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SUMMARY

Test Report No.: VX-TR-20-0260 Copy No.: 1

DETERMINATION OF THE VIRUCIDAL ACTIVITY – HUMAN CORONAVIRUS TEST (EN 14476) OF ARMOR 8



TITANIUM WORLD TECHNOLOGY SDN BHD (1032758-D)

No.16, Jalan Jalil Jaya 6 Block 5, Bukit Jalil 57000 Kuala Lumpur Malaysia





Product and Test Information:

Sample Name:	ARMOR 8		
Test Method:	EN 14476:2013+A1:2015 (E) Chemical disinfectants and antiseptics – Quantitative suspension test for the evaluation of virucidal activity in the medical area – Test method and requirements (phase 2, step 1)		
Product appearance:	Clear solution		
Product info:	ARMOR 8 is a unique self-applied nanotechnology coating that deodorises, sanitises, and eliminates germs, bacterias, and viruses through the process of photocatalysis. Armor 8 is an easy solution to keep you and your family safe and healthy.		
Test organism(s):	Human coronavirus, strain 229E, ATCC VR-740		
Concentration/contact time:	100.00 %* / 5 and 30 minutes		
Loading:	0.30 g/L bovine albumin solution		
Test temperature:	20 °C ± 1 °C		
Incubation period:	5 days, 36 °C ± 1 °C		
Testing method:	Quantal test		
Inactivation method:	Immediate dilution Molecular sieving using MicroSpinTM S 400 HR		

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Test results:

Table A: Evaluation of the virucidal activity of ARMOR 8 on test strains according to EN 14476

Product: ARMOR 8 Loading: 0.30 g/L bovine albumin solution

Test strain: Human coronavirus ATCC VR-740

Virus control, V _C	Cytotoxicity effect, CE	
V _{C1} : 6.38 ± 0.25	CE ₁ : 1.50 ± 0.00	
V _{C2} : 6.00 ± 0.38	CE ₂ : 1.50 ± 0.00	

Test concentration (%) / contact time (min)	First assay, N _{a1}	Second assay, N _{a2}	Average reduction
100.00*/5	N _{a1} : ≤1.50 ± 0.00 lg R ₁ : ≥4.88 ± 0.25	N _{a2} : 1.63 ± 0.25 lg R ₂ : 4.38 ± 0.45	lg R: ≥4.63 ± 0.36
100.00* / 30	N _{a1} : ≤1.50 ± 0.00 lg R ₁ : ≥4.88 ± 0.25	N _{a1} : 1.63 ± 0.25 lg R ₁ : 4.38 ± 0.45	lg R: ≥4.63 ± 0.36

Table B: Control tests and method validation for Table A

Test strain	Cell susceptibility control	Suppression efficiency control	Reference test for virus inactivation
Human coronavirus ATCC VR-740	A: 6.25 ± 0.33 A _{PBS} : 6.50 ± 0.00	B: 5.38 ± 0.25 Vc: 5.88 ± 0.37	C _{30:} ≥4.00 ± 0.00 C _{60:} ≥4.00 ± 0.00
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Note:

TCID ₅₀ :	The dilution of the virus suspension that induces a cytopathic effect (CPE) in 50 % of cell culture units.
СРЕ:	The morphological alteration of cells and/or their destruction caused by the cytopathic effect of virus multiplication.
V _C :	$\log_{10} TCID_{50}$ per ml in the viral test suspension at the beginning and at the maximum contact time.
N _a :	$\log_{10} TCID_{50}$ per ml in the test mixture at the end of the contact time.
CE:	The morphological alteration of cells caused by the cytotoxicity effect of the product test solution.
A:	log_{10} TCID $_{50}$ per ml in the cell susceptibility control as compared to PBS
B:	$\log_{10} \ TCID_{50}$ per ml in the suppression efficiency control as compared to the virus control
C:	log ₁₀ TCID ₅₀ per ml in the reference test for virus inactivation after 30 and 60 minutes (5 and 15 minutes for vacciniavirus)

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Table C: Summary of the log reductions of the quantitative suspension test according to EN 14476

Test strain	Test concentration (%) / contact time (min)	Log reduction (TCID50/ml)	Associated risk [†]
Human coronavirus ATCC VR-740	100.00* / 5	≥4.63 ± 0.36	Minimal risk of false acceptance
	100.00* / 30	≥4.63 ± 0.36	Minimal risk of false acceptance

^{*} The product can only be tested at 80.00 % concentration or less, as some dilution always occurs when test organisms and interfering substance are added.

