

Report For: Canada Masq Corporation
4-10 Brodie Drive
Richmond Hill, Ontario
L4B 3K8
Phone: 647-628-9615
Email: anthonyz@canadamasq.com

Laboratory #: 864813-21
Report Date: July 1, 2021
Received Date: June 14, 2021

Attention: Anthony Zhao
Specimen: #1: Respirator Black/White Flat Fold

TEST REPORT

One specimen, consisting of respirators, identified as Black/White Flat Fold, was submitted to CMTL for assessment of particulate filter efficiency, airflow resistance and mechanical strength of headstrap or head harness properties to evaluate acceptability with Health Canada performance criteria for filtering facepiece respirators (Date published: 2020-08-25, Date modified: 2021-02-02).



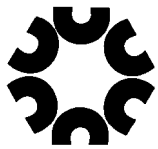
This report is subject to the following terms and conditions: 1. This report relates only to the specimen provided and there is no representation or warranty that it applies to similar substances or materials or the bulk of which the specimen is a part. 2. The content of this report is for the information of the customer identified above only and it shall not be reprinted, published or disclosed to any other party except in full. Prior written consent from Cambridge Materials Testing Limited is required. 3. The name Cambridge Materials Testing Limited shall not be used in connection with the specimen reported on or any substance or materials similar to that specimen without the prior written consent of Cambridge Materials Testing Limited. 4. Neither Cambridge Materials Testing Limited nor any of its employees shall be responsible or held liable for any claims, loss or damages arising in consequence of reliance on this report or any default, error or omission in its preparation or the tests conducted. 5. Specimens are retained 6 months, test reports and test data are retained 7 years from date of final test report and then disposed of, unless instructed otherwise in writing. 6. When making a statement of conformity to a specification or standard the report will make the statement of conformity based on the absolute value of the test result. Test Report Template Revision August 20, 2019

Per Stephen Brown
Authorized By Stephen Brown
Per Derek Wild
Technician, Derek Wild



Requirement for Filtering Facepiece Respirators per
Health Canada National Standard Specifications for Respirators during COVID-19:
Guidance for Canadian Manufacturers, Date published: 2020-08-25, Date modified: 2021-02-02

Characteristic	Barrier	Summary Results
Particulate Filter Efficiency (%)	≥95	Pass
Mechanical Headstrap Strength, Observations and Proof Load (Newtons)	≥20	Pass
Airflow (Inhalation) Resistance, mmH ₂ O (Pa)	≤35 (343)	Pass
Airflow (Exhalation) Resistance, mmH ₂ O (Pa)	≤25 (245)	Pass



PARTICULATE FILTER EFFICIENCY

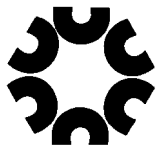
Fourteen submitted specimens were evaluated for particulate filter efficiency based on the TEB-APR-STP-00059 test procedure, with exceptions based on appropriate NRC deviations for maximum expected particle loading being used to filter ambient air in a hospital (medical) or other non-industrial setting.

Seven of the specimens were conditioned (C) within a CSZ environmental control chamber for 25±1 hour at a 85±5% relative humidity and 38°C ± 2.5°C, then tested within 10 hours of extraction from the chamber as indicated in NIOSH standard procedure TEB-APR-STP-0059. A remaining seven additional specimens were also tested to evaluate the effect of no conditioning (U), based on NRC recommendations, to assess composition of filtering material.

The particulate filter efficiency was performed on a TSI 8130A automated filter tester, and challenged for 4-minutes under unidirectional airflow at 85 L/min ± 4 L/min with an aerosol of sodium chloride (NaCl) particles. The particles were generated by an aerosol generator and neutralized to their Boltzmann equilibrium state. The particles were considered to have an average count median diameter of 0.075 ± 0.020 micrometers and a geometric standard deviation not exceeding 1.86. This was equivalent to approximately 7.5 mg of NaCl loading.

RESULTS

Specimen #	Conditioned	Flow Rate	Initial Filter Resistance (mmH ₂ O)	Maximum Allowable Leakage (%)	Initial Leakage (%)	Maximum Leakage (%)	Particulate Filtration Efficiency (%)	Requirement (≥95%)	
								Result	Overall Result
1-1	C	85	9.81	5.00	0.26	0.27	99.73	Pass	Pass
1-2	C	85	9.88	5.00	0.55	0.55	99.45	Pass	
1-3	C	85	9.48	5.00	0.42	0.42	99.58	Pass	
1-4	C	85	10.11	5.00	0.42	0.42	99.58	Pass	
1-5	C	85	10.38	5.00	0.55	0.59	99.41	Pass	
1-6	C	85	9.52	5.00	0.21	0.21	99.79	Pass	
1-7	C	85	10.42	5.00	0.57	0.58	99.42	Pass	
1-8	U	85	9.95	5.00	0.18	0.18	99.82	Pass	
1-9	U	85	9.81	5.00	0.19	0.19	99.81	Pass	
1-10	U	85	10.03	5.00	0.12	0.12	99.88	Pass	
1-11	U	85	10.14	5.00	0.10	0.10	99.90	Pass	
1-12	U	85	9.95	5.00	0.50	0.51	99.49	Pass	
1-13	U	85	9.52	5.00	0.24	0.24	99.76	Pass	
1-14	U	85	9.88	5.00	0.15	0.15	99.85	Pass	



MECHANICAL HEADSTRAP STRENGTH

Ten submitted specimens were subjected to proof load testing in accordance with Health Canada National Standard Specifications for Respirators during COVID-19: Guidance for Canadian Manufacturers (Date published: 2020-08-25, Date modified: 2021-02-02). Testing was performed by donning the mask body on to a head form. A proof load of 10 N was then applied to the elastomeric strap for 10 seconds. The proof load was then removed and the specimens were examined for failure. Testing machine was operated in accordance with ASTM A370-20 paragraph 8 with a test speed of 75mm/min.

