

Conversion of Existing UVB Phototherapy Units to UVC Germicidal Chambers for N95 Decontamination: Lessons Learned

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To the Editor:

SHORTAGES OF PERSONAL PROTECTIVE EQUIPMENT (PPE) during the COVID-19 pandemic have necessitated development of novel decontamination procedures to facilitate N95 mask reuse.¹ Ultraviolet (UV) germicidal irradiation has demonstrated efficacy against viruses, including SARS-CoV-2, at 1 J/cm² and a peak UVC germicidal wavelength of 254 nm.² The following question arises: how can UVC best be delivered in the field to enable mask reuse? UVA and UVB phototherapy units are available in dermatology clinics worldwide, making UVC conversion an attractive option for N95 decontamination. Daavlin 4Series phototherapy units have been successfully converted into a UVC device currently under FDA consideration for commercial purchase for \$4,000 per unit.¹ Even if approved, this device is limited by cost, size, and its ability to deliver UV irradiation from one side only. Efforts to convert existing phototherapy units into UVC devices are, therefore, warranted, particularly in times of high demand or in low-resource settings. We explored direct conversion of a full body phototherapy unit for UVC decontamination by replacement of UVB bulbs with UVC bulbs currently available in the United States.

Between May and June 2020, we conducted a systematic search of existing phototherapy units available in dermatology practices in the United States based on manufacturers listed by the National Psoriasis Foundation.³ Based on its market share, we focused on converting the wavelength of the Daavlin 3SeriesX UVB photobooth. We then conducted a systematic search for existing UVC bulbs from major lighting manufacturers that closely matched specifications

of the UVB bulb used in the Daavlin 3SeriesX booth (Philips® UVB NarrowBand TL 100W/01).^{4,5} Specific search criteria included bulb length, pin form factor, and wattage (Fig. 1).

Our search identified the Philips TL-D 95W HO bulb as the most promising replacement option currently marketed in the United States based on the aforementioned criteria. However, lack of concordance in some key parameters poses challenges. The first is a difference in bulb length, with the Daavlin 3SeriesX requiring 70 inch (177.8 cm) bulbs and the closest UVC substitute bulb measuring 60 inches (152.4 cm). The second is a difference in electrical pinout (R17D recessed double contact vs. G13 medium bi-pin). A third challenge is that Daavlin 3SeriesX booths use special ballasts, making it difficult to establish the electrical compatibility of substitute bulbs. An adapter could be fabricated to interconvert pins and extend shorter bulbs, but this likely voids electrical safety/product certifications, precluding simple UVB to UVC conversion.

Our investigation suggests that there is no convenient solution for retrofitting UVC bulbs into the Daavlin 3SeriesX UVB photobooth using inventories of commercially available bulbs. Given the likelihood of lasting PPE shortages throughout the pandemic, maximizing the utility of these pre-existing UV devices requires one or more of the following (1) manufacturing germicidal UVC bulbs compatible with existing phototherapy units, (2) developing engineered solutions for retrofitting available UVC bulbs into the existing sockets on Daavlin phototherapy units, and (3) identifying phototherapy units from other manufacturers that might more easily accommodate UVC

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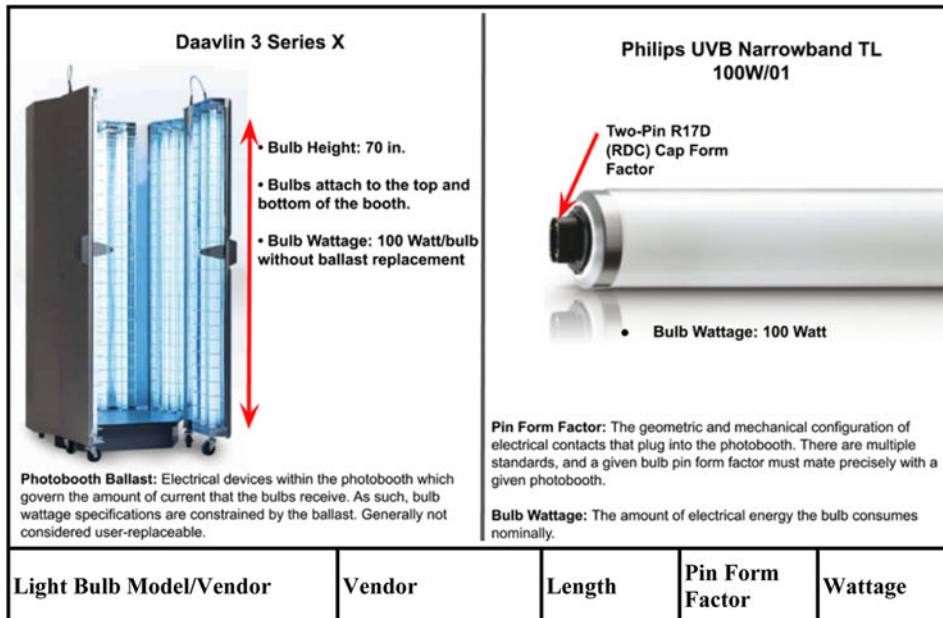


FIG. 1. The parameters, and their respective definitions, of the Daavlin 3SeriesX UVB4 photobooth and its original bulb, the Philips® UVB NarrowBand TL 100W/013. Search of UVC bulb candidates. Commercially available bulb parameters compatible with our design requirements are depicted in green, and incompatible ones are depicted in red. The original bulb corresponding to the Daavlin 3SeriesX UVB photobooth is shown in the top row, with all valid parameters. The most promising candidate bulb with the closest matching parameters is highlighted in orange.

bulbs. In all three cases, collaboration with manufacturers is required; we call on them to collaborate with interested academic medical centers.

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