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- (71) Applicant (for all designated States except US): **THE TRUSTEES OF COLUMBIA UNIVERSITY IN THE CITY OF NEW YORK** [—/US]; 412 Low Memorial Library, 535 West 116th Street, New York, New York 10027 (US).
- (72) Inventors; and
- (75) Inventors/Applicants (for US only): **GERHARD, Randers-Pehrson** [—/US]; 113 Hawkes Avenue, Ossining, US 10562 (US). **BRENNER, David Jonathan** [GB/US]; 139 West 82nd St., Apt. 9G, New York, New York 10024 (US). **BIGELOW, Alan** [US/US]; 117 East 77th Street, Apt. 5C, New York, New York 10075 (US).
- (74) Agents: **ABELEV, Gary** et al.; Dorsey & Whitney LLP, 51 West 52nd Street, New York, New York 10019 (US).

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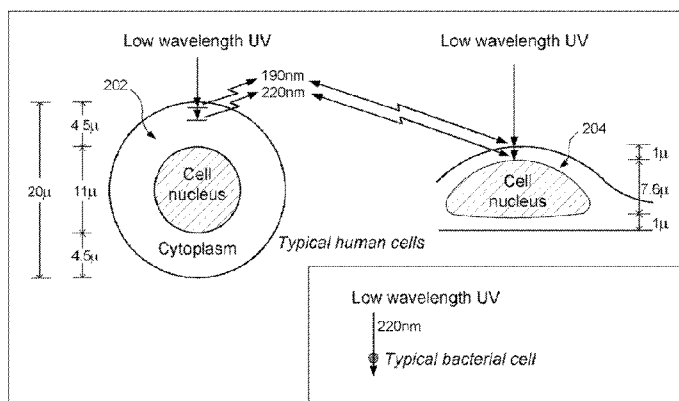


FIG. 2

(57) Abstract: Apparatus and methods for generating at least one radiation can be provided, which can selectively kill and/or affect at least one bacteria are provided. For example, the radiation source can generate at least one radiation having one or more wavelengths in a range of about 190 nanometers (nm) to about 230 nm, and the apparatus can be configured to prevent the radiation from having any wavelength that is outside of the range from being transmitted.

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APPARATUS, METHOD, AND SYSTEM FOR SELECTIVELY AFFECTING AND/OR KILLING BACTERIA

CROSS-REFERENCE TO PRIOR APPLICATION(S)

5 [0001] This application claims priority to U.S. Provisional Application Serial No. 61/450,038, filed on March 7, 2011, which is incorporated by reference herein in its entirety.

FIELD OF THE DISCLOSURE

10 [0002] Exemplary embodiments of the present disclosure relate to selectively affecting and/or killing bacteria, and more specifically to exemplary apparatus, methods and systems which can use an ultraviolet radiation to selectively affecting and/or killing bacteria while not harming human cells.

BACKGROUND INFORMATION

15 [0003] It has been estimated that between 2% and 5% of clean surgeries result in surgical site infections (SSI). Patients who develop SSI can be 60% more likely to spend time in an ICU, can be 5 times as likely to be readmitted, can have a mortality rate twice that of noninfected patients, can have an average of 7 days additional length of hospital stay, and can have an average of about \$3,000 additional costs. It has been estimated that about 40-
20 60% of SSIs can be preventable (*see, e.g., Barie PS, Eachempati SR. Surgical site infections. Surg Clin North Am 2005;85(6):1115-35, viii-ix*).

[0004] It has been approximately 50 years since Deryl Hart and colleagues at Duke University showed that ultraviolet (UV) irradiation of surgical wounds can be a highly effective methodology for reducing surgical wound infection rates (*see, e.g., Hart D. Bactericidal ultraviolet radiation in the operating room. Twenty-nine-year study for control of infections. J Am Med Assoc 1960;172:1019-28*). However, UV radiation can be a hazard
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both to the patient and to the operating team, and the use of additional clothing, hoods, and eye shields for protection can be both cumbersome and costly, preventing widespread use of the technique.

[0005] UV radiation can be a very efficient bactericidal agent, and the mechanisms by which it mutates and kills bacteria, as well as human cells, are well established (*see, e.g., Mitchell DL, Nairn RS. The biology of the (6-4) photoproduct. Photochem Photobiol 1989;49(6):805-19; Witkin EM. Ultraviolet mutagenesis and inducible DNA repair in Escherichia coli. Bacteriol Rev 1976;40(4):869-907; Koch-Paiz CA, Amundson SA, Bittner ML, Meltzer PS, Fornace AJ, Jr. Functional genomics of UV radiation responses in human cells. Mutat. Res. 2004;549(1-2):65-78; and Harm W. Biological effects of ultraviolet radiation. Cambridge, UK: Cambridge University Press, 1980*). Ultraviolet germicidal irradiation (UVGI) has been used to break down microorganisms in food, air, and water purification. UVGI typically uses a short wavelength of UV, typically in the UVB or UVC range, to destroy nucleic acids in small organisms, removing their reproductive capabilities. UV irradiation (including UVGI) is typically produced with low-pressure mercury lamps, which can produce a range of UV wavelengths, ranging from UVA (wavelengths 400 to 320 nm) to UVB (wavelengths 320 to 290 nm) to UVC (wavelengths 290 to 100 nm). Figure 1 shows the spectrum of UV wavelengths emitted from a typical mercury UV lamp. UVGI is typically produced by mercury-vapor lamps that emit at around 254 nm. However, UVGI lamps may be harmful to humans and other life forms, and are typically shielded or in environments where exposure is limited.

[0006] UV lamps can also facilitate a UV emission from an excited molecule complex (e.g., an exciplex, such as either krypton-bromine or krypton-chlorine), using arrangements called excilamps. The basic theory behind exciplex UV emission was developed in the 1970s (*see, e.g., Lorents DC. A model of rare-gas excimer formation and*

decay and its application to vuv lasers. *Radiat. Res.* 1974;59(2):438-40; and Measures RM. Prospects for developing a laser based on electrochemiluminescence. *Appl Opt* 1974;13(5):1121-33). The first excimer lasers were made in the 1980s and they are now in common use, for example, e.g., in LASIK ophthalmic surgery (see, e.g., Pallikaris IG, Papatzanaki ME, Stathi EZ, Frenschock O, Georgiadis A. *Laser in situ keratomileusis. Lasers Surg Med* 1990;10(5):463-8). Current excimer lasers, however, are typically not feasible for wound sterilization both in terms of beam size (e.g., excimer laser beams are very narrow) and their high cost. In the past, an excimer lamp (excilamp) has been developed in Russia (see, e.g., Sosnin EA, Oppenlander T, Tarasenko VF. *Applications of capacitive and barrier discharge excilamps in photoscience. J. Photochem. Photobiol C: Photochem. Rev.* 2006;7:145-63), which can produce a wide high-intensity beam of single-wavelength UV radiation. These lamps can be small, inexpensive (e.g., ~\$1,000), high powered (e.g., wound irradiation time can be a few seconds), and long-lived (e.g., 1,000 to 10,000 hours). Certain papers (see, e.g., Sosnin EA, Avdeev SM, Kuznetzova EA, Lavrent'eva LV. *A bacterial barrier-discharge KrBr Excilamp. Instr. Experiment. Tech.* 2005;48:663-66; Matafonova GG, Batoev VB, Astakhova SA, Gomez M, Christofi N. *Efficiency of KrCl excilamp (222 nm) for inactivation of bacteria in suspension. Lett Appl Microbiol* 2008;47(6):508-13; and Wang D, Oppenlander T, El-Din MG, Bolton JR. *Comparison of the disinfection effects of vacuum-UV (VUV) and UV light on Bacillus subtilis spores in aqueous suspensions at 172, 222 and 254 nm. Photochem Photobiol* 2010;86(1):176-81) have been published on their bactericidal properties (as expected they are highly efficient), but the concept that these lamps will kill bacteria but not human cells is not described.

