

# ULTRAVIOLET AIR AND SURFACE TREATMENT

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ULTRAVIOLET germicidal irradiation (UVGI) uses short-wave ultraviolet (UVC) energy to inactivate viral, bacterial, and fungal organisms so they are unable to replicate and potentially cause disease. UVC energy disrupts the deoxyribonucleic acid (DNA) of a wide range of microorganisms, rendering them harmless (Brickner et al. 2003; CIE 2003). Early work established that the most effective UV wavelength range for inactivation of microorganisms is between 220 and 280 nm, with peak effectiveness near 265 nm. The standard source of UVC in commercial systems is low-pressure mercury vapor lamps, which emit mainly near-optimal 253.7 nm UVC. Use of germicidal ultraviolet (UV) lamps and lamp systems to disinfect room air and air streams dates to about 1900 (Reed 2010). Riley (1988) and Shechmeister (1991) wrote extensive reviews of UVC disinfection. Application of UVC is becoming increasingly frequent as concerns about indoor air quality increase. UVC is now used as an engineering control to interrupt the transmission of pathogenic organisms, such as *Mycobacterium tuberculosis* (TB), influenza viruses, mold, and potential bioterrorism agents (Brickner et al. 2003; CDC 2002, 2005; GSA 2010; McDevitt et al. 2008; Rudnick et al. 2009).

UVC lamp devices and systems are placed in air-handling systems and in room settings for the purpose of air and surface disinfection (Figure 1). Control of bioaerosols using UVC can improve indoor air quality (IAQ) and thus enhance occupant health, comfort, and productivity (ASHRAE 2009; Menzies et al. 2003). Detailed descriptions of UVGI components and systems are given in Chapter 17 of the 2016 *ASHRAE Handbook—HVAC Systems and Equipment*. Upper-air (also commonly called upper-room) devices are installed in occupied spaces to control bioaerosols (e.g., suspended viruses, bacteria, fungi contained in droplet nuclei) in the space. In-duct systems are installed in air-handling units to control bioaerosols in recirculated air that may be collected from many spaces, and to control microbial growth on cooling coils and other surfaces. Keeping the coils free of biofilm buildup can help reduce pressure drop across the coils and improve heat exchanger efficiency (therefore lowering the energy required to move and condition the air), and eliminates one potential air contamination source that could degrade indoor air quality. UVC is typically combined with conventional air quality control methods, including dilution ventilation and particulate filtration, to optimize cost and energy use (Ko et al. 2001).

This chapter discusses these common approaches to the application of UVC products. It also surveys the most recent UVC design guidelines, standards, and practices and discusses energy use and economic considerations for the application of UVC systems. Photocatalytic oxidations (PCOs), another UV-based HVAC application, are not discussed in this chapter, but are addressed in Chapter 47 of this volume.

The preparation of this chapter is assigned to TC 2.9, Ultraviolet Air and Surface Treatment.

## 1. FUNDAMENTALS

Ultraviolet energy is electromagnetic radiation with a wavelength shorter than that of visible light and longer than x-rays (Figure 2). The International Commission on Illumination (CIE 2003) defines the UV portion of the electromagnetic spectrum as radiation having wavelengths between 100 and 400 nm. The UV spectrum is further divided into UVA (wavelengths of 400 to 315 nm), UVB (315 to 280 nm), UVC (280 to 200 nm), and vacuum UV (VUV; 200 to 100 nm) (IESNA 2000). The optimal wavelength for inactivating microorganisms is 265 nm (Figure 3), and the germicidal effect decreases rapidly if the wavelength is not optimal.

### UV Dose and Microbial Response

This section is based on Martin et al. (2008).

UVGI inactivates microorganisms by damaging the structure of nucleic acids and proteins at the molecular level, making them incapable of reproducing. The most important of these is DNA, which is responsible for cell replication (Harm 1980). The nucleotide bases (pyrimidine derivatives thymine and cytosine, and purine derivatives guanine and adenine) absorb most of the UV energy responsible for cell inactivation (Diffey 1991; Setlow 1966). Absorbed UV photons can damage DNA in a variety of ways, but the most significant damage event is the creation of pyrimidine dimers, where two adjacent thymine or cytosine bases bond with each other, instead of across the double helix as usual (Diffey 1991). In general, the DNA molecule with pyrimidine dimers is unable to function properly, resulting in the organism's inability to replicate or even its death (Diffey 1991; Miller et al. 1999; Setlow 1997; Setlow and Setlow 1962). An organism that cannot reproduce is no longer capable of causing disease.

UVGI effectiveness depends primarily on the UV dose ( $D_{UV}$ ,  $\mu\text{J}/\text{cm}^2$ ) delivered to the microorganisms:

$$D_{UV} = It \quad (1)$$

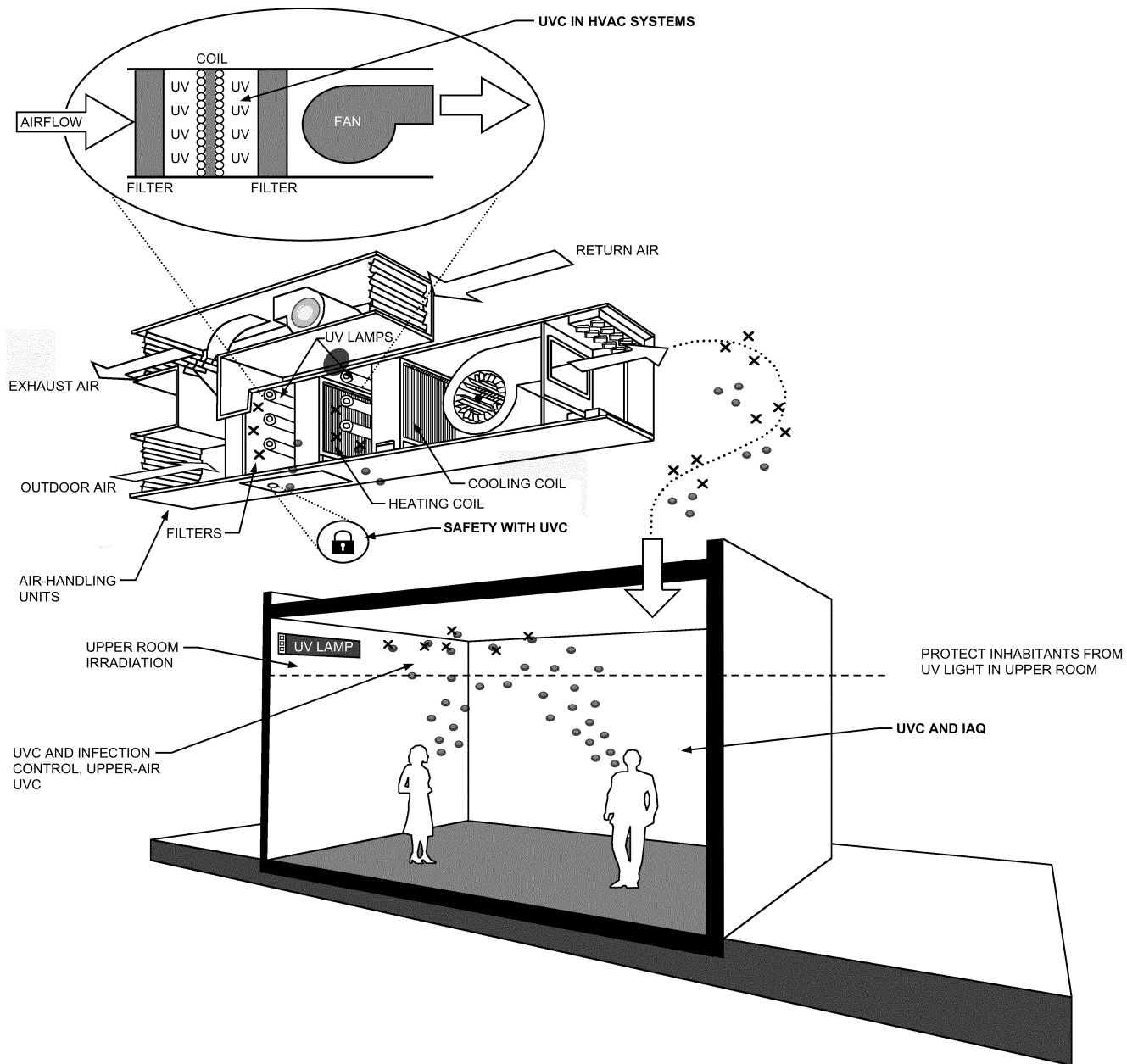
where  $I$  is the average irradiance in  $\mu\text{W}/\text{cm}^2$ , and  $t$  is the exposure time in seconds (note that  $1 \text{ J} = 1 \text{ W/s}$ ). Although Equation (1) appears quite simple, its application can be complex (e.g., when calculating the dose received by a microorganism following a tortuous path through a device with spatial variability in irradiance). The dose is generally interpreted as that occurring on a single pass through the device or system. Although the effect of repeated UV exposure on microorganisms entrained in recirculated air may be cumulative, this effect has not been quantified, and it is conservative to neglect it.

The survival fraction  $S$  of a microbial population exposed to UVC energy is an exponential function of dose:

$$S = e^{-kD_{UV}} \quad (2)$$

where  $k$  is a species-dependent inactivation rate constant, in  $\text{cm}^2/\mu\text{J}$ . The resulting single-pass inactivation rate  $\eta$  is the complement of  $S$ :

$$\eta = 1 - S \quad (3)$$



**Fig. 1 Potential Applications of UVC to Control Microorganisms in Air and on Surfaces**  
(ASHRAE 2009)

and is a commonly used indicator of overall UVC effectiveness, representing the percentage of the microbial population inactivated after one pass through the irradiance field(s).

Inactivation rate constants ( $k$ -values) are species-dependent and relate the susceptibility of a given microorganism population to UV radiation (Hollaender 1943; Jensen 1964; Sharp 1939, 1940). Measured  $k$ -values for many species of viruses, bacteria, and fungi have been published in the scientific literature and previously summarized (Brickner et al. 2003; Kowalski 2009; Philips 2006). As shown in Figure 4, bacteria are generally more susceptible to UVC energy than fungi, but this is not always the case (see Chapter 17 of the 2016 *ASHRAE Handbook—HVAC Systems and Equipment*). It is more difficult to generalize when it comes to viruses. Reported  $k$ -values for different species of microorganisms vary over several orders of magnitude. Consequently, choosing which  $k$ -value to use

for UVC system design is often difficult and confusing. The variation in reported  $k$ -values makes generalizing the use of Equation (2) particularly complicated for heterogeneous microbial populations. Even accurately determining  $S$  for one specific microorganism can be difficult, because the reported  $k$ -values for the same species sometimes differ significantly.

Variations in published  $k$ -values may relate to differences in conditions under which the UV irradiance of the microbial population was conducted (in air, in water, or on surfaces), the methods used to measure the irradiance level, and errors related to the microbiological culture-based measurements of microbial survival (Martin et al. 2008). Because no standard methods are currently available for the determination of inactivation rate constants, care is necessary when applying values reported in the literature to applications under different environmental conditions.

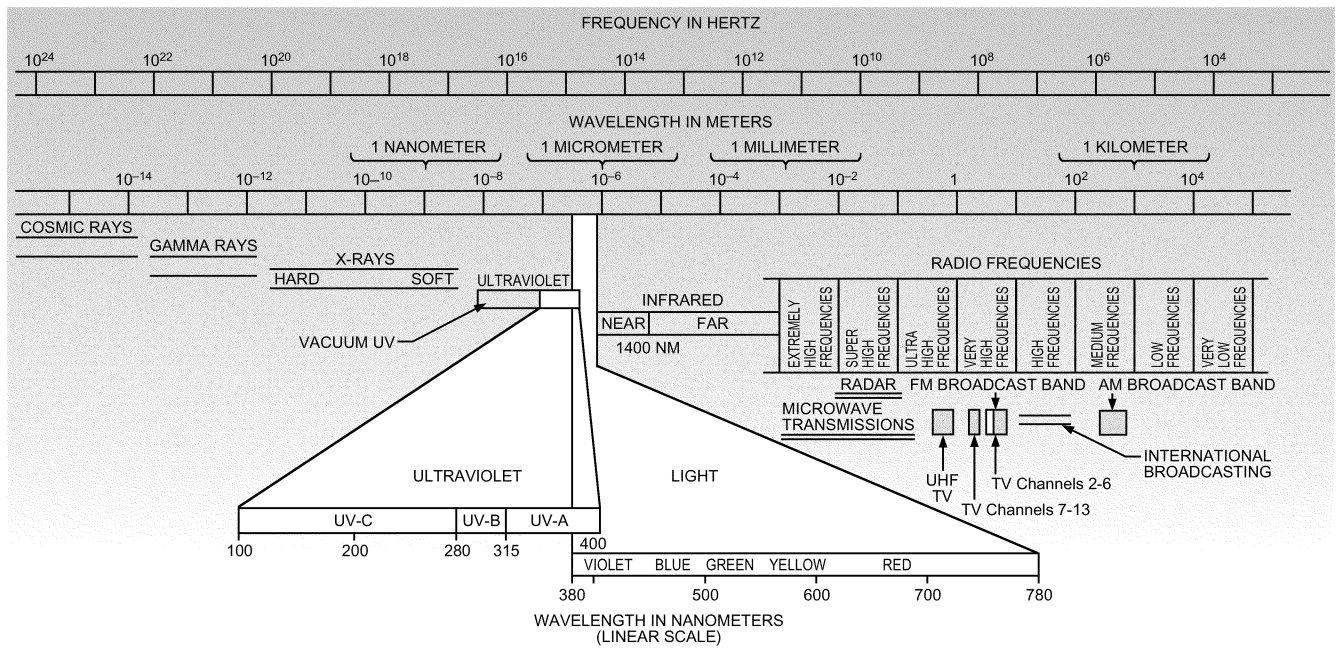


Fig. 2 Electromagnetic Spectrum (IESNA 2000)

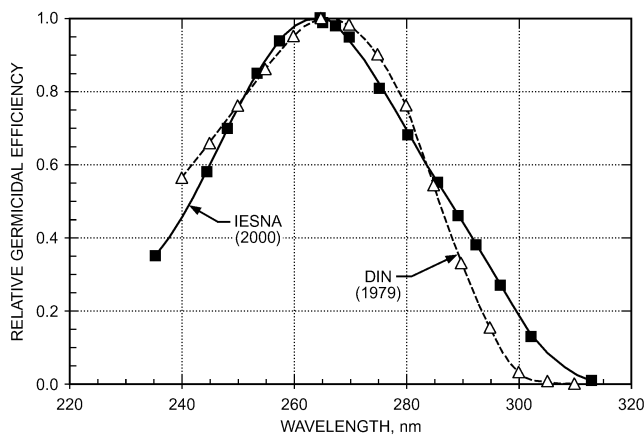


Fig. 3 Standardized Germicidal Response Functions

**UV Inactivation of Biological Contaminants**

The focus of this chapter is application of UVC energy to inactivate microorganisms, specifically bacteria, fungi, and viruses on surfaces and in air streams. The application of UVC for upper-air treatment generally applies to pathogenic bacteria and viruses. Under some circumstances, these pathogens have the potential to be transmitted throughout the HVAC system.

