

Customer Name	MedicAir DentAir Ltd
Customer Address	The Barns, Hilltop Farm, Lyne Lane, Chertsey, Surrey, KT16 0AW
Contact	Alex Hallwood
Sample Description	MedicAir FOZKYGB-04 Air Purifier
Number of Samples	1
Date of Receipt	17 June 2021
ASC Code	ASC004167
Report Number	ASCR092499
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# 1. Purpose

To assess the MedicAir FOZKYGB-04 air purifier in removing airborne *Escherichia* virus MS2 in a 28.5 m<sup>3</sup> environmental test chamber.

## 2. Test Item Description

One MedicAir FOZKYGB-04 air purifier was received by airmid healthgroup on 17 June 2021 in good condition (Figure 2.1).





### 3. Materials and Methods

#### 3.1. 28.5 m<sup>3</sup> Environmental Test Chamber

Testing was conducted in a state-of-the-art 28.5 m<sup>3</sup> test chamber purpose-built to comply with the American Society for Testing and Materials (ASTM) standard. The chamber features include HEPA filtered supply air and an ability to maintain selected temperature and humidity levels at a wide range of air change rates. The chamber was constructed using powder-coated stainless steel with all materials complying with low volatile organic compound (VOC) emission requirements.

The chamber is sealable from the exterior environment with access via an anteroom with interlocking doors and complies with cleanroom standards. The air change rate within the chamber can be controlled within a range of 0.06 to 20 air changes per hour.

#### 3.2. Bacteriophage MS2 (MS2)

Bacteriophage MS2 (MS2) is a non-enveloped positive-sense single stranded RNA virus that infects *Escherichia coli* and some other closely related bacteria but has not been shown to infect eukaryotes. However, several steps of MS2 replication, including viral gene expression and genome replication, share many similarities with positive-sense RNA viruses that infect eukaryotes. Thus, MS2 is often used as a surrogate for other viruses such as Norovirus, including studies where MS2 has been aerosolised [1] and where viral inactivation by ultraviolet light has been assessed [2,3]. Based on the requirements for aerosolisation, the use of ultraviolet light as the antiviral technology and its suitability as a surrogate for some human viral pathogens, MS2 was used as the challenge microorganism in this study.



### 4. Protocol

#### 4.1. Part One: Test Preparation

- 4.1.1. The environmental test chamber is operated at 20 air changes per hour.
- 4.1.2. The walls, floor and ceiling of the environmental test chamber are washed.
- 4.1.3. Once dry, they are wiped with an isopropanol-based wipe.
- 4.1.4. A UV-C light sterilises the surface for 1 hour before testing.

#### 4.2. Part Two: Equipment setup and chamber conditioning

- 4.2.1. The UV light is turned off. Sterile sampling equipment filled with virus collection media is set up on retort stands.
- 4.2.2. Sampling pumps located outside the chamber are calibrated to a specified flow rate for virus sampling.
- 4.2.3. Virus nebulisation equipment is set up in four locations in the test chamber.
- 4.2.4. The test chamber is conditioned to  $20^{\circ}$ C (±  $3^{\circ}$ C) and 50 % relative humidity (± 5 %).

#### 4.3. Part Three: Testing

- 4.3.1. The ventilation system on the environmental test chamber is shut down to minimise the number of air changes during the test.
- 4.3.2. Background air samples are taken in duplicate for 10 minutes.
- 4.3.3. MS2 virus is loaded into the four nebulisers.
- 4.3.4. A ceiling fan and floor fan is operated. The ceiling fan is turned off after nebulisation.
- 4.3.5. The virus is nebulised into the test chamber over 20 minutes.
- 4.3.6. When nebulisation is complete, triplicate air samples are taken for 10 min (t= -10 to 0 minutes).
- 4.3.7. After the air samples have been taken, the air purifier is operated remotely in MAX mode with UV on and remains operating for the duration of the test. In the control runs, no air purifier is operated.
- 4.3.8. Additional air samples are taken in triplicate at the following time points:
  - t = + 5 to 15 minutes
  - t = + 20 to 30 minutes
  - t = + 50 to 60 minutes

#### 4.4. Part Four: Post-study

- 4.4.1. The floor fan is stopped.
- 4.4.2. Air samples are collected and transferred to the laboratory for plaque assay analysis.
- 4.4.3. Sampling equipment is removed and bagged for autoclave sterilisation.
- 4.4.4. The UV light is operated inside the test chamber to decontaminate all surfaces.
- 4.4.5. After UV sterilisation the chamber is operated at 20 air changes per hour.

