

ASHRAE Position Document on Infectious Aerosols

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The ASHRAE Position Document on Infectious Aerosols was developed by the Society's Environmental Health Position Document Committee formed on April 24, 2017, with Erica Stewart as its chair.

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Note: ASHRAE's Technology Council and the cognizant committee recommend revision, reaffirmation, or withdrawal every 30 months.

Note: ASHRAE position documents are approved by the Board of Directors and express the views of the Society on a specific issue. The purpose of these documents is to provide objective, authoritative background information to persons interested in issues within ASHRAE's expertise, particularly in areas where such information will be helpful in drafting sound public policy. A related purpose is also to serve as an educational tool clarifying ASHRAE's position for its members and professionals, in general, advancing the arts and sciences of HVAC&R.

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ABSTRACT

The pathogens that cause infectious diseases are spread from a primary host to secondary hosts via several different routes. Some diseases are known to spread by infectious aerosols; for other diseases, the route of transmission is uncertain. The risk of pathogen spread, and therefore the number of people exposed, can be affected both positively and negatively by the airflow patterns in a space and by heating, ventilating, and air-conditioning (HVAC) and local exhaust ventilation (LEV) systems. ASHRAE is the global leader and foremost source of technical and educational information on the design, installation, operation, and maintenance of these systems. Although the principles discussed in this position document apply primarily to buildings, they may also be applicable to other occupancies, such as planes, trains, and automobiles.

ASHRAE will continue to support research that advances the knowledge base of indoor airmanagement strategies aimed to reduce occupant exposure to infectious aerosols. Chief among these ventilation-related strategies are dilution, airflow patterns, pressurization, temperature and humidity distribution and control, filtration, and other strategies such as ultraviolet germicidal irradiation (UVGI). While the exact level of ventilation effectiveness varies with local conditions and the pathogens involved, ASHRAE believes that these techniques, when properly applied, can reduce the risk of transmission of infectious diseases through aerosols.

To better specify the levels of certainty behind ASHRAE's policy positions stated herein, we have chosen to adopt the Agency for Healthcare Research and Quality (AHRQ) rubric for expressing the scientific certainty behind our recommendations (Burns et al. 2011). These levels of certainty, as adapted for this position document, are as follows:

Evidence Level	Description
Α	Strongly recommend; good evidence
В	Recommend; at least fair evidence
С	No recommendation for or against; balance of benefits and harms too close to justify a recommendation
D	Recommend against; fair evidence is ineffective or the harm outweighs the benefit
E	Evidence is insufficient to recommend for or against routinely; evidence is lacking or of poor quality; benefits and harms cannot be determined

ASHRAE's position is that facilities of all types should follow, as a minimum, the latest published standards and guidelines and good engineering practice. ANSI/ASHRAE Standards 62.1 and 62.2 (ASHRAE 2019a, 2019b) include requirements for outdoor air ventilation in most residential and nonresidential spaces, and ANSI/ASHRAE/ASHE Standard 170 (ASHRAE 2017a) covers both outdoor and total air ventilation in healthcare facilities. Based on risk assessments or owner project requirements, designers of new and existing facilities could go beyond the minimum requirements of these standards, using techniques covered in various ASHRAE publications, including the ASHRAE Handbook volumes, Research Project final reports, papers and articles, and design guides, to be even better prepared to control the dissemination of infectious aerosols.

EXECUTIVE SUMMARY

With infectious diseases transmitted through aerosols, HVAC systems can have a major effect on the transmission from the primary host to secondary hosts. Decreasing exposure of secondary hosts is an important step in curtailing the spread of infectious diseases.

Designers of mechanical systems should be aware that ventilation is not capable of addressing all aspects of infection control. HVAC systems, however, do impact the distribution and bio-burden of infectious aerosols. Small aerosols may persist in the breathing zone, available for inhalation directly into the upper and lower respiratory tracts or for settling onto surfaces, where they can be indirectly transmitted by resuspension or fomite contact.

Infectious aerosols can pose an exposure risk, regardless of whether a disease is classically defined as an "airborne infectious disease." This position document covers strategies through which HVAC systems modulate aerosol³ distribution and can therefore increase or decrease exposure to infectious droplets,⁴ droplet nuclei,⁵ surfaces, and intermediary fomites⁶ in a variety of environments.

This position document provides recommendations on the following:

- The design, installation, and operation of heating, ventilating, and air-conditioning (HVAC) systems, including air-cleaning, and local exhaust ventilation (LEV) systems, to decrease the risk of infection transmission.
- Non-HVAC control strategies to decrease disease risk.
- Strategies to support facilities management for both everyday operation and emergencies.

Infectious diseases can be controlled by interrupting the transmission routes used by a pathogen. HVAC professionals play an important role in protecting building occupants by interrupting the indoor dissemination of infectious aerosols with HVAC and LEV systems.

COVID-19 Statements

Separate from the approval of this position document, ASHRAE's Executive Committee and Epidemic Task Force approved the following statements specific to the ongoing response to the COVID-19 pandemic. The two statements are appended here due to the unique relationship between the statements and the protective design strategies discussed in this position document:

Statement on airborne transmission of SARS-CoV-2: Transmission of SARS-CoV-2 through the air is sufficiently likely that airborne exposure to the virus should be controlled. Changes to building operations, including the operation of heating, ventilating, and air-conditioning systems, can reduce airborne exposures.

Statement on operation of heating, ventilating, and air-conditioning systems to reduce SARS-CoV-2 transmission: Ventilation and filtration provided by heating, ventilating, and air-conditioning systems can reduce the airborne concentration of SARS-CoV-2 and thus

¹ Different HVAC systems are described in ASHRAE Handbook—HVAC Systems and Equipment (ASHRAE 2020).