[0007] Thus, there may be a need to address at least some of the deficiencies and/or issues that to date remained with respect to the above-described conventional systems and methods.

SUMMARY OF EXEMPLARY EMBODIMENTS

[0008] Accordingly, exemplary embodiments of the apparatus, methods and systems can be provided that can address at least some of such deficiencies. For example, the
5 exemplary embodiments of the apparatus, methods and systems can use an ultraviolet radiation to selectively affecting and/or killing bacteria while not harming human cells.

[0009] In particular, in certain exemplary embodiments of the present disclosure, a UV irradiator, e.g., the excilamp, can be provided which can effect and/or kill bacteria, without being harmful to human cells. The exemplary system, method and apparatus takes
10 into consideration that bacteria are typically physically much smaller than human cells, and thus, an appropriately chosen UV wavelength (e.g., around 207 nm to 220 nm) preferably penetrates and kills bacteria, but preferably would not be able to penetrate into the biologically sensitive nucleus of human cells. Irradiating a wound with this exemplary tailored UV radiation, for example, can therefore provide the advantages of UV
15 bacteriological sterilization, while being safe for patient and staff, and preferably not requiring protective clothing/hoods/eye shields, or the like. According to another exemplary embodiment of the present disclosure, the room air (as opposed to the wound), or surfaces (e.g., walls, floors, ceiling, countertops, furniture, fixtures, etc.) can be exposed to this exemplary UV lamp in hospital environments.

[0010] According to further exemplary embodiments of the present disclosure, it is possible to provide exemplary UV lamps that can emit at a single wavelength, in contrast to standard mercury UV lamps which typically emit over a wide range of wavelengths. The exemplary lamps can include UV emitted from an excited molecule complex (e.g., an exciplex, such as either krypton-bromine or krypton-chlorine), called excilamps, and can be
25 modified in accordance with certain exemplary embodiments of the present disclosure to

produce UV having a single wavelength, thus, facilitating modifying the UV irradiation to have enough energy to penetrate and kill bacteria, but not enough range to penetrate to the nucleus of human cells. This can be performed based on certain exemplary embodiments, e.g., using one or more modulators, wavelength-effecting masks, etc.

5 [0011] Certain exemplary embodiments of the present disclosure can be tested, for example, in an *in-vitro* (laboratory) human skin system (*see, e.g., Belyakov OV, Mitchell SA, Parikh D, Randers-Pehrson G, Marino S, Amundson SA, et al. Biological effects in unirradiated human tissue induced by radiation damage up to 1 mm away. Proc. Natl. Acad. Sci. U. S. A. 2005;102:14203-08; and Su Y, Meador JA, Geard CR, Balajee AS. Analysis of*
10 *ionizing radiation-induced DNA damage and repair in threedimensional human skin model system. Exp Dermatol 2010;19(8):e16-22*), in an *in-vitro* wound infection model (*see, e.g., Giannini GT, Boothby JT, Sabelman EE. Infected wound model development of an in vitro biomaterial-protected wound infection model to study microbial activity and antimicrobial treatment through microdialysis. Adv Skin Wound Care 2010;23(8):358-64*), in a clinically
15 relevant mouse model of surgical wound infection (*see, e.g., McLoughlin RM, Solinga RM, Rich J, Zaleski KJ, Cocchiaro JL, Risley A, et al. CD4+ T cells and CXC chemokines modulate the pathogenesis of Staphylococcus aureus wound infections. Proc. Natl. Acad. Sci. U.S. A. 2006;103(27):10408-13*), in a nude mouse model for *in-vivo* safety standards, in large animal studies, or in studies in the clinic. The exemplary excilamp wound irradiation can
20 facilitate a practical and inexpensive approach to significantly reducing surgical site infections.

[0012] According to certain exemplary embodiments of the present disclosure, a UV radiation at approximately 207 nm to 220 nm can be provided, for example, that can differentially damage and/or kill methicillin-resistant *Staphylococcus aureus* (MRSA),
25 relative to human cells. Although a conventional germicidal UV lamp can be approximately

equally efficient at killing MRSA and human cells, by contrast, the exemplary 207 to 220 nm wavelength UV from excilamps can be approximately 5,000 times more efficient at killing MRSA relative to human cells.

[0013] According to certain exemplary embodiments of the present disclosure, apparatus and method for generating at least one radiation can be provided. According to certain exemplary embodiments, the exemplary apparatus and/or method can selectively kill and/or affect at least one bacteria. For example, a radiation source first arrangement configured to generate at least one radiation having one or more wavelengths provided in a range of about 190 nanometers (nm) to about 230 nm, and at least one second arrangement configured to substantially prevent the at least one radiation from having any wavelength that is outside of the range can be provided. The radiation can be configured to selectively affect or destroy at least one bacteria on or within a body, while substantially avoiding harm to cells of the body. The radiation source can include a krypton-bromine lamp or a krypton-chlorine lamp excilamp. Additionally, the radiation source first arrangement can be further configured to generate the at least one radiation having a single wavelength provided in the range, and the at least one second arrangement can be further configured to prevent the radiation from having any wavelength other than the single wavelength. The single wavelength can be about 207 nm, and/or about 222 nm. Further, the second arrangement can include at least one of a chemical filter or a dielectric.

[0014] According to yet another exemplary embodiment, systems and methods can be provided for generating at least one radiation. For example, e.g., using a radiation source first arrangement or another arrangement, it is possible to generate the radiation(s) having one or more wavelengths provided in a range of about 190 nanometers (nm) to about 230 nm. Further, it is possible to, using at least one second arrangement and/or the same arrangement,

to substantially prevent the radiation(s) from having any wavelength that is outside of the range.

[0015] The radiation(s) can be configured to selectively affect or destroy at least one bacteria on or within a body, while substantially avoiding harming to any of cells of the body.

5 The radiation source can include an excilamp, a krypton-bromine lamp and/or a krypton-chlorine lamp. The radiation source first arrangement can be further configured to generate the radiation(s) having a single wavelength provided in the range, and the second arrangement(s) can be further configured to prevent the radiation(s) from having any wavelength other than the single wavelength. The single wavelength can be about 206 nm,
10 207 nm, and/or 222 nm. The second arrangement can include a chemical filter and/or a dielectric.

[0016] These and other objects, features and advantages of the present disclosure will become apparent upon reading the following detailed description of embodiments of the present disclosure in conjunction with the appended claims.

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BRIEF DESCRIPTION OF THE DRAWINGS

[0017] Further objects, features and advantages of the present disclosure will become apparent from the following detailed description taken in conjunction with the accompanying Figures showing illustrative embodiments of the present disclosure, in which:

20 [0018] Figure 1 is a graph of an exemplary spectrum of UV wavelengths generated by a typical mercury UV lamp;

[0019] Figure 2 is an illustration of an exemplary penetration of low wavelength UV radiation with respect to human cells and bacteria in accordance with an exemplary embodiment of the present disclosure;

[0020] Figure 3 is an illustration of an exemplary excilamp which can provide the UV radiation with at a single wavelength or in a particular range of wavelengths in accordance with an exemplary embodiment of the present disclosure;

[0021] Figure 4 is a graph of the exemplary spectral distributions of the UV radiation
5 generated by excilamps in accordance with certain exemplary embodiments of the present disclosure;

[0022] Figure 5 is an exemplary block diagram of an apparatus according to particular exemplary embodiments of the present disclosure;

[0023] Figures 6(a) and 6(b) are spectral graphs of exemplary excilamps according to
10 certain exemplary embodiments of the present disclosure;

[0024] Figure 7 is a graph of human cell survival with respect to ultra violet fluence, according to certain exemplary embodiments of the present disclosure; and

[0025] Figure 8 is a graph of MRSA survival with respect to an excilamp fluence according to certain exemplary embodiments of the present disclosure.

