



**Laboratory**  
of **Viruses** Contaminants  
of **Water** and **Food**



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### Summary Report

Evaluation of the efficiency of the air disinfection unit WADU-02, WELLIS (Wellis Co., Ltd.) against Human Respiratory Syncytial Virus under wet conditions

Report nº: 20191212\_3

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**REPORT Núm. 2019121203**

**Applicant:** RECO PLANT Co., Ltd  
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### **Product evaluation**

**Product Description:** Air Disinfection unit  
**Model number:** WADU-02  
**Brand:** WELLIS  
**Manufacturer:** Wellis Co., Ltd.  
**Issue date:** 12/9/2019

### **Summary**

The effectiveness of the WELLIS WADU-02 air disinfection unit for the disinfection of viruses was measured against Human Respiratory Syncytial Virus (RSV). The inactivation or decay of infectious RSV was quantified using cell culture (TCID<sub>50</sub> assay) and all tests were done in duplicate. Wet viral suspensions were exposed to the disinfection unit in order to test virus stability over time. Control viral suspensions, not exposed to the disinfection unit, were tested in parallel. The disinfection treatment was able to reduce 99% the initial concentration of RSV after 2 hour of treatment.

### **Experimental procedure**

RSV is an enveloped RNA virus with surface proteins that mediate RSV infection of human airway epithelial cells. RSV is the leading viral cause of acute lower respiratory tract infections, including bronchiolitis and pneumonia, among infants and young children globally. RSV can survive for many hours on hard surfaces such as tables and crib rails. It typically lives on soft surfaces such as tissues and hands for shorter amounts of time. It is usually transmitted through droplets from the cough or sneeze that contact with eyes, nose, or mouth, or by direct contact with a contaminated surface. For this test RSV strain A2 (ATCC® VR-1540TM) was produced in Hep2 cells (ATCC® CCL-23TM).

This experiment was performed for RSV under wet conditions. The air disinfection unit was stored in a metacrilate box (0,064 m<sup>3</sup>). All experiments were conducted at room temperature. One-hundred microliter droplets were disposed over small pieces of glass and placed inside or outside the box (control) as it is shown in picture 1.



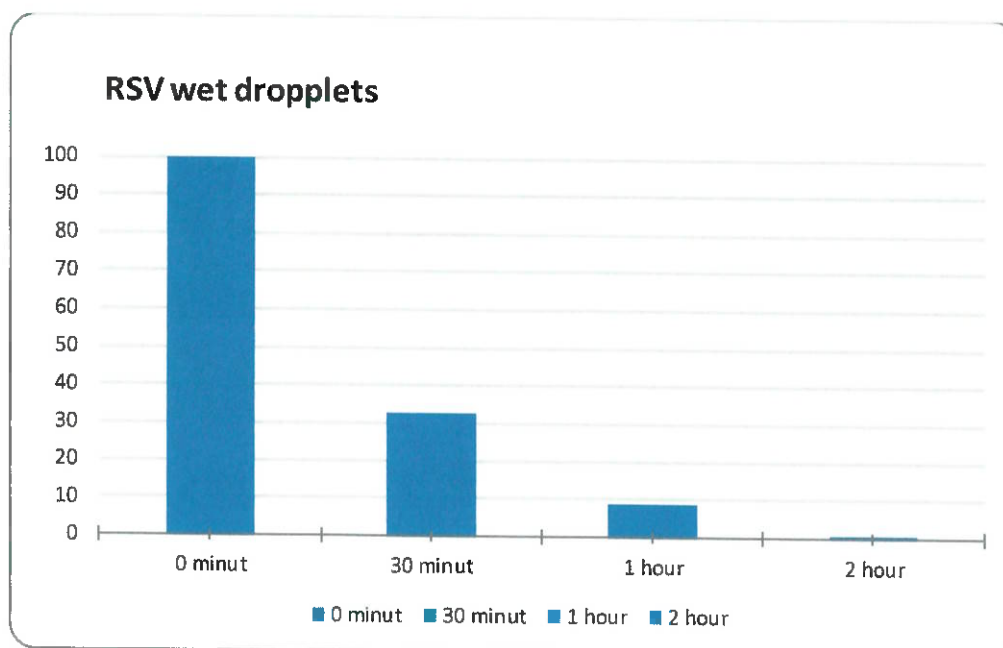
**Picture 1: Droplets disposed over small glass pieces.**

At each testing time viruses were recovered, from the glass surfaces, in culture medium (MEM) and the number of infectious viral particles were quantified by TCID<sub>50</sub> in Hep2 cells.

The inactivation effectiveness of the air disinfection unit over RSV wet suspensions are summarized in table 1 and figure 1. Viral quantities are expressed in logarithms

	Time	No treatment	Air disinfection unit (ozone + <i>d</i> -limomene)	log <sub>10</sub> decay	% of decay
WET	0 minut	3,58E+05	3,58E+05		
	30 minut	2,60E+05	8,08E+04	0,48	67%
	1 hour	2,82E+05	5,00E+04	1,26	91%
	2 hour	1,77E+05	6,55E+03	2,19	99%

**Table 1: RSV concentration decay over time under wet conditions.**



**Figure 1: Percentages in RSV concentration over time under wet conditions.**

### Conclusion

The equipment, significantly reduce the concentration of RSV in wet droplets. This virus presented in 2 hours a total decay of  $2,19\log_{10}$ . The efficiency of WELLIS WADU-02 on aerosols receiving equivalent doses could be expected to be at least equivalent.

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