² An object (such as a dish or a doorknob) that may be contaminated with infectious organisms and serve in their transmission.

An aerosol is a system of liquid or solid particles uniformly distributed in a finely divided state through a gas, usually air. They are small and buoyant enough to behave much like a gas.

⁴ In this document, *droplets* are understood to be large enough to fall to a surface in 3–7 ft (1–2 m) and thus not become aerosols.

Droplet nuclei are formed from droplets that become less massive by evaporation and thus may become aerosols.

⁶ Fomite transmission is a form of indirect contact that occurs through touching a contaminated inanimate object such as a doorknob, bed rail, television remote, or bathroom surface.

the risk of transmission through the air. Unconditioned spaces can cause thermal stress to people that may be directly life threatening and that may also lower resistance to infection. In general, disabling of heating, ventilating, and air-conditioning systems is not a recommended measure to reduce the transmission of the virus.

1. THE ISSUE

The potential for airborne dissemination of infectious pathogens is widely recognized, although there remains uncertainty about the relative importance of the various disease transmission routes, such as airborne, droplet, direct or indirect contact, and multimodal (a combination of mechanisms). Transmission of disease varies by pathogen infectivity, reservoirs, routes, and secondary host susceptibility (Roy and Milton 2004; Shaman and Kohn 2009; Li 2011). The variable most relevant for HVAC design and control is disrupting the transmission pathways of infectious aerosols.

Infection control professionals describe the chain of infection as a process in which a pathogen (a microbe that causes disease) is carried in an initial host or reservoir, gains access to a route of ongoing transmission, and with sufficient virulence finds a secondary susceptible host. Ventilation, filtration, and air distribution systems and disinfection technologies have the potential to limit airborne pathogen transmission through the air and thus break the chain of infection.

Building science professionals must recognize the importance of facility operations and ventilation systems in interrupting disease transmission. Non-HVAC measures for breaking the chain of infection, such as effective surface cleaning, contact and isolation precautions mandated by employee and student policies, and vaccination regimens, are effective strategies that are beyond the scope of this document. Dilution and extraction ventilation, pressurization, airflow distribution and optimization, mechanical filtration, ultraviolet germicidal irradiation (UVGI), and humidity control are effective strategies for reducing the risk of dissemination of infectious aerosols in buildings and transportation environments.

Although this position document is primarily applicable to viral and bacterial diseases that can use the airborne route for transmission from person to person, the principles of containment may also apply to infection from building reservoirs such as water systems with *Legionella spp.* and organic matter containing spores from mold (to the extent that the microorganisms are spread by the air). The first step in control of such diseases is to eliminate the source before it becomes airborne.

2. BACKGROUND

ASHRAE provides guidance and develop standards intended to mitigate the risk of infectious disease transmission in the built environment. Such documents provide engineering strategies for reducing the risk of disease transmission and therefore could be employed in a variety of other spaces, such as planes, trains, and automobiles.

This position document covers the dissemination of infectious aerosols and indirect transmission by resuspension but not direct-contact routes of transmission. *Direct contact* generally refers to bodily contact such as touching, kissing, sexual contact, contact with oral secretions or skin lesions and routes such as blood transfusions or intravenous injections.

2.1 Airborne Dissemination

Pathogen dissemination through the air occurs through droplets and aerosols typically generated by coughing, sneezing, shouting, breathing, toilet flushing, some medical procedures, singing, and talking (Bischoff et al. 2013; Yan et al. 2018). The majority of larger emitted droplets are drawn by gravity to land on surfaces within about 3–7 ft (1–2 m) from the source (see Figure 1). General dilution ventilation and pressure differentials do not significantly influ-

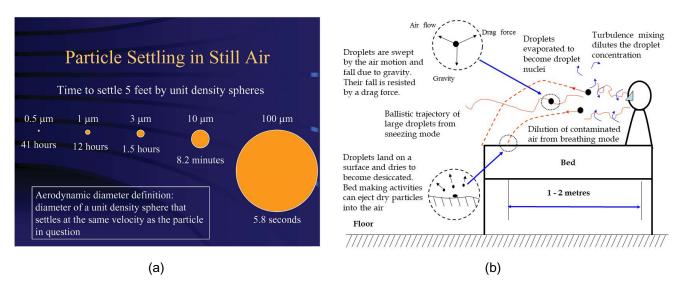


Figure 1 (a) Comparative settling times by particle diameter for particles settling in still air (Baron n.d.) and (b) theoretical aerobiology of transmission of droplets and small airborne particles produced by an infected patient with an acute infection (courtesy Yuguo Li).

ence short-range transmission. Conversely, dissemination of smaller infectious aerosols, including droplet nuclei resulting from desiccation, can be affected by airflow patterns in a space in general and airflow patterns surrounding the source in particular. Of special interest are small aerosols (<10 μ m), which can stay airborne and infectious for extended periods (several minutes, hours, or days) and thus can travel longer distances and infect secondary hosts who had no contact with the primary host.

Many diseases are known to have high transmission rates via larger droplets when susceptible individuals are within close proximity, about 3–7 ft (1–2 m) (Nicas 2009; Li 2011). Depending on environmental factors, these large (100 μ m diameter) droplets may shrink by evaporation before they settle, thus becoming an aerosol (approximately <10 μ m). The term *droplet nuclei* has been used to describe such desiccation of droplets into aerosols (Siegel et al. 2007). While ventilation systems cannot interrupt the rapid settling of large droplets, they can influence the transmission of droplet nuclei infectious aerosols. Directional airflow can create clean-to-dirty flow patterns and move infectious aerosols to be captured or exhausted.