[0026] Throughout the drawings, the same reference numerals and characters, unless
15 otherwise stated, are used to denote like features, elements, components, or portions of the illustrated embodiments. Moreover, while the present disclosure will now be described in detail with reference to the figures, it is done so in connection with the illustrative embodiments and is not limited by the particular embodiments illustrated in the figures and
20 the accompanying claims.

DETAILED DESCRIPTION OF EXEMPLARY EMBODIMENTS

[0027] UV radiations of different wavelengths can have different abilities to penetrate
into cells. Typically, the higher the wavelength, the more penetrating the radiation, and the
25 lower the wavelength, the less penetrating the radiation. For example, UV radiation with a

low wavelength of about 200 nm, for example, while able to pass through water quite efficiently, can be heavily absorbed in the outer part of a human cell (the cytoplasm, see, for example, Figure 2), and may not have enough energy to reach the biologically sensitive cell nucleus.

5 [0028] The limited penetrating ability of ~200 nm UV radiation can be used, as shown in Figure 2, because bacteria are typically physically far smaller than human cells. Specifically, a typical bacterial cell is less than about 1 μm (micrometer) in diameter, whereas human cells are typically about 10 to 30 μm across, depending on their type and location.

[0029] In particular, Figure 2 shows a typical human cell nucleus having a spherical
10 geometry 202 or a flattened geometry 204, illustrating the penetration into a human cell of UV radiation with wavelength of around 200 nm. As shown in Figure 2, essentially no UV of this wavelength preferably reaches the cell nucleus 202, 204, which contains the radiation-sensitive DNA. Accordingly, UV radiation of this wavelength would typically not be harmful to human cells or thus to humans. In addition to this geometric reason, there can be a
15 biological reason why UV with a wavelength around 200 nm UV will typically not be harmful to humans. At about 185 nm and below, UV can be very efficiently absorbed by oxygen, producing ozone and oxidative damage. Above about 240 nm, UV can be very efficient at producing oxidative DNA base damage (*see, e.g., Kvam E, Tyrrell RM. Induction of oxidative DNA base damage in human skin cells by UV and near visible radiation. Carcinogenesis 1997;18(12):2379-84; and Pattison DI, Davies MJ. Actions of ultraviolet light on cellular structures. EXS 2006(96):131-57*). Thus, 200 nm wavelength UV can be in
20 a narrow UV “safety window”. In contrast, because bacteria are typically physically much smaller in size than human cells, UV radiation with wavelength around 200 nm can penetrate through, and therefore kill, bacteria.

[0030] According to exemplary embodiments of the present disclosure, it is possible to utilize one or more UV excilamps which, in contrast to standard UV lamps, can produce UV radiation at a specific wavelength — e.g., around 200 nm. UV radiation around such exemplary wavelength (e.g., a single wavelength or in a range of certain wavelengths as described herein) can penetrate and kill bacteria, but preferably would not penetrate into the nucleus of human cells, and thus, can be expected to be safe for both patient and staff.

Exemplary Excilamp UV Irradiator

[0031] The exemplary excilamp technology can utilize certain exemplary concepts which were developed at the Institute of High Current Electronics (IHCE) in Tomsk, Siberia (see, e.g., Sosnin EA, Oppenlander T, Tarasenko VF. *Applications of capacitive and barrier discharge excilamps in photoscience. J. Photochem. Photobiol C: Photochem. Rev.* 2006;7:145-63.). Additional exemplary excilamps that can be utilized with the exemplary embodiments of the present disclosure may be available from Heraeus Noblelight in Germany. The IHCE lamps, an exemplary embodiment of such lamp 302 is shown in Figure 3, can be small, rugged, cost ~\$1,000, and can be made to produce a variety of single wavelength UV radiations. Based on the considerations above, exemplary embodiments of the present disclosure can use, for example, a krypton-bromine lamp excilamp, which can produce UV at about 207 nm, or a krypton-chlorine lamp (Figure 3), which can produce UV at about 222 nm. Another exemplary excilamp can produce UV at about 206 nm. The exemplary spectrum of these lamps are shown in the graph of Figure 4. As shown in Figure 4, a spectral distribution 402 can be produced by a krypton-bromine lamp, and spectral distribution 404 was produced by a krypton-chlorine lamp. Additionally, according to further exemplary embodiments of the present disclosure, certain exemplary features can be included (e.g., spectrum filtering elements such as multilayer dielectric filters or chemical filters) to

remove unwanted wavelengths, or those wavelengths that are outside of the preferable range of wavelengths. For example, absorption and/or reflective elements can be provided between the lamp and the irradiated surface to filter unwanted wavelengths, such as, e.g., a band-pass filter, a long-wavelength blocking filter. In one exemplary embodiment, the absorptive material can be fluorescent, such that it emits visible light when it absorbs UV radiation to provide an indication that the lamp is operating. Alternatively or in addition, other gases can be added to the lamp to suppress unwanted wavelengths. For example, adding argon to the krypton-chlorine lamp can suppress generation of the 228 nm UV.

[0032] The typical power output of the air-cooled excilamps can be about 7.5 to 20 mW/cm², although higher power can be obtained in a water-cooled system. At about 20mW/cm², only a few seconds of exposure can deliver about 20 mJ/cm², which can be a typical bactericidal dose.

[0033] Exemplary embodiments of the present disclosure can provide an excilamp, emitting about a 207 nm or about a 222 nm single wavelength UV radiation, to differentially kill bacteria while sparing adjacent human cells. Further, the wavelength(s) of the UV radiation according to further exemplary embodiments of the present disclosure can be in the range of about 190 nanometers (nm) to about 230 nm. Exemplary experiments implementing embodiments of the present disclosure can include: an *in-vitro* (laboratory) 3-D human skin system (see, e.g., Belyakov OV, Mitchell SA, Parikh D, Randers-Pehrson G, Marino S, Amundson SA, et al. *Biological effects in unirradiated human tissue induced by radiation damage up to 1 mm away. Proc. Natl. Acad. Sci. U. S. A. 2005;102:14203-08*; and Su Y, Meador JA, Geard CR, Balajee AS. *Analysis of ionizing radiation-induced DNA damage and repair in threedimensional human skin model system. Exp Dermatol 2010;19(8):e16-22*); a nude mouse model for *in-vivo* safety standards; an *in-vitro* wound infection model (see, e.g., Giannini GT, Boothby JT, Sabelman EE. *Infected wound model development of an in vitro*

biomaterial-protected wound infection model to study microbial activity and antimicrobial treatment through microdialysis. *Adv Skin Wound Care* 2010;23(8):358-64); a clinically relevant mouse model of surgical wound infection (see, e.g., McLoughlin RM, Solinga RM, Rich J, Zaleski KJ, Cocchiaro JL, Risley A, et al. *CD4+ T cells and CXC chemokines modulate the pathogenesis of Staphylococcus aureus wound infections. Proc. Natl. Acad. Sci. U.S. A.* 2006;103(27):10408-13); larger animal model studies; studies in a clinic. According to yet another exemplary embodiment of the present disclosure, excilamp irradiation can be provided for a wound, a room air and/or surfaces (e.g., walls, floors, ceiling, countertops, furniture, fixtures, etc.), which can facilitate a practical and inexpensive approach to significantly reducing surgical wound infection.

