to the inlet fitting
   f. Observe readings are within limits stated on the calibration cylinder

3. To attach probe to sampling line, follow these steps:

   a. Grasp the probe handle by the top two sections [the large section (cap) with the MSA logo and the center section (base) with the label].
   b. Unscrew lower section (guard) from the label section.
   c. Feed male end of the sample line through the guard and screw into the exposed connector ring on the probe.
   d. Screw the guard back onto the base.

PERFORMANCE CHECKLIST

STS Line Item 4.5.2.6.4: Combustible Gas Meters MSA Passport® Personal Alarm

Proficiency Code:	3c
PC Definition:	Can do all parts of the task. Needs only a spot check of completed work. Can identify why and when the task must be done and why each step is needed.

DID THE TRAINEE…		YES	NO
1. Turn on Passport in clean, fresh air environment?			
2. Observe readings to verify no gas present?			
3. Check battery condition?			
4. Perform calibration check?			
5. Attach sampling lines and related equipment, if available and collecting a sample from a remote or inaccessible location?			
6. Expose instrument to environment?			
7. Record meter readings?			
8. Utilize DOEHRS or equivalent.			
Did the trainee successfully complete the task?			

TRAINEE NAME (PRINT) TRAINER NAME (PRINT)

STS Line Item 4.5.2.6.4: Combustible Gas Meters
MSA Sirius® Multigas Detector*

TRAINER GUIDANCE

Proficiency Code:	3c
PC Definition:	Can do all parts of the task. Needs only a spot check of completed work. Can identify why and when the task must be done and why each step is needed.
Prerequisites:	None
Training References:	MSA Sirius® Multigas Detector Operating Manual
Additional Supporting References:	http://www.msanorthamerica.com/catalog/product16577.html http://www.msanorthamerica.com/
CDC Reference:	4B051
Training Support Material:	MSA Sirius® Multigas Detector with fully charged battery/power source Calibration gas
Specific Techniques:	Conduct hands-on training and evaluation of calibration and operation of equipment with verification of steps.
Criterion Objective:	Given a Sirius® Multigas Detector, demonstrate how to operate it successfully completing all the checklist items with no trainer assistance.

Notes:

***** The Sirius® Multigas Detector is designed to detect gases and vapors in air only and to detect only specified toxic gases for which a sensor is installed.

Use only Teflon sampling lines for reactive gases such as chlorine ($CL_{2)}$, phosphine ($PH_{3)}$, ammonia ($NH_{3)}$, hydrogen cyanide (HCN), and for semivolatile organic compounds such as gasoline and jet fuels. Do not use silicone tubing or sampling lines. The operating manual contains additional warnings and acceptable usage limits for the unit as well as a discussion of how the unit functions.

TASK STEPS

Turning ON the Sirius® Multigas Detector

1. Press the **Power ON** button.[1]
    1. Perform Fresh Air Set Up Option for automatic zero adjustment of the Sirius® Multigas Detector sensors.[2]

Verifying Pump Operation

1. Turn **ON** the Sirius® Multigas Detector.[3]
2. Once gas readings are displayed, plug the free end of the sampling line or probe.[4]
3. Check the pump before each day's use.
4. Press the **RESET/▼** button to reset the alarm and restart the pump. [8]

Clearing an Alarm

1. Correct any flow blockage.
2. Press the **RESET/▼** button. The Pump will now restart.

Conducting a Pre-Operational Check

The pre-operational check is simple and should only take about one minute. Perform this check before each day's use for each installed sensor.

1. Turn **ON** the Sirius® Multigas Detector in clean, fresh air.
2. Verify that readings indicate no gas is present.
3. Attach regulator (supplied with calibration kit) to the cylinder.
4. Connect tubing (supplied with calibration kit) to the regulator.
5. Attach other end of tubing to the instrument.
6. Open the valve on the regulator, if so equipped.
7. Determine that the reading on the Sirius® Multigas Detector display is within the limits stated on the calibration cylinder or limits pre-determined by your flight.
8. If necessary, change cylinder to introduce other calibration gases.
9. If readings are not within these limits, the Sirius® Multigas Detector requires recalibration.[6]

Conducting a Calibration Check

1. Turn **ON** the Sirius® Multigas Detector in clean, fresh air.
2. Verify that readings indicate no gas is present.
3. Attach regulator (supplied with calibration kit) to the cylinder.
4. Connect tubing (supplied with calibration kit) to the regulator.
5. Attach other end of tubing to the instrument.
6. Open the valve on the regulator, if so equipped.[7]

Performing Recalibration (if necessary)

1. Turn **ON** the instrument and verify that battery has sufficient life.
2. Wait until the Measure Gases page appears.
3. Push and hold the **RESET/▼**button until **CAL ZERO?** flashes on the display.
4. Push the **ON-OFF/ACCEPT** button to zero the instrument.[8]
5. Connect the appropriate calibration gas (MSA recommends 100ppm isobutylene) to the instrument by connecting one end of the tubing to the pump inlet on the instrument and the other end of tubing to the cylinder regulator (supplied in the calibration kit).*
6. Open the valve on the regulator, if so equipped.
7. Push the **ON-OFF/ACCEPT** button to calibrate (span) the instrument.[9]
8. Remove the tubing from the instrument.