As shown in Table 1, infectious diseases can be transmitted by a variety of means. UVC is effective against microorganisms in the air that flows through the UVC irradiation field and on irradiated surfaces.

As shown in Table 2 and Figure 4, viruses and vegetative bacteria are the generally most susceptible to UV inactivation, followed by Mycobacteria, bacterial spores, and finally fungal spores. Within each group, an individual species may be significantly more resistant or susceptible, so this ranking should be used only as a general guideline. Note that the spore-forming bacteria and fungi also have vegetative forms, which are markedly more susceptible to

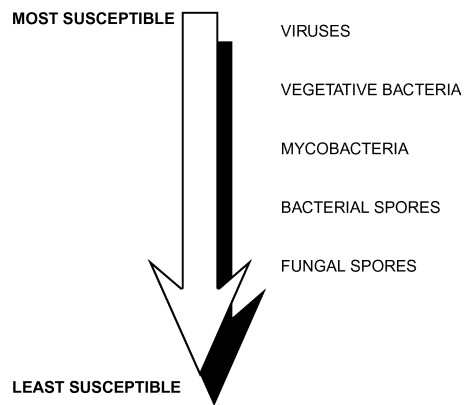


Fig. 4 General Ranking of Susceptibility to UVC Inactivation of Microorganisms by Group

inactivation than are the spore forms. Viruses are a separate case. As a group, their susceptibility to inactivation is even broader than for the bacteria or fungi.

**2. TERMINOLOGY**

Just as it is customary to express the size of aerosols in micrometers and electrical equipment’s power consumption in watts, regardless of the prevailing unit system, it is also customary to express total UVC output, UVC irradiance and fluence, and UVC dose using SI units.

Multiply I-P	By	To Obtain SI
Btu/ft <sup>2</sup> (International Table)	1135.65	μJ/cm <sup>2</sup>
Btu/h · ft <sup>2</sup>	315.46	μW/cm <sup>2</sup>
To Obtain I-P	By	Divide SI

**Burn-in time.** Period of time that UV lamps are powered on before being put into service, typically 100 h.

Table 1 Modes of Disease Transmission

Exposure	Examples
Direct contact with an infected individual	Touching, kissing, sexual contact, contact with oral secretions, or contact with open body lesions Usually occurs between members of the same household/close friends/family
Indirect contact with a contaminated surface (fomite)	Doorknobs, handrails, furniture, washroom surfaces, dishes, keyboards, pens, phones, office supplies, children's toys
Droplet contact	Infected droplets contact surfaces of eye, nose, or mouth Droplets containing microorganisms generated when an infected person coughs, sneezes, or talks Droplets are too large to be airborne for long periods of time, and quickly settle out of air
Airborne droplet nuclei (residue from evaporated droplets) or other particles containing microorganisms $\sim 5 \mu\text{m}$	Size allows them to remain airborne for long periods of time Organisms generally hardy (capable of surviving for long periods of time outside the body, resistant to drying) Organisms enter the upper and lower respiratory tracts
Fecal-oral	Usually associated with organisms that infect the digestive system Microorganisms enter via ingestion of contaminated food/water and shed in feces Lack of proper hygienic and sanitation practices
Vectorborne	Transmission through animals Bite, feces of a vector, contact with outside surface of a vector (e.g., a fly)

Table 2 Representative Members of Organism Groups

Organism Group	Member of Group
Vegetative Bacteria	<i>Staphylococcus aureus</i>
	<i>Streptococcus pyogenes</i>
	<i>Escherichia coli</i>
	<i>Pseudomonas aeruginosa</i>
	<i>Serratia marcescens</i>
Mycobacteria	<i>Mycobacterium tuberculosis</i>
	<i>Mycobacterium bovis</i>
	<i>Mycobacterium leprae</i>
Bacterial Spore	<i>Bacillus anthracis</i>
	<i>Bacillus cereus</i>
	<i>Bacillus subtilis</i>
Fungal Spores	<i>Aspergillus versicolor</i>
	<i>Penicillium chrysogenum</i>
	<i>Stachybotrys chartarum</i>
Viruses	Influenza viruses
	Measles
	SARS
	Smallpox

**Cutaneous damage.** Any damage to the skin, particularly that caused by exposure to UVC energy.

**Disinfection.** Compared to sterilization, a less lethal process of inactivating microorganisms.

**Droplet nuclei.** Residual viable microorganisms in air, following evaporation of surrounding moisture. These microscopic particles are produced when an infected person coughs, sneezes, shouts, or sings. The particles can remain suspended for prolonged periods and can be carried on normal air currents in a room and beyond to adjacent spaces or areas receiving exhaust air.

**Erythema (actinic).** Reddening of the skin, with or without inflammation, caused by the actinic effect of solar radiation or artificial optical radiation. See CIE (2011) for details. (Nonactinic erythema can be caused by various chemical or physical agents.)

**Exposure.** Being subjected to infectious agents, irradiation, particulates, or chemicals that could have harmful effects.

**Fluence.** Radiant flux passing from all directions through a unit area, often expressed as  $\text{J}/\text{m}^2$ ,  $\text{J}/\text{cm}^2$ , or  $(\mu\text{W} \cdot \text{s})/\text{cm}^2$ .

**Irradiance.** Power of electromagnetic radiation incident on a surface per unit surface area, typically reported in microwatts per square centimeter ( $\mu\text{W}/\text{cm}^2$ ). See CIE (2011) for details.

***Mycobacterium tuberculosis.*** The namesake member of the *M. tuberculosis* complex of microorganisms, and the most common cause of tuberculosis (TB) in humans. In some instances, the species name refers to the entire *M. tuberculosis* complex, which includes *M. bovis*, *M. africanum*, *M. microti*, *M. canettii*, *M. caprae*, *M. pinipediti*, and others.

**Ocular damage.** Any damage to the eye, particularly that caused by exposure to UV energy.

**Permissible exposure time (PET).** Calculated time period that humans, with unprotected eyes and skin, can be exposed to a given level of UV irradiance without exceeding the NIOSH recommended exposure limit (REL) or ACGIH Threshold Limit Value® (TLV®) for UV radiation.

**Personal protective equipment (PPE).** Protective clothing, helmets, goggles, respirators, or other gear designed to protect the wearer from injury from a given hazard, typically used for occupational safety and health purposes.

**Photokeratitis.** Defined by CIE (1993) as corneal inflammation after overexposure to ultraviolet radiation.

**Photokeratoconjunctivitis.** Inflammation of cornea and conjunctiva after exposure to UV radiation. Exposure to wavelengths shorter than 320 nm is most effective in causing this condition. The peak of the action spectrum is approximately 270 nm. See CIE (1993) for details. Note that different action spectra have been published for photokeratitis and photoconjunctivitis (CIE 1993); however, the latest studies support the use of a single action spectrum for both ocular effects.

**Radiometer.** An instrument used to measure radiometric quantities, particularly UV irradiance or fluence.

**Threshold Limit Value® (TLV®).** An exposure level under which most people can work consistently for 8 h a day, day after day, without adverse effects. Used by the ACGIH to designate degree of exposure to contaminants. TLVs can be expressed as approximate milligrams of particulate per cubic meter of air ( $\text{mg}/\text{m}^3$ ). TLVs are listed either for 8 h as a time-weighted average (TWA) or for 15 min as a short-term exposure limit (STEL).

**Ultraviolet radiation.** Optical radiation with a wavelength shorter than that of visible radiation. (See CIE [1987] for details.) The range between 100 and 400 nm is commonly subdivided into

UVA: 315 to 400 nm  
UVB: 280 to 315 nm  
UVC: 200 to 280 nm  
Vacuum UV 100 to 200 nm

**Ultraviolet germicidal irradiation (UVGI).** Ultraviolet radiation that inactivates microorganisms. UVC energy is generated by germicidal lamps that kill or inactivate microorganisms by emitting radiation predominantly at a wavelength of 253.7 nm.

**UV dose.** Product of UV irradiance and specific exposure time on a given microorganism or surface, typically reported in millijoules per square centimeter ( $\text{mJ}/\text{cm}^2$ ).

**Wavelength.** Distance between repeating units of a wave pattern, commonly designated by the Greek letter lambda ( $\lambda$ ).

### 3. UVGI AIR TREATMENT SYSTEMS

#### Design Guidance

Early guidelines published by General Electric (Buttolph and Haynes 1950), Philips (1985), and Westinghouse (1982) are still used by many system designers today. First et al. (1999), Kowalski (2003, 2006, 2009), NIOSH (2009), and Riley et al. (1976) made meaningful advances in the analysis and modeling of UVGI systems that improved guidance for system design, yet no consensus guidelines exist that comprehensively address all aspects of UVGI system design required to ensure desired performance.

UVC system design today relies on performance data from lamp, ballast, and fixture manufacturers and the experience of system designers. Many equipment manufacturers have methods for estimating the UV dose delivered, which may include using tabulated data charts, mathematical modeling, and complex formulas. Like most HVAC components, UVC systems are often oversized to ensure performance. This oversizing, though conservative, can potentially increase equipment and utility costs, and may result in less energy-efficient systems.

Although application support for UVC technologies is growing and many successful systems have been installed, “the most important needs in the area of UVGI are industry standards to rate devices and installations, as well as guidance for installation and maintenance” (EPA 2017). ASHRAE Technical Committee 2.9, Ultraviolet Air and Surface Treatment, was created in 2003 (initially as a Task Group, converted to a standing Technical Committee in 2007) in part to address these deficiencies by initiating research programs, preparing Handbook chapters, and serving as the cognizant committee for developing the needed standards. So far, two new ASHRAE standards have been developed that provide end users with ratings of equipment performance and aid UVC system designers in selecting appropriate components:

- ASHRAE *Standard* 185.1, Method of Testing UV-C Lights for Use in Air-Handling Units or Air Ducts to Inactivate Airborne Microorganisms, establishes a test method for evaluating the efficacy of UVC lights for their ability to inactivate airborne microorganisms installed inside general ventilation systems.
- ASHRAE *Standard* 185.2, Method of Testing Ultraviolet Lamps for Use in HVAC&R Units or Air Ducts to Inactivate Microorganisms on Irradiated Surfaces, establishes a similar test method to measure the intensity of ultraviolet lamps on irradiated surfaces under typical HVAC&R operating conditions.

Work is ongoing to initiate round-robin testing between laboratories that can potentially conduct testing on UVC devices according to these new standards. Such testing will generate critical data on the repeatability of the testing methods and identify issues that must be addressed in updates to the standards.

For any application, the ability of UVC to inactivate microorganisms is a function of dose. **Dose** is the length of time of exposure multiplied by the irradiance measured in  $\mu\text{W}/\text{cm}^2$  (see Chapter 17 in the 2016 *ASHRAE Handbook—HVAC Systems and Equipment* for more details). A key difference between surface decontamination and airborne inactivation of organisms is exposure time. In a duct system, exposure time is on the order of seconds or fractions of seconds because of the rapid movement of air through the duct. Therefore, the irradiance must be sufficiently high to provide the dose necessary to inactivate the pathogen in seconds or a fraction of a second, depending upon the configuration and characteristics of the UVC system.

As mentioned previously, organisms differ in their susceptibility to UVC inactivation. Depending on the application, a public health or medical professional, microbiologist, or other individual with knowledge of the threat or organisms of concern should be consulted during the design process.

#### Upper-Air UVC Devices (Fixtures)

The primary objective of upper-air UVC placement and use is to interrupt the transmission of airborne infectious pathogens within the indoor environment. The source of these infectious organisms may be infected humans, animals, or bioaerosols introduced for terrorism purposes. Humans are the predominant sources of airborne agents that infect people (ACGIH 1999). The measles and influenza viruses and the tuberculosis bacterium are three important infectious organisms known to be transmitted indoors by means of air shared, by any means, between infected and susceptible persons. Studies of person-to-person outbreaks indicate at least two transmission patterns: within-room exposure such as in a congregate space, and transmission beyond a room through corridors and by entrainment in ventilation ductwork, through which air is then recirculated throughout the building. ASHRAE also provides guidance on protecting buildings from extraordinary incidents in which a bio-terror agent is aerosolized into a building (ASHRAE 2003).

UVC is used, in combination with other environmental controls, to protect building occupants in all areas of concern (Brickner et al. 2003; Kowalski and Bahnfleth 2003). Since the 1930s (Riley and O’Grady 1961; Wells 1955) and continuing to the present day (First et al. 2007a, 2007b; Miller et al. 2002; Xu et al. 2003), numerous experimental studies have demonstrated the efficacy of upper-air UVC. Additionally, evidence of effectiveness has been established for inactivation of tuberculosis (Escombe et al. 2009; Mphahlele et al. 2015), reducing measles transmission in a school, and the interruption of influenza transmission within a hospital (McLean 1961).

