

National Institute for Occupational Safety and Health National Personal Protective Technology Laboratory P.O. Box 18070 Pittsburgh, PA 15236

Procedure No. TEB-APR-STP-0512-CBRN

Revision: 0.1

Date: 9 February 2009

## DETERMINATION OF CBRN ACID GASES (CYANOGEN CHLORIDE) SERVICE LIFE TEST, LOOSE-FITTING POWERED AIR PURIFYING RESPIRATORS (PAPR) STANDARD TEST PROCEDURE (STP)

## 1. <u>PURPOSE</u>

This test establishes the procedure for ensuring that the level of protection provided by the *CBRN Acid Gases* (*Cyanogen Chloride*) Service Life Test, Loose-Fitting Powered Air Purifying Respirators Standard Test Procedure submitted for Approval, Extension of Approval, or examined during Certified Product Audits, meet the minimum certification standards set forth in 42 CFR Part 84, Subpart G, 84.63(a)(c)(d); Volume 60, Number 110, June 8, 1995 and the Statement of Standard for Chemical, Biological, Radiological, and Nuclear (CBRN) Powered Air Purifying Respirator (PAPR).

## 2. <u>GENERAL</u>

This STP describes *Determination of CBRN Acid Gases (Cyanogen Chloride) Service Life Test, Loose-Fitting Powered Air-Purifying Respirators Standard Testing Procedure* in sufficient detail that a person knowledgeable in the appropriate technical field can select equipment with the necessary resolution, conduct the test and determine whether the product passes the test.

## 3. EQUIPMENT AND MATERIAL

- 3.1. The list of necessary test equipment and materials follows:
  - 3.1.1. Miller Nelson Research Model 401 Flow-Temperature-Humidity Control System or equivalent. This system is an automated system to control the airflow, temperature and humidity of an air supply for an operating system. Laboratory air and distilled water are supplied to the unit. The unit output is air of the variable volume/flow dependant on the size of unit (10% of max flow to max flow in Lpm  $\pm$  2%), and relative humidity (10-98  $\pm$  3%) and temperature (20-30  $^{0}$ C  $\pm$  0.3%).
  - 3.1.2. EdgeTech Dew Prime II Hygrometer, Model 2000 or equivalent. A microprocessor based programmable chilled mirror dew point hygrometer. The hygrometer uses the dew point and ambient temperature to calculate the relative humidity. Ambient temperature range is: -50°C to 130°C  $\pm$  0.2°C; relative humidity is 1% to 95%  $\pm$  0.5%
  - 3.1.3. Labview software developed for NIOSH Service Life Testing.

Approvals: First Level	Second Level	Third Level	Fourth Level

Revision: 0.1

California Analytical Instruments, Model 1314 Photoacoustic Multigas Monitor with UA0985 optical filter or equivalent. This monitor uses photoacoustic infrared detection to measure gases that absorb infrared light. A filter is used to select the wavelength specific for the gas being detected. Range for cyanogen chloride is 1000 to 0.1 ppm  $\pm 0.01$  ppm. Two detector are used, one for challenge concentration and one for downstream concentration.

- 3.1.4. Mass Flow Controller, Brooks Instruments, variable flow rate depending on use.
- 3.1.5. Read out and Control Electronics, Brooks Instruments, Model 0154. Power supply and controller for the Brooks Mass Flow Controller or equivalent. The flow controllers are an integrally mounted control valve module with which stable gas flows can be achieved. Various flow rates are used with an accuracy of ± 0.7 % of rate and ± 0.2 % full scale.
- 3.1.6. Dry Test Meter, American Meter Co., model and size depending on air flow rate to be measured. Must have NIST traceable calibration certificate.
- 3.1.7. Cyanogen chloride cylinder, 99 % purity, with regulator and purge valve.
- 3.1.8. Kin-Tek 491MB Basic Gas Standards Generator with cyanogen chloride permeation tubes that are certified to NIST standard or equivalent.
- 3.2. Test fixture for mounting cartridges.
- 3.3. The test chamber consisting of an air tight box with door opening lined with gasket material. Two fittings located on the test chamber for the introduction of the test concentration and for the exit of the test fixture. This fixture is not commercially available.
- 3.4. Airflow measurement equipment as described in standard test procedure RCT-APR-