Measuring Gas Concentrations[10]

1. Expose instrument to environment
2. Calculate response factor

Resetting Short Term Exposure Limits (STELs) [11]

1. Access the STEL page.
2. Press the RESET/▼ button

Resetting the Time Weighted Average (TWA)[12]

1. Access the TWA page.
2. Press the RESET/▼ button.

Recording data

1. Utilize DOEHRS or equivalent.

LOCAL REQUIREMENTS:

NOTES:

1. The instrument displays the following information:
    a. A self-test:
        a. Audible alarm sounds
        b. Alarm LEDs illuminate
        c. Display backlight illuminates
        d. Pump activates
        e. Software version displays
        f. Internal diagnostics.
    b. Alarm setpoints:
        a. Low
        b. High
        c. STEL (if activated)
        d. TWA (if activated)
    c. Calibration gas (expected calibration gas values)

d. Time and date (if data logging option installed)

e. Last CAL date (if data logging option installed) —The Sirius® Multigas Detector is equipped with a "last successful calibration date" feature. The date shown is the last date that all installed sensors were successfully calibrated. **LAST CAL** is displayed with this date in the following format: **MM/DD/YY**

f. Instrument warm-up period

g. Fresh Air Setup (FAS) option.

2. Persons responsible for the use of the Sirius® Multigas Detector must determine whether or not the Fresh Air Setup option should be used. The user's abilities, training and normal work practices must be considered when making this decision.

   Warning: Do not activate the Fresh Air Setup unless you are certain you are in fresh, uncontaminated air; otherwise, inaccurate readings can occur which can falsely indicate that a hazardous atmosphere is safe. If you have any doubts as to the quality of the surrounding air, do not use the Fresh Air Setup feature. Do not use the Fresh Air Setup as a substitute for daily calibration checks. The calibration check is required to verify span accuracy. Failure to follow this warning can result in serious personal injury or death.

   To perform a Fresh Air Setup, push the ON/OFF button while **ZERO?** is flashing. The Fresh Air Setup (FAS) has limits. If a hazardous level of gas is present, the Sirius® Multigas Detector ignores the FAS command and goes into alarm.

   Once the instrument self check is complete, **ZERO?** flashes for 10 seconds.

   If no buttons are pushed, the **ZERO?** automatically stops flashing after the 10 seconds have expired and the FAS is not performed.

   To immediately skip the FAS, push the **RESET/▼** button.

3. The pump motor will start fast and then slows down as the instrument adjusts the power to run the pump.

4. If the pump motor shuts down and an alarm sounds, **PUMP ALARM** will flash on the display and the readings on the display may change. When the pump inlet, sample line or probe is blocked, the pump alarm must activate. If the alarm does not activate, check the sample line and probe for leaks. Once leak is fixed, re-check pump alarm by blocking flow.

5. **Warning:** Perform a blocked flow test before each day's use. Do not use the pump, sample line, or probe unless the pump alarm activates when the flow is blocked. Lack of an alarm is an indication that a sample may not be drawn to the sensors, which could cause inaccurate readings. Failure to follow the above can result in serious personal injury or death.

   During operation, a pump alarm may occur when the flow system is blocked, pump is inoperative, and sample lines are attached or removed.

   When the instrument is in a gas alarm, the pump alarm may not display until gas alarm is cleared.

6. The presence of other calibration gases may cause the PID to under range, indicated by dashes for the displayed **VOC** reading.

7. The reading on the Sirius® Multigas Detector display should be within the limits stated on the calibration cylinder or limits which are predetermined by the user. If necessary, change cylinder to introduce other calibration gases. If readings are not within these limits, the Sirius Multigas Detector requires recalibration.

8. Instrument must be in fresh air to perform the zero. **CAL ZERO** flashes. To skip the Zero procedure and move directly to the calibration span procedure, push the **RESET/▼** button. If no button is pushed for 30 seconds, the instrument returns to the Measure mode. Once the zeros are set, **CAL SPAN?** flashes

9. **CAL SPAN** flashes for approximately 90 seconds. If autocalibration sequence passes, the instrument beeps three times and returns to the Measure mode. To skip calibration and return to the Measure mode, push the **RESET/▼** button. If no button is pushed for 30 seconds, it will return to the Measure page.

10. **Warning:** Never let the end of the sampling line touch or go under any liquid surface. If liquid is sucked into the instrument, readings will be inaccurate and the instrument could be damaged. We recommend the use of an MSA Sample Probe (P/N 10042621, 10042622, 10040589, or equivalent) containing a special membrane filter, permeable to gas but impermeable to water, to prevent such an occurrence.

    The Sirius® Multigas Detector can be equipped to detect combustible gases in the atmosphere.

    a. Alarms sound when concentrations reach:

        i. Alarm Setpoint or
        ii. 100% LEL (Lower Explosive Limit), 5% CH4.

    b. When the combustible gas indication reaches the Alarm Setpoint:

        i. Alarm sounds
        ii. Alarm lights flash
        iii. % LEL or CH4 flag above the concentration flashes.

    c. To silence the alarm, press the RESET/▼ button. The alarm will stay silent if the alarm condition has cleared.

    d. When the combustible gas indication reaches 100% LEL or 5% CH4, the LockAlarm™ circuit locks the combustible gas reading and alarm and:

        i. Alarm sounds
        ii. Alarm lights flash
        iii. 100 (or 5.00 in CH4 mode) appears on the display and flashes.

    e. This alarm cannot be reset with the RESET/▼ button. After moving to a safe, fresh-air environment, reset the alarm by turning OFF the instrument and turning it ON again.