3. PRACTICAL IMPLICATIONS FOR BUILDING OWNERS, OPERATORS, AND ENGINEERS

Even the most robust HVAC system cannot control all airflows and completely prevent dissemination of an infectious aerosol or disease transmission by droplets or aerosols. An HVAC system's impact will depend on source location, strength of the source, distribution of the released aerosol, droplet size, air distribution, temperature, relative humidity, and filtration. Furthermore, there are multiple modes and circumstances under which disease transmission occurs. Thus, strategies for prevention and risk mitigation require collaboration among designers, owners, operators, industrial hygienists, and infection prevention specialists.

3.1 Varying Approaches for Facility Type

Healthcare facilities have criteria for ventilation design to mitigate airborne transmission of infectious diseases (ASHRAE 2013, 2017a, 2019a; FGI 2010); however, infections are also transmitted in ordinary occupancies in the community and not only in industrial or healthcare occupancies. ASHRAE provides general ventilation and air quality requirements in Standards 62.1, 62.2, and 170 (ASHRAE 2019a, 2019b, 2017a); ASHRAE does not provide specific requirements for infectious disease control in homes, schools, prisons, shelters, transportation, or other public facilities.

In healthcare facilities, most infection control interventions are geared at reducing direct or indirect contact transmission of pathogens. These interventions for limiting airborne transmission (Aliabadi et al. 2011) emphasize personnel education and surveillance of behaviors such as hand hygiene and compliance with checklist protocols and have largely been restricted to a relatively small list of diseases from pathogens that spread only through the air. Now that microbiologists understand that many pathogens can travel through both contact and airborne routes, the role of indoor air management has become critical to successful prevention efforts. In view of the broader understanding of flexible pathogen transmission modes, healthcare facilities now use multiple modalities simultaneously (measures that are referred to as infection control bundles) (Apisarnthanarak et al. 2009, 2010a, 2010b; Cheng et al. 2010). For example, in the cases of two diseases that clearly utilize airborne transmission, tuberculosis and measles, bundling includes administrative regulations, environmental controls, and personal protective equipment protocols in healthcare settings. This more comprehensive approach is needed to control pathogens, which can use both contact and airborne transmission pathways. Similar strategies may be appropriate for non-healthcare spaces, such as public transit and airplanes, schools, shelters, and prisons, that may also be subject to close contact of occupants.

Many buildings are fully or partially naturally ventilated. They may use operable windows and rely on intentional and unintentional openings in the building envelope. These strategies create different risks and benefits. Obviously, the airflow in these buildings is variable and unpredictable, as are the resulting air distribution patterns, so the ability to actively manage risk in such buildings is much reduced. However, naturally ventilated buildings can go beyond random opening of windows and be engineered intentionally to achieve ventilation strategies and thereby reduce risk from infectious aerosols. Generally speaking, designs that achieve higher ventilation rates will reduce risk. However, such buildings will be more affected by local outdoor air quality, including the level of allergens and pollutants within the outdoor air, varying temperature and humidity conditions, and flying insects. The World Health Organization has published guidelines for naturally ventilated buildings that should be consulted in such projects (Atkinson et al. 2009).

3.2 Ventilation and Air-Cleaning Strategies

The design and operation of HVAC systems can affect infectious aerosol transport, but they are only one part of an infection control bundle. The following HVAC strategies have the potential to reduce the risks of infectious aerosol dissemination: air distribution patterns, differential room pressurization, personalized ventilation, source capture ventilation, filtration (central or local), and controlling temperature and relative humidity. While UVGI is well researched and validated, many new technologies are not (ASHRAE 2018). (Evidence Level B)

Ventilation with effective airflow patterns (Pantelic and Tham 2013) is a primary infectious disease control strategy through dilution of room air around a source and removal of infectious

agents (CDC 2005). However, it remains unclear by how much infectious particle loads must be reduced to achieve a measurable reduction in disease transmissions (infectious doses vary widely among different pathogens) and whether these reductions warrant the associated costs (Pantelic and Tham 2011; Pantelic and Tham 2012). (Evidence Level B)

Room pressure differentials and directional airflow are important for controlling airflow between zones in a building (CDC 2005; Siegel et al. 2007) (Evidence Level B). Some designs for airborne infection isolation rooms (AIIRs) incorporate supplemental dilution or exhaust/capture ventilation (CDC 2005). Interestingly, criteria for AIIRs differ substantially between regions and countries in several ways, including air supply into anterooms, exhaust from space, and required amounts of ventilation air (Fusco et al. 2012; Subhash et al. 2013). A recent ASHRAE Research Project found convincing evidence that a properly configured and operated anteroom is an effective means to maintain pressure differentials and create containment in hospital rooms (Siegel et al. 2007; Mousavi et al. 2019). Where a significant risk of transmission of aerosols has been identified by infection control risk assessments, design of AIIRs should include anterooms. (Evidence Level A)

The use of highly efficient particle filtration in centralized HVAC systems reduces the airborne load of infectious particles (Azimi and Stephens 2013). This strategy reduces the transport of infectious agents from one area to another when these areas share the same central HVAC system through supply of recirculated air. When appropriately selected and deployed, single-space high-efficiency filtration units (either ceiling mounted or portable) can be highly effective in reducing/lowering concentrations of infectious aerosols in a single space. They also achieve directional airflow source control that provides exposure protection at the patient bedside (Miller-Leiden et al. 1996; Mead and Johnson 2004; Kujundzic et al. 2006; Mead et al. 2012; Dungi et al. 2015). Filtration will not eliminate all risk of transmission of airborne particulates because many other factors besides infectious aerosol concentration contribute to disease transmission. (Evidence Level A)

