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STULV20AA1474-1	Measurement of antivition activation by means of	ral activity of PHOEBE treate visible light	d surface after	
Sponsor	COLOROBBIA Consulting Srl			
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	ITALY			
REFERENCE TEST METHOD	ISO 21702:2019 - Measurement of antiviral activity on plastics and other non- porous surfaces (method modified for the specific type and size of surface and process)			
Test Item				
PRODUCT NAME	PHOEBE filter			
MATRIX OF THE PRODUCT	treated ceramic surface			
Ватсн	Not applicable	CODE	MTNT000012	
MANUFACTURING DATE	Not applicable	EXPIRY DATE	Not applicable	
MANUFACTURER	COLOROBBIA Consulting Srl			
ACTIVE INGREDIENT	Colorobbia Photocatalyst			
MATERIAL ITEM ALIQUOT	LV-MAT-FOV7-20-101-0877:a			
PARCEL REGISTRATION N.	IP-LV-2020101-AEJ	RECEIVING DATE	April 10 th 2020	
STORAGE CONDITIONS	Room temperature			
Analysis Starting Date	May 27 th 2020	ANALYSIS ENDING DATE	June 01 st 2020	
EXPERIMENTAL CONDITION	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	4400		
TEST TEMPERATURE SPECIMEN	Room temperature (25±1°C) at ≥90%RH 2x6 cm specimen			
DESCRIPTION VIRAL INOCULUM	400 µl of viral inoculum with known viral titre - were applied onto each specimer evenly distributed. The inoculum was left adsorbing and drying onto the porous ceramic specimen at room temperature and under biosafety hood.			
PRODUCT APPLICATION	Visibile light LEDs are switched on in order to activate the treated surface in the spectra range of wave length from 400 to 600nm			
VOLUME APPLIED	NA NA			
CONTACT TIME	4 hours (±5 minutes)			
INACTIVATION OF PRODUCT RESIDUES	Dilution-neutralization in cell culture medium (no detoxification needed)			
INCUBATION TEMPERATURE	37°C ± 1°C (with 5% CO ₂)			



BioPharma Product Testing

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TEST VIRUS	Bovine Coronavirus (BCoV) - strain S379 Riems	
CELL LINE	PT cells (Ovis aries). Code: CCLV-RIE 11	
	Check of cytotoxicity of the test item The test item was not cytotoxic, i.e. its contribution in terms of CPE was not visible in the test.	
	Assay of viral infectivity (virus titration) The minimum titre of the starting viral suspension was sufficiently high to at least enable a theoretical viral titre reduction of 4 LogTCID ₅₀ .	
	Check of viral recovery (untreated and treated surface) The dose of infectious particles recovered immediately after inoculation from the untreated test specimens was within the range of 5 to 6LogTCID ₅₀ . The dose of infectious particles recovered from each untreated test specimen after contacting for 24 h wasnot higher than 3LogTCID ₅₀ .	
VALIDITY AND EFFICACY CRITERIA	Check of host cells susceptibility to virus and suppression of antiviral activity (neutralization) The difference of the average value of $TCID_{50}$ among the cellular cultures treated with the treated samples or untreated samples and then with the viral inoculum and the ones treated only with the viral inoculum (negative control) was $\leq 0.5 \text{ Log}TCID_{50}$.	
· 70 m · .	Accuracy of virus control among the three replicas The maximum difference of the value of $TCID_{50}$ among the cellular cultures treated with the viral inoculum recovered from the 3 different untreated specimen was ≤ 0.5 Log.	
	Antiviral efficacy The LogTCID $_{50}$ reduction factor (R) was calculated as per ISO 21702 :2019 standard, i.e. subtracting the average LogTCID $_{50}$ of treated specimen (At) from the average LogTCID $_{50}$ of untreated specimen (Ut) at the chosen contact time (4 hours). The LogTCID $_{50}$ was calculated by the Spearman-Karber method.	
	Bovine coronavirus is used as a surrogate virus for SARS-related viruses as it belongs to the same Betacoronavirus 1 genus and showed similar susceptibility to WHO formulations in published studies.	



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	Cytotoxicity			
RESULTS	PT cells (Ovis aries) cell destruction	≤0.50 (Log)		
	Log reductions at the different contact times			
		4 hours		
	Bovine coronavirus (Betacoronavirus 1)	Average		
		≥3.22±0.200 (Log)		
		≥99.9%		
	See Annex N.1 for the detail	of the test results		
Conclusions	The antiviral treatment causes a complete viral contact time in the adopted test conditions.	titre reduction after 4 hours of		
Annex	N. 1: Raw Data Elaboration			
TEST FACILITY MANAGER	Mi Caust 16	106/2		

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The test results relate only to the tested items. Sampling, except specific indication on test report, is always intended to be made by the Sponsor. Characterization of the test sample is under Sponsor responsibility.