To determine a response factor for a target chemical, perform the following procedure:

1. Calibrate the Sirius Detector using isobutylene as the span gas.
2. On the monitor, set the sample gas name to isobutylene.
3. Apply a known concentration of the target chemical to the monitor and note the concentration reported in the display.
4. The response factor for the target chemical relative to isobutylene:

$$\text{RF target gas} = \frac{\text{Actual known concentration}}{\text{Concentration reported by instrument}}$$

For example:

A monitor is calibrated on isobutylene, and has isobutylene defined as the sample gas. When sampling 106 ppm of benzene in air, the instrument reports a concentration of 200 ppm. In this example, the response factor for benzene relative to isobutylene would be:

$$RF\ benz = \frac{106 \text{ ppm known conc. benzene}}{200 \text{ ppm reported}} = 0.53$$

When surveying, if benzene is selected as the sample gas in the Response Factor page, and 0.53 is entered into the monitor as the response factor, the instrument would use this response factor to automatically correct the displayed concentration into PPM benzene.

If a chemical has a response factor between zero and one, the monitor has a higher detector response for this chemical than isobutylene. If the response factor is greater than one, the monitor has a lower detector response for this chemical than isobutylene.

11. The STEL alarm is calculated over a 15-minute exposure. Calculation examples are as follows:

• Assume the detector has been running for at least 15 minutes:

• 15-minute exposure of 35 PPM:

$$\frac{(15 \text{ minutes} \times 35 \text{ PPM})}{15 \text{ minutes}} = 35 \text{ PPM}$$

• 10-minute exposure of 35 PPM

• 5-minute exposure of 15 PPM:

$$\frac{(10 \text{ minutes} \times 35 \text{ PPM}) + (5 \text{ minutes} \times 15 \text{ PPM})}{15 \text{ minutes}} = 28 \text{ PPM}$$

12. The TWA alarm is calculated over an eight-hour exposure. Calculation examples are as follows:

• 1-hour exposure of 50 PPM:

$$\frac{(1 \text{ hour} \times 50 \text{ PPM}) + (7 \text{ hours} \times 0 \text{ PPM})}{8 \text{ hours}} = 6.25 \text{ PPM}$$

• 4-hour exposure of 50 PPM
4-hour exposure of 100 PPM:

$$\frac{(4 \text{ hours} \times 50 \text{ PPM}) + (4 \text{ hours} \times 100 \text{ PPM})}{8 \text{ hours}} = 75 \text{ PPM}$$

• 12-hour exposure of 100 PPM:

$$\frac{(12 \text{ hours} \times 100 \text{ PPM})}{8 \text{ hours}} = 150 \text{ PPM}$$

NOTE: The accumulated reading is always divided by eight hours.

MSA Sirius® MultiGas Detector

PERFORMANCE CHECKLIST

STS Line Item 4.5.2.6.4: Combustible Gas Meters
MSA Sirius® Multigas Detector

Proficiency Code:	3c
PC Definition:	Can do all parts of the task. Needs only a spot check of completed work. Can identify why and when the task must be done and why each step is needed.

DID THE TRAINEE…		YES	NO
TURNING ON THE SIRIUS® MULTIGAS DETECTOR			
1. Press the **Power ON** button?			
2. Perform Fresh Air Set Up Option for automatic zero adjustment of the Sirius® MultiGas Detector sensors?			
VERIFYING PUMP OPERATION			
1. Turn ON the Sirius® MultiGas Detector?			
2. Once gas readings were displayed, plug the free end of the sampling line or probe?			
3. Check the pump before use?			
4. Press the RESET/ ▼ button to reset the alarm and restart the pump?			
CLEARING AN ALARM			
1. Correct any flow blockage?			
2. Press the RESET/ ▼ button to restart the pump?			
CONDUCTING A PRE-OPERATIONAL CHECK			
1. Turn ON the Sirius® Multigas Detector in clean, fresh air?			
2. Verify that readings indicate no gas is present?			
3. Attach regulator (supplied with calibration kit) to the cylinder?			
4. Connect tubing (supplied with calibration kit) to the regulator?			
5. Attach other end of tubing to the instrument?			

6. Open the valve on the regulator, if so equipped?			
7. Determine that the reading on the Sirius® Multigas Detector display is within the limits stated on the calibration cylinder or limits pre-determined by your flight?			
8. If necessary, change cylinder to introduce other calibration gases?			
9. If readings are not within these limits, the Sirius® Multigas Detector requires recalibration?			
CONDUCTING A CALIBRATION CHECK			
1. Turn ON the Sirius Multigas Detector in clean, fresh air?			
2. Verify that readings indicate no gas is present?			
3. Attach regulator (supplied with calibration kit) to the cylinder?			
4. Connect tubing (supplied with calibration kit) to the regulator?			
5. Attach other end of tubing to the instrument?			
6. Open the valve on the regulator, if so equipped?			
PERFORMING RECALIBRATION (IF NECESSARY)			
1. Turn **ON** the instrument and verify that battery has sufficient life?			
2. Wait until the Measure Gases page appears?			
3. Push and hold the **RESET**/ ▼ button until **CAL ZERO?** flashes on the display?			
4. Push the **ON-OFF**/**ACCEPT** button to zero the instrument?			
5. Connect the appropriate calibration gas to the instrument by connecting one end of the tubing to the pump inlet on the instrument and the other end of tubing to the cylinder regulator (supplied in the calibration kit)?			
6. Open the valve on the regulator, if so equipped?			
7. Push the **ON-OFF**/**ACCEPT** button to calibrate (span) the instrument?			
8. Remove the tubing from the instrument?			
MEASURING GAS CONCENTRATIONS			
1. Expose instrument to environment?			
2. Calculate a response factor?			
RESETTING SHORT TERM EXPOSURE LIMITS (STELS)			

1. Access the STEL page			
2. Press the RESET/ ▼ button			
RESETTING THE TIME WEIGHTED AVERAGE (TWA)			
1. Access the TWA page			
2. Press the RESET/ ▼ button			
RECORDING DATA			
1. Utilize DOEHRS or equivalent.			
Did the trainee successfully complete the task?			

