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Germicidal Efficacy and Mammalian Skin Safety of 222-nm UV Light

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INTRODUCTION

The use of ultraviolet (UV) light for inactivating bacteria and viruses is well established (1, 2). However, UV radiations emitted by typical germicidal lamps with a peak emission at 254 nm represent a human health hazard, causing skin cancer (3, 4) and cataracts (5, 6).

We have developed an approach to kill bacteria without harming human cells in skin tissue models (7) and mouse skin in vivo (8) that employs single-wavelength UVC light generated by inexpensive filtered excilamps (9). The approach is based on the limited penetration distance of UVC light in the wavelength range of 200–222 nm in biological samples. Specifically, while far-UVC light has enough range to traverse microbes that are much smaller in size than human cells [less than 1 μm in diameter (10, 11), compared to the diameter of typical human cells ranging from about 10–25 μm (10)], it is strongly absorbed by the proteins in the cytoplasm of human cells (12, 13) and is drastically attenuated before reaching the human cell nucleus. It follows that far-UVC light is not able to penetrate the stratum corneum of skin and reach the underlying critical basal cells or melanocytes (4).

Another organ especially sensitive to UV damage is the lens; however, the lens is positioned distal to the cornea that is sufficiently thick [~500 μm (14)]. Therefore, penetration of far-UVC ~200-nm light through the cornea to the lens is predicted to be essentially zero (15).

The potential use of UVC light for microbe sterilization purposes in the presence of humans paves the way to numerous clinical applications, including reduction of surgical site infections (SSI) that are the second most common healthcare-associated infections resulting in readmissions, prolonged hospital stays, increased morbidity and mortality, and an overall higher medical cost (16, 17). A key factor contributing to the severity of SSI is the incidence of drug-resistant bacteria such as methicillin-resistant Staphylococcus aureus (MRSA) (18, 19).

To address the issue of reducing SSI, we have developed an approach that involves the use of inexpensive excimer lamps that, appropriately filtered, emit monoequivalent wavelengths in the far-UVC range. A crucial property of
UVC-mediated germicidal killing is that it is essentially independent of acquired drug resistance (20, 21).

We have previously shown that 207-nm light emitted by a filtered krypton-bromine (Kr-Br) excilamp has bactericidal efficacy while being minimally cytotoxic to human cells in a 3D skin tissue model in vitro (7) and in a hairless mouse skin model in vivo (8). Thus, continuous exposure of the wound to far-UVC light during surgery may inactivate the microbes alighting directly onto the surgical wound from the air. If proven to be safe to eyes as well as skin, continuous operation of far-UVC light would not require the use of cumbersome protective clothing, hoods and eye shields for the surgical staff and the patient (22, 23).

Here we extended those studies to a filtered krypton-chlorine (Kr-Cl) excimer lamp that produces essentially monoenergetic UV light at 222 nm and established that a wavelength window exist in the far-UVC region (from 200–222 nm) that inactivates bacteria efficiently but is not cytotoxic or mutagenic to mammalian cells.

We describe measurements of MRSA survival and of typical UV-induced premutagenic DNA lesions in a 3D human skin model immediately after exposure to different fluences of 222-nm light. We compared the results to those measured in samples exposed to similar fluences from a typical germicidal lamp emitting at 254 nm.

We further tested eight cellular and molecular endpoints relevant to skin damage in vivo in dorsal skin of hairless mice 48 h after exposure to 222- or 254-nm light used as positive control (8), at a fluence at which the 254-nm light is predicted to produce a significant decrease in SSI rates (22).

In agreement with our previous results using 207-nm light (7, 8), here we showed that 222-nm light has similar antimicrobial properties as a conventional germicidal lamp but without causing mammalian skin damage.

The finding that a far-UVC wavelength window (200–225 nm) is differentially cytotoxic to bacteria relative to mammalian cells is novel and can be used for various applications that would require the presence of humans, including room sterilization and reduction of surgical site infections, without the need of additional personal protective equipment.