Conclusion

ARMOR 8 showed the required virus reduction of \geqslant 4.0 log10 against test strain Human coronavirus ATCC VR-740 in accordance with EN 14476:2013+A1:2015 (E) at 100.00 %* concentration after 5 and 30 minutes under the stated condition. According to the simple acceptance decision rule†, there is a minimal risk of false acceptance. This result clearly to show that percent reduction of strain Human coronavirus ATCC is greatly reduce between 99.99% and 99.999% within 5 minutes of exposure time (please check Appendix A for better understanding).

References

- 1. Photocatalytic disinfection using titanium dioxide: spectrum and mechanism of antimicrobial activity, Applied Microbiology and Biotechnology 90(6):1847-68, (2011).
- 2. Photocatalytic inactivation of influenza virus by titanium dioxide thin film, Photochemical and Photobiological Sciences, 11: 1293-1298, (2012).
- 3. Understanding the antimicrobial mechanism of TiO2-based nanocomposite films in a pathogenic bacterium, 4: 4134-4143, (2014).
- 4. Nanomaterials for alternative antibacterial therapy, International Journal of Nanomedicine, 12: 8211–8225, (2017).
- 5. Contaminant-activated visible light photocatalysis, Scientific Reports, volume 8: Article number: 1894 (2018).
- 6. Photocatalysis could be used to inactivate coronaviruses, Photonics Media, USA 2020, (https://www.photonics.com/Articles/Photocatalysis Could Be Used to Inactivate/a65 761).

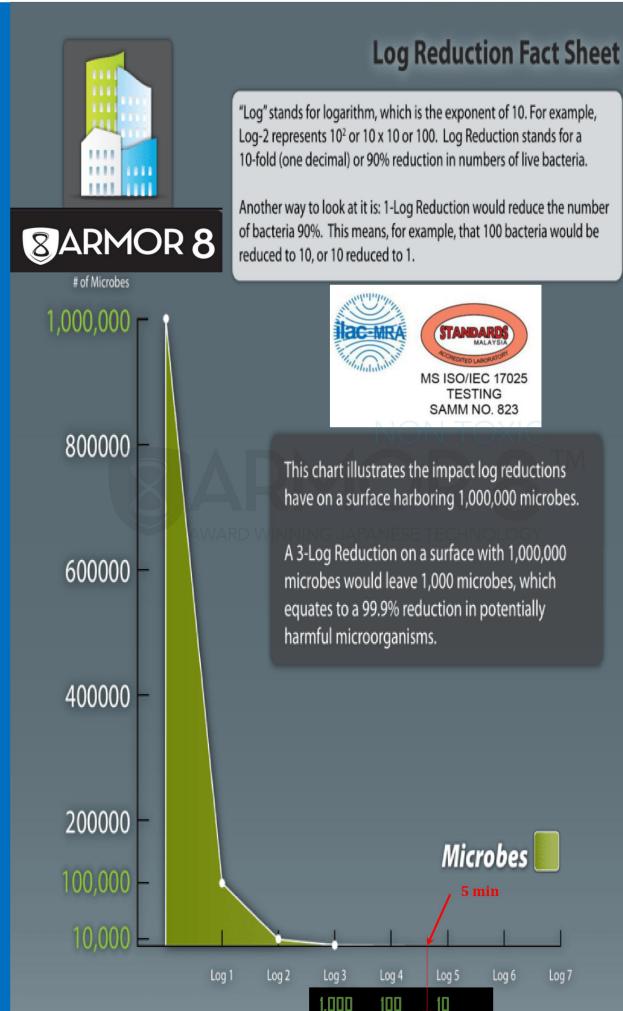
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 $[\]dagger$ The decision rule applied is simple acceptance rule with no guard band and up to 50 % risk of false acceptance or rejection. This rule has been determined by the laboratory and agreed with the client prior to testing.

APPENDIX A





99.9%

99.99%

99.999%

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