TRAINEE NAME (PRINT)

TRAINER NAME (PRINT)

STS Line Item 4.5.2.10: Calculate equivalent OEELs*

TRAINER GUIDANCE

Proficiency Code:	3c
PC Definition:	Can do all parts of the task. Needs only a spot check of completed work. Can identify why and when the task must be done and why each step is needed.
Prerequisites:	None
Training References:	• *Patty's Industrial Hygiene and Toxicology*, 6.1 Brief and Scala Model and 6.2 OSHA Model. • American Conference of Governmental Industrial Hygienist (ACGIH), *TLVs® and BEIs®* (guidebook), most current edition.
Additional Supporting References:	• Air Force Manual (AFMAN) 48-146, *Occupational and Environmental Health Program Management*, 9 Oct 2012, Attachment 4. • Air Force Manual (AFMAN) 48-155, *Occupational and Environmental Health Exposure Controls*, 1 Oct 2008, Chapter 1.
CDC Reference:	4B051
Training Support Material:	• Sampling data. • American Conference of Governmental Industrial Hygienist (ACGIH), *TLVs® and BEIs®* (guidebook), most current edition. • Calculator.
Specific Techniques:	Conduct hands-on training and evaluation.
Criterion Objective:	Given a sampling data scenario, calculate equivalent OEELs successfully completing all checklist items with no trainer assistance.

Notes:
*****OEELs are based on the assumption that exposure occurs for an 8-hour period after which the body is no longer exposed but allowed to recover for the next 16 hours. Where the worker is exposed for more than 8-hours in a day, these assumptions do not hold true. Numerous biological factors come into play when adjusting the OEEL. The booklet produced each year by the American Conference of Governmental Industrial Hygienist (ACGIH), Threshold Limit Values (TLVs) and Biological Exposure Indices (BEIs), should be consulted to ensure it is appropriate to adjust the limit. For example, it is unnecessary to adjust limits where they are based on odor. Although limits can be adjusted downwards to accommodate longer periods of exposure, standards can never be adjusted upwards to accommodate shorter periods of exposure.

TASK STEPS

1. Select appropriate OEEL.
2. Determine total number of hours worked.[1]
3. Select the appropriate equation.[2]
4. Substitute hours worked per day (h) or week (hw) into the equation.
5. Solve to determine equivalent OEEL.

LOCAL REQUIREMENTS:

NOTES:

1. Most often exposure limits are based on the conventional 8-hour day and 40-hour week. When you encounter non-standard (extended) work shifts, OEELs may be adjusted to account for the longer exposures. Adjustments can be made for both daily and weekly exposures.

2. Formulas:

Brief & Scala Model:

The Brief and Scala model is the preferred method for a variety of reasons:

- Easy to use.
- Takes into account increased hours of exposure and decreased recovery time.
- Most conservative model.
- Results in the greatest reduction of the exposure limit.

Daily adjustment of exposure limit:

$$Adjusted\ exposure\ standard\ (TWA) = \left\{\frac{8}{h} x\ \left(\frac{24-h}{16}\right)\right\}\ x\ listed\ TWA$$

Where h = hours worked per day.

Weekly adjustment of limit:

$$Weekly\ Reduction\ Factor\ = \left\{\frac{40}{hw} \times \left(\frac{168-hw}{128}\right)\right\}$$

Where hw = hours worked per week

$$Adjusted\ Exposure\ Limit\ = 8\ hr\ OEEL\ \times Weekly\ Reduction\ Factor$$

OSHA Model:

Daily adjustment of exposure limit:

$$Adjusted\ OEEL_{daily} = OEEL \times \frac{8\ hr}{h}$$

Where *h* = total hours worked.

Weekly adjustment of exposure limit:

$$Adjusted\ OEEL_{weekly} = OEEL \times \frac{40\ hr}{h}$$

Where h = total hours worked.

TRAINEE REVIEW QUESTIONS

STS Line Item 4.5.2.10: Calculate equivalent OEELs

1. Calculate the equivalent OEEL using the Brief & Scala method for an employee that worked a 12-hour shift during which you collected samples for a contaminant that has an OEEL of 0.05 mg/m^3.
2. Calculate the equivalent OEEL using the OSHA method for an employee that worked five (5) 12-hour shifts which you collected samples for a contaminant that has an OEEL of 200 ppm.

PERFORMANCE CHECKLIST

STS Line Item 4.5.2.10: Calculate equivalent OEELs

Proficiency Code:	3c
PC Definition:	Can do all parts of the task. Needs only a spot check of completed work. Can identify why and when the task must be done and why each step is needed.

DID THE TRAINEE…	YES	NO
1. Select appropriate OEEL?		
2. Determine total number of hours worked?		
3. Select the appropriate equation?		
4. Substitute hours worked per day (h) or week (hw) into the equation?		
5. Solve to determine equivalent OEEL?		
Did the trainee successfully complete the task?		

TRAINEE NAME (PRINT)

TRAINER NAME (PRINT)

ANSWERS

1. Calculate the equivalent OEEL using the Brief & Scala method for an employee that worked a 12-hour shift during which you collected samples for a contaminant that has an OEEL of 0.05 mg/m^3.