MATERIALS AND METHODS

UV Lamps

We used an excimer lamp based on a krypton-chlorine (Kr-Cl) gas mixture that emits principally at 222 nm. The lamp (High Current Electronics Institute, Tomsk, Russia) was air cooled with a 6,000-mm² exit window (24). A custom bandpass filter (Omega Optical, Brattleboro, VT) was used to remove essentially all but the dominant 222-nm wavelength emission.

A UV spectrometer (Photon Control, BC, Canada) sensitive in the wavelength range from 200–360 nm was used to characterize the wavelength spectra emitted by the excimer lamp, and a deuterium lamp standard with a NIST-traceable spectral irradiance (Newport Corp, Stratford, CT) was used to calibrate the UV spectrometer.

Studies were also carried out with a conventional mercury germicidal lamp (Hygare, Atlantic Ultraviolet Corporation, Hauppauge, NY) with peak emission at 254 nm and used as positive control. An ILT1400 sensor equipped with a SEL220 detector (International Light Technologies, Peabody, MA) was used to measure the fluence rate from both lamps. For acute exposures (MRSA and 3D tissues), the 222- and 254-nm lamps were positioned 9 and 99 cm, respectively, from the sample at which corresponded a power density of 0.036 mJ/cm². For chronic exposures of mice, delivery of 157 mJ/cm² in a 7 h period was obtained by locating the 222- and the 254-nm lamps 41 and 205 cm from the samples, respectively; the corresponding power densities were 0.0062 mJ/cm² for the 222-nm lamp and 0.0008 mJ/cm² in the case of the conventional germicidal lamp.

Figure 1 shows the measured spectra emitted from the Kr-Cl excimer lamp that, together with the main excimer emission at 222 nm, also includes lower fluences of higher wavelength light (i.e. ~237 and ~258 nm). To remove these more penetrating higher wavelengths and therefore potentially more harmful to human cells, we used a custom bandpass filter (Omega Optical, Brattleboro, VT). As shown in the inset in Fig. 1, the filtered lamp effectively emitted only the characteristic single wavelength of 222 nm.

IN VITRO STUDIES

MRSA Cell Survival

We used methicillin-resistant S. aureus (MRSA USA300, multilocus sequence type 8, clonal complex 8, staphylococcal cassette chromosome mec type IV), which is a major cause of both community and nosocomial infections (25–27). Survival of MRSA as a function of UVC fluence was assessed immediately after exposure with the standard colony forming unit (CFU) assay, as previously reported (7).

UV-Associated Premutagenic DNA Lesions in a 3D Human Skin Model

We used a 3D human skin model EpiDerm-FT (MatTek Corp., Ashland, MA) consisting of stratum corneum and 8–12 human cell layers to reproduce human epidermis and dermis (28). We measured induction of the two most abundant premutagenic DNA lesions in the epidermis,
cyclobutane pyrimidine dimers (CPD) and pyrimidine-pyrimidone 6-4 photoproducts (6-4PP) (4), as a function of UVC fluence in 3D human skin constructs immediately after exposure using the immunohistochemical approach previously described (7).

**IN VIVO STUDIES**

**Mouse Irradiations**

We used six- to eight-week-old male hairless mice (SKH1-Elite strain 477, Charles River Laboratories, Stone Ridge, NY); the strain has UV action spectra for histological, physical and visible changes similar to those of human skin (29, 30). Moreover, the typical thickness of the stratum corneum of SKH1 mice (5–10 μm) (31, 32) is comparable to that of human skin (5–20 μm) (33).

One group of three mice was exposed to fluence of 157 mJ/cm² from 222 nm light delivered in a 7 h period by a filtered Kr-Cl excimer lamp while another group of three mice was sham irradiated to zero UV fluence. Mice exposed to 157 mJ/cm² delivered in 7 h period from 254 nm light represented the positive controls (8).

All animal procedures were carried out in accordance with federal guidelines and protocols approved by the Columbia University Medical Center IACUC.