Various upper-air UVC devices are designed to generate a controlled UVC field above the heads of occupants and to minimize UVC in the lower, occupied area of the room. Settings appropriate to upper-air UVC placement include congregate spaces, where unknown and potentially infected persons may share the same space with uninfected persons (e.g., a medical waiting room or homeless shelter). Common corridors potentially used by unknown infected persons in a medical facility would also benefit from upper-air UVGI fixtures. Upper-air UVC also covers situations where untreated recirculated air might enter an occupied space (see Figures 5 and 6 for illustrations of upper-air pathogen control using UVC). Upper-air UVC is very effective in areas with no, or minimal, ventilation; 2 air changes per hour (ach) equivalency, up to normal recommended levels of 6 ach can be achieved. Ventilation patterns (natural and mechanical) should promote good air mixing in the space equipped with UVC so that infectious microorganisms encounter the UVC zone and are inactivated, thus reducing the risk of exposure of occupants to airborne infectious agents. Recent studies that have used natural ventilation and UVC have shown that upper-air UVC is an effective, low-cost intervention for use in TB infection control (Escombe et al. 2009; Mphahlele et al. 2015).

Upper-air UVC devices are designed and installed to irradiate only air in the upper part of the room (Figures 7 and 8). Parameters

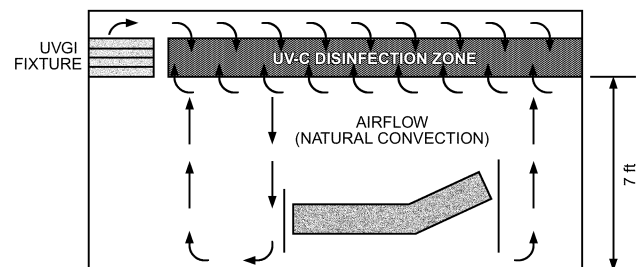
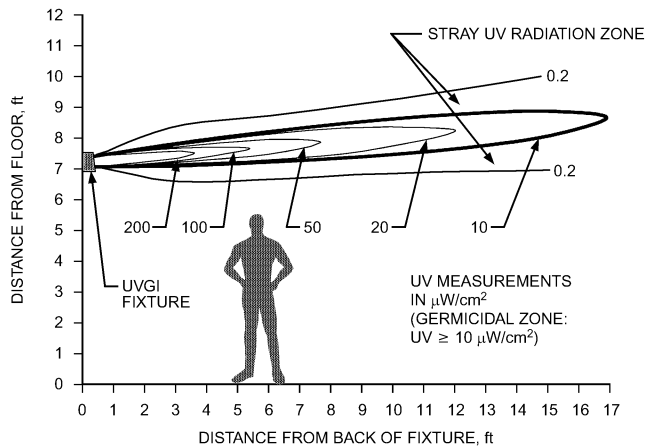


Fig. 5 Typical Elevation View of Upper-Air UVC Applied in Hospital Patient Room



**Fig. 6 Typical Elevation View Showing UVC Placed above Heads of Room Occupants for Safety**



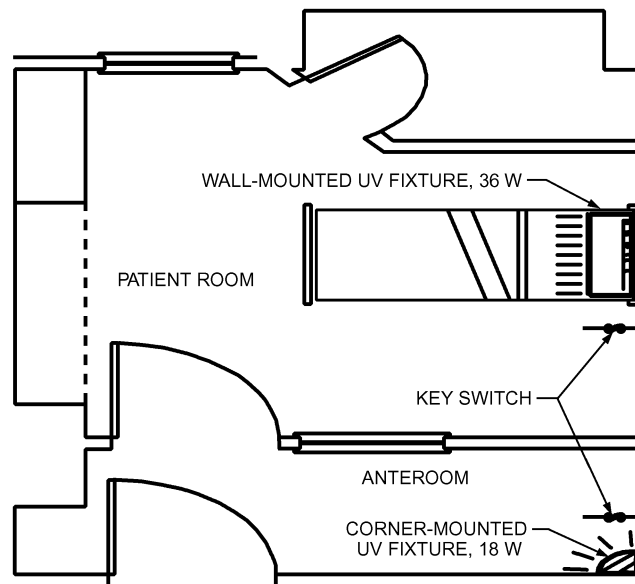
**Fig. 7 Upper-Air UVC Treating Congregate Setting**  
(TUSS Project, St. Vincent's Hospital, New York City)

for UVC effectiveness include room configuration, UV fixture placement, and the adequacy of air currents in bringing contaminated air into the upper UV zone. UVC devices should be placed appropriately spaced to accommodate the area, shape, and height of the space in which air is to be disinfected. Figures 9 to 11 show examples of upper-air fixture placement. An upper-air computer-based tool can calculate the average fluence in the upper room (Rudnick et al. 2012; Vincent et al. 2013; Zhang et al. 2012). Additionally, computational fluid dynamics (CFD) is being used to understand the interaction between airflow and upper-air UVC (Gilkeson and Noakes 2013; Xu et al. 2013; Zhu et al. 2013).

Upper-air UVC devices typically use low-pressure UVC lamps in tubular and compact shapes and accommodate a variety of electrical wattages and voltages. Beyond lamp size, shape, and ballasts, fixtures are available in open or restricted energy distribution, depending on the physical space to be treated. UVC fixtures are selected based on the floor-to-ceiling height. Ceiling heights above 10 ft may allow for more open fixtures, which may be more efficient because they may allow for a larger irradiation zone. For occupied spaces with lower ceilings (less than 10 ft), various louvered upper-air UVC devices (wall-mount, pendant, and corner-mount) are available for use in combinations and are mounted with at least 7 ft from the floor to the bottom of the fixture. The fixture should be mounted so that its UV energy is distributed parallel to the plane of the ceiling. Device construction and placement prevent excessive



**Fig. 8 Upper-Air UVC Devices in Naturally Ventilated Corridor of TB Facility in Brazil**  
(Centers for Disease Control and Prevention)



**Fig. 9 Suggested Layout of UVC Fixtures for Patient Isolation Room**  
(First et al. 1999)

ultraviolet energy from striking occupants below. For example, in high-risk areas such as corridors of infectious disease wards, a maximum UV irradiation of  $0.4 \mu\text{W}/\text{cm}^2$  at eye level is an acceptable engineering guide (Coker et al. 2001). No long-term health effects of UVC exposure at these levels in the lower occupied part of rooms are known. Figure 5 shows a typical elevation and corresponding UV levels, and Figure 6 illustrates typical UVC energy distribution in a room.

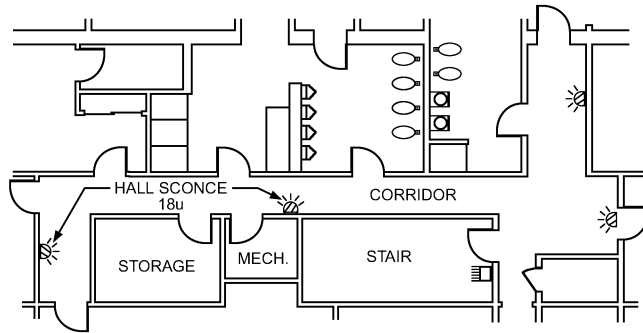
Application guidance with placement criteria for UV equipment is provided by Boyce (2003), CDC (2005), CIE (2003), Coker et al. (2001), First et al. (1999), IESNA (2000), and NIOSH (2009). An example of the guidance provided by Coker et al. is shown in Table 3. Additionally, manufacturer-specific advice on product operation and placement should be followed. A new computer-aided lighting software program is being modified to help automate the placement of fixtures, and to calculate the uniformity and average UV provided (Brickner et al. 2009). Upper-air UVC fixtures that are typically used in developed countries are often cost-prohibitive for use in less

**Table 3 Suggested UVC Fixture Mounting Heights**

	Wall-Mounted Fixtures*		Ceiling-Mounted Figures*	
	Corner Mount	Wall Mount	Pendant	Pendant with Fan
Beam pattern	90°	180°	360°	360°
Minimum ceiling height	2.44 m	2.44 m	2.89 m	2.89 m
Fixture mounted height	2.1 m	2.1 m	2.4 m	2.4 m
Ideal UV-C intensity for effective disinfection	> 10 $\mu\text{W}/\text{cm}^2$	> 10 $\mu\text{W}/\text{cm}^2$	> 10 $\mu\text{W}/\text{cm}^2$	> 10 $\mu\text{W}/\text{cm}^2$

Source: Coker et al. (2001)

\*Appropriately designed UV fixtures are available for all locations. Only the most commonly used have been included in the table.



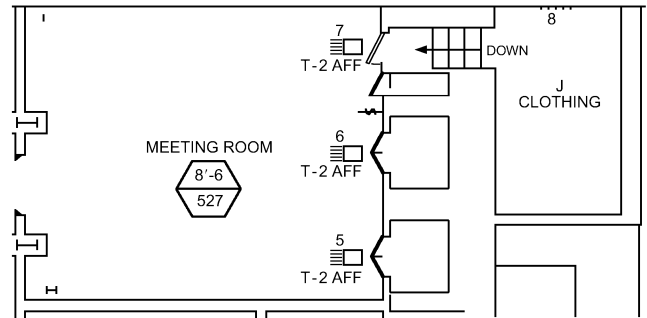
**Fig. 10 Upper-Air UVC Devices with 180° Emission Profile Covering Corridors**  
(First et al. 1999)

developed parts of the world. International guidance is needed to understand best practice for UVC application in the developing world where extensive drug-resistant TB is an increasing global threat (Nardell et al. 2013).

Some upper-air installations rely on air convection and mixing to move air from the lower to the upper portion of the room, where it can be irradiated and airborne microorganisms inactivated (Kethley and Branc 1972). The overall effectiveness of upper-air UVC systems improves significantly when the air in the space can be well mixed. Although convection air currents created by occupants and equipment can provide adequate air circulation in some settings, mechanical ventilation systems that maximize air mixing are preferable. If mechanical ventilation is not possible, fans can be placed in the room to enhance mixing. Many fixtures incorporate a safety switch that breaks the circuit when fixtures are opened for servicing and should contain baffles or louvers appropriately positioned to direct UV irradiation to the upper air space. Baffles and louvers must never be bent or deformed.

A UVC installation that produces a maintained, uniform distribution of UV irradiance averaging between 30 and 50  $\mu\text{W}/\text{cm}^2$  is effective in inactivating most airborne droplet nuclei containing mycobacteria, and is presumably effective against viruses as well (First et al. 2007a, 2007b; Miller et al. 2002; Xu et al. 2003). Beyond UVC irradiance, effectiveness of upper-air UVC is related to air mixing, relative humidity, and the inherent characteristics of the pathogenic organisms being addressed (Ka et al. 2004; Ko et al. 2000; Rudnick 2007). Effectiveness can improve greatly with well-mixed air (First et al. 2007a, 2007b; Miller et al. 2002; Riley and Permutt 1971; Riley et al. 1971), so ventilation systems that maximize air mixing receive the greatest benefit from upper-air UVC. Relative humidity should be less than 60%; levels over 80% rh may reduce effectiveness (Kujundzic et al. 2007; Xu et al. 2003).

Depending on the disinfection goals, upper-air devices should be operated similarly to in-duct UVC systems. Systems designed to reduce or eliminate the spread of airborne infectious diseases in buildings with continuous occupancy and/or with immunocompromised populations should be operated 24 h per day, 7 days per week.



SPACE USAGE	ROOM DIMENSIONS			UVGI FIXTURE COVERAGE			
	L ft	W ft	H ft	MH ft	A ft <sup>2</sup>	V ft <sup>3</sup>	(N) WM
CONFERENCE/OPEN OFFICE ROOM	8.9	5.5	2.6	2.2	49	117.4	(3) 8.5 W

*Design Concept: Congregate setting, high occupancy, shared air with adjacent auditorium (multipurpose room). Dropped ceiling for an open office plan. Look for long path lengths, evenly space fixtures along one wall.*

*L = length, W = width, H = floor-to-ceiling height, A = area covered, V = room volume, MH = mounting height above finished floor, N = number of UVGI fixtures, W = nominal wattage, WM = wall mounted, CM = ceiling mounted*

**Fig. 11 Example Upper-Air UVC Layout for A Meeting Room**

Upper-air systems designed for improved indoor air quality installed in more traditional commercial buildings may be operated intermittently, or powered on during hours of normal building occupancy and powered off when the facility is empty. This may provide acceptable indoor air quality during periods of building occupancy, simultaneously saving energy and requiring less frequent lamp replacements. However, intermittent operation must be factored into the initial system design because cycling UV lamps on and off may negatively affect lamp and ballast performance and life.

**In-Duct UVC Systems: Airstream Disinfection**

The principal design objective for an in-duct UVC air disinfection system is to distribute UV energy uniformly in all directions throughout the length of the duct or air-handling unit (AHU) to deliver the appropriate UV dose to air moving through the irradiated zone with minimum system power. Enhancing the overall reflectivity of the inside of the air handler can improve UVC system performance by reflecting UVC energy back into the irradiated zone, thus increasing the effective UV dose. Using materials such as aluminum or other highly reflective materials can increase reflectivity. Properly designed in-duct UV air disinfection systems are also able to maintain the cleanliness of cooling coil surfaces and condensate pans, when the UV lamps are installed in close proximity to this equipment. On the other hand, systems designed specifically for coil and condensate pan applications may not be adequate for proper air disinfection.

Design dose is a function of the design-basis microbe (i.e., the targeted microorganism with the smallest *k*-value) and the desired

level of disinfection. Generally, single-pass inactivation efficiencies are specified, analogous to the specification of a particulate filter MERV rating. In some cases, the design disinfection level may be a true performance specification based on the exposure in an occupied space. Determining this value requires analysis of the entire system that is used to determine the single-pass performance. Which approach is selected depends on the type of application. Laboratory/hospital installations are more likely to have specific, identified targets than, for example, school or office installations. The required average irradiance for a typical in-duct system is on the order of 1000 to 10,000  $\mu\text{W}/\text{cm}^2$ , but it could be higher or lower depending on the application requirements.