A:

$$Adjusted\ exposure\ standard\ (TWA) = \left\{\frac{8}{12} x\ \left(\frac{24-12}{16}\right)\right\}\ x\ 0.05$$

$$Adjusted\ exposure\ standard\ (TWA) = \left(\frac{8}{12} x\ \frac{12}{16}\right)\ x\ 0.05$$

$$Adjusted\ exposure\ standard\ (TWA) = \frac{96}{192}\ x\ 0.05$$

$$Adjusted\ exposure\ standard\ (TWA) = \ 0.5\ x\ 0.05$$

$$Adjusted\ exposure\ standard\ (TWA) = \ 0.025\ mg/m^3$$

(Source: Note #2 of this QTP)

2. Calculate the equivalent OEEL using the OSHA method for an employee that worked five (5) 12-hour shifts which you collected samples for a contaminant that has an OEEL of 200 ppm.

A:

$$Adjusted\ OEEL_{weekly}\ =\ 200\ ppm\ \times \frac{40\ hr}{60\ hr}$$

$$Adjusted\ OEEL_{weekly}\ = 200\ ppm\ \times\ 0.67\ \text{(rounded)}$$

$$Adjusted\ OEEL_{weekly}\ = 134\ ppm$$

(Source: Note #2 of this QTP)

STS Line Item 4.5.2.11: Convert raw concentrations (i.e., grams to mg/m^3)*

TRAINER GUIDANCE

Proficiency Code:	3c
PC Definition:	Can do all parts of the task. Needs only a spot check of completed work. Can identify why and when the task must be done and why each step is needed.
Prerequisites:	None
Training References:	American Conference of Governmental Industrial Hygienist (ACGIH), *TLVs® and BEIs®* (guidebook), most current edition.
Additional Supporting References:	None
CDC Reference:	4B051
Training Support Material:	Sample results. Calculator.
Specific Techniques:	Conduct hands-on training and evaluation.
Criterion Objective:	Given sampling results, convert concentrations successfully completing all checklist items with no trainer assistance.

Notes:
*****When you receive your air sampling results back from the analytical laboratory, in most cases you will need to calculate the TWA and compare the results to the applicable OEEL. For those cases where the laboratory uses the air volume you provided and calculates the concentration in milligrams per cubic meter (mg/m3) or parts per million (ppm), at normal temperature and pressure (NTP), no corrections are required; you can simply compare your results directly to the applicable OEEL as long as the OEEL is in the same units. If the units are not the same, you will need to use one of the formulas provided below to convert to the same units as the OEEL: mg/m3 or ppm.

In situations where the laboratory reported your results as a weight of contaminant collected on the sample media, such as milligrams (mg) or micrograms (µg), these results cannot be compared to the OEEL. When this happens, you must first determine the volume of air drawn through the sample then calculate the mass/volume concentration in mg/m3.

Remember, the units must be the same in order to make a comparison.

TASK STEPS

1. Select the appropriate equation.[1]
2. Substitute the known variables into the appropriate equation.
3. Solve to convert given results to desired units.

LOCAL REQUIREMENTS:

NOTES:

1. Conversion Formulas:

mg/m3 to ppm:

$$mg/m^3 = \frac{Molecular\ Weight\ (MW)\ of\ substance \times Sample\ results\ in\ ppm}{24.45}$$

Where 24.45 = a constant and represents the molar volume of air liters at NTP conditions (25°C and 760 torr).

ppm to mg/m3:

$$ppm = \frac{24.45\ \times\ Sample\ results\ in\ mg/m^3}{Molecular\ Weight\ (MW)\ of\ substance}$$

Where 24.45 = a constant and represents the molar volume of air liters at NTP conditions (25°C and 760 torr).

volume:

$$Volume\ (liters) =\ flow\ rate\ (lpm) \times time$$

liters to cubic meters:

$$Volume\ (meters^3) = \frac{liters}{1} \times \frac{1\ meter^3}{1000\ liters}$$

mass/volume concentration in mg/m3:

$$Concentration\ (mg/m^3) = \frac{mass\ reported\ (mg)}{volume\ (m^3)}$$

TRAINEE REVIEW QUESTIONS

STS Line Item 4.5.2.11: Convert raw concentrations (i.e., grams to mg/m^3)

1. You just received your air sample results from the analytical laboratory. You were sampling for benzene. The sample results were returned as 1.5 mg/m^3 but the OEEL for benzene is listed as ppm, so you will need to convert the concentration in order to compare it to the standard.
2. You just received your air sample results from the analytical laboratory. You were sampling for methyl ethyl ketone (MEK). The sample results were returned as 42 ppm but the OEEL for MEK is listed as mg/m^3 so you will need to convert the concentration in order to compare it to the standard.
3. Your sample flow rate was 1.25 liter/minute (lpm) and your total sampling time was 90 minutes, what is your sample volume?

4. Using the sample volume from Question #3, convert the liters to cubic meters (m^3).

5. Your sample results have returned from the laboratory, which reported the results as a mass of 18 mg. Using the converted sample volume from Question #4, what are your results in mg/m^3?

PERFORMANCE CHECKLIST

STS Line Item 4.5.2.11: Convert raw concentrations (i.e., grams to mg/m^3)

Proficiency Code:	3c
PC Definition:	Can do all parts of the task. Needs only a spot check of completed work. Can identify why and when the task must be done and why each step is needed.

DID THE TRAINEE…	YES	NO
1. Select the appropriate equation?		
2. Substitute the known variables into the appropriate equation?		
3. Solve to convert given results to desired units?		
Did the trainee successfully complete the task?		

TRAINEE NAME (PRINT)

TRAINER NAME (PRINT)

ANSWERS

1. You just received your air sample results from the analytical laboratory. You were sampling for benzene. The sample results were returned as 1.5 mg/m^3 but the OEEL for benzene is listed as ppm, so you will need to convert the concentration in order to compare it to the standard.