#### 4. TESTING REQUIREMENTS AND CONDITIONS

- 4.1. Prior to beginning any testing, all measuring equipment to be used must have been calibrated in accordance with the manufacturer's calibration procedure and schedule. At a minimum, all measuring equipment utilized for this testing must have been calibrated within the preceding 12 months using a method traceable to the National Institute of Standards and Technology (NIST).
- 4.2. Any laboratory using this procedure to supply certification test data to NIOSH will be subject to the provisions of the NIOSH Supplier Qualification Program (SQP). This program is based on the tenets of *ISO/IEC 17025*, the NIOSH Manual of Analytical Methods and other NIOSH guidelines. An initial complete quality system audit and follow on audits are requirements of the program. Additional details of the Program and its requirements can be obtained directly from the Institute.

- 4.3. Precision and accuracy (P&A) must be determined for each instrument in accordance with laboratory procedures and NIOSH/NPPTL guidance. Sound practice requires, under *NIOSH Manual of Analytical Methods*, demonstrating a tolerance range of expected data performance of a plus or minus 25% of a 95% confidence interval of the stated standard requirement. NIOSH/NPPTL P&A tolerance can be higher but not lower.
- 4.4. Compressed gas cylinders must meet all applicable Department of Transportation requirements for cylinder approval as well as retesting / requalification.
- 4.5. Normal laboratory safety practices must be observed. This includes safety precautions described in the current NIOSH Pittsburgh Health and Safety Program.
  - 4.5.1. Safety glasses, lab coats and hard-toe shoes must be worn at all times.
  - 4.5.2. Workbenches must be maintained free of clutter and non-essential test equipment.
  - 4.5.3. When handling any broken glass laboratory equipment, lab technicians and personnel must wear special gloves, which protect against lacerations or punctures.
- 4.6. Please refer to Material Safety Data Sheets and the NIOSH Health and Safety Manual for the proper protection and care in handling, storing, and disposing of the chemicals and gases used in this procedure.

#### 5. PROCEDURE

Note: Reference Section 3 for equipment, model numbers and manufacturers. For calibration purposes use those described in the manufacturer's operation and maintenance manuals.

- 5.1. Follow individual instruction manuals for set up and maintenance of equipment used in this procedure prior to beginning testing. Malfunctioning equipment must be repaired or replaced and properly set up and calibrated before starting all tests.
- 5.2. Calibrate both photoacoustic monitors for cyanogen chloride using the Kin-Tek gas generator and cyanogen chloride permeation tubes. The challenge monitor is calibrated for the challenge concentration of 150 ppm and the downstream monitor is calibrated for the breakthrough concentration of 2 ppm. Make any necessary adjustments to the Labview software so that the concentration display on the software agrees with the detector.
- 5.3. Set up test equipment as shown in Figure 1. The humidity reading controlled by the Miller Nelson system and monitored the Dew Point Hygrometer. The sample pickup for the hygrometer is place into the air stream via a tee after the Miller Nelson and before the introduction point of challenge agent. The thermocouple for the hygrometer is placed in the challenge gas stream immediately before the test chamber.

- 5.4. Verify the following equipment is on:
  - 5.4.1. Air and water supplies.
  - 5.4.2. Miller Nelson Unit.
- 5.5. Establish the correct humidity and temperature as per the test standard in paragraph 6.3.
- 5.6. Set the required airflow for the system. Individual cartridges will be tested by dividing 170 Lpm by the number of air purifying elements on the system. See appendix 8.2. Verify the airflow from the test fixture using the appropriate dry test meter.
- 5.7. Weigh and record initial weight of the test cartridge on Test Data Sheet.
- 5.8. Take and record initial airflow measurements in accordance with RCT-APR-012. See section 84.1152 (b) Title 42, Code of Federal Regulations, Part 84 for minimum airflow requirements.
- 5.9. Mount cartridge onto test fixture and place in testing chamber.
- 5.10. Ensure that the nitrogen gas is ready for immediate flushing of regulator through the purge valve.
- 5.11. Turn on the cyanogen chloride cylinder.
- 5.12. Establish the test concentration of 150 ppm  $\pm$  15 ppm cyanogen chloride.
- 5.13. Once the cyanogen chloride concentration has been established, testing may begin.
- 5.14. Direct challenge concentration airflow into test chamber.
- 5.15. Start timer. Airflow out of the fixture is directed into the breakthrough detector. Monitor and record the upstream and downstream temperatures of the air stream throughout testing. Record breakthrough values and times.
- 5.16. Run test until breakthrough of 2 ppm is observed or selected service life time is surpassed. Record this data on the test data sheet.
- 5.17. Direct challenge concentration airflow out of test chamber.
- 5.18. Weigh and record final weight of the test cartridge on Test Data Sheet.
- 5.19. Take and record final airflow measurements in accordance with RCT-APR-012. See section 84.1152 (b) Title 42, Code of Federal Regulations, Part 84 for minimum airflow requirements.
- 5.20. Allow clean air to purge through system for 5 minutes.