In-duct air disinfection systems should be designed to have the desired single-pass inactivation level under worst-case conditions of air temperature and velocity in the irradiated zone. The worst-case performance reflects the combined effect of the number/power of UVC fixtures; air residence time, which is inversely proportional to air velocity; and lamp/ballast characteristics, including wind chill effect and depreciation (as discussed in Chapter 17 of the 2016 *ASHRAE Handbook—HVAC Systems and Equipment*). Lee et al. (2009) showed that it may be advantageous to use simulation to determine the design condition, given the complex interactions between air temperature, velocity, and lamp performance. Lamps may be located anywhere in an air conveyance system; however, some locations provide more efficiency and potentially greater benefit. In most cases, the lowest maximum velocity in a system occurs inside an air-handling unit. For this reason, and because it provides the ability to treat air from many spaces and simultaneously irradiate cooling coils and condensate pans, this is a very common choice, although systems may also be located in air distribution ducts.

Because they are typically installed in air handling units, most in-duct systems are designed for an air velocity of around 500 ft/min. At this velocity, an irradiance zone 8 ft in length achieves a 1 s exposure. As a rule of thumb, in-duct systems should be installed in a location that can provide a minimum of 0.25 s of UV exposure; otherwise, system cost and power consumption will be excessive. UVC devices are most often located downstream of the heating/cooling coils. However, in some cases, mounting fixtures upstream of the coil may result in lower in-duct temperatures, resulting in a more optimum lamp performance temperature and more cost-effective disinfection. The trade-off is reducing the effectiveness of disinfection of the cooling coil and forgoing irradiation of the drain pan that lamps mounted downstream of the coil provide.

In-duct air disinfection systems designed to reduce the spread of airborne infectious diseases (e.g., tuberculosis, influenza) in buildings with continuous occupancy and/or with immunocompromised populations (e.g., hospitals, prisons, homeless shelters) should be operated on a continuous basis. However, properly designed systems installed in more traditional commercial buildings (e.g., offices, retail) can be operated intermittently, or powered on during hours of normal building occupancy and powered off when the facility is empty. This may save energy costs and require less frequent lamp replacement while providing acceptable indoor air quality during periods of occupancy. However, the effect of intermittent operation on lamp and ballast life must be factored into the design analysis: cycling reduces the operating hours to failure of hot cathode lamps. In-duct UVC should always be used in combination with proper filtration. Filters may help to protect UV lamps from dust and debris accumulation which may reduce UV output over time, and filters enhance the overall air cleaning capabilities of the system.

### Studies of Airstream Disinfection Effectiveness

Laboratory studies (e.g., RTI 2005; VanOsdell and Foarde 2002) conclusively demonstrate the ability of commercially available equipment to achieve a high level of disinfection of moving airstreams. These studies have generally involved tests with surrogates

rather than actual infectious disease agents, but it can be assumed that an infectious agent with a  $k$ -value similar to an experimental surrogate will be similarly inactivated. Previous field studies showed clinical effectiveness (i.e., reduced incidence of infection) (Nagy et al. 1954; Rentschler and Nagy 1940), but similar recent studies are lacking. Although pilot studies have begun (Bierman and Brons 2007; Rudnick et al. 2009), further recorded field studies are needed to benchmark installed system performance. Many UV airstream disinfection systems have been installed in hospital environments to help reduce pathogens by complementing conventional dilution/filtration systems.

## 4. HVAC SYSTEM SURFACE TREATMENT

### Coil and Drain Pan Irradiation

Conditions in HVAC systems can promote the growth of bacteria and mold-containing biofilms on damp or wet surfaces such as cooling coils, drain pans (Levetin et al. 2001), plenum walls, humidifiers, fans, energy recovery wheels, and filters. Locations in and downstream of the cooling coil section are particularly susceptible because of condensation and carryover of moisture from coil fins. Cooling coil fouling by biofilms may increase coil pressure drop and reduce airflow and heat exchange efficiency (Montgomery and Baker 2006). Filters capture bacteria, mold, and dust, which may lead to microbial growth in damp filter media. As the growth proliferates, a filter's resistance to airflow can increase. This can result in more frequent filter changeouts and increased exposure to microbes for maintenance workers and building occupants. As airflow and coil performance degrades, so does the air quality in occupied spaces (Kowalski 2006).

Conventional methods for maintaining air-handling system components include chemical and mechanical cleaning, which can be costly, difficult to perform, and dangerous to maintenance staff and building occupants. Vapors from cleaning agents can contribute to poor air quality, chemical runoff contributes to groundwater contamination, and mechanical cleaning can reduce component life. Furthermore, system performance can begin to degrade again shortly after cleaning, as microbial growth reappears or reactivates.

UVC can be applied to HVAC systems, typically in air-handling units, to complement conventional system maintenance procedures (Bahnfleth 2011) and has been shown to be effective in reducing air-side pressure drop and increasing air-side heat transfer coefficient of wetted cooling coils (Bahnfleth 2017). A large dose can be delivered to a stationary surface with a low UVC irradiance because of the essentially infinite exposure time, making it relatively easy to cost-effectively prevent the growth of bacteria and mold on system components. In contrast to air disinfection irradiance levels, which may exceed 1000  $\mu\text{W}/\text{cm}^2$ , coil surface irradiance levels on the order of 1  $\mu\text{W}/\text{cm}^2$  can be effective (Kowalski 2009), although 50 to 100  $\mu\text{W}/\text{cm}^2$  is more typical. Using reflectors to focus lamp output on surfaces may reduce the power required for surface treatment, but at the expense of reducing air treatment effectiveness. Potential advantages of UVC surface treatment include keeping surfaces clean continuously rather than periodically restoring fouled surfaces, no use of chemicals, lower maintenance cost, and potentially better HVAC system performance.

Lamps can be installed to target problematic components such as cooling coils, condensate pans, or filters (Figure 12), or applied to give broad distribution of UVC energy over an entire enclosure (e.g., mixing box/plenum) that might have microbial activity. Like in-duct air-treatment equipment, systems for surface treatment in air-handling units should be designed to withstand moisture and condensate and selected to operate over a full range of system operating conditions.



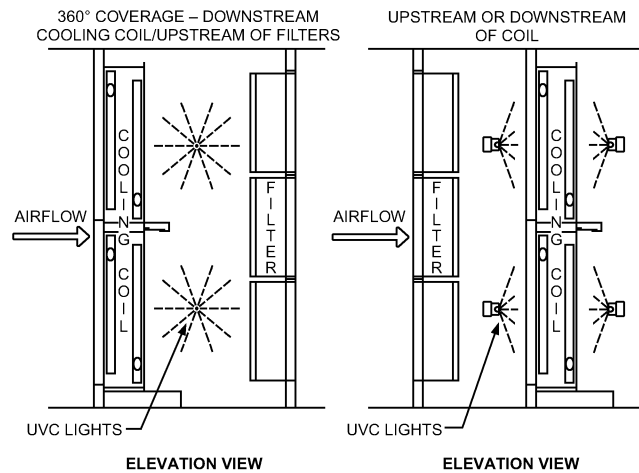


Fig. 12 Section View of Typical HVAC Surface Treatment Installations

### Alternative and Complementary Systems

ASHRAE (2014) identifies the following demonstrated ways of reducing airborne infectious disease transmission:

- UVC
- Dilution, personalized, and source capture ventilation
- In-room airflow control
- Room pressure differentials
- Filtration

From one perspective, these may be viewed as distinct, mutually exclusive alternatives for bioaerosol control. In principle, ventilation alone, filtration alone, or UVC alone can yield the same level of control of a given contaminant source. However, in most cases, multiple modes of air quality control are used in the same system, often as a result of code requirements. For example, air quality codes for commercial buildings based on ASHRAE *Standard* 62.1 minimally require both dilution ventilation and particulate filtration at prescribed levels.

When used in combination with other mandatory air treatment modes, UVC provides an incremental benefit. For example, if a particulate filter removes 85% of a given agent in an incoming air-stream and a UVC system with a single-pass efficiency of 85% for the same contaminant is installed in series with it, the combined filter/UVC system would have a combined single-pass capture and inactivation efficiency of approximately 98% (i.e., the incremental benefit of adding an 85% efficient device is only 13%). Situations involving ventilation, filtration, and UVC can be evaluated quantitatively by analyzing the entire system.

An example of this type of analysis was given by Nazaroff and Wechsler (2009) for several common arrangements of air cleaners in combination with ventilation. The performance of an air cleaner added to a system with ventilation is defined in terms of an effectiveness  $\epsilon$ , which is the difference in contaminant concentration in a space of interest caused by adding an air cleaner and the concentration that would exist without the air cleaner:

$$\epsilon = \frac{C_{baseline} - C_{control}}{C_{baseline}} \quad (4)$$

where  $C_{baseline}$  is the concentration without the air cleaner and  $C_{control}$  is the concentration after addition of the air cleaner. This performance measure would show, for example, that adding UVC to a system with a low ventilation rate would have a higher effectiveness (i.e., greater impact) than adding the same device to the same

system with a higher ventilation rate. The extension of this concept to multiple-space systems and multiple air cleaners and air cleaner types is straightforward. System designers can use such methods to obtain more accurate cost/benefit estimates and to optimize the characteristics and placement of air cleaners.

Even in the absence of the constraints imposed by building codes, the system designer should consider the potential benefits of combining air treatment methods. For example, the cost of particulate filters and their negative impact on fan energy use increase in inverse relation to the sizes of particles to be controlled (i.e., filters for smaller particles tend to be more expensive and have higher pressure drop than filters for larger particles). On the other hand, many larger microorganisms that may be resistant to UVC, such as some fungal spores, can be captured effectively by filters of moderately high efficiency and cost (Kowalski 2009). In addition, using UVC to suppress microbial growth on filters that capture but do not kill is a potential complementary use of these two technologies. Ultimately, the decision to use or not use one of the available, effective microbial control methods should be based on a complete analysis that considers overall performance goals for air quality, impact on energy use, and economic factors. Such an analysis is illustrated for a typical air disinfection system by Lee et al. (2009), as discussed in the following section.

## 5. ENERGY AND ECONOMIC CONSIDERATIONS

The major costs of owning and operating a UVC system include initial equipment and installation costs, maintenance costs (primarily lamp replacement), and energy cost (direct cost of lamp operation plus impact on heating and cooling energy consumption). For a given system, these costs are relatively straightforward to estimate. The benefits of a UVC system are not so easily quantified. Energy use is of concern, and it is also the major operating cost component of most systems. Considerations of energy conservation measures inevitably lead to the issue of cost effectiveness. Therefore, it is appropriate to discuss energy use in conjunction with its economic impact.

Air treatment systems and room surface disinfection systems have the objective of improving the safety, health, and productivity of building occupants through reduced incidence of infectious disease and sick building complaints. Although many studies exist to support claims of UVC's effectiveness in these applications, it is difficult to express the resulting benefits in economic terms. A conservative approach to economic evaluation is to compare the costs of alternative approaches such as dilution ventilation and particulate filtration that have the same effectiveness.

When alternative systems are compared with UVC, all associated costs must be carefully estimated. Increased ventilation adds to heating- and cooling-coil loads and may also affect fan energy use. Particle filtration systems have their own associated installation and maintenance costs and may significantly increase air-side pressure drop and, therefore, fan energy consumption.

Cooling-coil treatment systems have the two-fold objectives of maintaining coil performance and minimizing energy use by reducing air-side flow resistance and increasing the overall heat transfer coefficient relative to a conventionally maintained, mechanically and chemically cleaned coil.

Field studies in the United States (Bahnfleth and Firrantello 2017; Firrantello and Bahnfleth 2017a) and Singapore (Wang et al. 2016a, 2016b) in hot, humid climates report significant improvements in air-side pressure drop and heat transfer coefficient. A system in Tampa, FL, experienced a 22% reduction in pressure drop and 15% increase in air-side heat transfer coefficient after less than two months of surface treatment system operation. Similar results were obtained from a system in Singapore. Improvement in heat transfer coefficient of the Singapore system cooling coil (Wang et

al. 2016b) resulted in a chilled-water flow rate reduction of 8.0 to 11.9% and an increase in chilled-water temperature difference of 0.7 to 1.1°F. Changes in performance in drier climates were less dramatic, as indicated by a laboratory study in Colorado (Luongo et al. 2017; Luongo and Miller 2016) and field data from a system in State College, PA (Bahnfleth and Firrantello 2017). As in the case of air disinfection systems, costs to install and operate coil treatment systems are easily estimated, but though there are many reports of significant improvement in performance, there are relatively few peer-reviewed studies documenting its real-world performance (summarized by Bahnfleth 2017).

Economic analysis of UVC coil treatment based on field measurements (Firrantello and Bahnfleth 2017b; Wang 2017) indicates that energy consumption of germicidal lamps is less than corresponding savings in fan, chiller, and pump energy. However, annual energy savings vary greatly between hot, humid climates where coils are continuously wet and temperate ones in which coils may be dry or inactive for several months per year. Thus, cost effectiveness of coil treatment based on energy savings alone is not certain. Economic performance appears much more favorable when reductions in maintenance cost and improvements in air quality are included in the analysis. Firrantello and Bahnfleth (2017c) modeled effects of air disinfection by a coil treatment system on sick leave for six typical buildings in 16 climate zones. They found that, although typical sizing practices for coil UVC systems only reduced illness-related costs by 3.5%, the monetized value of this improvement was 20 times the energy cost to operate the system.

### Upper-Air UVC Devices

The effectiveness of upper-air UVC performance has often been described in terms of equivalent air changes per hour (ach): that is, by the rate of outside airflow measured in room volumes per hour that would achieve the same reduction of microbial air contamination in a well-mixed space. Riley et al.'s (1976) study of UVGI efficacy found that one 17 W UVC lamp covering 200 ft<sup>2</sup> produced 10 equivalent ach versus a natural die-off of 2 ach when a surrogate for tuberculosis was released in the room. The UVC lamp took less than 20 min to inactivate the bioaerosol, versus over 30 minutes for a natural die-off. In a bioaerosol room study, McDevitt et al. (2008) showed seasonal variations of between 20 to 1000 equivalent ach for a surrogate for smallpox. Ko et al. (2001) modeled the cost of using three air-cleansing strategies to control transmission of tuberculosis in a medical waiting room. They calculated a present value per avoided tuberculin skin test conversion (evidence of infection) of \$1708 for increased ventilation, \$420 for HEPA filtration, and \$133 for upper-air UVC: that is, UVC was less expensive by a factor of 3 to 13. Another metric is cost to provide a typical level of treatment per unit of floor area. The estimated health care benefit, typical of such analyses, was much larger than the cost: roughly \$40/ft<sup>2</sup> per year.