The entire ultraviolet (UV) spectrum can kill or inactivate microorganisms, but UV-C energy (in the wavelengths from 200 to 280 nm) provides the most germicidal effect, with 265 nm being the optimum wavelength. The majority of modern UVGI lamps create UV-C energy at a near-optimum 254 nm wavelength. UVGI inactivates microorganisms by damaging the structure of nucleic acids and proteins with the effectiveness dependent upon the UV dose and the susceptibility of the microorganism. The safety of UV-C is well known. It does not penetrate deeply into human tissue, but it can penetrate the very outer surfaces of the eyes and skin, with the eyes being most susceptible to damage. Therefore, shielding is needed to prevent direct exposure to the eyes. While ASHRAE Position Document on Filtration and Air Cleaning (2018) does not make a recommendation for or against the use of UV energy in air systems for minimizing the risks from infectious aerosols, Centers for Disease Control and Prevention (CDC) has approved UVGI as an adjunct to filtration for reduction of tuberculosis risk and has published a guideline on its application (CDC 2005, 2009). (Evidence Level A)

Personalized ventilation systems that provide local exhaust source control and/or supply 100% outdoor, highly filtered, or UV-disinfected air directly to the occupant's breathing zone (Cermak et al. 2006; Bolashikov et al., 2009; Pantelic et al. 2009, 2015; Licina et al. 2015a, 2015b) may offer protection against exposure to contaminated air. Personalized ventilation may be effective against aerosols that travel both long distances as well as short ranges (Li 2011).

In addition to UVGI, optical radiation in longer wavelengths as high as 405 nm is an emerging disinfection technology that may also have useful germicidal effectiveness.

Personalized ventilation systems, when coupled with localized or personalized exhaust devices, further enhance the overall ability to mitigate exposure in breathing zones, as seen from both experimental and computational fluid dynamics (CFD) studies in healthcare settings (Yang et al. 2013, 2014, 2015a, 2015b; Bolashikov et al. 2015; Bivolarova et al. 2016). However, there are no known epidemiological studies that demonstrate a reduction in infectious disease transmission. (Evidence Level B)

Advanced techniques such as computational fluid dynamics (CFD) analysis, if performed properly with adequate expertise, can predict airflow patterns and probable flow paths of airborne contaminants in a space. Such analyses can be employed as a guiding tool during the early stages of a design cycle (Khankari 2016, 2018a, 2018b, 2018c).

3.3 Temperature and Humidity

HVAC systems are typically designed to control temperature and humidity, which can in turn influence transmissibility of infectious agents. Although HVAC systems can be designed to control relative humidity (RH), there are practical challenges and potential negative effects of maintaining certain RH set points in all climate zones. However, while the weight of evidence at this time (Derby et al. 2016), including recent evidence using metagenomic analysis (Taylor and Tasi 2018), suggests that controlling RH reduces transmission of certain airborne infectious organisms, including some strains of influenza, this position document encourages designers to give careful consideration to temperature and RH.

In addition, immunobiologists have correlated mid-range humidity levels with improved mammalian immunity against respiratory infections (Taylor and Tasi 2018). Mousavi et al. (2019) report that the scientific literature generally reflects the most unfavorable survival for microorganisms when the RH is between 40% and 60% (Evidence Level B). Introduction of water vapor to the indoor environment to achieve the mid-range humidity levels associated with decreased infections requires proper selection, operation, and maintenance of humidification equipment. Cold winter climates require proper building insulation to prevent thermal bridges that can lead to condensation and mold growth (ASHRAE 2009). Other recent studies (Taylor and Tasi 2018) identified RH as a significant driver of patient infections. These studies showed that RH below 40% is associated with three factors that increase infections. First, as discussed previously, infectious aerosols emitted from a primary host shrink rapidly to become droplet nuclei, and these dormant yet infectious pathogens remain suspended in the air and are capable of traveling great distances. When they encounter a hydrated secondary host, they rehydrate and are able to propagate the infection. Second, many viruses and bacteria are anhydrous resistant (Goffau et al. 2009; Stone et al. 2016) and actually have increased viability in low-RH conditions. And finally, immunobiologists have now clarified the mechanisms through which ambient RH below 40% impairs mucus membrane barriers and other steps in immune system protection (Kudo et al. 2019). (Evidence Level B)

This position document does not make a definitive recommendation on indoor temperature and humidity set points for the purpose of controlling infectious aerosol transmission. Practitioners may use the information herein to make building design and operation decisions on a case-by-case basis.

3.4 Emerging Pathogens and Emergency Preparedness

Disease outbreaks (i.e., epidemics and pandemics) are increasing in frequency and reach. Pandemics of the past have had devastating effects on affected populations. Novel microor-

ganisms that can be disseminated by infectious aerosols necessitate good design, construction, commissioning, maintenance, advanced planning, and emergency drills to facilitate fast action to mitigate exposure. In many countries, common strategies include naturally ventilated buildings and isolation. Control banding is a risk management strategy that should be considered for applying the hierarchy of controls to emerging pathogens, based on the likelihood and duration of exposure and the infectivity and virulence of the pathogen (Sietsema 2019) (Evidence Level B). Biological agents that may be used in terrorist attacks are addressed elsewhere (USDHHS 2002, 2003).

4. CONCLUSIONS AND RECOMMENDATIONS

Infectious aerosols can be disseminated through buildings by pathways that include air distribution systems and interzone airflows. Various strategies have been found to be effective at controlling transmission, including optimized airflow patterns, directional airflow, zone pressurization, dilution ventilation, in-room air-cleaning systems, general exhaust ventilation, personalized ventilation, local exhaust ventilation at the source, central system filtration, UVGI, and controlling indoor temperature and relative humidity. Design engineers can make an essential contribution to reducing infectious aerosol transmission through the application of these strategies. Research on the role of airborne dissemination and resuspension from surfaces in pathogen transmission is rapidly evolving. Managing indoor air to control distribution of infectious aerosols is an effective intervention which adds another strategy to medical treatments and behavioral interventions in disease prevention.

