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Central Laboratory of the Intitute (SÚJCHBO, v.v.i.)

TESTING REPORT NO. 1/20/LBMO - Brno

	A	ssessment (of Decont	amination Effect of F	iltration M	aterial			
Order No.				1/20					
Order acce	epted	on:		10 June, 2020					
Ordering	g Par	ty							
Name/Lega				RESPILON Group s.r.	0.				
Registered	office			Jaselská 14, 602 00 Bri					
Testing dor	ne at:			Laboratory of Biologic Brno, National Institute					
Proxy:				-					
Party Ac	cepti	ng the Orde	r						
Testing done at:				Laboratory of Biological Monitoring and Protection in					
				Brno, National Institute for NBC Protection					
Testing dor	ne on:			28 July, 2020					
Tester:				Mgr. Kateřina Rosenbergová, PhD.					
Used metho	ods:								
Testing rep	ort iss	ued on:		23 September, 2020					
Issuer:				Mgr. Oldřich Kubíček, CSc.					
			ate Office fo	r Nuclear Safety – Prague (SÚJB)				
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Subject of Testing:

Bactericidal and viricidal effect of tested material

Testing procedure – Description

Internal procedure of LBMO – **No.:** Cultivation assessment of microbicidal effect of protective material B-MIKR-01.

Nanomaterial (material with active copper layer):

S.No. 1 Virus and bacteria deactivating layer – Respilon VKB01



Bacterial cultures:

Staphylococcus aureus (strain CCM 299) Streptococcus agalactiae (strain CCM 6187) Escherichia coli (strain CCM 3954) Pseudomonas aeruginosa (strain CCM 1960)

strains from the Czech Collection of Microorganisms at Masaryk University Brno - Faculty of Science

Virus culture:

influenza virus (A/sw/Brno/12) isolate from the Department of Infectious Diseases and Microbiology at Faculty of Veterinary Medicine – Brno

Systematic procedure:

The material was cut into 1x1 cm squares. Bacterial or viral culture respectively was applied on one square and covered with a second square of the material. The incubation of the samples was performed at room temperature for 1 min, 5 min, 30 min, and 180 min. The cultures were then eluted into distilled water.

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Bacterial cultures were then plated on blood agar (BA) plates and incubated for 24 hours at 37 °C.

7-day-old chicken embryos were infected with the virus culture and incubated for 3 days at $37 \, ^{\circ}\text{C}$.

Results:

No - full growth

Yes - 100% growth inhibition

NT - not tested

Staphylococcus aureus concentration 1,07 x $10^8 \pm 0,21$ x 10^8 CFU/ml

Growth inhibition (100 %) corresponds to the decrease in viability of at least $\underline{1,07 \times 10^6}$ times

			Incubation	period		
sample	1 min	5 min	30 min	180 min		
	Growth inhibition (100 %)					
1	no	no	no	yes		

Streptococcus agalactiae concentration 1,1 x $10^8 \pm 0.9$ x 10^7 CFU/ml

Growth inhibition (100 %) corresponds to the decrease in viability of at least $\underline{1,1} \times \underline{10^6}$ times

			Incubation	period		
sample	1 min	5 min	30 min	180 min		
•	Growth inhibition (100 %)					
1	no	no	yes	NT		

E. coli concentration 1,32 x $10^8 \pm 0,18$ x 10^8 CFU/ml

Growth inhibition (100 %) corresponds with the decrease in viability of at least $\underline{1,32 \times 10^6}$ times

			Incubation	period		
sample	1 min	5 min	30 min	180 min		
•	Growth inhibition (100 %)					
1	no	no	no	yes		

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Pseudomonas aeruginosa concentration 9,4 x $10^7 \pm 0.5$ x 10^7 CFU/ml

Growth inhibition (100 %) corresponds to the decrease in viability of at least 9.4×10^5 times

			Incubation	period	
sample	1 min	5 min	30 min	180 min	
			Growth inhibiti	on (100 %)	
1	no	no	no	yes	

Titer of influenza virus 1:256

	Incubation period					
sample	1 min	5 min	30 min	180 min		
•	Growth inhibition (100 %)					
1	no	no	yes	NT		

Growth inhibition (100 %) corresponds with the decrease in viability of at least 256 times

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Appendixes: No	
	End of Report

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