- 5.21. Repeat steps 5.5 through 5.21 for each test described in section 6.3.
- 5.22. When all tests are completed turn off cyanogen chloride cylinder, set temperature and humidity to zero and allow air to pass through the system for 30 minutes.

#### 6. PASS OR FAIL CRITERIA

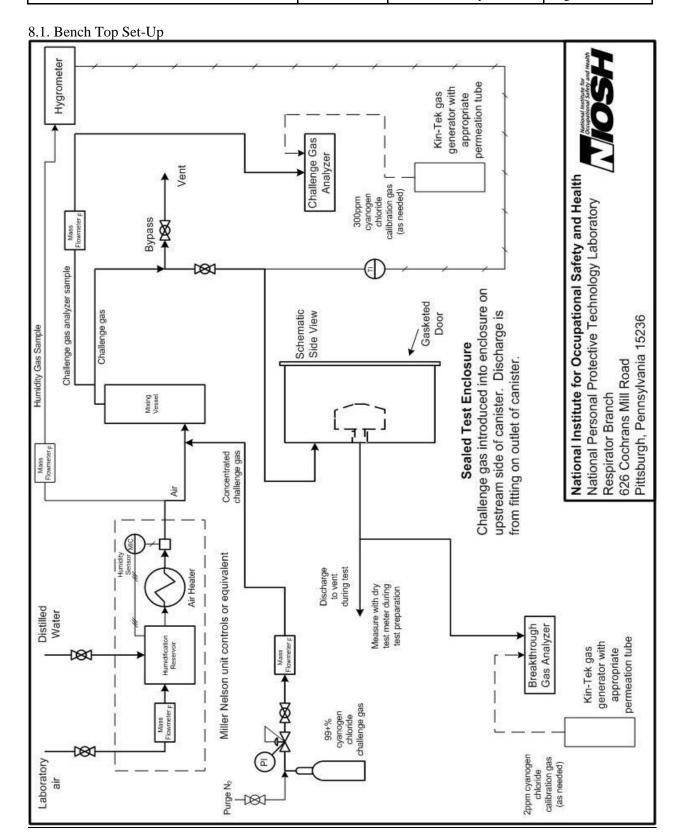
- 6.1. The criterion for passing this test is set forth in 42 CFR Part 84, Subpart G, 84.63(a)(c)(d): Volume 60, Number 110, June 8, 1995.
- 6.2. This test establishes the standard procedure for ensuring that:
  - 84.63 Test requirements; general.
  - (a) Each respirator and respirator component shall when tested by the applicant and by the Institute, meet the applicable requirements set forth in subparts H through L of this part.
  - (c) In addition to the minimum requirements set forth in subparts H through L of this part, the Institute reserves the right to require, as a further condition of approval, any additional requirements deemed necessary to establish the quality, effectiveness, and safety of any respirator used as protection against hazardous atmospheres.
  - (d) Where it is determined after receipt of an application that additional requirements will be required for approval, the Institute will notify the applicant in writing of these additional requirements, and necessary examinations, inspections, or tests, stating generally the reasons for such requirements, examinations, inspections, or tests.
- 6.3. Acid Gases (Cyanogen Chloride) Test for CBRN Cartridges
  - 6.3.1. Airflow measurements of system must be taken before and after each test.
  - 6.3.2. Three cartridges will be tested at the determined, continuous air flow, 25 %  $\pm$  2.5 % relative humidity (RH), 25 °C  $\pm$  2.5 °C and 150 ppm cyanogen chloride.
  - 6.3.3. Three cartridges will be tested at the determined, continuous air flow, 80 %  $\pm$  2.5 % relative humidity (RH), 25 °C  $\pm$  2.5 °C and 150 ppm cyanogen chloride.
  - 6.3.4. Minimum service life will be 15, 30, 45, 60, 90 or 120 minutes as per manufacturer request. End of service life concentration is 2 ppm cyanogen chloride.