All testing to the above method is conducted in triplicate for both active test and inactive controls



## 5. Results

Table 5.1 summarises the Log<sub>10</sub> values of the MS2 plague-forming units per cubic meter of air (PFU/m<sup>3</sup>) measured inside the environmental test chamber at each time point. The results of three inactive control runs (no air purifier) and three active test runs (air purifier operating) are presented, the average of which is graphed in Figure 5.1.

Table 5.1. Average Plaque Forming Units/m <sup>3</sup> for MedicAir active test and inactive control runs (Log <sub>10</sub> )								
Time (minute)	Control				Test			
	Run 1	Run 2	Run 3	Average (n=3)	Run 1	Run 2	Run 3	Average (n=3)
-10 to 0	8.0	10.2	8.2	9.7	8.1	8.4	8.1	8.2
+ 5 to 15	8.0	10.1	8.0	9.7	6.7	6.3	6.6	6.5
+20 to 30	7.6	10.0	7.6	9.5	4.8	4.3	4.8	4.7
+50 to 60	7.6	10.0	7.6	9.5	3.6	4.3	3.7	4.0

<LOD: Less than Limit of detection: 3.66 PFU/m<sup>3</sup> (Log10)



The measured starting concentration of MS2 was similar between each of the active test and inactive control runs. In the inactive control (no air purifier), there was a 0.2 log reduction due to natural decay after 60 minutes.

In the active test runs, the concentration of airborne MS 2 reduced by on average 8.2 log to 4.0 Log<sub>10</sub> MS2 PFU/m<sup>3</sup>.





### 6. Discussion

Our environmental test chamber assessment demonstrated that, when challenged with MS2, the MedicAir FOZKYGB-04 air purifier could reduce the average airborne concentration of the virus from 8.2 to 4.0 Log<sub>10</sub> PFU/m<sup>3</sup> after 60 minutes of operation. A reduction was observed in each of the triplicate active test runs.

The triplicate inactive control runs (no air purifier) did not show the same scale of reduction, the average concentration of MS2 decreased from 9.7 to 9.5 Log<sub>10</sub> PFU/m<sup>3</sup>.

Calculating the percentage reduction based on the PFU/m<sup>3</sup> results there was a greater than 99.9% reduction of airborne MS2 within 20 - 30 minutes of the air purifier operating.



## 7. References

- Tung-Thompson G, Libera DA, Koch KL, de los Reyes FL III, Jaykus L-A (2015) Aerosolization of a Human Norovirus Surrogate, Bacteriophage MS2, during Simulated Vomiting. PLoS ONE 10(8): e0134277. <u>hilps://doi.org/10.1371/journal.pone.0134277</u>
- 2) G.W. Park, K.G. Linden, M.D. Sobsey (2010) Inactivation of murine norovirus, feline calicivirus and echovirus 12 as surrogates for human norovirus (NoV) and coliphage (F+) MS2 by ultraviolet light (254 nm) and the effect of cell association on UV inactivation. Letters in Applied Microbiology (Volume 52, Issue 2, Pages 162-167). <u>hiips://doi.org/10.1111/j.1472-765X.2010.02982.x</u>
- Jung Eun Lee, GwangPo Ko (2013) Norovirus and MS2 inactivation kinetics of UV-A and UV-B with and without TiO2. Water Research (Volume 47, Issue 15, Pages 5607-5613). <u>hiips://doi.org/10.1016/j.watres.2013.06.035</u>

## 8. Appendix

	Table	e 8.1. Average PFU	/m <sup>3</sup> recovered fro	m test run sample	S
Time (minute)	Test 1	Test 2	Test 3	Average	% Reduction
0	115,151,515	224,242,424	113,787,879	115,151,515	N/A
15	4,469,697	2,060,606	3,681,818	4,469,697	97.7
30	66,667	21,212	65,152	66,667	>99.9
60	1,515	18,182	4,545	1,515	>99.9

	Table 8.2. Average PFU/m <sup>3</sup> recovered from control run samples						
Time (minute)	Control 1	Control 2	Control 3	Average	% Reduction		
0	90,117,994	14,132,475,194	160,766,962	4,794,453,383	N/A		
15	91,445,428	13,301,153,124	90,265,487	4,494,288,013	6.3		
30	39,233,038	10,297,666,935	37,463,127	3,458,121,033	27.9		
60	35,840,708	10,351,300,617	36,725,664	3,474,622,329	27.5		

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