4.1 ASHRAE's Positions

- HVAC design teams for facilities of all types should follow, as a minimum, the latest published standards and guidelines and good engineering practice. Based on risk assessments or owner project requirements, designers of new and existing facilities could go beyond the minimum requirements of these standards, using techniques covered in various ASHRAE publications, including the ASHRAE Handbook volumes, Research Project final reports, papers and articles, and design guides, to be even better prepared to control the dissemination of infectious aerosols.
- Mitigation of infectious aerosol dissemination should be a consideration in the design of all facilities, and in those identified as high-risk facilities the appropriate mitigation design should be incorporated.
- The design and construction team, including HVAC designers, should engage in an integrated design process in order to incorporate the appropriate infection control bundle in the early stages of design.
- Based on risk assessments, buildings and transportation vehicles should consider designs that promote cleaner airflow patterns for providing effective flow paths for airborne particulates to exit spaces to less clean zones and use appropriate air-cleaning systems. (Evidence Level A)
- Where a significant risk of transmission of aerosols has been identified by infection control risk assessments, design of AIIRs should include anterooms. (Evidence Level A)

- Based on risk assessments, the use of specific HVAC strategies supported by the evidence-based literature should be considered, including the following:
 - Enhanced filtration (higher minimum efficiency reporting value [MERV] filters over code minimums in occupant-dense and/or higher-risk spaces) (Evidence Level A)
 - Upper-room UVGI (with possible in-room fans) as a supplement to supply airflow (Evidence Level A)
 - Local exhaust ventilation for source control (Evidence Level A)
 - Personalized ventilation systems for certain high-risk tasks (Evidence Level B)
 - Portable, free-standing high-efficiency particulate air (HEPA) filters (Evidence Level B)
 - Temperature and humidity control (Evidence Level B)
- Healthcare buildings⁸ should consider design and operation to do the following:
 - Capture expiratory aerosols with headwall exhaust, tent or snorkel with exhaust, floor-to-ceiling partitions with door supply and patient exhaust, local air HEPA-grade filtration.
 - Exhaust toilets and bed pans (a must).
 - Maintain temperature and humidity as applicable to the infectious aerosol of concern.
 - Deliver clean air to caregivers.
 - Maintain negatively pressurized intensive care units (ICUs) where infectious aerosols may be present.
 - Maintain rooms with infectious aerosol concerns at negative pressure.
 - Provide 100% exhaust of patient rooms.
 - Use UVGI.
 - Increase the outdoor air change rate (e.g., increase patient rooms from 2 to 6 ach).
 - Establish HVAC contributions to a patient room turnover plan before reoccupancy.
- Non-healthcare buildings should have a plan for an emergency response. The following modifications to building HVAC system operation should be considered:
 - Increase outdoor air ventilation (disable demand-controlled ventilation and open outdoor air dampers to 100% as indoor and outdoor conditions permit).
 - Improve central air and other HVAC filtration to MERV-13 (ASHRAE 2017b) or the highest level achievable.
 - Keep systems running longer hours (24/7 if possible).
 - Add portable room air cleaners with HEPA or high-MERV filters with due consideration to the clean air delivery rate (AHAM 2015).
 - Add duct- or air-handling-unit-mounted, upper room, and/or portable UVGI devices in connection to in-room fans in high-density spaces such as waiting rooms, prisons, and shelters.
 - Maintain temperature and humidity as applicable to the infectious aerosol of concern.
 - Bypass energy recovery ventilation systems that leak potentially contaminated exhaust air back into the outdoor air supply.
- Design and build inherent capabilities to respond to emerging threats and plan and practice for them. (Evidence Level B)

⁸ It is assumed that healthcare facilities already have emergency response plans.

4.2 ASHRAE's Commitments

- Address research gaps with future research projects, including those on the following topics:
 - Investigating and developing source generation variables for use in an updated ventilation rate procedure
 - Understanding the impacts of air change rates in operating rooms on patient outcomes
 - Determining the effectiveness of location of supply, return, and exhaust registers in patient rooms
 - Conducting controlled interventional studies to quantify the relative airborne infection control performance and cost-effectiveness of specific engineering strategies, individually and in combination, in field applications of high-risk occupancies
 - Evaluating and comparing options to create surge airborne isolation space and temporary negative pressure isolation space and the impacts on overall building operation
 - Understanding the appropriate application of humidity and temperature control strategies across climate zones on infectious aerosol transmission
 - Investigating how control banding techniques can be applied to manage the risk of infectious aerosol dissemination
- Partner with infection prevention, infectious disease, and occupational health experts and building owners to evaluate emerging control strategies and provide evidence-based recommendations.
- Educate stakeholders and disseminate best practices.
- Create a database to track and share knowledge on effective, protective engineering design strategies.
- Update standards and guidelines to reflect protective evidence-based strategies.