RESULTS

Specimen #	Observations	Result
1-1	There was no evidence of breakage, tearing, separation from the point of fixation to the respirator body, permanent deformation or other obvious loss of function in the securing mechanism.	Pass
1-2	There was no evidence of breakage, tearing, separation from the point of fixation to the respirator body, permanent deformation or other obvious loss of function in the securing mechanism.	Pass
1-3	There was no evidence of breakage, tearing, separation from the point of fixation to the respirator body, permanent deformation or other obvious loss of function in the securing mechanism.	Pass
1-4	There was no evidence of breakage, tearing, separation from the point of fixation to the respirator body, permanent deformation or other obvious loss of function in the securing mechanism.	Pass
1-5	There was no evidence of breakage, tearing, separation from the point of fixation to the respirator body, permanent deformation or other obvious loss of function in the securing mechanism.	Pass
1-6	There was no evidence of breakage, tearing, separation from the point of fixation to the respirator body, permanent deformation or other obvious loss of function in the securing mechanism.	Pass
1-7	There was no evidence of breakage, tearing, separation from the point of fixation to the respirator body, permanent deformation or other obvious loss of function in the securing mechanism.	Pass
1-8	There was no evidence of breakage, tearing, separation from the point of fixation to the respirator body, permanent deformation or other obvious loss of function in the securing mechanism.	Pass
1-9	There was no evidence of breakage, tearing, separation from the point of fixation to the respirator body, permanent deformation or other obvious loss of function in the securing mechanism.	Pass
1-10	There was no evidence of breakage, tearing, separation from the point of fixation to the respirator body, permanent deformation or other obvious loss of function in the securing mechanism.	Pass



AIRFLOW (INHALATION) RESISTANCE

Twenty submitted specimens were evaluated for airflow (inhalation) resistance based on TEB-APR-STP-0007 using a TSI 8130A automated filter tester considered by NIOSH to be an acceptable pressure drop measurement.

Tests were performed with the salt generator turned -off under no loading conditions. Using hot-melt glue the filtering facepiece respirators were sealed onto flat plates with joint for connection to the resistance apparatus for measurements of pressure drop.

RESULTS

Specimen #	Maximum Allowable Resistance (mmH ₂ O) Inhalation	Actual Resistance (mmH ₂ O) Inhalation	Requirement (≤35)	
			Result	Overall Result
1-1	35	9.9	Pass	Pass
1-2	35	9.6	Pass	
1-3	35	9.9	Pass	
1-4	35	10.0	Pass	
1-5	35	9.7	Pass	
1-6	35	9.3	Pass	
1-7	35	9.7	Pass	
1-8	35	9.9	Pass	
1-9	35	10.2	Pass	
1-10	35	9.7	Pass	
1-11	35	9.9	Pass	
1-12	35	10.0	Pass	
1-13	35	9.5	Pass	
1-14	35	9.6	Pass	
1-15	35	9.5	Pass	
1-16	35	9.6	Pass	
1-17	35	9.8	Pass	
1-18	35	9.5	Pass	
1-19	35	9.4	Pass	
1-20	35	9.6	Pass	



AIRFLOW (EXHALATION) RESISTANCE

Twenty submitted specimens were evaluated for airflow (inhalation) resistance based on TEB-APR-STP-0003 using a TSI 8130A automated filter tester considered by NIOSH to be an acceptable pressure drop measurement.

Tests were performed with the salt generator turned off under no loading conditions. Using hot-melt glue, the filtering facepiece respirators were sealed onto flat plates and mounted in reverse with joint for connection to the resistance apparatus for measurements of pressure drop.

RESULTS

Specimen #	Maximum Allowable Resistance (mmH ₂ O) Exhalation	Actual Resistance (mmH ₂ O) Exhalation	Requirement (≤25)	
			Result	Overall Result
1-1	25	9.7	Pass	Pass
1-2	25	9.6	Pass	
1-3	25	9.7	Pass	
1-4	25	9.6	Pass	
1-5	25	9.6	Pass	
1-6	25	9.4	Pass	
1-7	25	9.5	Pass	
1-8	25	9.6	Pass	
1-9	25	9.8	Pass	
1-10	25	9.7	Pass	
1-11	25	9.8	Pass	
1-12	25	9.8	Pass	
1-13	25	9.5	Pass	
1-14	25	9.7	Pass	
1-15	25	9.6	Pass	
1-16	25	9.7	Pass	
1-17	25	9.9	Pass	
1-18	25	9.6	Pass	
1-19	25	9.5	Pass	
1-20	25	9.7	Pass	