### In-Duct Air Disinfection

Bahnfleth et al. (2009) and Lee et al. (2009) used simulation to investigate the energy use and operating cost of in-duct UVC air treatment applied upstream or downstream of the cooling coil in a cooling-only variable-air-volume system located in New York and compared it with equivalent added particulate filter. A representative MERV 12 filter was estimated to provide the same performance as UVC designed for 85% single-pass inactivation under design conditions. They computed not only the costs associated with the alternatives considered, but also estimated the health benefit using a method based on the Wells-Riley equation as applied by Fisk et al. (2005). They found that locating the UVC system upstream of the cooling coil in the normally warmer mixed-air section of the air-handling unit reduced its required size by roughly 50% relative to a downstream location using typical in-duct lamp characteristics.

Annual energy cost at an average electric rate of \$0.10/kW·h (\$0.03/1000 Btu) was approximately \$0.02/ft<sup>2</sup> for the downstream location and \$0.01/ft<sup>2</sup> for the upstream location, whereas the additional MERV 12 filter cost \$0.10/ft<sup>2</sup>. Annualized life-cycle cost, including installation and maintenance, was \$0.74/ft<sup>2</sup> for the downstream location, \$0.38/ft<sup>2</sup> for the upstream location, and \$1.79/ft<sup>2</sup> for MERV 12 filtration. The drawback to the more economically advantageous upstream UVC location is that it is considered a less favorable location for cooling coil irradiation, which many air treatment systems are designed to do as a benefit of increased airflow and heat exchange efficiency and reduced coil cleaning.

### Upper-Air Versus In-Duct

Economic factors clearly favor an upper-air fixturing when the building being treated with UVC has no air distribution system. When a recirculating central air distribution system is present, a choice becomes possible between upper-air devices, which must be distributed throughout occupied spaces, and in-duct systems, which can be centralized. As noted in the preceding discussion of in-duct systems, an annual operating cost of \$0.01 to 0.02/ft<sup>2</sup> is possible at an electric rate of \$0.10/kW·h (\$0.03/1000 Btu). The same study (Lee et al. 2009) estimated an installed cost for equipment of \$0.13 to 0.25/ft<sup>2</sup>. By comparison, a typical upper-air system might cost more than \$2/ft<sup>2</sup> to install and more than \$0.10/ft<sup>2</sup> to operate, based on typical sizing procedures and current equipment costs. This comparison seems to strongly favor in-duct systems where they are applicable, but is based on an assumption of equal performance that may not be valid. In a health care setting, controlling transmission of airborne pathogens at their source would suggest an upper-room approach. However, where feasible, a whole-building approach to UV should be considered.

### Cooling Coil Surface Treatment

Cooling coil surface treatment is done as an alternative to periodic mechanical and chemical cleaning of coils. By suppressing the formation of biofilms and mold growth on coils, coil irradiation should reduce air-side pressure drop, increase heat transfer coefficient, and reduce both fan and refrigeration system energy consumption. Several studies have documented the ability of coil irradiation to reduce microbial growth (Levetin et al. 2001; Shaughnessy et al. 1998). No peer-reviewed studies have yet been published to document the effect of coil irradiation on energy consumption, but there are many strong anecdotal reports of its effectiveness. As noted previously, the U.S. General Services Administration has sufficient confidence in this application to include it in its mechanical requirements (GSA 2018).

## 6. ROOM SURFACE TREATMENT

Environmental contamination in health care settings and transmission of health-care-associated pathogens to patients occurs most frequently via contaminated hands of health care workers and transmission of pathogens to patients (Boyce 2010). A primary concern in health care settings has been reducing nosocomial infections and finding new approaches for these environments to help eliminate infections from hospital settings. Hospital-acquired infections generate a high financial burden for the health care industry and the consumer. In the United States, an estimated 1.7 million hospital-acquired infections occur annually, leading to about 100,000 deaths (U.S. HHS 2009). UVC for surface disinfection, particularly in health care settings, has been applied to reduce the number of microorganisms on surfaces, and consequently UVC should contribute to a reduction in these healthcare-acquired infections (HAIs). Scientific studies have shown reductions in viable infectious agents on surfaces after UV exposure. However, further evidence of reductions in HAIs is needed, as well as a method to test

various portable UVC devices being used for “whole-room” decontamination.

Various portable UVC devices are available for hospitals, which can be easily moved into patient rooms, surgical suites, ICUs, and other critical areas that need surface and air disinfection during a terminal cleaning process or when a patient is diagnosed with a disease transferred by pathogens. Some of the pathogens of interest and their reduction in health care settings are multidrug resistant, such as methicillin-resistant *Staphylococcus aureus* (MRSA), *Clostridium difficile*, *Acinetobacter baumannii*, and vancomycin-resistant *Enterococci* (VRE). These pathogens can be inactivated by proper application of UVC energy. A study by Rastogi et al. (2007) investigated the efficacy of UVC disinfection of *Acinetobacter baumannii* on contaminated surfaces relevant to medical treatment facilities. The UVC exposure to surfaces resulted in  $\geq 4$ -log (CFU) reduction in viable cells of *A. baumannii*.

UVC fixtures can also be installed in surgical suites to disinfect surfaces and air between or during procedures. A 19-year study on UVC during orthopedic surgery showed that 47 infections occurred following 5980 joint replacements. The infection rates for total hip replacements decreased from 1.03% to 0.72% ( $p = 0.5407$ ), and for total knee replacements from 2.20% to 0.5% ( $p < 0.0001$ ). The study concluded that UVC appears to be an effective way to lower the risk of infection in the operating room during total joint replacement (Ritter et al. 2007). Safety precautions must be followed when applying UVC during surgery to protect workers from accidental exposure (see the following discussion of intensity of source) or upper air fixtures may be used as discussed previously. Tools used in healthcare applications can be irradiated with UVC for simple surface disinfection. However, UV irradiation should never replace sterilization of surgical instruments.

UVC surface disinfection could also be applied in schools, morgues, nursing homes, and homeless shelters: surfaces can be irradiated with fixed or portable in-room UVC fixtures that serve as part of the room’s disinfection methodology.

Application of UVC to any surface is based on the UV dose delivered to the surface. The dose ( $\mu\text{J}/\text{cm}^2$ ) of UVC needed to disinfect a surface depends on the selected target and desired disinfection level. Different microorganisms require various levels of UVC energy for inactivation (see Figure 4). Vegetative forms of bacteria tend to be more susceptible to UVC energy than spore-forming microorganisms. UVC irradiates all line-of-sight objects and into shadowed areas (e.g., tables, chairs, surgical equipment, objects) through reflection, so the desired level of disinfection can be achieved, even on surfaces which are not directly irradiated. Different materials absorb and reflect UVC energy at different rates, depending on the overall reflectivity of the materials, irradiation time, and intensity. UVC surface disinfection should only be applied as an adjunct to normal surface cleaning procedures of the facility. No living organisms, including animals and plants, should be in the room when UVC is used. It should be noted that most organic compound-based materials degrade when exposed to UVC energy.

The same principles as for in-duct applications apply here. There are two primary methods of UVC delivery: direct (line of sight) and indirect (reflection). Most surface applications use a direct source, where the source (typically a mercury vapor lamp) is contained in an assembly designed to direct the UVC energy at a particular surface or in a particular direction with no impedance to the energy beam. In an indirect application, the energy is reflected onto a surface using a reflective material. The reflected UVC energy can be measured to determine accurately when a given amount of the UVC dose has been delivered to the desired target.

The basics of determining the radiant energy levels to a surface are as follows:

**Length of exposure.** When disinfecting surfaces, it must be first determined if the target is moving or stationary. This helps to determine if there are any limiting factors associated with the length of exposure time. In most surface disinfection applications, time is relative to intensity, meaning that increasing the intensity of the source can decrease the exposure time necessary. It is important to remember that microorganisms vary, requiring a higher or lower intensity for inactivation, depending on their structure (Brickner et al. 2003).

**Intensity of source.** UVC lamp and equipment manufacturers normally provide the intensity of a given source (lamp or fixture) at a given distance. A distance correction factor may be needed when calculating a desired dose or intensity for a surface. UVC energy follows the same inverse square law for intensity as visible energy and other electromagnetic sources: the amount of energy at the surface is measured in proportion to the square of the distance from the energy’s source (UVC lamp), assuming no loss through scattering or absorption. Temperature and airflow corrections may also be necessary, depending on the location of the application. The intensity of a source is given in power per unit area (i.e.,  $\mu\text{W}/\text{cm}^2$ ).

**Distance from source to surface.** In a point irradiation application, the distance is relatively easy to calculate. Calculating time requirements and intensity levels for a three-dimensional object or space is more complex. The varying distances from the source are the first challenge, because the object itself creates a shadowing effect, and any shadows from the local environment must be taken into consideration. However, portable devices are available that effectively measure the reflected dose from shadow areas and offer quantifiable results.

Studies on in-room UVC disinfection devices have shown that UVC can be successfully applied to reduce microbiological loads of surfaces located in shadow areas in addition to line of sight (Rutala 2009). The reductions were up to 4-log for organisms such as MRSA, VRE, *Acinetobacter*, and *C. difficile*. Furthermore, it was concluded that UV room decontamination with the test device reduced colony counts of pathogens by greater than 99.9% within 20 min. Note that, depending on the portable or stationary UVC device, performance could greatly differ with respect to irradiation time, because overall dose delivered to surfaces is the critical measure of portable device performance.

## 7. SAFETY

### Hazards of Ultraviolet Radiation to Humans

UVC is a low-penetrating form of UV compared to UVA or UVB. Measurements of human tissue show that 4 to 7% of UVC (along with a wide range of wavelengths, 250 to 400 nm) is reflected (Diffey 1983) and absorbed in the first 2  $\mu\text{m}$  of the stratum corneum (outer dead layer of human skin), thus minimizing the amount of UVC transmitted through the epidermis (Bruls 1984).

Although UV is far more energetic than the visible portion of the electromagnetic spectrum, it is invisible to humans. Therefore, exposure to ultraviolet energy may result in transient corneal inflammation, which can go unnoticed.

Ocular damage generally begins with **photokeratitis** (inflammation of the cornea) but can also result in **photokeratoconjunctivitis** (inflammation of the conjunctiva [ocular lining]). Symptoms, which may not be evident until several hours after exposure, may include an abrupt sensation of sand in the eyes, tearing, and eye pain, possibly severe. These symptoms usually appear within 6 to 12 h after UV exposure, and resolve fully within 24 to 48 h. Acute overexposure to UVC radiation may cause some incapacity due to eye discomfort, but this generally abates after several days, leaving no permanent damage.

**Cutaneous damage** consists of erythema, a reddening of the skin akin to sunburn (but without tanning). The maximum effect of

erythema occurs at a wavelength of 296.7 nm in the UVB band. UVC radiation at a wavelength of 253.7 nm is less effective in causing erythema. Because ultraviolet radiation is carcinogenic, questions have been raised concerning open-air UVC systems. The International Commission on Illumination (CIE) completed a review of UVC photocarcinogenesis risks from germicidal lamps using basic biophysical principles: because of the attenuation provided by the stratum corneum and epithelial tissues of the skin, upper-air disinfection can be safely used without significant risk for long-term delayed effects such as skin cancer (CIE 2010).

### Sources of UV Exposure

UVC energy does not normally penetrate through solid substances and is attenuated by most materials. Quartz glass, soda barium glass, and TFPE plastic have high transmissions for UVC radiation.

UVC energy can reflect from most metals and several types of painted and nonpainted surfaces; however, a surface's ability to reflect visible light cannot be used to indicate its UV reflectance. The fact that a blue glow can be observed on a metal surface from an operating low-pressure UV fixture lamp could indicate the presence of UV, and a measurement should be performed to ensure there is no exposure risk. The lack of reflected blue light clearly indicates the absence of UV energy. Note that ultraviolet energy is invisible to the normal human eye; however, it follows the same optical path as the visible blue light spectrum generated by the UVC lamp.

Well-designed and commissioned UVC installations, education of maintenance personnel, signage, and use of safety switches can help to avoid overexposure. During commissioning and before operation of the UVC installation, hand-held radiometers with sensors tuned to read the specific 254 nm wavelength should be used to measure stray UVC energy and should be used in upper-air systems.

### Exposure Limits

In 1972, the Centers for Disease Control and Prevention (CDC) and National Institute for Occupational Safety and Health (NIOSH) published a **recommended exposure limit (REL)** for occupational exposure to UV radiation. The REL is intended to protect workers from the acute effects of UV exposure, although photosensitive persons and those exposed concomitantly to photoactive chemicals might not be protected by the recommended standard.

Exposures exceeding CDC/NIOSH REL levels require that workers use personal protective equipment (PPE), which consists of eyewear and clothing known to be nontransparent to UVC penetration and which covers exposed eyes and skin.

UV inspection, maintenance, and repair workers typically do not remain in one location during their workday, and therefore are not exposed to UV irradiance levels for 8 h. Threshold Limit Value® (TLV®) consideration should be based on real-time occupancy of spaces treated by UVC (ACGIH 2007; Sliney 2013). This recommendation is supported by UV monitoring data from First et al. (2005), which showed that peak meter readings poorly predict actual exposure of room occupants.

### Evidence of Safety

During the height of the tuberculosis resurgence in the United States in the 1990s, the Tuberculosis Ultraviolet Shelter Study (TUSS), a double-blind, placebo-controlled field trial of upper-air UVC, was conducted at 14 homeless shelters in six U.S. cities from 1997 to 2004 (Brickner et al. 2000). Following available recommended placement, installation, and maintenance guidelines, each building in the study was evaluated for treatment with upper-air UVC fixtures. At the conclusion of the study, the safety of room occupants was evaluated using data from a total of 3,611 staff and homeless study subjects regarding eye and skin irritation. Analysis showed no statistically significant difference in the number of reports of symptoms between the

active and placebo periods. There was one definite instance of UV-related photokeratoconjunctivitis (from eye overexposure). This occurred from a placement of an elevated bunk bed in a dormitory where a single bed had been used when the UV fixtures were first installed. By moving the UV fixture, this incident was resolved (Brickner and Vincent 2013). This study demonstrated that, with careful application, side effects of UV overexposure can be avoided. Because of the enclosed nature of in-duct UVC systems, with careful adherence to safety guidelines, these systems should not result in UV exposure.