5. REFERENCES

- AHAM. 2015. ANSI/AHAM AC-1-2015, *Method For Measuring Performance Of Portable Household Electric Room Air Cleaners*. Washington, DC: Association of Home Appliance Manufacturers.
- Aliabadi, A.A., S.N. Rogak, K.H. Bartlett, and S.I. Green. 2011. Preventing airborne disease transmission: Review of methods for ventilation design in health care facilities. *Advances in Preventive Medicine*. Article ID 12406. DOI: 10.4061/2011/124064.
- Apisarnthanarak, A., P. Apisarnthanarak, B. Cheevakumjorn, and L.M. Mundy. 2009. Intervention with an infection control bundle to reduce transmission of influenza-like illnesses in a Thai preschool. *Infection Control and Hospital Epidemiology* 30(9):817–22. DOI: 10.1086/599773.
- Apisarnthanarak, A., P. Apisarnthanarak, B. Cheevakumjorn, and L.M. Mundy. 2010a. Implementation of an infection control bundle in a school to reduce transmission of influenza-like illness during the novel influenza A 2009 H1N1 pandemic. *Infection Control and Hospital Epidemiology* 31(3):310–11. DOI: 10.1086/651063.
- Apisarnthanarak, A., T.M. Uyeki, P. Puthavathana, R. Kitphati, and L.M. Mundy. 2010b. Reduction of seasonal influenza transmission among healthcare workers in an intensive care unit: A 4-year intervention study in Thailand. *Infection Control and Hospital Epidemiology* 31(10):996–1003. DOI: 10.1086/656565.
- ASHRAE. 2009. Indoor Air Quality Guide: Best Practices for Design, Construction and Commissioning. Atlanta: ASHRAE.

- ASHRAE. 2013. HVAC Design Manual for Hospitals and Clinics, 2d ed. Atlanta: ASHRAE.
- ASHRAE. 2017a. ANSI/ASHRAE/ASHE Standard 170-2017, Ventilation of Health Care Facilities. Atlanta: ASHRAE.
- ASHRAE. 2017b. ANSI/ASHRAE Standard 52.2-2017, Method of Testing General Ventilation Air- Cleaning Devices for Removal Efficiency by Particle Size. Atlanta: ASHRAE.
- ASHRAE. 2018. ASHRAE Position Document on Filtration and Air-Cleaning. Atlanta: ASHRAE. www.ashrae.org/file%20library/about/position%20documents/filtration-and-air-cleaning-pd.pdf.
- ASHRAE. 2019a. ANSI/ASHRAE Standard 62.1-2019, Ventilation for Acceptable Indoor Air Quality. Atlanta: ASHRAE.
- ASHRAE. 2019b. ANSI/ASHRAE Standard 62.2-2019, Ventilation and Acceptable Indoor Air Quality in Low-Rise Residential Buildings. Atlanta: ASHRAE.
- ASHRAE. 2020. ASHRAE Handbook—HVAC Systems and Equipment. Atlanta: ASHRAE.
- Atkinson J., Y. Chartier, C.L. Pessoa-Silva, P. Jensen, and W.H. Seto. 2009. *Natural Ventilation for Infection Control in Health-Care Settings*. Geneva: World Health Organization. www.who.int/water_sanitation_health/publications/natural_ventilation/en.
- Azimi, P., and B. Stephens. 2013. HVAC filtration for controlling infectious airborne disease transmission in indoor environments: Predicting risk reductions and operational costs. *Building and Environment* 70:150–60.
- Baron, P. n.d. Generation and Behavior of Airborne Particles (Aerosols). Presentation published at CDC/NIOSH Topic Page: Aerosols, National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention, Public Health Service, U.S. Department of Health and Human Services, Cincinnati, OH. www.cdc.gov/niosh/topics/aerosols/pdfs/Aerosol_101.pdf.
- Bischoff, W.E., K. Swett, I. Leng, and T.R. Peters. 2013. Exposure to influenza virus aerosols during routine patient care. *Journal of Infectious Diseases* 207(7):1037–46. DOI: 10.1093/infdis/jis773.
- Bivolarova, M.P., A.K. Melikov, C. Mizutani, K. Kajiwara, and Z.D. Bolashikov. 2016. Bed-integrated local exhaust ventilation system combined with local air cleaning for improved IAQ in hospital patient rooms. *Building and Environment* 100:10–18.
- Bolashikov, Z.D., A.K. Melikov, and M. Krenek. 2009. Improved performance of personalized ventilation by control of the convection flow around occupant body. *ASHRAE Transactions* 115(2):421–31.
- Bolashikov, Z.D., M. Barova, and A.K. Melikov. 2015. Wearable personal exhaust ventilation: Improved indoor air quality and reduced exposure to air exhaled from a sick doctor. Science and Technology for the Built Environment 21(8):1117–25.
- Burns, P.B., R.J. Rohrich, and K.C. Chung. 2011. Levels of evidence and their role in evidence-based medicine. *Plast Reconstr Surg* 128(1):305–10.
- CDC. 2005. Guidelines for Preventing the Transmission of Mycobacterium tuberculosis in Health-Care Settings. Morbidity and Mortality Weekly Report (MMWR) 54(RR17):1–140. Atlanta: Centers for Disease Control and Prevention. www.cdc.gov/mmwr/preview/mmwrhtml/rr5417a1.htm.
- CDC. 2009. Environmental Control for Tuberculosis: Basic Upper-Room Ultraviolet Germicidal Irradiation Guidelines for Healthcare Settings. Atlanta: Centers for Disease Control and Prevention. www.cdc.gov/niosh/docs/2009-105/pdfs/2009-105.pdf.

