**From:** **Gary Cohen** gcohen@iuva.org
**Subject:** EUAs for Decontamination and Bioburden Reduction Systems for Surgical Masks and Respirators
**Date:** September 23, 2020 at 2:26 PM
**To:** jeff.shuren@fda.hhs.gov
**Cc:** Denise.Hinton@fda.hhs.gov, Judith.McMeekin@fda.hhs.gov, Richard Martinello richard.martinello@yale.edu, Malley, Jim jim.malley@unh.edu, Troy Cowan troy@visionbasedconsulting.us, Ron Hofmann ron.hofmann@utoronto.ca, CDRH-COVID19SurgicalMasks@fda.hhs.gov

Dear Dr. Shuren,
On behalf of the International Ultraviolet Association (IUVA), we respectfully submit the attached "FDA N95 Mask Emergency Use Authorization Requirements – Questions for the FDA on the Requirements for UV Testing." In addition, please find a supporting cover letter from Dr. Richard A. Martinello, MD, Yale School of Medicine/Yale New Haven Hospital; and James P. Malley, Jr., Ph.D., University of New Hampshire—who helped lead this effort for IUVA. Please contact me if we may provide further information; we welcome the opportunity to discuss the use of UV with the FDA.

Thank you,

gary

Gary Cohen
Executive Director
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