

Room G8, LB Pavilion 902 W 10TH Avenue
Vancouver, BC, Canada V5Z 1M9

Lab Phone 604 875 4111 Ext 67418
Email PPETesting@vch.ca
Website <http://www.vch.ca/PPETesting>



Test Report for Medical Masks – ATSM F1862 Resistance to Penetration by Synthetic Blood

Prepared for:

Client Name: Layfield Canada Ltd.

Client Contact: Jessica Dominguez

Email Address: Jessica.Dominguez@layfieldgroup.com

Job ID: 220914-LFC-A

Quote ID: PTL-Q-2022-1021

Date: 9/28/2022

Report Version: 2

Room G8, Ground Floor, Leon Blackmore Pavilion
902 W 10TH Avenue, Vancouver, BC, Canada V5Z 1M9

TEL: 604-875-4111 Ext. 67418 | FAX: 604-875-5918 | EMAIL: PPETesting@vch.ca

[The VCH PPE Testing Laboratory's ISO 17025 Accreditations can be found on the Standard Council of Canada's website.](#)

Contents

Executive Summary.....	3
Materials	3
Methods.....	4
Results.....	5
References	6

Test Report for Medical Masks – ASTM F1862 Resistance to Penetration by Synthetic Blood

Client Name: Layfield Canada Ltd.
 Client Contact: Jessica Dominguez
 Email Address: Jessica.Dominguez@layfieldgroup.com
 Job ID: 220914-LFC-A
 Quote Number: PTL-Q-2022-1021
 Date: 9/28/2022
 Report Version: 2

Executive Summary

Thirty-two M6060 respirator samples were tested for resistance to penetration by synthetic blood in accordance with ASTM F1862/F1862-M-17 at a test pressure of 160 mm Hg. All testing were performed in the VCH PPE testing laboratory. Calibration of instruments and equipment were verified to be current, and within operation specifications prior to use. Samples were catalogued upon receipt and inventoried after visual inspection.

Two samples exhibited penetration of synthetic blood on the interior of the respirator when tested at a pressure of 160 mm Hg (Table 4), meeting the acceptable quality limit of 4% for level 3 barrier specified in ASTM F1862/F1862-M-17 and ASTM F2100-21.

Materials

Table 1. Sample and testing info.

Sample Name	Number of Samples	Analysis ID	Date Samples Received	Sample Prep and Test Date
M6060	32	220914-LFC-A-1	9/14/2022	9/15/2022

Methods

Test Standard: ASTM F2100-21¹

Test: ASTM F1862/F1862-M-17²

Test Apparatus: *Conditioning environment: 21 ± 5 °C, 85 ± 5 % relative humidity*

Apparatus: ASTM 1862 Facemask Blood Penetration Test Apparatus

Procedures:

Testing were performed in accordance with ASTM F1862/F1862-M-17. The distance from the target area to the tip of the cannula is 30.5 cm. Visual inspection of the inner surface of the respirator for detecting the presence of synthetic blood penetration was performed within 10 seconds of testing. All equipment had a valid calibration at the time of testing.

Sample conditioning environment: 21 ± 5 °C, 85 ± 5 % relative humidity; samples were conditioned for a minimum of four hours prior to testing.

Equipment:

Table 2. Equipment calibration due dates.

Equipment	Calibration Due Date
ASTM 1862 Facemask Blood Penetration Test Apparatus	N/A
Precision Balance (C021452358)	June 30 2023
Climate Chamber (W819.0119)	October 15 2022
Timer (210779774)	September 13 2023

Lab Environmental Conditions:

Table 3. Laboratory environmental conditions.

Date	Temperature (°C)	Relative Humidity (%)
9/15/2022	21.3	56

Results

Table 4. ASTM F1862/F1862-M-17 test results for M6060 N95-Style Respirator tested at a pressure of 160 mm Hg.