- Cermak, R., A.K. Melikov, L. Forejt, and O. Kovar. 2006. Performance of personalized ventilation in conjunction with mixing and displacement ventilation. *HVAC&R Research* 12(2):295–311.
- Cheng, V.C., J.W. Tai, L.M. Wong, J.F. Chan, I.W. Li, K.K. To, I.F. Hung, K.H. Chan, P.L. Ho, and K.Y. Yuen. 2010. Prevention of nosocomial transmission of swine-origin pandemic influenza virus A/H1N1 by infection control bundle. *Journal of Hospital Infection* 74(3):271–77. DOI: 10.1016/j.jhin.2009.09.009.
- Derby, M., S. Eckels, G. Hwang, B. Jones, R. Maghirang, and D. Shulan. 2016. *Update the Scientific Evidence for Specifying Lower Limit Relative Humidity Levels for Comfort, Health and IEQ in Occupied Spaces*. ASHRAE Research Report 1630. Atlanta: ASHRAE.
- Dungi, S.R., U. Ghia, K.R. Mead, and M. Gressel. 2015. Effectiveness of a local ventilation/fil-tration intervention for health-care worker exposure reduction to airborne infection in a hospital room. Paper no. CH-15-C017. 2015 ASHRAE Winter Conference—Papers [download].
- FGI. 2010. Guidelines for Design and Construction of Health Care Facilities. St Louis, MO: Facility Guidelines Institute.
- Fusco, F.M., S. Schilling, G. De Iaco, H.R. Brodt, P. Brouqui, H.C. Maltezou, B. Bannister, R. Gottschalk, G. Thomson, V. Puro, and G. Ippolito. 2012. Infection control management of patients with suspected highly infectious diseases in emergency departments: Data from a survey in 41 facilities in 14 European countries. *BMC Infectious Diseases* January 28:12–27.
- de Goffau, M.C., X. Yang, J.M. van Dijl, and H.J. Harmsen. 2009. Bacterial pleomorphism and competition in a relative humidity gradient. *Environmental Microbiology* 11(4):809–22. DOI: 10.1111/j.1462-2920.2008.01802.x.
- Khankari, K. 2016. Airflow path matters: Patient room HVAC. ASHRAE Journal 58(6.
- Khankari, K. 2018a. Analysis of spread index: A measure of laboratory ventilation effectiveness. Paper no. HO-18-C043. 2018 ASHRAE Annual Conference—Papers [download].
- Khankari, K. 2018b. CFD analysis of hospital operating room ventilation system part I: Analysis of air change rates. *ASHRAE Journal* 60(5).
- Khankari, K. 2018c. CFD analysis of hospital operating room ventilation system part II: Analyses of HVAC configurations. *ASHRAE Journal* 60(6).
- Kudo, E., E. Song, L.J. Yockey, T. Rakib, P.W. Wong, R.J. Homer, and A. Iwasaki. 2019. Low ambient humidity impairs barrier function, innate resistance against influenza infection. *PNAS* 116(22):10905–10. https://doi.org/10.1073/pnas.1902840116.
- Kujundzic, E., F. Matalkah, D.J. Howard, M. Hernandez, and S.L. Miller. 2006. Air cleaners and upper-room air UV germicidal irradiation for controlling airborne bacteria and fungal spores. *Journal of Occupational and Environmental Hygiene* 3:536–46.
- Lax, S., and J.A. Gilbert. 2015. Hospital-associated microbiota and implications for nosocomial infections. *Trends in Molecular Medicine* 21(7):427-32. www.sciencedirect.com/science/article/abs/pii/S147149141500074X.
- Lax, S., D. Smith, N. Sangwan, K. Handley, P. Larsen, M. Richardson, S. Taylor, E. Landon, J. Alverdy, J. Siegel, B. Stephens, R. Knight, and J.A. Gilbert. 2017. Colonization and Succession of Hospital-Associated Microbiota. Sci Transl Med. 9(391):eaah6500. DOI: 10.1126/scitranslmed.aah6500. www.ncbi.nlm.nih.gov/pmc/articles/PMC5706123.

- Licina, D., A. Melikov, C. Sekhar, and K.W. Tham. 2015a. Human convective boundary layer and its interaction with room ventilation flow. *Indoor Air* 25(1):21–35. DOI:10.1111/ina.12120.
- Licina, D., A. Melikov, J. Pantelic, C. Sekhar, and K.W. Tham. 2015b. Human convection flow in spaces with and without ventilation: Personal exposure to floor-released particles and cough-released droplets. *Indoor Air* 25(6):672–82. DOI:10.1111/ina.12177.
- Li, Y. 2011. The secret behind the mask. (Editorial.) Indoor Air 21(2):89-91.
- Mead, K., and D. Johnson. 2004. An evaluation of portable high-efficiency particulate air filtration for expedient patient isolation in epidemic and emergency response. *Annals of Emergency Medicine* 44(6):635–45.
- Mead, K.R., A. Feng, D. Hammond, and S. Shulman. 2012. *In-depth report: Expedient methods for surge airborne isolation within healthcare settings during response to a natural or manmade epidemic*. EPHB Report no. 301-05f. Cincinnati, OH: U.S. Department of Health and Human Services, Public Health Service, Centers for Disease Control and Prevention, National Institute for Occupational Safety and Health. www.cdc.gov/niosh/surveyreports/pdfs/301-05f.pdf.
- Miller-Leiden, S., C. Lobascio, J.M. Macher, and W.W. Nazaroff. 1996. Effectiveness of inroom air filtration for tuberculosis control in healthcare settings. *Journal of the Air & Waste Management Association* 46:869–82.
- Mousavi, E., R. Lautz, F. Betz, and K. Grosskopf. 2019. *Academic Research to Support Facility Guidelines Institute & ANSI/ASHRAE/ASHE Standard 170*. ASHRAE Research Project CO-RP3. Atlanta: ASHRAE.
- Nicas, M., and R.M. Jones. 2009. Relative contributions of four exposure pathways to influenza infection risk. *Risk Analysis* 29:1292–303.
- Pantelic, J., and K.W. Tham. 2011. Assessment of the ability of different ventilation systems to serve as a control measure against airborne infectious disease transmission using Wells-Riley approach. IAQ 2010: Airborne Infection Control—Ventilation, IAQ, and Energy [CD]. Atlanta: ASHRAE.
- Pantelic, J., G.N. Sze-To, K.W. Tham, C.Y. Chao, and Y.C.M. Khoo. 2009. Personalized ventilation as a control measure for airborne transmissible disease spread. *Journal of the Royal Society Interface* 6(suppl_6):S715–S726.
- Pantelic, J., and K.W. Tham. 2012. Assessment of the mixing air delivery system ability to protect occupants from the airborne infectious disease transmission using Wells-Riley approach. HVAC&R Research 18(4):562–74.
- Pantelic, J., and K.W. Tham. 2013. Adequacy of air change rate as the sole indicator of an air distribution system's effectiveness to mitigate airborne infectious disease transmission caused by a cough release in the room with overhead mixing ventilation: A case study. HVAC&R Research 19(8):947–61.
- Pantelic, J., K.W. Tham, and D. Licina. 2015. Effectiveness of a personalized ventilation system in reducing personal exposure against directly released simulated cough droplets. *Indoor Air* 25(6):683–93.
- Roy, C.J., and D.K. Milton. 2004. Airborne transmission of communicable infection—The elusive pathway. *New England Journal of Medicine* 350:17.
- Shaman, J., and M. Kohn. 2009. Absolute humidity modulates influenza survival, transmission, and seasonality. *Proceedings of the National Academy of Sciences* 106(0):3243–48.