   A:

$$ppm = \frac{24.45 \times Sample\ results\ in\ mg/m^3}{Molecular\ Weight\ (MW)\ of\ substance}$$

$$ppm = \frac{24.45 \times 1.5\ mg/m^3}{78.11}$$

$$ppm = 0.47\ (rounded)$$

(Source: Note 2 of this training module)

2. You just received your air sample results from the analytical laboratory. You were sampling for methyl ethyl ketone (MEK). The sample results were returned as 42 ppm but the OEEL for MEK is listed as mg/m^3 so you will need to convert the concentration in order to compare it to the standard.

A:

$$mg/m^3 = \frac{Molecular\ Weight\ (MW)\ of\ substance \times Sample\ results\ in\ ppm}{24.45}$$

$$mg/m^3 = \frac{M72.10 \times 42\ ppm}{24.45}$$

$$mg/m^3 = 113\ (rounded)$$

(Source: Note 1 of this training module)

3. Your sample flow rate was 1.25 liter/minute (lpm) and your total sampling time was 90 minutes, what is your sample volume?

   A:

$$Volume\ (liters) = flow\ rate\ (lpm) \times time$$

$$Volume\ (liters) = 1.25\ lpm \times 90\ mins$$

$$Volume = 112.5\ liters$$

(Source: Note 3 of this training module)

4. Using the sample volume from Question #3, convert the liters to cubic meters (m^3).

   A:

$$Volume\ (meters^3) = \frac{liters}{1} \times \frac{1\ meter^3}{1000\ liters}$$

$$Volume\ (meters^3) = \frac{90\ liters}{1} \times \frac{1\ meter^3}{1000\ liters}$$

$$Volume\ (meters^3) = 0.09\ m^3$$

(Source: Note 4 of this training module)

5. Your sample results have returned from the laboratory, which reported the results as a mass of 18 mg. Using the converted sample volume from Question #4, what are your results in mg/m^3?

   A:

$$Concentration\ (mg/m^3) = \frac{mass\ reported\ (mg)}{volume\ (m^3)}$$

$$Concentration\ (mg/m^3) = \frac{18\ mg}{0.09\ m^3}$$

$$Concentration\ (mg/m^3) = 200\ mg/m^3$$

(Source: Note 5 of this training module)

STS Line Item 4.5.2.13: Calculate upper and lower confidence limits

TRAINER GUIDANCE

Proficiency Code:	3c
PC Definition:	Can do all parts of the task. Needs only a spot check of completed work. Can identify why and when the task must be done and why each step is needed.
Prerequisites:	None
Training References:	ACGIH TLV Booklet, 2008 29 CFR 1910.1000(d)
Additional Supporting References:	*Fundamentals of Industrial Hygiene*, 5th edition http://www.cdc.gov/niosh/docs/2003-154/
CDC Reference:	4B051
Training Support Material:	Air sampling results Calculator *NIOSH Manual of Analytical Methods* ACGIH TLV Booklet
Specific Techniques:	Conduct hands-on training and evaluation.
Criterion Objective:	Given air sampling results and references, calculate upper and lower confidence limits successfully completing all checklist items with no trainer assistance.
Notes: See Notes Section for formulas.	

TASK STEPS

1. Obtain laboratory results of analyte(s) sample
2. Determine the standard concentration of the analyte(s) and the sampling procedure information[1]
3. Identify the sampling and analytical error (SAE) factor for the analyte(s) [2]
4. Calculate the upper confidence limit (UCL) [3]
5. Calculate the lower confidence limit (LCL) [4]
6. Recognize when the OEEL could be exceeded[5]
7. Utilize OEHMIS (DOEHRS or equivalent), as applicable.

LOCAL REQUIREMENTS:

NOTES:

1. The standardized concentration of an analyte takes into consideration random fluctuations of its presence throughout the sampling process. This variation is expressed by the National Institute for Occupational Safety and Health (NIOSH) as S_{rT} which is an equivalent term indicating the measure of precision. It can be found in the *NIOSH Manual of Analytical Methods*. The standardized concentration, therefore, can be expressed by the following formula where Y equals standardized concentration and X equals the full period sampling result obtained in step one:

$$Y = \frac{X}{OEEL}$$

The occupational exposure limit can be found in the ACGIH TLV Booklet. It is expressed as either a time-weighted average (TWA) or as a short-term exposure limit (STEL). The sampling results obtained will determine which value to use based on the sampling procedure.

2. The SAE is expressed by multiplying the precision measurement obtained in the *NIOSH Manual of Analytical Methods* by the statistical constant 1.645. In other words, an SAE is found by the S_{rT} multiplied by 1.645, e.g., the SAE of Stoddard solvent is 0.082 (S_{rT} = 0.05 x 1.645).

$$SAE = S_{rT} \times 1.645$$

3. The formula to calculate the upper confidence level (UCL) is the following:

$$UCL = Y + SAE$$

If the sampling process consists of multiple consecutive samples instead of one continuous sample, the formula is different:

$$UCL = Y + \frac{SAE\sqrt{(T_1X_1)^2 + (T_2X_2)^2 + \ldots (T_nX_n)^2}}{OEEL\ (T_1 + T_2 + \ldots T_n)}$$

T_n = Time of each sample

X_n = Concentration of each sample

4. The formula to calculate the lower confidence limit (LCL) is as follows:

$$LCL = Y - SAE$$

Again, if the sampling process consists of multiple consecutive samples instead of one continuous sample, the formula is different:

$$LCL = Y - \frac{SAE\sqrt{(T_1X_1)^2 + (T_2X_2)^2 + \ldots (T_nX_n)^2}}{OEEL\ (T_1 + T_2 + \ldots T_n)}$$

T_n = Time of each sample
X_n = Concentration of each sample

5. If the TWA is already above the OEEL, there is no need to calculate the UCL; however, an LCL should be calculated. If the SAE results in a UCL of more than 1, it is not clear whether the exposure was in compliance. An error of method could make it unclear as to if the exposure was too high.