Because in-duct UVC systems are installed inside air-handling units or ventilation ductwork, typical building occupants are not expected to be exposed to UV energy. On the other hand, building facilities workers and maintenance personnel are at risk of high UV exposures with in-duct systems. To minimize the risk to these workers, UVC systems should be designed with specific safety features and all workers that could potentially work around the UV fixtures should receive UV-specific training.

### Safety Design Guidance

**Upper-air systems** should have on/off switches and an electrical disconnect device on the louvers. If UV radiation measurements at the time of initial installation exceed the recommended exposure limit, all highly UV-reflecting materials should be removed, replaced, or covered. UV-absorbing paints containing titanium oxide can be used on ceilings and walls to minimize reflectance in the occupied space.

Warning labels must be posted on all upper-air UV fixtures to alert personnel to potential eye and skin hazards. Damaged or illegible labels must be replaced as a high priority. Warning labels must contain the following information:

- Wall sign for upper-air UVC  
**Caution:** Ultraviolet energy. Switch off lamps before entering upper room.
- General warning posted near UVC lamps.  
**Caution:** Ultraviolet energy. Protect eyes and skin.

Upper-air UVC fixtures can vary widely in their luminaire efficiency factors, which rates the performance of emitted UVC from a fixture. Zhang et al. (2012) developed a protocol and performed gonioradiometric measurements (i.e., measuring both radiance and irradiance at concurrent angles) for upper-air UVGI fixtures, which is now being used to test total UVC fixture output (Leuschner and Salie 2013). These gonioradiometric measurements are reported in standard IES format compatible with computer-aided design (CAD) lighting software adapted for use with upper room UVC devices (Rudnick et al. 2012; Vincent et al. 2013).

**In-duct systems** should be fully enclosed and sealed to prevent leakage of UV radiation to unprotected persons or materials outside of the HVAC equipment. The fifth edition of UL *Standard* 1995, which carries a November 2019 compliance date, requires that no opening permit leakage of UVC greater than  $0.1 \mu\text{W}/\text{cm}^2$ , and that points of intentional access to UV sources must be equipped with an interlocking mechanism that deenergize the UV source. All access panels or doors to the lamp chamber and panels or doors to adjacent chambers where UV radiation may penetrate or be reflected should be interlocked and have warning labels posted in appropriate languages. Labels should be placed on the outside of each panel or door, in a prominent location visible to people accessing the system. At a minimum, the labels should state

- General warning posted near UVGI/UVC lamps  
**Caution:** Ultraviolet energy. Protect eyes and skin.
- Multilingual warning posted on the door of air handlers where UVC is present in ductwork.

**Caution:** Ultraviolet energy in duct. Do not switch off safety button or activate lamps with door open.

Lamp chambers should have door safety interlock switches and electrical disconnect devices. Disconnection devices must be able to be locked or tagged out, and should be located outside the lamp chamber, next to the chamber's primary access panel or door. Switches should be wired in series so that opening any switched access deenergizes the system. It is recommended that on/off switches for UV lamps not be located in the same location as general room lighting; instead, they should be in a location that only authorized persons can access and should be locked or password protected to ensure that they are not accidentally turned on or off.

The lamp chamber should have one or more viewports of UVC-absorbing materials. Viewports should be sized and located to allow an operating UV system to be viewed from outside of the HVAC equipment.

## 8. INSTALLATION, START-UP, AND COMMISSIONING

The operating instructions and advice of UVC system designers and lamp manufacturers should always be followed to ensure the proper operation of any UVGI/UVC system. It is important to operate any such system within the temperature and relative humidity ranges considered during the system design process. The following section presents some general guidelines for initially verifying and maintaining adequate system performance.

### Upper-Air UVC Devices

Those responsible for the commissioning process should inspect fixture placement and eye level irradiance measurements using a 254 nm selective radiometer. UVC levels can be measured with a UV radiometer directly facing the device at eye height at various locations in a room and must be taken in the same location each time. UVC measurements should be taken at eye level (between 5.5 and 6.0 ft) at compass points from each fixture. Check reflective surfaces (e.g., TVs, monitors). CAD software can be used to preview safety of UVGI/UVC upper room installations (Vincent et al. 2013). Incorporate readings into final commissioned drawings. If the readings indicate an eye-level exposure that exceeds the 8 h TLV for UVC of  $6 \mu\text{J}/\text{cm}^2$ , the UV systems must be deactivated until adjustments can be made or the manufacturer can be contacted. Measurements should be made at initial installation, whenever new UV lamps are installed (newer lamp designs may provide increased irradiance), and whenever modifications are made to the UVC device or room (e.g., adjusting fixture height, relocating or repositioning louvers, adding UV-absorbing or -reflecting materials, changing room dimension or modular partition height).

### In-Duct UVC Systems

Installation, start-up, and commissioning of in-duct UVGI systems are straightforward. Those responsible for installation should ensure that the system is installed as designed and that all lamps, ballasts, and/or fixtures are the same as included in the final design. Take care to ensure that all safety interlocks and view ports are installed in appropriate positions and functional. Once the UV lamps are powered on, ensure that all lamps are burning. Unfortunately, there are no good methods for in situ testing of in-duct system performance, so relying on final design parameters is essential to ultimate system performance.

## 9. MAINTENANCE

All UVC systems require periodic inspection, maintenance, and lamp replacement to ensure proper system performance. Whenever maintenance is performed on UVC systems, the appropriate safety

guidelines outlined elsewhere in this chapter should be carefully followed.

### Material Degradation

UVC energy can be detrimental to most organic materials. If the UVC is not applied properly and sensitive materials are not shielded or substituted, degradation can occur. However, the degradation may not be enough to cause failure of the material if UVC only penetrates micrometers into the material before the degradation plateaus off, leaving a still fully functional material, as found by ASHRAE research project RP-1509, sponsored by TC 2.9 (Kauffman 2010). Air filters are known to be sensitive to degradation by UVC, especially those made from synthetic materials. Glass fibers by themselves are unaffected by UV exposure, but binding materials in glass fiber filters may be degraded. As a general rule, synthetic air filters should not be exposed to UVC.

Lower doses, or those typically sized for cooling coil surface treatment, of UVC exposure to organic materials resulted in much slower rates of degradation (Kauffman 2017). Although UVC photodegradation is of concern, with the selection of the proper material or metallic shielding of components, the problem is significantly reduced, and components can be expected to meet product design life. As a simple, practical approach, it is wise to shield all organic material components within about 5 ft of the UV lamp. Some indoor plants do not tolerate prolonged UVC exposure and should not be hung higher in the room where upper air UVC devices are installed.

### Visual Inspection

Maintenance personnel should routinely perform periodic visual inspection of the UVC lamp assembly. Typically, a viewing port or an access door window is sufficient for in-duct applications. Closer visual inspection may be required for upper-air systems because a single burned-out lamp in a multilamp fixture may not be apparent from the lower room. Personal protective measures are required for this close-up inspection.

Any burned-out or failing lamps should be replaced immediately. If lamps become dirty in dusty environments, they should be cleaned with a lint-free cloth and isopropyl alcohol. Care should be taken to ensure no film remains on the surface of the lamps after cleaning. This film could reduce UV output from the lamp. Complete lamp fixtures should be replaced whenever they are visibly damaged or in accordance with manufacture warranty guidelines.

### Radiometer

Another means of monitoring UVC lamps is with a stationary or portable radiometer. These are generally used to monitor the "relative" output of the UVC system by measuring the UV intensity produced by the lamps. Caution is needed when using a radiometer in critical applications, because these devices are intended only to give a relative indication of the lamp output unless measured identically each time with a calibrated instrument. Radiometer sensors can degrade over time with constant exposure to UV. If accurate measurements of UV intensity are required, a calibrated laboratory radiometer should be used, and readings must always be taken in the exact position each time as the readings are extremely sensitive to inverse square law losses and gains.

### Lamp Replacement

UVC lamps should be replaced at the end of their useful life, based on equipment manufacturer recommendations or radiometer measurements. Where applicable, it may be prudent simply to change lamps annually (8760 h when lamps are run continuously) to ensure that adequate UV energy is supplied by a given system. Lamps can operate after their useful life, but at reduced performance, and require regular measurement to ensure that a maintained level of UVC is being generated. A blue visible light emitted from

the lamp does *not* indicate that UVC is present. The typical rated life of UVC lamps is 9000 h of operation. Switching lamps on and off too often may lead to early lamp failure, depending on the ballast type used. Consult the lamp manufacturer for specific information on expected lamp life and effects of switching.

### Lamp and Ballast Disposal

UVC lamps should be treated in the same manner as other mercury-containing devices, such as fluorescent lamps. Some lamps may need to be treated as hazardous waste and not discarded with regular waste, although low mercury lamps may be an exception; however, check state and local codes for proper determination. The U.S. EPA's universal waste regulations allow users to treat mercury lamps as regular waste for transport to a recycling facility (EPA 2018). This simplified process was developed to promote recycling. The National Electrical Manufacturers Association maintains an online list of companies claiming to recycle or handle used mercury lamps (NEMA 2009). The most stringent of local, state, or federal regulations for disposal should be followed.

UVC systems currently depend on the use of an electronic ballast to provide the UV lamp with power; however, many older systems used magnetic ballasts instead. Magnetic ballasts manufactured before 1979 contain polychlorinated biphenols (PCB) in the dielectric of their capacitors (EPA 2017). Recycling is the best way to dispose of all magnetic ballasts. The process allows the reuse of copper and aluminum wire, steel laminations, and steel cases, and disposes of capacitors and potting compound as hazardous waste in high-temperature incinerators.

Failed electronic ballasts should be treated as electronic waste. Many lamp and ballast recyclers are expanding their businesses and becoming certified to accept electronic waste. Some recyclers now accept both lamps and electronic ballasts.

### Personnel Safety Training

Workers should be provided with as much training as necessary, including health and safety training, and some degree of training in handling lamps and materials. Workers should be made aware of hazards in the work area and trained in precautions to protect themselves. Training topics include the following:

- UVC exposure hazards
- Electrical safety
- Lock-out/tag-out (for in-duct units)
- Health hazards of mercury
- Rotating machinery (for in-duct units)
- Slippery condensate pans (for in-duct units)
- Sharp unfinished edges (for in-duct units)
- Confined-space entry (if applicable) (for in-duct units)
- Emergency procedures

Workers expected to clean up broken lamps should be trained in proper protection, cleanup, and disposal.

No personnel should be subject to direct UV exposure, but if exposure is unavoidable, personnel should wear protective clothing (no exposed skin), protective eyewear, and gloves. Most types of eyewear, including prescription glasses, may be sufficient to protect eyes from UV, but not all offer complete coverage. Standard-issue safety goggles or clear full-face masks may be the best alternative.

If individual lamp operating conditions must be observed, this should preferably be done using the view port or window(s).

During maintenance, renovation, or repair work in rooms with upper-air UV systems, all UVC devices must be deactivated before personnel enter the upper part of the room.

For in-duct systems, access to lamps should be allowed only when lamps are deenergized. The lamps should be turned off before air-handling unit (AHU) or fan shutdown to allow

components to cool and/or to purge any ozone in the lamp chamber (if ozone-producing lamps are used). If AHUs or fans are deenergized first, the lamp chamber should be opened and allowed to ventilate for several minutes. Workers should always wear protective eyewear and puncture-resistant gloves for protection in case a lamp breaks.

Access to the lamp chamber should follow a site-specific lock-out/tag-out procedure. Do not rely on panel and door safety switches as the sole method to ensure lamp deenergizing. Doors may be inadvertently closed, or switches may be inadvertently contacted, resulting in unexpected lamp activation.

If workers enter the condensate area of equipment, the condensate pan should be drained and any residual water removed.

In general, avoid performing readings with the fan running and workers inside an AHU (e.g., only to test for output reduction caused by air cooling). Tests of this nature should be instrumented and monitored from outside the equipment.

### Lamp Breakage

If workers break a lamp, they should warn all other workers to exit the HVAC equipment area. Panels or doors should be left open and any additional lamp chamber access points should also be opened. Do not turn air-handling unit fans back on. After 15 min, workers may reenter the HVAC equipment to begin lamp clean-up.

If a lamp breaks in a worker's hand, the worker should not exit the HVAC equipment with the broken lamp. The worker should carefully set the broken lamp down, and then exit the space. When possible, try not to set the broken lamp in any standing condensate water. Follow standard ventilation and reentry procedures.

Cleanup requires special care because of mercury drop proliferation and should be performed by trained workers. As a minimum, workers should wear cut-resistant gloves, as well as safety glasses to protect eyes from glass fragments. Large bulb pieces should be carefully picked up and placed in an impervious bag. HEPA-vacuum the remaining particles, or use other means to avoid dust generation.

## REFERENCES

ASHRAE members can access *ASHRAE Journal* articles and ASHRAE research project final reports at [technologyportal.ashrae.org](http://technologyportal.ashrae.org). Articles and reports are also available for purchase by nonmembers in the online ASHRAE Bookstore at [www.ashrae.org/bookstore](http://www.ashrae.org/bookstore).

ACGIH. 1999. *Bioaerosols: Assessment and control*, Ch. 9: Respiratory infections—Transmission and environmental control, by E.A. Nardell and J.M. Macher. American Conference on Governmental Industrial Hygienists, Cincinnati, OH.

ACGIH. 2007. *TLVs® and BEIs®*. American Conference of Governmental Industrial Hygienists, Cincinnati, OH.

ASHRAE. 2016. Ventilation for acceptable indoor air quality. ANSI/ASHRAE *Standard* 62.1-2016

ASHRAE. 2015. Method of testing UV-C lights for use in air-handling units or air ducts to inactivate airborne microorganisms. ANSI/ASHRAE *Standard* 185.1-2015.

ASHRAE. 2014. Method of testing ultraviolet lamps for use in HVAC&R units or air ducts to inactivate microorganisms on irradiated surfaces. ANSI/ASHRAE *Standard* 185.2-2014.