[0034] In an exemplary experiment implementing certain exemplary embodiments of the present disclosure, an exemplary test bench was developed for gathering, e.g., exemplary preliminary sterilization results from exemplary UV light sources. For example, the exemplary test bench can include: a) a light-tight box; b) a shutter control; c) a filter holder; and d) adjustable exposure parameters for time, distance, and wavelength (e.g., 207 nm KrBr excilamp, 222 nm, KrCl excilamp, and 254 nm standard germicidal lamp). Additionally, exemplary custom filters can be designed to eliminate higher-wavelength components in the excilamp emission spectra to provide optimal single-wavelength exposure. A UV spectrometer and deuterium lamp (e.g., for equipment calibration) can be used to validate the filter effectiveness, as shown, for example, in Figures 6(a) and 6(b), which illustrate the normalized spectra comparing excilamp emission (red --- 602a and 602b) with filtered excilamp emission (blue -- 604a and 604b) for both KrBr and KrCl excilamps. This exemplary test bench facilitated, for example, a generation of biological findings of filtered excilamp exposure to both bacteria and healthy human cells, which are described below. In turn, the exemplary biological testing experience has provided details regarding exemplary

parameters for developing filtered KrBr and KrCl excilamps into optimal devices for clinical applications.

Exemplary Biological Results

[0035] Described below are certain exemplary experiments implementing certain
5 exemplary embodiments of the present disclosure. The exemplary experiments investigated, for example, whether UV from exemplary filtered excilamps can be effective at killing bacteria while sparing normal human cells.

[0036] In the exemplary experiment, human fibroblasts were, for example, exposed to about 3 mJ/cm² from a standard germicidal UV lamp (e.g., 254 nm), and their survival was
10 less than about 10⁻⁴. By contrast, when they were exposed to fluences as high as 150 mJ/cm² from the exemplary filtered KrBr or KrCl excilamp (e.g., 207 and 222 nm, respectively), their survival was in the range from about 1 to about 10⁻¹ (see Figure 7). Indeed, Figure 7 shows an exemplary graph indicating a clonogenic survival of normal human skin fibroblasts (AG1522) exposed to UV from exemplary filtered KrBr (207 nm) or KrCl (222 nm)
15 excilamps, or from a conventional germicidal lamp (254 nm).

[0037] In the exemplary experiment, bactericidal killing efficacy of the exemplary excilamps was tested, for example, on methicillin resistant *Staphylococcus aureus* (MRSA). MRSA can be the cause of about 25% of surgical site infection and can be associated with approximately 20,000 deaths per year in the United States, mostly healthcare related. MRSA
20 and antibiotic-susceptible *S. aureus* are typically equally susceptible to UV from conventional germicidal lamps. (See, e.g., Conner-Kerr TA, Sullivan PK, Gaillard J, Franklin ME, Jones RM. The effects of ultraviolet radiation on antibiotic-resistant bacteria in vitro. *Ostomy Wound Manage.* 1998 ;44(10):50-6). The exemplary results are shown, for example, in Figure 8, indicating that at an excilamp fluence of about 100 mJ/cm², an MRSA survival
25 level of 10⁻⁴ can be achieved. For example, Figure 8 shows an exemplary graph of MRSA

(strain US300) inactivation after exposure to UV from the exemplary filtered KrBr or KrCl excilamps (207 nm and 222 nm, respectively).

[0038] Comparing the exemplary results in Figures 7 and 8, the exemplary filtered excilamp UV radiation at 207 nm and at 222 nm can differentially effect and/or kill MRSA relative to the human cells. For example, at exemplary filtered excilamp fluences of about 5 100 mJ/cm², the survival level of human cells is, for example, in the range of about 0.1 to 1, while the survival level of MRSA is in the range of about 10⁻⁴. Such exemplary findings are in considerable contrast to the situation for conventional germicidal UV lamps (GUVL), which is roughly equally efficient at killing bacteria and human cells. For example, for a 10 conventional germicidal UV lamp, at a UV fluence for which a GUVL produces a bacterial survival of 10⁻⁴, the human cell survival from the GUVL is about 0.3 x 10⁻⁴, a human cell survival advantage of 0.3. With the exemplary excilamp at 207 or 222 nm, at a UV fluence for which the exemplary 207 or 222 nm filtered excilamp produces a bacterial survival of 10⁻⁴, the human cell survival by the exemplary filtered excilamps is in the range of about 0.1 to 1, 15 a human cell survival advantage in the range of 5,000.

[0039] Figure 5 shows an exemplary block diagram of an exemplary embodiment of a system according to the present disclosure. For example, exemplary procedures in accordance with the present disclosure described herein can be performed by or controlled using a UV generation source 580 and/or hardware processing arrangement and/or a 20 computing arrangement 510, separately and in conjunction with one another. Such exemplary processing/computing arrangement 510 can be, e.g., entirely or a part of, or include, but not limited to, a computer/processor 520 that can include, e.g., one or more microprocessors, and use instructions stored on a computer-accessible medium (e.g., RAM, ROM, hard drive, or other storage device).

[0040] As shown in Figure 5, e.g., a computer-accessible medium 530 (e.g., as described herein above, a storage device such as a hard disk, floppy disk, memory stick, CD-ROM, RAM, ROM, etc., or a collection thereof) can be provided (e.g., in communication with the processing arrangement 510). The computer-accessible medium 530 can contain
5 executable instructions 540 thereon. In addition or alternatively, a storage arrangement 550 can be provided separately from the computer-accessible medium 530, which can provide the instructions to the processing arrangement 510 so as to configure the processing arrangement to execute certain exemplary procedures, processes and methods, as described herein above, for example.

10 [0041] Further, the exemplary processing arrangement 510 can be provided with or include an input/output arrangement 570, which can include, e.g., a wired network, a wireless network, the internet, an intranet, a data collection probe, a sensor, etc. As shown in Figure 5, the exemplary processing arrangement 510 can be in communication with an exemplary display arrangement 560, which, according to certain exemplary embodiments of the present
15 disclosure, can be a touch-screen configured for inputting information to the processing arrangement in addition to outputting information from the processing arrangement, for example. Further, the exemplary display 560 and/or a storage arrangement 550 can be used to display and/or store data in a user-accessible format and/or user-readable format.

[0042] The foregoing merely illustrates the principles of the disclosure. Various
20 modifications and alterations to the described embodiments will be apparent to those skilled in the art in view of the teachings herein. It will thus be appreciated that those skilled in the art will be able to devise numerous systems, arrangements, and procedures which, although not explicitly shown or described herein, embody the principles of the disclosure and can be thus within the spirit and scope of the disclosure. In addition, all publications and references
25 referred to above can be incorporated herein by reference in their entireties. It should be

understood that the exemplary procedures described herein can be stored on any computer accessible medium, including a hard drive, RAM, ROM, removable disks, CD-ROM, memory sticks, etc., and executed by a processing arrangement and/or computing arrangement which can be and/or include a hardware processors, microprocessor, mini, 5 macro, mainframe, etc., including a plurality and/or combination thereof. In addition, certain terms used in the present disclosure, including the specification, drawings and claims thereof, can be used synonymously in certain instances, including, but not limited to, e.g., data and information. It should be understood that, while these words, and/or other words that can be synonymous to one another, can be used synonymously herein, that there can be instances 10 when such words can be intended to not be used synonymously. Further, to the extent that the prior art knowledge has not been explicitly incorporated by reference herein above, it can be explicitly being incorporated herein in its entirety. All publications referenced can be incorporated herein by reference in their entirety.

