

EVALUATION OF VIRUCIDE ACTIVITY OF A UV DEVICE

Aim of the Study

The aim of this study is to determine the virucidal activity of the device BAIRY Air Purifier, produced for RIDI Leuchten GmbH, against SARS-CoV2.

The aforementioned device consists of a system with porous filter separated by a support with 4 UV lights.

Virus was positioned on the filter in three position irradiated with UV light.

Method

On the three positions are $100\mu L$ of a viral suspension and positioned in correspondence with UV light.

After a contact time of 5′, 1h, 4h we tested the residual virus activity by evaluating by Tissue Culture Infective Dose 50% (TCDI₅₀).

Name of product	BAIRY Air Purifier
Period of analysis	14/01/21 – 17/01/21
Temperature of incubation	37°C
Identification of Viral strain	SARS-CoV-2_COV2019 ITALY/INMI1
Contact time	See protocol



Contact Time Protocol

5 min	3 repetitions
1 hour	3 repetitions
4 hour	3 repetitions

All repetitions were tested for SARS-CoV-2 concentration by $TCDI_{50}$ using VERO E6 C1008 (ATCC CRL-1586) cell line.

Results

Suspension virus used $10^{6.25}$ TCID₅₀ /mL (6.25 expressed by Log)

Value of Log $TCDI_{50} = 1.50$ (Note that this value means total viral inactivation)

	Results	
Cytotoxicity	No Cytotoxicity	
	observed	

	Time	Media Log TCDI ₅₀	% reduction against control
T0		4.75	
TEST	5 min	1.50	99.94
	1 hour	1.50	99.94
	4 hour	1.50	99.94



Conclusions

The purpose of the study was to determine the virucidal efficacy of the device BAIRY Air Purifier, produced for RIDI Leuchten GmbH, against SARS-CoV-2_COV2019 ITALY/INMI1 at a contact time of 5 minutes, 1 hour and 4 hours at room temperature.

The evaluated test device demonstrated an average of 3.25 Log10 reduction in viral titer (99.94% reduction) after exposition of 5 minutes of exposure.

The Plate Recovery Control (T0) demonstrated a viral titer of 4.75 log10 TCID50 per 1 ml.

No test substance cytotoxicity was detected.

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