

Antimicrobial proof-of-concept: a novel device combining UV light and ozone for human skin antisepsis



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INTRODUCTION

- · Hand hygiene (HH) remains a cornerstone for the prevention of microbial transmission, both in the community and healthcare setting
- While hand-washing with plain soap and warm water remains the "gold standard" for HH, the process is less than standardized in practice (10 sec to 2 min) and the microbial reduction variable (0.5 to 3.3 log reduction)
- · Alcohol-based hand rubs (ABHR) and wipes offer an ostensibly more rapid and convenient alternative with satisfactory efficacy against vegetative microbes (up to 3 log reduction), but the effect against Clostridioides difficile spores limits their use
- Ultraviolet (UV) light has well-established antimicrobial effect (AME), including sporicidal activity
- Likewise, ozone (O₃) has potent AME and is commonly utilized in the field of dentistry
- In this study, we report proof-of-concept data on the AME of a novel investigational device, HyLuxO3 (GMI, LLC, Nashville, TN, USA; Fig. 1), that is engineered to synergistically combine UV-C light energy and high velocity O₃ airflow to not only decontaminate environmental surfaces and fomites, but also to safely achieve human skin antisepsis within regulatory limits

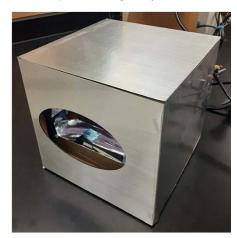


Figure 1: HvLuxO3 prototype device, 222 nm UV-C light beams are directed centrally from above and below at a fixed UV irradiance (0.6 mW/cm2), [O₃] output (0.1 ppm), and sample-to-device distance (5 cm). The only user-defined variable for this prototype device is exposure time.

METHODS

- HyLuxO3 was tested on LB agar to titrate device variables to ascertain intensities for optimal AME: later testing was performed on VITRO-SKIN (Florida Suncare Testing, Bunnell, FL), a human glabrous skin surrogate
- ATCC strains of MRSA, Staphylococcus epidermidis, Klebsiella pneumoniae, Escherichia coli, Pseudomonas aeruginosa, and Candida albicans were used to test AME vs. vegetative microbes; Bacillus atrophaeus spores were used as a surrogate for C. difficile
- Tested variables included time under device, [O₃], airflow velocity, 222 and/or 254 nm UV light, sample distance from UV lamp, and UV beam width. Positive controls were used to calculate log-kill curves for AME

| wicrobe (exposure time) | CFC inoculated onto | Cr C anter | Cro alter | Log-Kill (illean) |
|-------------------------|---------------------|-----------------|--------------------|-------------------|
| | plate | exposure (mean) | exposure (std dev) | |
| B. atrophaeus (5 sec) | 2,000,000 | 88 | 52 | 4.36 |
| B. atrophaeus (30 sec) | 2,000,000 | 14 | 8.7 | 5.14 |
| B. atrophaeus (60 sec) | 2,000,000 | 6.7 | 5.6 | 5.47 |
| C. albicans (5 sec) | 160,000 | 2.1 | 1.2 | 4.88 |
| C. albicans (30 sec) | 160,000 | 0.4 | 0.5 | 5.19 |
| C. albicans (60 sec) | 160,000 | 0 | 0 | 5.20 |
| E. coli (5 sec) | 2,000,000 | 7.3 | 5.6 | 5.44 |
| E. coli (30 sec) | 2,000,000 | 1.3 | 1.4 | 6.19 |
| E. coli (60 sec) | 2,000,000 | 1.0 | 1.3 | 6.30 |
| K. pneumoniae (5 sec) | 3,200,000 | 18 | 13 | 5.26 |
| K. pneumoniae (30 sec) | 3,200,000 | 5.2 | 3.8 | 5.79 |
| K. pneumoniae (60 sec) | 3,200,000 | 2.0 | 1.9 | 6.20 |
| MRSA (5 sec) | 16,000,000 | 42 | 24 | 5.58 |
| MRSA (30 sec) | 16,000,000 | 7.3 | 12 | 6.34 |
| MRSA (60 sec) | 16,000,000 | 1.3 | 1.4 | 7.09 |
| P. aeruginosa (5 sec) | 2,400,000 | 17 | 9.7 | 5.14 |
| P. aeruginosa (30 sec) | 2,400,000 | 7.9 | 4.4 | 5.48 |
| P. aeruginosa (60 sec) | 2,400,000 | 3.1 | 1.8 | 5.89 |
| S. epidermidis (5 sec) | 4,000,000 | 13 | 5.3 | 5.48 |
| S. epidermidis (30 sec) | 4,000,000 | 1.8 | 3.8 | 6.35 |
| S. epidermidis (60 sec) | 4,000,000 | 0.9 | 2.1 | 6.60 |

- Log-kill data for the HyLuxO3 device against spores and vegetative microbes as a function of exposure time are presented in Table 1
- 4.36 log-kill against spores was achieved in 5 sec; a >5 log-kill was achieved by extending sample exposure to 30 sec (Fig. 2)
- >5 log-kill against all vegetative bacteria was achieved in 5 sec; ≥6 log-kill was achieved by extending sample exposure to 30-60 sec



Figure 2: HvLuxO3 device AME vs. B. atrophaeus spores. Plates were inoculated with 2.0 x 106 CFU. Positive control (bottom) was not exposed; the 5 sec (left), 30 sec (top), and 60 sec (right) plates were exposed to the device for their respective time durations.

CONCLUSIONS

- HyLuxO3 combines 222 nm UV-C light and O₃ to achieve >4 logkill against spores in 5 sec and >5 log-kill against spores in 30 sec
- This device showed a >5 log-kill against common pathogenic vegetative microbes in 5 sec and >6 log-kill in 30-60 sec.
- These AME results rival those of hand-washing and ABHRs by 2-4 logs while decreasing the necessary exposure time from minutes to seconds. Moreover, similar efficacy was shown on agar as on VITRO-SKIN, a well-studied glabrous human skin surrogate
- · Studies on human hands are needed to confirm the efficacy and safety of HyLuxO3 under OSHA and EPA regulations

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