15

What Is Claimed Is:

1. An apparatus for generating at least one radiation, comprising:
a radiation source first arrangement configured to generate the at least one radiation having one or more wavelengths provided in a range of about 190 nanometers (nm) to about
5 230 nm; and
at least one second arrangement configured to substantially prevent the at least one radiation from having any wavelength that is outside of the range.
2. The apparatus of claim 1, wherein the at least one radiation is configured to selectively
10 affect or destroy at least one bacteria on or within a body, while substantially avoiding harm to cells of the body.
3. The apparatus of claim 1, wherein the radiation source includes an excilamp.
- 15 4. The apparatus of claim 3, wherein the excilamp includes at least one of a krypton-bromine lamp or a krypton-chlorine lamp.
5. The apparatus of claim 1, wherein the radiation source first arrangement is further configured to generate the at least one radiation having a single wavelength provided in the
20 range, and wherein the at least one second arrangement is further configured to prevent the at least one radiation from having any wavelength other than the single wavelength.
6. The apparatus of claim 5, wherein the single wavelength is about 207 nm.
- 25 7. The apparatus of claim 5, wherein the single wavelength is about 222 nm.

8. The apparatus of claim 1, wherein the at least one second arrangement includes at least one of a chemical filter or a dielectric.

5 9. A method for selectively killing or affecting at least one bacteria, comprising:

providing at least one radiation having at least one wavelength that is in a range of between about 190 nanometers (nm) to about 230 nm; and

causing a substantial prevention of the at least one radiation from having wavelengths outside of the range.

10

10. The method of claim 9, wherein the at least one radiation is configured to selectively effect or destroy the at least one bacteria on or within a body, while substantially avoiding harm to cells of the body.

15 11. The method of claim 9, wherein the at least one radiation is directed by an excilamp.

12. The method of claim 11, wherein the excilamp includes at least one of a krypton-bromine lamp or a krypton-chlorine lamp.

20 13. The method of claim 9, wherein the at least one radiation is provided having a single wavelength provided in the range, and wherein the prevention prevents the at least one radiation from having any wavelength other than the single wavelength.

14. The method of claim 13, wherein the single wavelength is about 207 nm.

25

15. The method of claim 13, wherein the single wavelength is about 222 nm.

16. The method of claim 0, wherein the causing procedure includes utilizing at least one of a chemical filter or a dielectric.

5

17. An apparatus for generating at least one radiation, comprising:

at least one arrangement configured to (i) provide the at least one radiation having one or more wavelengths provided in a range of about 190 nanometers (nm) to about 230 nm, and (ii) substantially prevent the at least one radiation from having any wavelength that is outside

10 of the range.