ASHRAE. 2003. Risk management guidance for health, safety, and environmental security under extraordinary incidents. *Report*, Presidential Ad Hoc Committee for Building Health and Safety under Extraordinary Incidents.

ASHRAE. 2009. *Indoor air quality guide: Best practices for design, construction, and commissioning*.

ASHRAE. 2014. *Position document on airborne infectious diseases*.

Bahnfleth, W. 2011. Cooling coil ultraviolet germicidal irradiation. *ASHRAE Journal* 53(4):70-72.

Bahnfleth, W. 2017. UVGI in air handlers. *ASHRAE Journal* 59(10):72-74.

Bahnfleth, W., and J. Firantello. 2017. Field measurement and modeling of UVC cooling coil irradiation for HVAC energy use reduction. ASHRAE Research Project RP-1738, *Final Report*.

- Bahnfleth, W., B. Lee, J. Lau, and J. Freihaut. 2009. Annual simulation of in-duct ultraviolet germicidal irradiation system performance. *Proceedings of Building Simulation 2009, The 11th International Building Performance Simulation Association Conference and Exhibition*, Glasgow.
- Bierman, A., and J. Brons. 2007. *Field evaluation of ultraviolet germicidal irradiation (UVGI) in an air duct system*. Lighting Research Center, RPI, Troy, NY. [www.lrc.rpi.edu/researchAreas/pdf/FieldEvaluationUVGIReport.pdf](http://www.lrc.rpi.edu/researchAreas/pdf/FieldEvaluationUVGIReport.pdf).
- Boyce, P. 2003. *Controlling tuberculosis transmission with ultraviolet irradiation*. Rensselaer Polytechnic Institute, Troy, NY.
- Boyce, J. 2010. When the patient is discharged: Terminal disinfection of hospital rooms. *Medscape.com*. [www.medscape.com/viewarticle/723217](http://www.medscape.com/viewarticle/723217) (requires free registered account).
- Brickner, P.W., and R.L. Vincent. 2013. Ultraviolet germicidal irradiation safety concerns: A lesson from the tuberculosis ultraviolet shelter study Murphy's law affirmed. *Photochemistry and Photobiology* 89 (4):819-821. [dx.doi.org/10.1111/php.12034](https://doi.org/10.1111/php.12034).
- Brickner, P.W., R.L. Vincent, E.A. Nardell, C. Pilek, W.T. Chaisson, M. First et al. 2000. Ultraviolet upper room air disinfection for tuberculosis control: An epidemiological trial. *Journal of Healthcare Safety Compliance & Infection Control* 4(3):123-131.
- Brickner, P.W., R.L. Vincent, M. First, E. Nardell, M. Murray, and W. Kaufman. 2003. The application of ultraviolet germicidal irradiation to control transmission of airborne disease: Bioterrorism countermeasure. *Public Health Report* 118(2):99-114.
- Brickner, P.W., et al. 2009. Computer aided design for UVGI. NYSERDA Project 9425. St. Vincent's Hospital, New York.
- Bruls, W. 1984. Transmission of human epidermis and stratum corneum as a function of thickness in the ultraviolet and visible wavelengths. *Journal of Photochemistry and Photobiology* 40:485-494.
- Buttolph, L.J., and H. Haynes. 1950. Ultraviolet air sanitation. *General Electric Report* LD-11.
- CDC. 2002. *Comprehensive procedures for collecting environmental samples for culturing Bacillus anthracis*. Centers for Disease Control and Prevention, Atlanta. [www.cdc.gov/niosh/topics/emres/unp-envsamp.html](http://www.cdc.gov/niosh/topics/emres/unp-envsamp.html).
- CDC. 2005. Guidelines for preventing the transmission of *Mycobacterium tuberculosis* in health-care settings. *Morbidity and Mortality Weekly Report (MMWR)* 37-38, 70-75.
- CIE. 2011. *International lighting vocabulary*. Commission Internationale de L'Éclairage, Vienna. CIE S 017/E:2011.
- CIE. 1993. CIE collection in photobiology and photochemistry. *Publications* 106/1 (Determining ultraviolet action spectra), 106/2 (Photokeratitis), and 106/3 (Photoconjunctivitis). Commission Internationale de L'Éclairage, Vienna.
- CIE. 2003. Ultraviolet air disinfection. *Publication* 155:2003. Commission Internationale de L'Éclairage, Vienna.
- CIE. 2010. UV-C photocarcinogenesis risks from germicidal lamps. *Publication* 187:2010. Commission Internationale de L'Éclairage, Vienna.
- Coker, A., E. Nardell, P. Fourie, W. Brickner, S. Parsons, N. Bhagwandin, and P. Onyebujoh. 2001. *Guidelines for the utilisation of ultraviolet germicidal irradiation (UVGI) technology in controlling the transmission of tuberculosis in health care facilities in South Africa*. South African Centre for Essential Community Services and National Tuberculosis Research Programme, Medical Research Council, Pretoria.
- Diffey, B.L. 1983. A mathematical model for ultraviolet optics in skin. *Physics in Medicine and Biology* 28:657-747.
- Diffey, B.L. 1991. Solar ultraviolet radiation effects on biological systems. *Physics in Medicine and Biology* 36(3):299-328.
- EPA. 2017. *Storage and disposal: Ballasts*. U.S. Environmental Protection Agency, Washington, D.C. [www3.epa.gov/region9/pcbs/ballast.html](http://www3.epa.gov/region9/pcbs/ballast.html).
- EPA. 2018. *Universal wastes*. U.S. Environmental Protection Agency, Washington, D.C. [www.epa.gov/hw/universal-waste](http://www.epa.gov/hw/universal-waste).
- Escombe, A.R., R.H. Gilman, M. Navincopa, E. Ticona, B. Mitchell, C. Noakes, C. Martínez, P. Sheen, R. Ramirez, W. Quino, A. Gonzalez, J.S. Friedland, and C.A. Evans. 2009. Upper-room ultraviolet light and negative air ionization to prevent tuberculosis transmission. *PLoS Med* 17(6).
- Firrantello, J., and W. Bahnfleth. 2017a. Field measurement and modeling of UVC cooling coil irradiation for HVAC energy use reduction (RP-1738)—Part 1: Field measurements. *Science and Technology for the Built Environment* 24(6):588-599.
- Firrantello, J., and W. Bahnfleth. 2017b. Field measurement and modeling of UVC cooling coil irradiation for HVAC energy use reduction (RP-1738)—Part 2: Energy, indoor air quality, and economic modeling. *Science and Technology for the Built Environment* 24(6):600-611.
- Firrantello, J., and W. Bahnfleth. 2017c. Simulation and monetization of collateral airborne infection risk improvements from UVGI for coil maintenance. *Science and Technology for the Built Environment* 24(2):135-148.
- First, M.W., E.A. Nardell, W.T. Chaisson, and R.L. Riley. 1999. Guidelines for the application of upper-room ultraviolet irradiation for preventing transmission of airborne contagion—Part 1: Basic principles. *ASHRAE Transactions* 105(1):869-876.
- First, M.W., R.A. Weker, S. Yasui, and E.A. Nardell. 2005. Monitoring human exposures to upper-room germicidal ultraviolet irradiation. *Journal of Occupational and Environmental Hygiene* 2:285-292.
- First, M.W., F.M. Rudnick, K. Banahan, R.L. Vincent, and P.W. Brickner. 2007a. Fundamental factors affecting upper-room ultraviolet germicidal irradiation—Part 1: Experimental. *Journal of Environmental Health* 4:1-11.
- First, M.W., K. Banahan, and T.S. Dumyahn. 2007b. Performance of ultraviolet light germicidal irradiation lamps and luminaires in long-term service. *LEUKOS* 3:181-188.
- Fisk, W.J., O. Seppanen, D. Faulkner, and J. Huang. 2005. Economic benefits of an economizer system: Energy savings and reduced sick leave. *ASHRAE Transactions* 111(2).
- Gilkeson, C.A., and C. Noakes. 2013. Application of CFD simulation to predicting upper-room UVGI effectiveness. *Photochemistry and Photobiology* 89(4):799-810. [onlinelibrary.wiley.com/doi/abs/10.1111/php.12013](https://onlinelibrary.wiley.com/doi/abs/10.1111/php.12013).
- GSA. 2018. *The facilities standards for the Public Buildings Service*. Public Buildings Service of the General Services Administration, Washington, D.C.
- Harm, W. 1980. *Biological effects of ultraviolet radiation*. Cambridge University Press, New York.
- Hollaender, A. 1943. Effect of long ultraviolet and short visible radiation (3500 to 4900 Å) on *Escherichia coli*. *Journal of Bacteriology* 46(6):531-541.
- IESNA. 2000. *The IESNA lighting handbook*, 9th ed., Ch. 5: Nonvisual effects of optical radiation. M.S. Rea ed. Illuminating Engineering Society of North America, New York, NY.
- Jensen, M.M. 1964. Inactivation of airborne viruses by ultraviolet irradiation. *Applied Microbiology* 12(5):418-420.
- Ka, M., H.A.B. Lai, and M.W. First. 2004. Size and UV germicidal irradiation susceptibility of *Serratia marcescens* when aerosolized from different suspending media. *Applied and Environmental Microbiology* (April):2021-2027.
- Kauffman, R.E. 2010. Study the degradation of typical HVAC materials, filters and components irradiated by UVC energy. ASHRAE Research Project RP-1509, *Final Report*.
- Kauffman, R.E. 2017. Study the HVAC system photodegradation caused by the low level UVC light irradiance used for coil maintenance and air stream disinfection. ASHRAE Research Project RP-1724, *Final Report*.
- Kethley, T.W., and K. Branc. 1972. Ultraviolet lamps for room air disinfection: Effect of sampling location and particle size of bacterial aerosol. *Archives of Environmental Health* 25(3):205-214.
- Ko, G., M.W. First, and H.A. Burge. 2000. Influence of relative humidity on particle size and UV sensitivity of *Serratia marcescens* and *Mycobacterium bovis* BCG aerosols. *Tubercle and Lung Disease* 80(4-5):217-228.
- Ko, G., H. Burge, E. Nardell, and K. Thompson. 2001. Estimation of tuberculosis risk and incidence under upper room ultraviolet germicidal irradiation in a waiting room in a hypothetical scenario. *Risk Analysis* 21(4):657-673.
- Kowalski, W.J. 2003. *Immune building systems technology*. McGraw-Hill, New York.
- Kowalski, W.J. 2006. *Aerobiological engineering handbook*. McGraw-Hill, New York.
- Kowalski, W. 2009. *Ultraviolet germicidal irradiation handbook*. Springer-Verlag, Berlin.
- Kowalski, W., and W. Bahnfleth. 2003. Immune building technology and bioterrorism defense. *HPAC Engineering* 75(1):57-62. [pennstate.pure.elsevier.com/en/publications/immune-building-technology-and-bioterrorism-defense](http://pennstate.pure.elsevier.com/en/publications/immune-building-technology-and-bioterrorism-defense).
- Kujundzic, E., M. Hernandez, and S.L. Miller. 2007. Ultraviolet germicidal irradiation inactivation of airborne fungal spores and bacteria in upper-room air and HVAC in-duct configurations. *Journal of Environmental Engineering Science* 6:1-9.