#### 7. RECORDS AND TEST SHEETS

- 7.1. All test data will be recorded on the DETERMINATION OF CYANOGEN CHLORIDE SERVICE LIFE test data sheet.
- 7.2. All videotapes and photographs of the actual test being performed, or of the tested equipment shall be maintained in the task file as part of the permanent record.
- 7.3. All equipment failing any portion of this test will be handled as follows:
  - 7.3.1. If the failure occurs on a new certification application, or extension of approval application, send a test report to the TEB Team Leader and prepare the hardware for return to the manufacturer.
  - 7.3.2. If the failure occurs on hardware examined under an off-the-shelf audit the hardware will be examined by a technician and the TEB Team Leader for cause. All equipment failing any portion of this test may be sent to the manufacturer for examination and then returned to NIOSH. However, the hardware tested shall be held at the testing laboratory until authorized for release by the Branch Chief, or the Branch Chief's designee, following the standard operating procedures outlined in *Procedure for Scheduling, and Processing Post-Certification Product Audits*, *RB-SOP-0005-00*.

#### 8. APPENDIXES

- 8.1. Bench Top Set-Up
- 8.2. Airflow calculations
- 8.3. Test Data Sheet



Procedure No. TEB-APR-STP-0512-CBRN	Revision: 0.1	Date: 9 February 2009	Page 8 of 10
-------------------------------------	---------------	-----------------------	--------------

# 8.2. Service life testing airflow calculations

# Loose-fitting PAPR

\_\_\_\_\_Number of filtering elements on a Loose-fitting PAPR System.

# 170 Lpm airflow

Airflow for service life test = 170 Lpm / number of filtering elements

Test Flow				
4 Cartridges	3 Cartridges	2 Cartridges	1 Cartridge	
42.5 Lpm	56.7 Lpm	85 Lpm	170 Lpm	

## 8.3. Test Data Sheet

Гask Nu	mber:						Company:				
est Date	e:		_ 7	#:	Test	Time:	19.0				
Canister Initial	Canister Final		Inhalation Resistanc Canister only		e Inhalation Resistance System		Exhalation Resistance System		Flow	Conc	
grams	grams	Initi in H2		Final in H2O	Initial in H2O	Final in H2O	Initial in H2O	Final in H2O	Lpm	ppm	%RI
									~		
Time				mperature	Temperature Upstream Remarks:						
STA								đ			
est Date	e:		_ #	<b>#</b> :	Test	Time:					
Canister Initial	Canister Final		Inhalation R		e Inhalation Resistance System		Exhalation Resistance System		Flow Conc	Conc	
grams	grams	Initia in H2		Final in H2O	Initial in H2O	Final in H2O	Initial in H2O	Final in H2O	Lpm	ppm	%Rl
Time Leakage		Leakage	Temperature Downstream Upstream		Remarks:						
Canister	e:	Inhala		#:		Time:	Exhalation				
Initial	Final	C	aniste	er only	S	ystem	Sys	tem	Flow	Conc	
grams	grams	Initia in H2		Final in H2O	Initial in H2O	Final in H2O	Initial in H2O	Final in H2O	Lpm	ppm	%Rl
Time		eakage		nperature	Temperature	Remarks					
			Do	wnstream	Upstream					<del>Marian</del> Andrews	
K					, -1,	Test Ope	erator				
										age (	of

Procedure No. TEB-APR-STP-0512-CBRN	Revision: 0.1	Date: 9 February 2009	Page 10 of 10
-------------------------------------	---------------	-----------------------	---------------

# **Revision History**

Revision	Date	Reason for Revision
0.0	21 November 2006	Original Issue
0.1	9 February 2009	Name changed in alignment with current naming convention, paragraphs immediately inferior to 4.5. renumbered to reflect their proper order in the procedure, minor editorial revision, no change to procedure