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## Synthetic Blood Penetration Resistance Final Report

Test Article:	3-Ply Face Mask / Lot #200402323	
Purchase Order:	AMYO04242020	
Study Number:	1293285-S01	
Study Received Date:	28 Apr 2020	
Testing Facility:	Nelson Laboratories, LLC	
	6280 S. Redwood Rd.	
	Salt Lake City, UT 84123 U.S.A.	
Test Procedure(s):	Standard Test Protocol (STP) Number:	STP0012 Rev 09
Deviation(s):	None	

**Summary:** This procedure was performed to evaluate surgical facemasks and other types of protective clothing materials designed to protect against fluid penetration. The purpose of this procedure is to simulate an arterial spray and evaluate the effectiveness of the test article in protecting the user from possible exposure to blood and other body fluids. The distance from the target area surface to the tip of the cannula is 30.5 cm. A test volume of 2 mL of synthetic blood was employed using the targeting plate method.

This test method was designed to comply with ASTM F1862 and ISO 22609 (as referenced in EN 14683:2019 and AS4381:2015) with the following exception: ISO 22609 requires testing to be performed in an environment with a temperature of  $21 \pm 5^{\circ}$ C and a relative humidity of  $85 \pm 10^{\circ}$ . Instead, testing was performed at ambient conditions within one minute of removal from the environmental chamber held at those parameters.

All test method acceptance criteria were met. Testing was performed in compliance with US FDA good manufacturing practice (GMP) regulations 21 CFR Parts 210, 211 and 820.

Number of Test Articles Tested:32Number of Test Articles Passed:31Test Side:OutsidePre-Conditioning:Minimum of 4 hours at 21 ± 5°C and 85 ± 5% relative humidity (RH)Test Conditions:21.0°C and 22% RH

**Results:** Per ASTM F1862 and ISO 22609, an acceptable quality limit of 4.0% is met for a normal single sampling plan when  $\geq$ 29 of 32 test articles show passing results.

Test Pressure: 120 mmHg (16.0 kP	a)	
Test Article Number	Synthetic Blood Penetration	
1-28, 30-32	None Seen	
29	Yes	
Sean Shepherd electronically approved for	09 May 2020 18:55 (+00:00)	
Study Director	James Luskin Study Completion Date and Time	
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