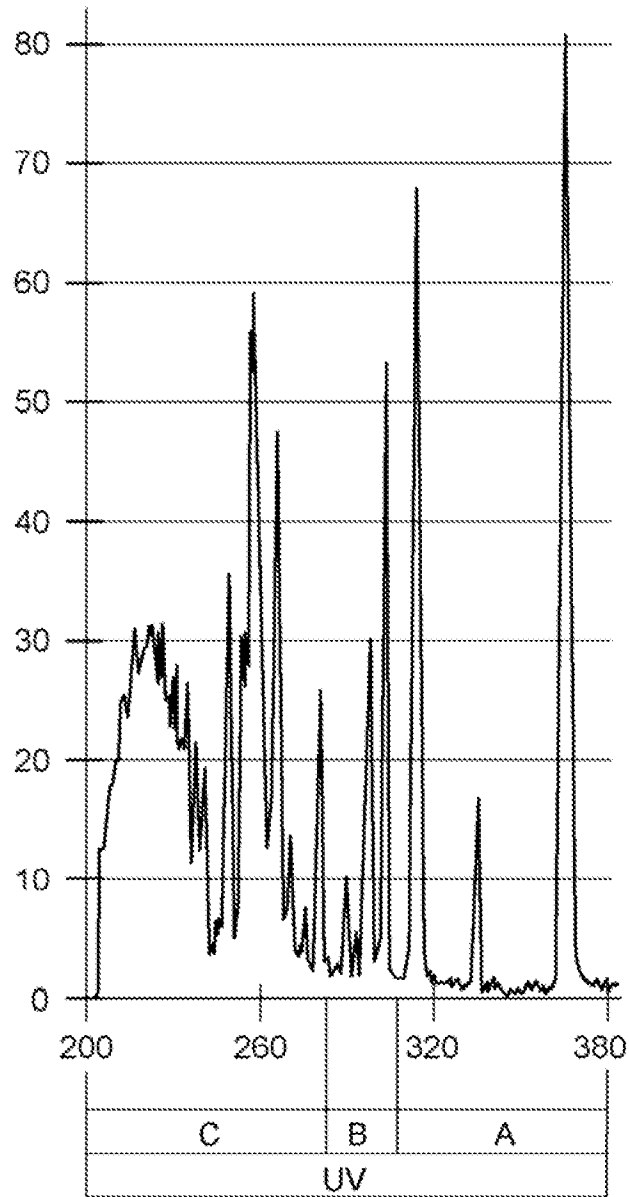


FIG. 1

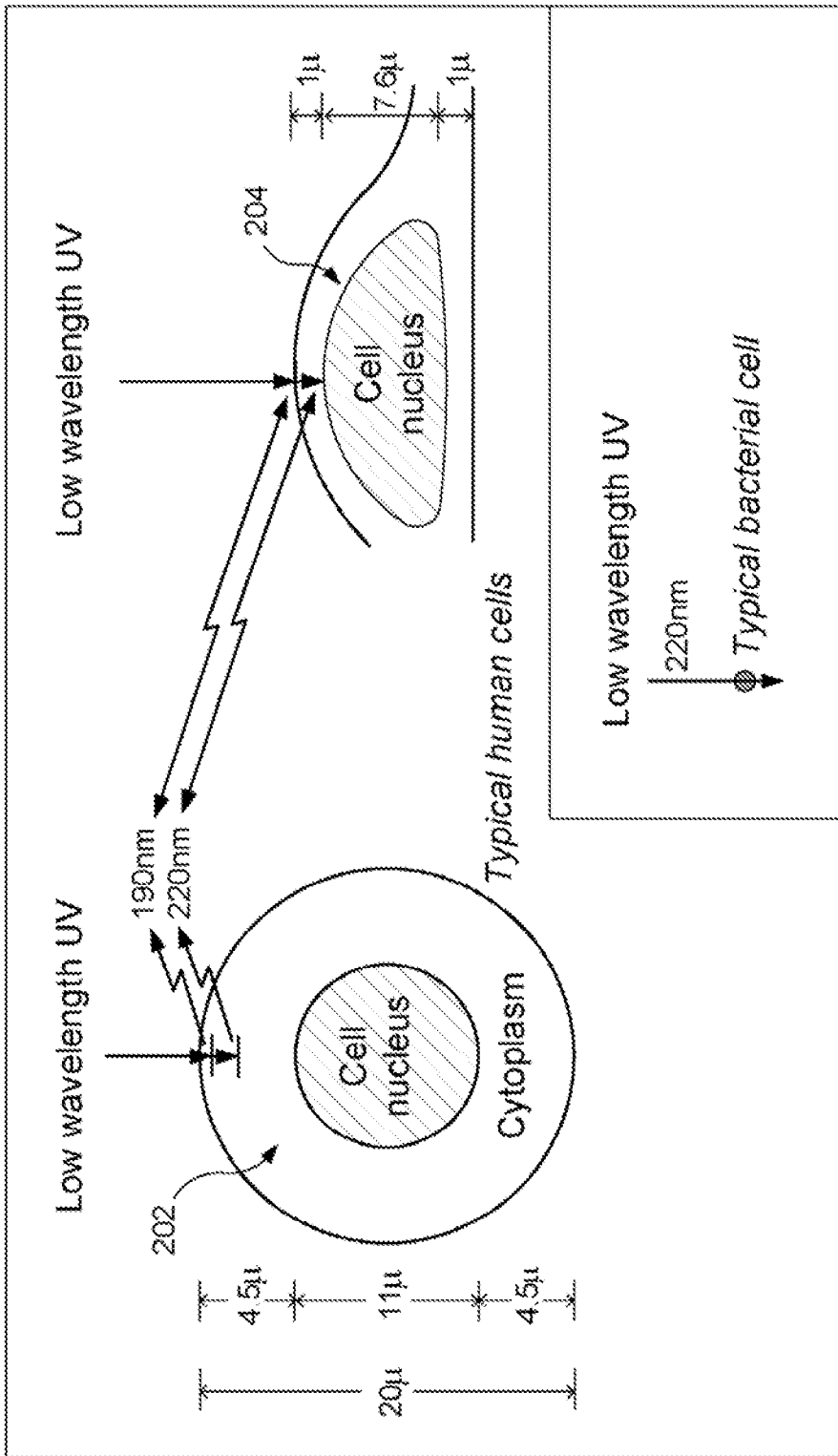
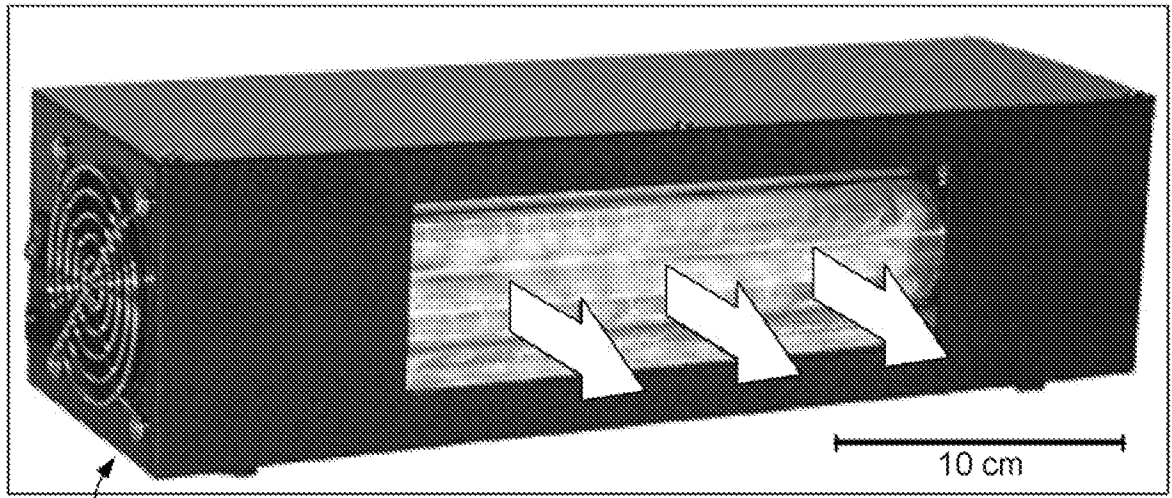
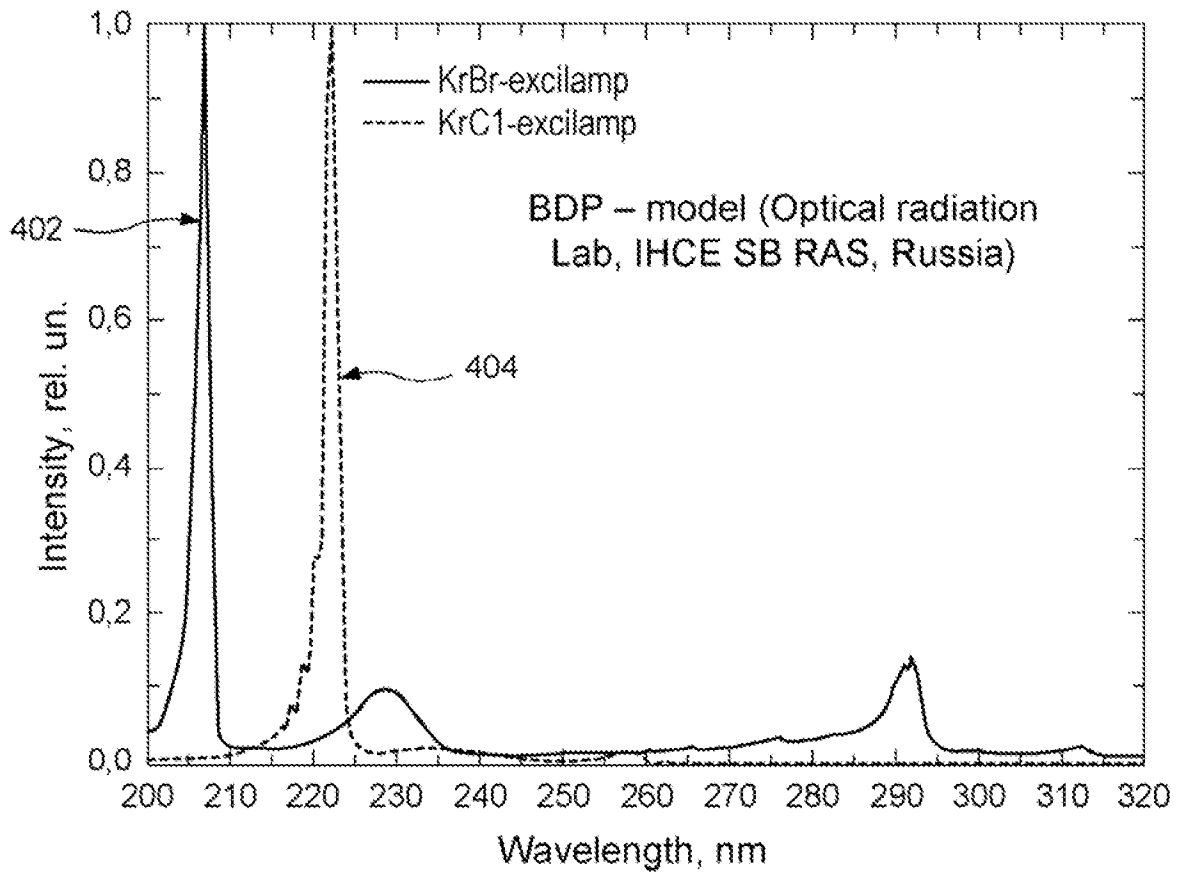