- Lee, B., W. Bahnfleth, and K. Auer. 2009. Life-cycle cost simulation of in-duct ultraviolet germicidal irradiation systems. *Proceedings of Building Simulation 2009, 11th International Building Performance Simulation Association Conference and Exhibition*, July, Glasgow.
- Leuschner, W., and F. Salie. 2013. Characterizing ultraviolet germicidal irradiance luminaires. *Photochemistry and Photobiology* 89(4):811-815. dx.doi.org/10.1111/php.12064.
- Levetin, E., R. Shaughnessy, C. Rogers, and R. Scheir. 2001. Effectiveness of germicidal UV radiation for reducing fungal contamination within air-handling units. *Applied and Environmental Microbiology* 67(8):3712-3715.
- Luongo, J., and S. Miller. 2016. Ultraviolet germicidal coil cleaning: Decreased surface microbial loading and resuspension of cell clusters. *Building and Environment* 105:50-55.
- Luongo, J., J. Brownstein, and S. Miller. 2017. Ultraviolet germicidal coil cleaning: Impact on heat transfer effectiveness and static pressure drop. *Building and Environment* 112:159-165.
- Martin, S.B., C. Dunn, J.D. Freihaut, W.P. Bahnfleth, J. Lau, and A. Nedeljkovic-Davidovic. 2008. Ultraviolet germicidal irradiation current best practices. *ASHRAE Journal* (August):28-36.
- McDevitt, J.J., D.K. Milton, S.N. Rudnick, and M.W. First. 2008. Inactivation of poxviruses by upper-room UVC light in a simulated hospital room environment. *PLoS ONE* 3(9):e3186. journals.plos.org/plosone/article?id=10.1371/journal.pone.0003186.
- McLean, R.L. 1961. General discussion: The mechanism of spread of Asian influenza. Presented at the International Conference of Asian Influenza, Bethesda, MD. *American Review of Respiratory Diseases* 83(2 Part 2): 36-38.
- Menzies, D., J. Popa, J. Hanley, T. Rand, and D. Milton. 2003. Effect of ultraviolet germicidal lights installed in office ventilation systems on workers' health and well being: Double-blind multiple cross over trial. *Lancet* 363:1785-1792.
- Miller, R.V., W. Jeffrey, D. Mitchell and M. Elasi. 1999. Bacterial responses to ultraviolet light. *American Society for Microbiology (ASM) News* 65(8): 535-541.
- Miller, S.L., M. Fennelly, M. Kernandez, K. Fennelly, J. Martyny, J. Mache, E. Kujundzic, P. Xu, P. Fabian, J. Peccia, and C. Howard. 2002. Efficacy of ultraviolet irradiation in controlling the spread of tuberculosis. *Final Report*, Centers for Disease Control, Atlanta, and National Institute for Occupational Safety and Health, Washington, D.C.
- Montgomery, R., and R. Baker. 2006. Study verifies coil cleaning saves energy. *ASHRAE Journal* 48(11):34-36.
- Mphaphlele, M., A.S. Dharmadhikari, P.A. Jensen, S.N. Rudnick, T.H van Reenen, M.A. Pagano, W. Leuschner, T.A. Sears, S.P. Milonova, M. van der Walt, A.C. Stoltz, K. Weyer, and E.A. Nardell. 2015. Controlled trial of upper room ultraviolet air disinfection: A basis for new dosing guidelines. *American Journal of Respiratory and Critical Care Medicine* 192(4):477-484.
- Nagy, R., G. Mouromseff, and F.H. Rixton. 1954. Disinfecting air with sterilizing lamps. *Heating, Piping, and Air Conditioning* 26(April):82-87.
- Nardell, E., R. Vincent, and D.H. Sliney. 2013. Upper-room ultraviolet germicidal irradiation (UVGI) for air disinfection: A symposium in print. *Photochemistry and Photobiology* 89(4):764-769. dx.doi.org/10.1111/php.12098.
- Nazaroff, W. and C. Weschler. 2009. Air cleaning effectiveness for improving indoor air quality: Open-path and closed-path configurations. *Proceedings of Healthy Buildings 2009*, Syracuse, NY, Paper 376.
- NEMA. 2009. *Lamprecycle.org: Environmental responsibility starts here*. National Electrical Manufacturers Association. www.lamprecycle.org/.
- NIOSH. 1972. Criteria for a recommended standard: Occupational exposure to ultraviolet radiation. *Publication* 73-11009. National Institute for Occupational Safety and Health, Washington, D.C.
- NIOSH. 2009. Environmental control for tuberculosis: Basic upper-room ultraviolet germicidal irradiation guidelines for healthcare settings. *NIOSH Publication* 2009-105. www.cdc.gov/niosh/docs/2009-105/default.html.
- Philips. 1985. *Germicidal lamps and applications*. Philips Lighting Division, Roosendaal, the Netherlands.
- Philips. 2006. *Ultraviolet purification application information*. Philips Lighting B.V., Roosendaal, the Netherlands.
- Rastogi, V.K., L. Wallace, and L.S. Smith. 2007. Disinfection of *Acinetobacter baumannii*-contaminated surfaces relevant to medical treatment facilities with ultraviolet C light. *Military Medicine* 172(11):1166.
- Reed, N.G. 2010. The history of ultraviolet germicidal irradiation for air disinfection. *Public Health Reports* 125:15-27.
- Rentschler, H.C., and R. Nagy. 1940. Advantages of bactericidal ultraviolet radiation in air conditioning systems. *HPAC* 12:127-130.
- Riley, R.L. 1988. Ultraviolet air disinfection for control of respiratory contagion. In *Architectural design and indoor microbial pollution*, pp. 179-197. Oxford University Press, New York.
- Riley, R.L., and F. O'Grady. 1961. *Airborne infection—Transmission and Control*. Macmillan, New York.
- Riley, R.L., and S. Permutt. 1971. Room air disinfection by ultraviolet irradiation of upper air: Air mixing and germicidal effectiveness. *Archives of Environmental Health* 22(2):208-219.
- Riley, R.L., S. Permutt, and J.E. Kaufman. 1971. Convection, air mixing, and ultraviolet air disinfection in rooms. *Archives of Environmental Health* 22(2):200-207.
- Riley, R.L., M. Knight, and G. Middlebrook. 1976. Ultraviolet susceptibility of BCG and virulent tubercle bacilli. *American Review of Respiratory Disease* 113:413-418.
- Ritter, M.A., E.M. Olberding, and R.A. Malinzak. 2007. Ultraviolet lighting during orthopaedic surgery and the rate of infection. *The Journal of Bone & Joint Surgery* 89:1935-1940.
- RTI. 2005. *Test/QA plan for biological inactivation efficiency by HVAC in-duct ultraviolet light air cleaners*. Research Triangle Institute, Research Triangle Park, NC.
- Rudnick, S. 2007. Fundamental factors affecting upper-room germicidal irradiation—Part 2: Predicting effectiveness. *Journal of Occupational and Environmental Hygiene* 4(5):352-362.
- Rudnick, S.N., M.W. First, R.L. Vincent, and P.W. Brickner. 2009. In-place testing of in-duct ultraviolet germicidal irradiation. *HVAC&R Research (now Science and Technology for the Built Environment)* 15(3).
- Rudnick, S.N., M.W. First, T. Sears, R.L. Vincent, P.W. Brickner, P.Y. Ngai, J. Zhang, R.E. Levin, K. Chin, R.O. Rahn, S.L. Miller, and E.A. Nardell. 2012. Spatial distribution of fluence rate from upper-room ultraviolet germicidal irradiation: Experimental validation of a computer-aided design tool. *HVAC&R Research (now Science and Technology for the Built Environment)* 18(4):774-794.
- Rutala, W. 2009. Disinfection and sterilization: Successes and failures. Presented at APIC Convention, June, Ft. Lauderdale, FL.
- Setlow, J.K. 1966. The molecular basis of biological effects of ultraviolet radiation and photoreactivation. *Current Topics in Radiation Research* 2:195-248.
- Setlow, R.B. 1997. DNA damage and repair: A photobiological odyssey. *Photochemistry and Photobiology* 65S:119S-122S.
- Setlow, R.B., and J.K. Setlow. 1962. Evidence that ultraviolet-induced thymine dimers in DNA cause biological damage. *Proceedings of the National Academy of Sciences* 48(7):1250-1257.
- Sharp, D.G. 1939. The lethal action of short ultraviolet rays on several common pathogenic bacteria. *Journal of Bacteriology* 37(4):447-460.
- Sharp, D.G. 1940. The effects of ultraviolet light on bacteria suspended in air. *Journal of Bacteriology* 39(5):535-547.
- Shaughnessy, R., E. Levetin, and C. Rogers. 1998. The effects of UV-C on biological contamination of AHUs in a commercial office building: preliminary results. *Proceedings of IAQ and Energy '98*, pp. 229-236.
- Shechmeister, I.L. 1991. Sterilization by ultraviolet radiation. In *Disinfection, sterilization and preservation*, pp. 535-565. Lea and Febiger, Philadelphia.
- Sliney, D. 2013. Balancing the risk of eye irritation from UV-C with infection from bioaerosols. *Photochemistry and Photobiology* 89(4):770-776. dx.doi.org/10.1111/php.12093.
- UL. 2015. Heating and cooling equipment. *ANSI/UL Standard* 1995. Underwriters Laboratories, Northbrook, IL.
- U.S. HHS. 2009. National healthcare quality report. U.S. Department of Health and Human Services. Agency for Healthcare Research and Quality (AHRQ) *Publication* 10-0003.
- VanOsdell, D., and K. Foorde. 2002. Defining the effectiveness of UV lamps installed in circulating air ductwork. *Final Report*, Air-Conditioning and Refrigeration Technology Institute 21-CR Project 61040030.
- Vincent, R.L., T. Sears, P.W. Brickner, and E.A. Nardell. 2013. Computer-aided design (CAD) for applying upper room UVGI fixtures to control airborne disease transmission. *Proceedings of the CIE*, Paris, CIE x038: 2013.
- Wang, Y. 2017. *Field study of ultraviolet germicidal irradiation systems for cooling coils in a hot and humid climate—Energy and disinfection analysis*. Ph.D. dissertation. National University of Singapore.



- Wang, Y., C. Sekhar, W. Bahnfleth, K-W Cheong, and J. Firrantello. 2016a. Effectiveness of an ultraviolet germicidal irradiation system in enhancing cooling coil energy performance in a hot and humid climate. *Energy and Buildings* 130(2016):321-329. dx.doi.org/10.1016/j.enbuild.2016.08.063.
- Wang, Y., C. Sekhar, W. P. Bahnfleth, K-W Cheong, and J. Firrantello. 2016b. Effects of an ultraviolet coil irradiation system on the air-side heat transfer coefficient and low  $\Delta T$  syndrome in a hot and humid climate. *Science and Technology for the Built Environment* 23(4):582-593.
- Wells, W.F. 1955. *Airborne contagion and air hygiene; an ecological study of droplet infections*. Cambridge: Published for the Commonwealth Fund by Harvard University Press.
- Westinghouse. 1982. Westinghouse sterilamp germicidal ultraviolet tubes. Westinghouse *Engineering Notes* A-8968.
- Xu, P., J. Peccia, P. Fabian, J.W. Martyny, K.P. Fennelly, M. Hernandez, and S.L. Miller. 2003. Efficacy of ultraviolet germicidal irradiation of upper-room air in inactivating airborne bacterial spores and mycobacteria in full-scale studies. *Atmospheric Environment* 37(3):405-419.
- Xu, P., N. Fisher, and S.L. Miller. 2013. Using computational fluid dynamics modeling to evaluate the design of hospital ultraviolet germicidal irradiation systems for inactivating airborne Mycobacteria. *Photochemistry and Photobiology* 89(4):792-798. dx.doi.org/10.1111/php.12062.
- Zhang J., R. Levin, R. Angelo, R. Vincent, P. Brickner, P. Ngai, and E. Nardell. 2012. A radiometry protocol for UVGI fixtures using a moving-mirror type goniometer. *Journal of Occupational and Environmental Hygiene* 9(3):140-148.
- Zhu, S., J. Srebric, S.N. Rudnick, R.L. Vincent, and E.A. Nardell. 2013. Numerical investigation of upper-room UVGI disinfection efficacy in an environmental chamber with a ceiling fan. *Photochemistry and Photobiology* 89:782-791. dx.doi.org/10.1111/php.12039.
- Dumyahn, T. and M.W. First. 1999. Characterization of ultraviolet upper room air disinfection devices. *American Industrial Hygiene Association Journal* 60:219-227.
- EPA. 2006. Biological inactivation efficiency of HVAC in-duct ultraviolet light devices. EPA/600/S-11/002. U.S. Environmental Protection Agency, Washington, D.C.
- Linnes, J.C., S.N. Rudnick, G.M. Hunt, J.J. McDevitt, and E.A. Nardell. 2013. Eggcrate UV: A whole ceiling upper-room ultraviolet germicidal irradiation system for air disinfection in occupied rooms. *Indoor Air* 24(2):116-124.
- Luckiesh, M. 1946. *Applications of germicidal, erythematous and infrared energy*. D. Van Nostrand, New York.
- Masschelein, W.J. 2002. *Ultraviolet light in water and wastewater sanitation*. R.G. Rice, ed. Lewis Publishers, New York.
- Miller, S.L., J. Linnes, and J. Luongo. 2013. Ultraviolet germicidal irradiation: Future directions for air disinfection and building applications. *Photochemistry and Photobiology* 89(4):777-781. dx.doi.org/10.1111/php.12080.
- Nardell, E.A., S.J. Bucher, P.W. Brickner, C. Wang, R.L. Vincent, K. Becan-McBride, M.A. James, M. Michael, and J.D. Wright. 2008. Safety of upper-room ultraviolet germicidal air disinfection for room occupants: Results from the tuberculosis ultraviolet shelter study. *Public Health Report* 123(1):52-60.
- NEHC. 1992. *Ultraviolet radiation guide*. Navy Environmental Health Center, Bureau of Medicine and Surgery, Norfolk, VA.
- NEMA. 2004. Performance testing for lighting controls and switching devices with electronic fluorescent ballasts. *Standard* 410-2004. National Electrical Manufacturers Association, Rosslyn, VA.
- Philips Lighting. 1992. *Disinfection by UV-radiation*. Eindhoven, the Netherlands.
- Rahn, R.O. 2013. Fluence measurements employing iodide/iodate chemical actinometry as applied to upper-room germicidal radiation. *Photochemistry and Photobiology* 89(4):816-818. dx.doi.org/10.1111/php.12094.
- RLW Analytics. 2006. Improving indoor environment quality and energy performance of California K-12 schools, project 3: Effectiveness of UVC light for improving school performance. *Final Report*, California Energy Commission Contract 59903-300.
- Scheir, R. and F.B. Fencl. 1996. Using UVGI technology to enhance IAQ. *Heating, Piping and Air Conditioning* 68:109-124.
- Siegel, J., I. Walker, and M. Sherman. 2002. Dirty air conditioners: Energy implications of coil fouling. *Proceedings of the ACEEE Summer Study on Energy Efficiency in Buildings*, pp. 287-299.
- Sylvania. 1982. Germicidal and short-wave ultraviolet radiation. *Sylvania Engineering Bulletin* 0-342.
- Vincent, R., and P. Brickner. 2008. Safety and UV exposure. *IAQ Applications* 9(3).
- WHO. 2006. Solar ultraviolet radiation: Global burden of disease from solar ultraviolet radiation. *Environmental Burden of Disease Series* 13. World Health Organization, Geneva. www.who.int/quantifying\_ehimpacts/publications/ebd13/en/.
- Witham, D. 2005. Ultraviolet—A superior tool for HVAC maintenance. *IUVA Congress, Tokyo*.

## BIBLIOGRAPHY

- Abshire, R.L., and H. Dunton. 1981. Resistance of selected strains of *Pseudomonas aeruginosa* to low-intensity ultraviolet radiation. *Applied Environmental Microbiology* 41(6):1419-1423.
- ASHRAE. 2017. Thermal environmental conditions for human occupancy. *ANSI/ASHRAE Standard* 55-2017.
- Bahnfleth, W.P., and W.J. Kowalski. 2004. Clearing the air on UVGI systems. *RSES Journal*, pp. 22-24.
- Bernstein, J.A., R.C. Bobbitt, L. Levin, R. Floyd, M.S. Crandall, R.A. Shalwitz, A. Seth, and M. Glazman. 2006. Health effects of ultraviolet irradiation in asthmatic children's homes. *Journal of Asthma* 43(4):255-262.
- Blatt, M.S., T. Okura, and B. Meister. 2006. Ultraviolet light for coil cleaning in schools. *Engineered Systems* (March):50-61.
- Bolton, J.R. 2001. *Ultraviolet applications handbook*. Photosciences, Ontario.
- Department of General Services. 2001. *Working with ultraviolet germicidal irradiation (UVGI) lighting systems: Code of safe practice*. County of Sacramento, CA.
- DIN. 1979. Optical radiation physics and illumination engineering. *Standard* 5031. German Institute for Standardization, Berlin.