- If the UCL < 1, it is with 95 percent confidence that an overexposure does not exist.
- If the LCL < 1 and the UCL> 1, it is not certain, so may be classified as a possible overexposure.
- If the LCL > 1, it is with 95 percent confidence that an overexposure exists.

TRAINEE REVIEW QUESTIONS

STS Line Item 4.5.2.13: Calculate upper and lower confidence limits

1. What do calculated UCL and LCL values represent?
2. What is the SAE of methylene chloride?
3. Calculate the UCL and LCL for a continuous sampling for hydrazine. The laboratory results were 0.06ppm TWA. Did an overexposure occur?

4. Calculate the UCL and LCL for multiple consecutive samples of vinyl acetate. The laboratory returned results of 9.2ppm over three hours sampling, 8.9 over three hours sampling, and 9.6 over two hours sampling. Did an overexposure occur?

PERFORMANCE CHECKLIST

STS Line Item 4.5.2.13: Calculate upper and lower confidence limits

Proficiency Code:	3c
PC Definition:	Can do all parts of the task. Needs only a spot check of completed work. Can identify why and when the task must be done and why each step is needed.

DID THE TRAINEE…	YES	NO
1. Obtain laboratory results of analyte(s) sample?		
2. Determine the standard concentration of the analyte(s) and the sampling procedure information?		
3. Identify the sampling and analytical error (SAE) factor for the analyte(s)?		
4. Calculate the upper confidence limit (UCL)?		
5. Calculate the lower confidence limit (LCL)?		
6. Recognize when the OEL could be exceeded?		
7. Utilize OEHMIS (DOEHRS or equivalent), as applicable?		
Did the trainee successfully complete the task?		

TRAINEE NAME (PRINT) TRAINER NAME (PRINT)

ANSWERS

1. What do calculated UCL and LCL values represent?

   A:
   The UCL and LCL incorporate SAE factors statistically in order to obtain the lowest (LCL) and highest (UCL) value that the true exposure could be, within a 95-percent confidence interval.

(Source: Fundamentals of Industrial Hygiene, 5th edition)

2. What is the SAE of methylene chloride?

   A:
   The SAE of Stoddard solvent is 0.082.

   $SAE = S_{rT} \times 1.645$

   $SAE = 0.05 \times 1.645$

   $SAE = 0.082$

(Source: Note #2 and NIOSH Manual of Analytical Methods)

3. Calculate the UCL and LCL for a continuous sampling for hydrazine. The laboratory results were 0.06ppm TWA. Did an overexposure occur?

$$Y = \frac{X}{OEEL}$$

$$Y = \frac{0.06}{0.1}$$

$$\underline{\boldsymbol{Y = 0.6}}$$

$$SAE = S_{rT} \times 1.645$$

$$SAE = 0.094 \times 1.645$$

$$\underline{\boldsymbol{SAE = 0.15}}$$

$$UCL = 0.6 + 0.15 = \underline{\boldsymbol{0.75}}$$

$$LCL = 0.6 - 0.15 = \underline{\boldsymbol{0.45}}$$

   An overexposure did not occur.

   (Source: Note #1 – 5, *NIOSH Manual of Analytical Methods,* and ACGIH TLV Booklet*)*

4. Calculate the UCL and LCL for multiple consecutive samples of vinyl acetate. The laboratory returned results of 9.2ppm over three hours sampling, 8.9 over three hours sampling, and 9.6 over two hours sampling. Did an overexposure occur?

   OEEL = 10 ppm
   S_{rT} = 0.018

$$Y = \frac{X}{OEEL}$$

$$Y = \frac{9.2}{10} = 0.92$$

$$SAE = S_{rT} \text{ x } 1.645$$

$$SAE = 0.018 \text{ x } 1.645 = 0.029$$

$$UCL = Y + \frac{SAE\sqrt{(T_1X_1)^2 + (T_2X_2)^2 + \ldots (T_nX_n)^2}}{OEEL\ (T_1 + T_2 + \ldots T_n)}$$

$$UCL = 0.92 + \frac{0.029\sqrt{(3x9.2)^2 + (3x8.9)^2 + (2x9.6)^2}}{10\ (3 + 3 + 2)}$$

$$UCL = 0.92 + \frac{0.029\sqrt{(27.6)^2 + (26.7)^2 + (19.2)^2}}{10\ (8)}$$

$$UCL = 0.92 + \frac{0.029\sqrt{761.76 + 712.89 + 368.64}}{80}$$

$$UCL = 0.92 + \frac{0.029\sqrt{1843.29}}{80}$$

$$UCL = 0.92 + \frac{0.029\ x\ 42.93}{80}$$

$$UCL = 0.92 + \frac{1.24}{80}$$

$$UCL = 0.92 + 0.015$$

$$UCL = 0.9355$$

$$LCL = Y - \frac{SAE\sqrt{(T_1X_1)^2 + (T_2X_2)^2 + \ldots (T_nX_n)^2}}{OEEL\ (T_1 + T_2 + \ldots T_n)}$$

$$LCL = 0.92 - \frac{0.029\sqrt{(3x9.2)^2 + (3x8.9)^2 + (2x9.6)^2}}{10\ (3 + 3 + 2)}$$

$$LCL = 0.92 - \frac{0.029\sqrt{(27.6)^2 + (26.7)^2 + (19.2)^2}}{10(8)}$$

$$LCL = 0.92 - \frac{0.029\sqrt{761.76 + 712.89 + 368.64}}{80}$$

$$LCL = 0.92 - \frac{0.029\sqrt{1843.29}}{80}$$

$$LCL = 0.92 - \frac{0.029\ x\ 42.93}{80}$$

$$LCL = 0.92 - \frac{1.24}{80}$$

$$LCL = 0.92 - 0.015$$

$$LCL = 0.905$$

A: An overexposure did not occur.