FIG. 2



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FIG. 3



Spectral distribution of KrBr and KrC1 excilamps in relative units

FIG. 4

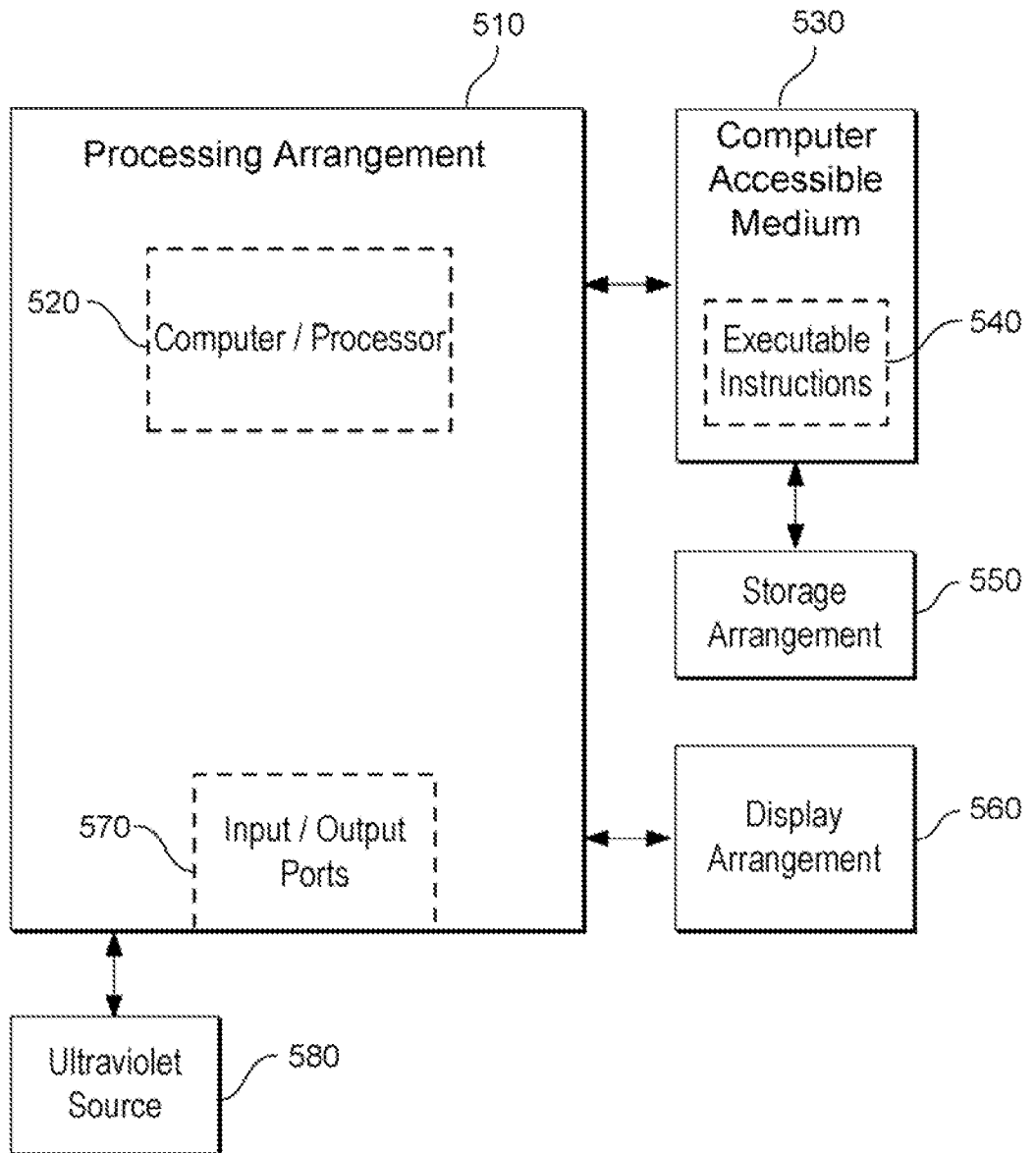


FIG. 5

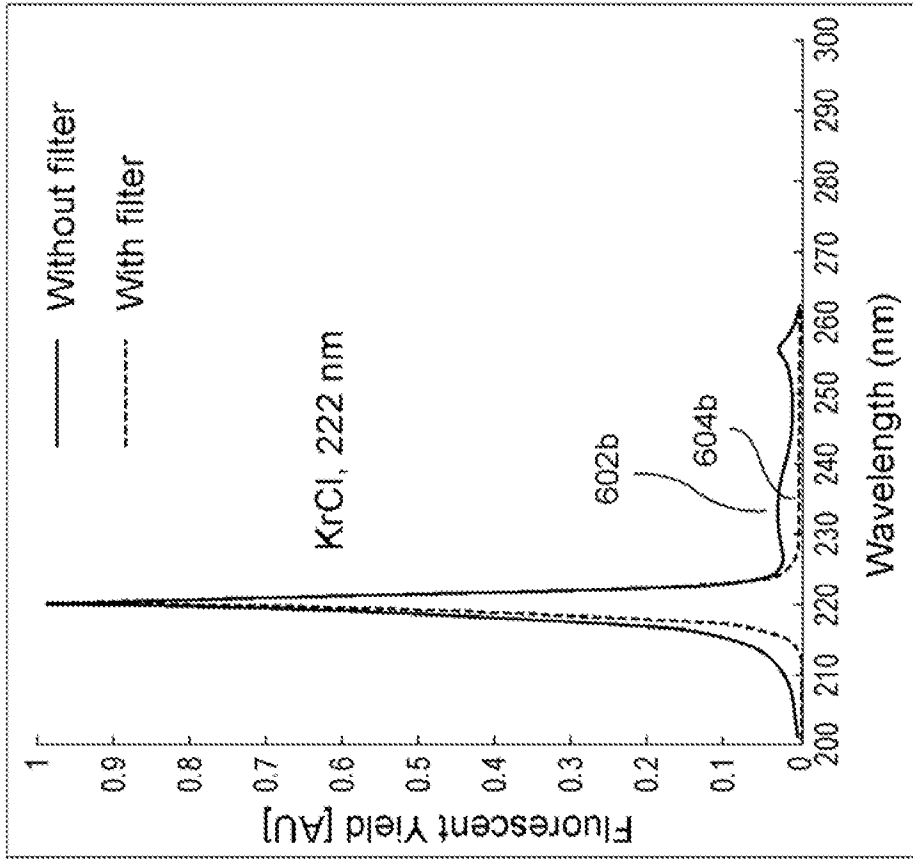


FIG. 6b

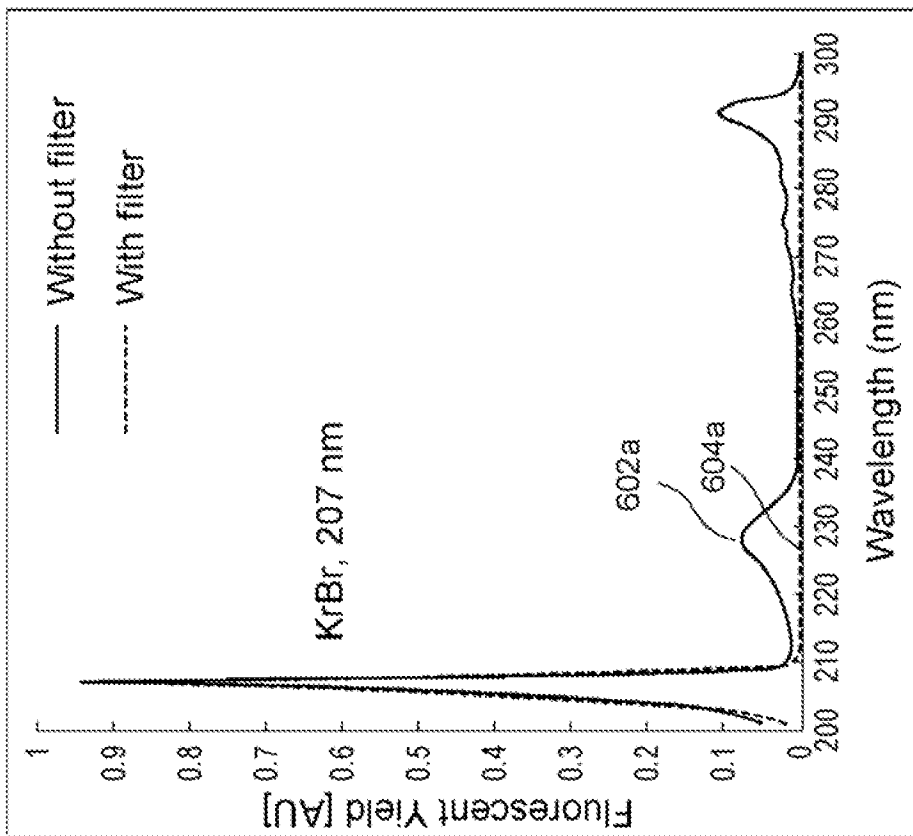


FIG. 6a

6 / 6

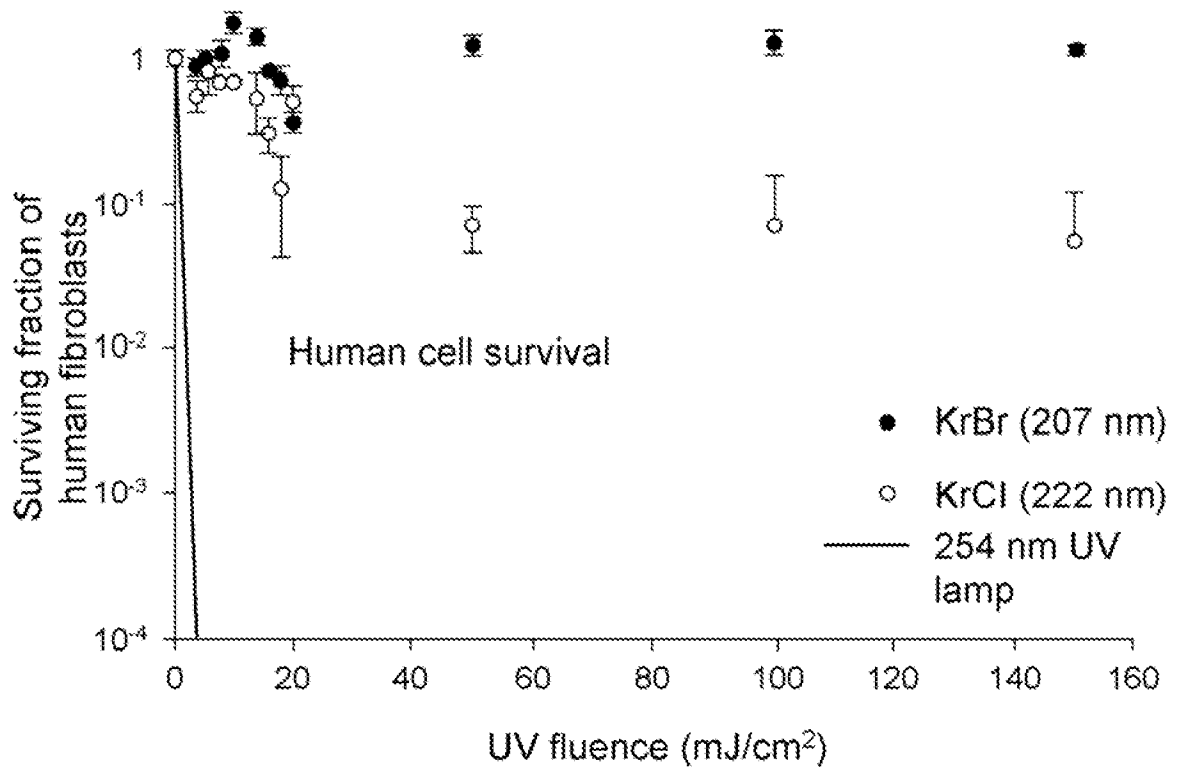


FIG. 7

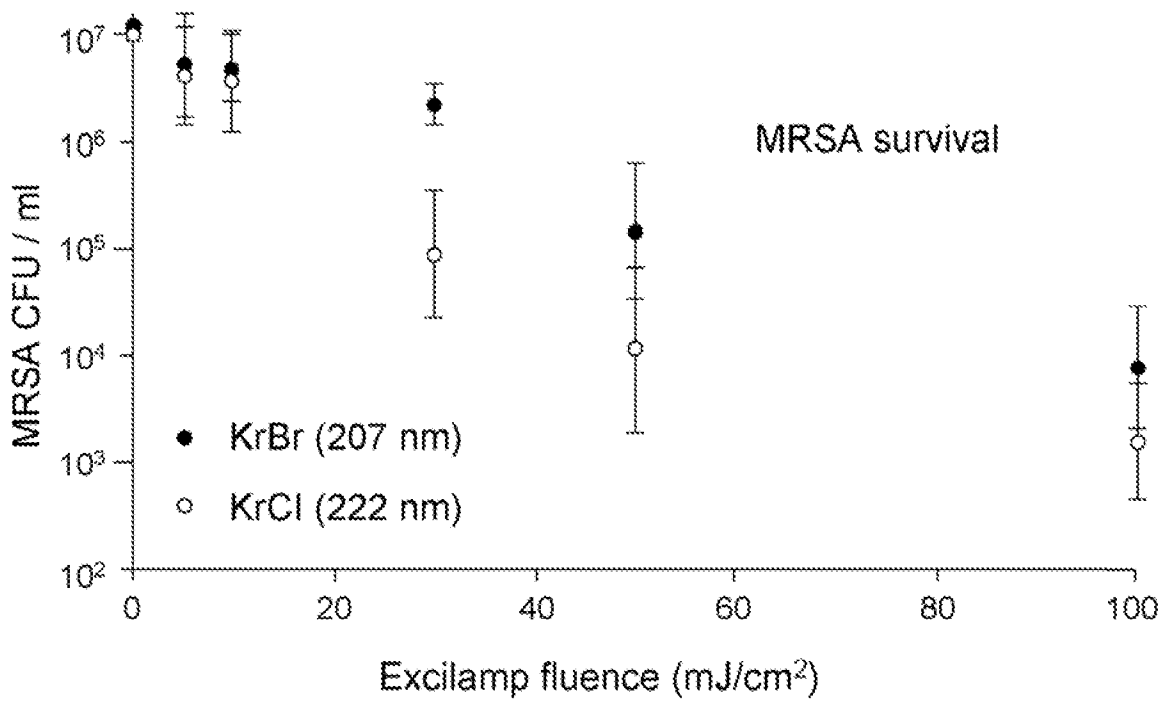


FIG. 8

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US 12/27963

A. CLASSIFICATION OF SUBJECT MATTER

IPC(8) - A61N 5/06 (2012.01)

USPC - 607/94; 422/2

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC(8) - A61N 5/06 (2012.01)

USPC - 607/94; 422/2

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched
IPC(8) - A61N 5/06 (2012.01), USPC - 607/94; 422/2, 250/492.1, 250/493.1, 607/88: keyword search, as below

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

USPTO PubWest (databases: PGPB,USPT,USOC,EPAB,JPAB), Google Scholar -- Search Terms: Excilamp, bacteria, prokaryote, 207 nm, 220 nm, 180, 190, 200, 210, 220, 230, nm, generate, produce, light, radiation, electromagnetic, filter, block, prevent, range, longer, shorter, above, below, kill, killing, destroy, sterilize, affect