**Mouse Skin Safety-Specific Endpoints**

Following the same protocols described in our previous mouse skin safety study (8), 48 h after UV exposure we measured the following endpoints: 1. Epidermal thickness measured in hematoxylin and eosin stained samples; 2. The percentage of proliferating keratinocytes expressing the Ki-67 antigen (34, 35); 3. Yields of UV-induced cyclobutane pyrimidine dimers (CPD) and 6-4 photoproducts (6-4PP); 4. The number of neutrophils and mast cells as markers of skin inflammation (36); 5. The expression of keratin (K) K6A as marker of skin differentiation (37, 38).

Tissues were examined with an Olympus IX70 microscope equipped with a Photometrics® PVCAM high-resolution, high-efficiency digital camera; Image-Pro Plus 6.0 software and Fiji/Image J software were used to analyze the images. For each mouse, each endpoint was measured in at least six randomly selected fields of view.

**Statistical Analysis**

Comparisons of mean values between treatment groups and controls were performed using Student’s t test, and comparison of proportions were assessed with standard χ² tests.

**IN VITRO RESULTS**

**MRSA Survival**

We measured MRSA survival immediately after exposure to different fluences of 222-nm light generated by a filtered Kr-Cl excimer lamp (Fig. 2). We compared the results to survival of MRSA exposed to similar fluences from a conventional germicidal UV lamp (254 nm) (7). As illustrated in Fig. 2, at relatively high fluences at which the 254-nm light is predicted to produce a significant decrease in SSI rates (1), 222-nm light kills MRSA almost as efficiently as a conventional germicidal UV lamp. However, 254-nm light is almost equally efficient at killing human cells (7).

**Induction of Premutagenic DNA Lesions in Human Skin**

Figure 3 shows the comparison of the induced yields of CPD (left panel) and 6-4PP (right panel) in 3D skin tissue models exposed to 222-nm or 254-nm light. Unlike conventional germicidal light (254 nm), exposure to the 222-nm light at any of the studied fluences did not induce yields of either lesions that was significantly higher than the sham-irradiated samples.

**IN VIVO RESULTS**

**Epidermal Thickness and Keratinocyte Proliferation**

At 48 h after exposure, fixed dorsal skin sections were stained with hematoxylin and eosin (H&E) for analysis of epidermal thickness (Fig. 4A). As shown by the typical H&E-stained cross-sections of dorsal skin of sham-exposed mice (top panel), of mice exposed to 157 mJ/cm² from 254-nm light (middle panel) or 222-nm light (bottom panel), we found that, unlike the 254-nm light, the epidermal thickness of skin of mice exposed to the 222-nm excimer lamp was not statistically different from skin of unexposed mice (P = 0.18) (Fig. 4B and Table 1).

Figure 4C shows representative cross-sectional images of skin samples comparing Ki-67 expression in sham-exposed mice (Fig. 4A, top panel), in skin of mice exposed to 157 mJ/cm² from 254-nm light (middle panel) or 222-nm light (bottom panel). We found that Ki-67 expression in keratinocytes of skin exposed to the 222-nm excimer lamp was not statistically different from control (P = 0.52) (Fig. 4D and Table 1). In contrast, the same fluence from the 254-nm germicidal lamp triggered keratinocytes hyper-proliferation (Ki-67 expression increased by 2.5-fold compared to controls, P < 0.0001) (Fig. 4D and Table 1); this was consistent with severe tissue hyperplasia induced by 254-nm light as measured by a ~3-fold increase in epidermal thickness (P < 0.0001) (Fig. 4B and Table 1).

**Premutagenic UV-Associated DNA Lesions**

Figure 5A shows representative cross-sectional images of sham-irradiated mice or skin of mice exposed to 157 mJ/cm² from 222-nm light or 254-nm light, comparing premutagenic skin lesions CPD (Fig. 5A, top panels) and 6-4PP (Fig. 5A, bottom panels). In agreement with our
previous findings (7, 8), we measured no increase in UV-induced DNA photodamage in 222-nm exposed skin relative to controls (P = 0.18 for CPD and P = 0.41 for 6-4PP). On the contrary, exposure to the same fluence from a conventional germicidal lamp (254 nm) caused a dramatic increase of both lesions relative to respective controls (P < 0.0001) (Fig. 5B and C, respectively, and Table 1).