(Source: Note #1-5, *NIOSH Manual of Analytical Methods*, and ACGIH TLV Booklet*)*

STS Line Item 4.5.2.16: Interpret air sample results

TRAINER GUIDANCE

Proficiency Code:	3c
PC Definition:	Can do all parts of the task. Needs help only a spot check of completed work. Can identify why and when the task must be done and why each step is needed.
Prerequisites:	None
Training References:	ESOH Service Center, Laboratory Sampling Guide https://hpws.afrl.af.mil/dhp/OE/ESOHSC/pages/index.cfm?id=742 AFMAN 48-146, *Occupational and Environmental Health Program Management*. 9 October 2012. NIOSH Pocket Guide to Chemical Hazards http://www.cdc.gov/niosh/npg/ (or equivalent chemical reference) AFMAN 48-155, *Occupational and Environmental Health Program* ACGIH TLV® Booklet
Additional Supporting References:	AFI48-145, *Occupational and Environmental Health Program.*15 September 2011
CDC Reference:	4B051
Training Support Material:	Sample Report Calculator
Specific Techniques:	Conduct 4.5.2.16 Interpret Air Sample Results in conjunction with 4.5.2.12 - Calculate Time-weighted Averages and 4.5.2.13 - Calculate Upper and Lower Confidence Limits.
Criterion Objective:	Given a calculator and a Laboratory Sample Report of air sampling results, interpret the data to identify discrepancies successfully completing all steps with NO trainer assistance.
Notes: Trainee should be able to read and understand a Laboratory Sample Report, and identify discrepancies, sample results, blanks, and be able to perform blank corrections. Know how to compare the final Time-weighted Average (TWA) results to the Occupational Environmental Exposure Limit (OEEL).	

TASK STEPS

1. Locate "Narrative Comments" on sample report.
2. Identify if data is valid.
3. Check for any comments that might affect your samples.
4. Review sample information for correctness. (Sample ID, Air Volume, Date Sampled, Analyte, Method Reference – ensure these match what you had sent in.)
5. Check for qualifiers.
6. Check dilution factors.
7. Identify sample(s).
8. Identify blank(s).
9. Check if contaminant is detected greater than the Reporting Limit (RL) on media blanks.
10. Determine if blank correction is necessary.
11. Perform blank correction (if necessary) IAW Laboratory Sampling Guide for all samples.
12. Calculate TWA IAW QTP 4.5.2.12.
13. Calculate upper and lower confidence limits IAW QTP 4.5.2.13.
14. Compare resulting TWA to OEEL and determine if limit was exceeded.

LOCAL REQUIREMENTS:

NOTES:

TRAINEE REVIEW QUESTIONS

STS Line Item 4.5.2.16: Interpret air sample results

1. What needs to be done if the laboratory report only lists a mass of constituent, and did not report a concentration for a sample?
2. Why does air flow rate have to be converted from Liters (L) to cubic meters (m^3)?
3. Why do we have to accomplish censored data analysis?

PERFORMANCE CHECKLIST

STS Line Item 4.5.2.16: Interpret air sample results

Proficiency Code:	3c
PC Definition:	Can do all parts of the task. Needs only a spot check of completed work. Can identify why and when the task must be done and why each step is needed.

DID THE TRAINEE…		YES	NO
1. Locate "Narrative Comments" on sample report?			
2. Identify if data is valid?			
3. Check for any comments that might affect your samples?			
4. Review sample information for correctness. (Sample ID, Air Volume, Date Sampled, Analyte, Method Reference – ensure these match what you had sent in.)?			
5. Check for qualifiers?			
6. Check dilution factors?			
7. Identify sample(s)?			
8. Identify blank(s)?			
9. Check if contaminant is detected greater than the Reporting Limit (RL) on media blanks?			
10. Determine if blank correction is necessary?			
11. Perform blank correction (if necessary) IAW Laboratory Sampling Guide for all samples?			
12. Calculate TWA IAW QTP 4.5.2.12?			
13. Calculate upper and lower confidence limits IAW QTP 4.5.2.13?			
14. Compare resulting TWA to OEEL and determine if limit was exceeded?			
Did the trainee successfully complete the task?			

TRAINEE NAME (PRINT) TRAINER NAME (PRINT)

ANSWERS

1. What needs to be done if the laboratory report only lists a mass of constituent, and did not report a concentration for a sample?

A: Concentration can be calculated by dividing the mass reported by the total volume of air sampled.

(Source: 4B051 CDC)

2. Why does air flow rate have to be converted from Liters (L) to cubic meters (m^3)?

A: OEELs and TWA calculations rely on volume in units of m^3; therefore, liters must be converted to cubic meters (m3).

(Source: 4B051 CDC)

3. Why do we have to accomplish censored data analysis?

A: When you ignore or omit the data your estimated mean will be biased high and by setting to zero, the estimated mean will be biased too low.

(Source: ESOH Service Center, Laboratory Sampling Guide)