Sample ID	Pass/Fail
220914-LFC-A-1-1	Pass
220914-LFC-A-1-2	Pass
220914-LFC-A-1-3	Pass
220914-LFC-A-1-4	Pass
220914-LFC-A-1-5	Fail
220914-LFC-A-1-6	Fail
220914-LFC-A-1-7	Pass
220914-LFC-A-1-8	Pass
220914-LFC-A-1-9	Pass
220914-LFC-A-1-10	Pass
220914-LFC-A-1-11	Pass
220914-LFC-A-1-12	Pass
220914-LFC-A-1-13	Pass
220914-LFC-A-1-14	Pass
220914-LFC-A-1-15	Pass
220914-LFC-A-1-16	Pass
220914-LFC-A-1-17	Pass
220914-LFC-A-1-18	Pass
220914-LFC-A-1-19	Pass
220914-LFC-A-1-20	Pass
220914-LFC-A-1-21	Pass
220914-LFC-A-1-22	Pass
220914-LFC-A-1-23	Pass
220914-LFC-A-1-24	Pass
220914-LFC-A-1-25	Pass
220914-LFC-A-1-26	Pass
220914-LFC-A-1-27	Pass
220914-LFC-A-1-28	Pass
220914-LFC-A-1-29	Pass
220914-LFC-A-1-30	Pass
220914-LFC-A-1-31	Pass
220914-LFC-A-1-32	Pass
Summary	30 Pass 2 Fail

Table 5. ASTM F2100 Medical Face Mask material Requirements by Performance Level¹ classification of barrier performance.

Characteristic	Level 1 Barrier	Level 2 Barrier	Level 3 Barrier
Resistance to penetration by synthetic blood	80 mm Hg	120 mm Hg	160 mm Hg

References

1. American Society for Testing and Materials. (2021). *Standard Specification for Performance of Materials Used in Medical Face Masks* (ASTM F2100 – 21). Retrieved from <https://www.astm.org/f2100-21.html>
2. American Society for Testing and Materials. (2017). *Standard Test Method for Resistance of Medical Face Masks to Penetration by Synthetic Blood (Horizontal Projection of Fixed Volume at a Known Velocity)* (ASTM F1862/F1862M – 17). Retrieved from https://www.astm.org/f1862_f1862m-17.html

These data are representative of only the samples tested. This report may be copied only in its entirety.

Reports (“**Reports**”) contain the results of testing the samples submitted by the client to whom the Report is addressed (“**Client**”). Client acknowledges the following:

- (a) Reports are limited in scope. Reports only relate to samples as received and tested by VCH at the time of testing, and not from any lot(s) from which the samples may have been taken, or any apparently identical or similar products. Reports reflect the test results as recorded by VCH at the time of testing and are subject to expected measurement variability.
- (b) The interpretation of the Reports and any decisions to be made on the basis of any information contained in the Reports are the sole responsibility of Client. VCH does not provide any interpretation of, or any opinion with respect to, the information contained in the Reports.
- (c) If a Report is amended by VCH, the amended Report will identify that it is an amendment to an existing Report and any change of information will be identified and where appropriate, the reason for the change will be included in the amended Report. Client must not amend or alter any Report or other information received from VCH relating to VCH or the Services provided to Client.
- (d) Reports are for the exclusive benefit of Client and may not be relied upon by any other party. Client may not use, reproduce or otherwise disseminate excerpted, partial, redacted or otherwise altered Reports without the prior review of such use and written approval by VCH. Client is prohibited from manipulating data and/or extrapolating from Reports statistics or conclusions that contradict or eclipse the empirical results of testing as reflected by the complete Report.
- (e) Client will not, without the prior written consent of VCH (which may be subject to conditions) use VCH’s name, logo, test results or Reports prepared by VCH in connection with any sale, marketing, advertising or in any way that may be interpreted as denoting product conformity or that may imply that VCH has approved a product. Client will not, under any circumstances, use VCH’s name, logo or any results or Report prepared by VCH in any manner which may cause harm to VCH’s reputation and/or business.

Prepared by: *Jesse Cooper*, MSc. VCH PPE Testing Lab Manager
Reviewed by: *Titus Wong*, MD, Medical Director

Version	Replaces	Change	Description	Approved by	Release Date
2	220914-LFC-A	Editorial	Reference to model name, M6060, added. Added statement referencing AQL of 4% met. Job ID updated on page 1 and 3 to 220914-LFC-A.	TW	9/28/2022

End of Report