**Skin Inflammation**

To assess UV-induced cell inflammation, we measured the density of the markers mast cells and neutrophils [i.e., myeloperoxidase (MPO)-positive cells] (Fig. 6). Although 254-nm light caused a statistically significant increase of both markers (P < 0.0005) (Fig. 6 and Table 1), the density of mast cells and neutrophils induced by the 222-nm light (Fig. 6A and B, respectively) was not statistically distinguishable from controls (P = 0.68 and P = 0.59, respectively).

**Skin Differentiation**

We measured keratinocyte differentiation as keratin (K) 6A expression in skin of mice exposed to 222-nm light and compared the results to K6A expression in sham-exposed mice and in mice exposed to 254-nm light used as positive control. Figure 7 illustrates typical cross-sectional images of sham-exposed mice (Fig. 7, top panel), skin of mice exposed to 157 mJ/cm² from 254-nm light (Fig. 7, middle panel) or from 222-nm light (Fig. 7, bottom panel). The expression level of K6A in 222-nm exposed skin was not statistically different from controls (P = 0.22) (Table 1). This is in contrast with a ~3-fold increase of newly synthesized K6A in samples exposed to light from a conventional germicidal lamp (P < 0.0001).
FIG. 4. Epidermal thickness and keratinocyte proliferation in mouse skin exposed to UVC light. Panel A: Representative cross-sectional images of H&E-stained mouse dorsal skin comparing the epidermal thickness in sham-exposed mice (top panel), in mice exposed to 157 mJ/cm² from 254-nm light (middle panel) or from 222-nm light (bottom panel). Panel B: Quantification of epidermal thickness. Panel C: Ki-67-positive keratinocytes (dark brown-stained cells) in typical cross-sections of skin of sham-exposed mice (top panel), of mice exposed to 254-nm light (middle panel) or to 222-nm light (bottom panel). Panel D: Quantification of the percentage of keratinocytes expressing Ki-67 antigen. Values represent the average ± SD of epidermal thickness measured in nine randomly selected fields of view per mouse (n = 3). *Values from elsewhere (8).

TABLE 1
Summary of the Combined Results for Each Endpoint

<table>
<thead>
<tr>
<th>Endpoint</th>
<th>Controls</th>
<th>222 nm</th>
<th>222 nm P value</th>
<th>254 nm*</th>
<th>254 nm P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Epidermal thickness (μm)</td>
<td>21.0 ± 0.8</td>
<td>20.1 ± 0.4</td>
<td>0.18</td>
<td>60.9 ± 7.4</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Ki-67 positive keratinocytes (%)</td>
<td>14.2 ± 2.6</td>
<td>15.8 ± 2.7</td>
<td>0.52</td>
<td>36.1 ± 7.7</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>CPD positive keratinocytes (%)</td>
<td>0.43 ± 0.01</td>
<td>0.47 ± 0.03</td>
<td>0.18</td>
<td>52.3 ± 5.6</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>6-4PP positive keratinocytes (%)</td>
<td>0.50 ± 0.06</td>
<td>0.45 ± 0.03</td>
<td>0.41</td>
<td>31.1 ± 1.4</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Density of mast cells (number/cm²)</td>
<td>28.5 ± 11.9</td>
<td>30.6 ± 14.2</td>
<td>0.68</td>
<td>86.6 ± 16.8</td>
<td>&lt;0.0005</td>
</tr>
<tr>
<td>Density of MPO positive cells (number/cm²)</td>
<td>24.2 ± 9.1</td>
<td>27.2 ± 5.7</td>
<td>0.59</td>
<td>202.4 ± 11.3</td>
<td>&lt;0.0005</td>
</tr>
<tr>
<td>K6A optical density (%)</td>
<td>15.9 ± 5.7</td>
<td>18.9 ± 6.7</td>
<td>0.22</td>
<td>45.8 ± 6.9</td>
<td>&lt;0.0001</td>
</tr>
</tbody>
</table>