- Siegel J.D., E. Rhinehart, M. Jackson, and L. Chiarello. 2007. 2007 Guideline for Isolation Precautions: Preventing Transmission of Infectious Agents in Healthcare Settings. Atlanta: Centers for Disease Control and Prevention, The Healthcare Infection Control Practices Advisory Committee.
- Sietsema, M., L. Radonovich, F.J. Hearl, E.M. Fisher, L.M. Brosseau, R.E. Shaffer, and L.M. Koonin. 2019. A control banding framework for protecting the US workforce from aerosol transmissible infectious disease outbreaks with high public health consequences. *Health Security* 17(2):124–32. http://doi.org/10.1089/hs.2018.0103.
- Stone, W., O. Kroukamp, D.R. Korber, J. McKelvie, and G.M. Wolfaardt. 2016. Microbes at surface-air interfaces: The metabolic harnessing of relative humidity, surface hygroscopicity, and oligotrophy for resilience. *Frontiers in Microbiology* 7:1563. DOI: 10.3389/fmicb.2016.01563.
- Subhash, S.S., G. Baracco, K.P. Fennelly, M. Hodgson, and L.J. Radonovich, Jr. 2013. Isolation anterooms: Important components of airborne infection control. *American Journal of Infection Control* 41(5):452–55. DOI: 10.1016/j.ajic.2012.06.004.
- Taylor, S., and M. Tasi. 2018. Low indoor-air humidity in an assisted living facility is correlated with increased patient illness and cognitive decline. *Proceedings, Indoor Air 2018* 744:1–8.
- USDHHS. 2002. Guidance for Protecting Building Environments from Airborne Chemical, Biological, or Radiological Attacks. NIOSH Publication No. 2002-139. Washington, DC: United States Department of Health and Human Services.
- USDHHS. 2003. Guidance for Filtration and Air-Cleaning Systems to Protect Building Environments from Airborne Chemical, Biological, or Radiological Attacks. NIOSH Publication No. 2003-136. Washington, DC: United States Department of Health and Human Services.
- Yan, J., M. Grantham, J. Pantelic, P.J.B. de Mesquita, B. Albert, F. Liu, S. Ehrman, D.K. Milton, and EMIT Consortium. 2018. Infectious virus in exhaled breath of symptomatic seasonal influenza cases from a college community. *Proceedings of the National Academy of Sciences* 115(5):1081–86. DOI: 10.1073/pnas.1716561115.
- Yang, J., C. Sekhar, D. Cheong Kok Wai, and B. Raphael. 2013. CFD study and evaluation of different personalized exhaust devices. *HVAC&R Research* 19(8):934–46.
- Yang, J., C. Sekhar, D. Cheong, and B. Raphael. 2014. Performance evaluation of an integrated personalized ventilation-personalized exhaust system in conjunction with two background ventilation systems. *Building and Environment* 78:103–10. DOI:10.1016/j.buildenv.2014.04.015.
- Yang, J., S.C. Sekhar, K.W. Cheong, and B. Raphael. 2015a. Performance evaluation of a novel personalized ventilation-personalized exhaust system for airborne infection control. *Indoor Air* 25(2):176–87. DOI:10.1111/ina.12127.
- Yang, J., C. Sekhar, D.K.W. Cheong, and B. Raphael. 2015b. A time-based analysis of the personalized exhaust system for airborne infection control in healthcare settings. *Science and Technology for the Built Environment* 21(2):172–78. DOI:10.1080/10789669.2014.976511.

6. BIBLIOGRAPHY

ASHRAE. 2000. ASHRAE Guideline 12-2000, Minimizing the Risk of Legionellosis Associated with Building Water Systems. Atlanta: ASHRAE.

- ASHRAE. 2010. ASHRAE Research Strategic Plan 2010–2018. Atlanta: ASHRAE. www.ashrae.org/technical-resources/research/research-strategic-plan.
- ASHRAE. 2018. ASHRAE Position Document on Limiting Indoor Mold and Dampness in Buildings. Atlanta: ASHRAE. www.ashrae.org/file%20library/about/position%20documents/ashrae---limiting-indoor-mold-and-dampness-in-buildings.pdf.
- ASHRAE. 2017. ANSI/ASHRAE Standard 55-2017, *Thermal Environmental Conditions for Human Occupancy*. Atlanta: ASHRAE.
- