PFE RAPID SCREENING ASSESSMENT RESULTS

Summary:

In response to the COVID-19 pandemic, the Therapeutic Goods Administration (TGA) is undertaking a post-market review of all face masks included in the Australian Register of Therapeutic Goods (ARTG) to ensure the quality and effectiveness of face masks supplied in Australia, including that they meet the legislative requirements for medical devices and perform as intended.

Testing was performed in accordance with the TGA in-house Rapid Screening standard operating procedure D20-3958656 (SOP) to assess the particulate filter efficiency of TGA registered respirators claiming compliance with standards used in other countries.

The post-market rapid assessment program was specifically developed to expeditiously quantify the filtration efficiency of respirators. The in-house rapid screening test methodology is based on a modified version of the 42 CFR Part 84 Approval of Respiratory Protective Devices.

Whilst most of the test parameters listed in the SOP are consistent with NIOSH Standard Test Procedure TEB-APR-STP-0059 (STP-0059), this modified test differs for respirator pre-conditioning, test duration and respirator mass loading. Respirators assessed to this modified test plan do not meet the requirements of STP-0059, and therefore cannot be considered equivalent to N95 respirators that were tested to STP-0059. The values reported are only to provide an indication of filter efficiency to ensure masks perform as intended.

Respirator filters were tested for particle penetration against a polydispersed, sodium chloride (NaCl) particulate aerosol. The aerosol was dried, charge neutralised and passed through the test article at a flow rate of 85 ± 4 litres per minute. Each respirator was tested for five minutes after maximum penetration was reached or ten minutes and the findings recorded.

This assessment used convenience sampling, a non-probability sampling technique whereby samples were drawn from a population based on their availability. Respirator filter specimens were then selected at random from the sample provided.

Prior to penetration testing, specimens underwent a visual inspection to qualitatively assess build and marking quality. The initial inhalation resistance and maximum particle penetration (%) for each individual respirator was then determined.

An ATI 100Xs Salt Aerosol Automated Filter Tester was used capable of efficiency measurements of up to 99.9995%. The tester produces a particle size distribution with a count median diameter of 0.075 ± 0.02 um and a geometric standard deviation <1.86. The mass median diameter is approximately $0.26 \mu m$, which is generally accepted as the most penetrating aerosol size.

Unless stated otherwise, specimens were tested under normal laboratory environmental conditions in the condition received. Data relating to the initial resistance does not take into account any bias due to specific mounting fixture used for testing.

Australian Government Department of Health Therapeutic Goods Administration

Laboratories Branch

Sample Details		
LIMS No.		
ARTG No	#N/A	
Label name	#N/A	
Certification claim	ed	
Batch No.	#N/A	
Expiry date	#N/A	

Equipment Used

Equipment Oseu		
ATI Salt Aerosol Automated Filter Tester 100XS		33211
Analytical balance LIMS No.	33203	
NaCl 4% batch No.		
NaCl 0.9% batch No.		
Reference filter media sheet batch No.		
Lest method used	Test method used D20-3958656 - Particulate Penetration Efficiency Assessment -Rapid Screening	

Testing		
Operator	Test date	
Room Temp/RH	Probe LIMS	33215

Sample Conditioning		
Enclosure LIMS	Probe LIMS	
Date/Time IN	Temp/RH IN	
Date/Time OUT	Temp/RH OUT	

certification stand	•	ment for respirators much item tested. Test dat was as follows:	•	•
Minimum Filter E	fficiency =	0.00%	Masks Tested =	0
Maximum Filter E	Efficiency =	0.00%	Total Fails =	0
Maximum Initial F	Resistance =	0.0	Certification claimed	= 0
Filter No	Initial Airflow Resistance (mmH 2 O)	Max Particle Penetration (%)	Filtration Efficiency (%)	Test Result
1			not tested	not tested
2			not tested	not tested
3		_	not tested	not tested
4			not tested	not tested
5			not tested	not tested
6			not tested	not tested
7			not tested	not tested
8			not tested	not tested
9			not tested	not tested
10			not tested	not tested
11			not tested	not tested
12			not tested	not tested
13			not tested	not tested
14			not tested	not tested
15	I		not tested	not tested
16			not tested	not tested
17	I		not tested	not tested
18			not tested	not tested
19			not tested	not tested
20			not tested	not tested
File Name =		RapidPF	E_Results_	
Comments & obs	ervations:			

LIMS Number		0
SAMPLE_ID	LAB_RESPONS	
2105002052	336593	

ST_NAME	TEXT7
s22	N/A

DATE1 28/01/2023 15:21