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X — Y	US 2007/0135874 A1 (BALA) 14 June 2007 (14.06.2007) abstract; para [0013]; [0039]; [0054]; [0012]; [0014]; [0058]	1, 2, 5, 8-10, 13, 16, 17 ----- 3, 4, 6, 7, 11, 12, 14, 15
Y	OPPENLANDER et al. "Mercury-free Vacuum-(VUV) and UV Excilamps: Lamps of the Future?" IUVA News; December 2005; Vol. 7, No. 4; pg. 16-20. Entire document, especially pg. 19, col 1, para 2-3; Table 1; pg. 16, col 2, para 2	3, 4, 6, 7, 11, 12, 14, 15
A	US 2010/0007492 A1 (RESSLER et al.) 14 January 2010 (14.01.2010)	1-17
A	US 6,376,972 B1 (TARASENKO et al.) 23 April 2002 (23.04.2002)	3, 4, 6, 7, 11, 12, 14, 15
A	US 2010/0028201 A1 (NEISTER) 4 February 2010 (04.02.2010)	1-17

 Further documents are listed in the continuation of Box C.

* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier application or patent but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search

21 June 2012 (21.06.2012)

Date of mailing of the international search report

11 JUL 2012

Name and mailing address of the ISA/US

Mail Stop PCT, Attn: ISA/US, Commissioner for Patents
P.O. Box 1450, Alexandria, Virginia 22313-1450

Facsimile No. 571-273-3201

Authorized officer:

Lee W. Young

PCT Helpdesk: 571-272-4300
PCT OSP: 571-272-7774