

William S. Lawrence, PhD

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Jeffrey Mullen **Chief Executive Officer** Dynamics Inc. 493 Nixon Road Cheswick, PA 15024

Dear Mr. Mullen:

My laboratory is located in the Galveston National Laboratory at the University of Texas Medical Branch, a facility constructed with funds from the National Institute of Health, National Institute of Allergy and Infectious Disease.

On August 14, 2020, we tested the Nanowave device developed by Dynamics Inc. to measure inactivation of aerosolized SARS-CoV-2. SARS-CoV-2 (WA-01 isolate) was aerosolized and moved through the Nanowave device at the maximum airflow capability of our laboratory. Virus was inactivated to the detection limits of the test. Details of the SARS-CoV-2 inactivation test are attached hereto as Appendix A.

Sincerely,

William S. Lawrence, Ph.D.

Assistant Professor, Microbiology & Immunology

Manager, Aerobiology Service Center University of Texas Medical Branch

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## APPENDIX A

## SARS-COV-2 Inactivation With Dynamics Inc. Nanowave Technology

Test	Test	Nebulizer	Biosampler 1	Biosampler 2	Average
	Duration		(Before Device)	(After Device)	Humidity
Nanowave device at 10 LPM when ON	10 min	2.5E+05	7.5E+03	<50	88.10%
Nanowave device at 30 LPM when ON	10 min	7.4E+04	NA	<50	55.23%
Nanowave device at 10 LPM when OFF	10 min	7.5E+05	5.0E+04	7.5E+03	82.06%
Nanowave device at 30 LPM when OFF	10 min	7.5E+05	NA	7.5E+03	69.46%

## **Test Notes**

- SARS-CoV-2 virus utilized was WA01 Strain.
- Biosampler 1 was placed before the Nanowave Device and is the control sample.
- Biosampler 2 was placed after the Nanowave Device.
- To achieve 30LPM, biosampler 1 was removed. Result was determined based on other correlated data.
- Values provided as plaque forming units / ml (PFU/ml).
- Humidity was not controlled but was sampled 123 times during each test.