* Values from elsewhere (8).
Here we showed that 222 nm is essentially equi-effective at killing antibiotic-resistant bacteria as conventional germicidal UV lamps (254 nm). However, compared to the latter, 222-nm light does not induce typical UV-associated premutagenic DNA lesions in human keratinocytes in a 3D human skin model and appears to be safe for skin of exposed hairless mice, as assessed by eight cellular and molecular endpoints associated with damaged skin. The results are consistent with previous in vitro and in vivo findings using a filtered krypton-bromine lamp that emits light at 207 nm (7, 8); this suggests that there is a range of wavelengths, specifically between 200 and 222 nm, which are equitoxic to bacteria as typical germicidal lamps emitting at 254 nm, but without associated skin damage risks.

The lack of induced damage is due to the limited penetration of far-UVC light in biological samples (39): while light in the 200-222 nm region can traverse microbes that are much smaller in size (<1 μm) (10, 11, 40) than a typical mammalian cell (~10–25 μm in diameter) (10), it cannot penetrate mammalian cells cytoplasm (7) as well as all tissues with a stratum corneum (8).

A central application of our approach is reduction of SSI, which still represent a major complication of surgical procedures (41). Current evidence suggests that the majority of SSI result from bacteria alighting directly onto the surgical wound from the air (22, 42–44). Based on our previous studies (7, 8) and on the preclinical results reported here, lamps emitting far-UVC light in the 200–222 nm range could potentially be used for continuous low-fluence/low-rate exposures during the course of a surgical procedure to inactivate bacteria before they penetrate into the interior.
At bactericidal fluences, far-UVC light cannot traverse tissues with a stratum corneum such as skin and lens of the eyes. The use of this light would therefore not require additional protective clothing for patients or medical staff and could become a standard in hospital environments to reduce SSI rates, particularly those due to drug-resistant pathogens. In addition, targeting bacteria as they alight onto the wound would prevent the formation of bacterial clusters or biofilms (45), which are difficult to eradicate and impede wound healing (46).

Other potential applications of far-UVC light is sterilization of any environment with a high likelihood of airborne-based pathogen transmission, including tuberculosis, smallpox, severe acute respiratory syndrome (SARS) and pandemic influenza, which collectively affects one billion people annually (47). Although upper-room UV-irradiation systems based on conventional broad-spectrum UV lamps (48) have long been considered for room sterilization (49–51), they cannot be widely used due to safety concerns relating to skin cancer and cataract risks (49, 52, 53).

Collectively, our studies suggest that far-UVC light (200–225 nm), unlike conventional UV germicidal lamps, has considerable promise to be a safe and inexpensive modality for SSI reduction, while being cytotoxic to both drug-resistant and drug-sensitive microbes (22).

FIG. 6. UVC-induced inflammation in mouse skin. Density of (panel A) mast cells and (panel B) cells expressing the myeloperoxidase (MPO) enzyme (i.e., neutrophils) in the epidermis of sham-exposed mice, of mice exposed to 157 mJ/cm² from 254-nm light or 222-nm light. Values represent the average ± SD of the number of cells/m² measured in six randomly selected fields of view per mouse (n = 3). *Value from elsewhere (8).

FIG. 7. Tissue differentiation in UVC-exposed mouse skin. Representative cross-sectional images of mouse dorsal epidermis expressing K6A (brown stained area) with relative quantification. Mice were sham exposed (top panel), exposed to 157 mJ/cm² from 254-nm light (middle panel) or 222-nm light (bottom panel). Values represent the average ± SD of the percentage of keratin optical density measured in nine randomly selected fields of view per mouse (n = 3). *Value from elsewhere (8).
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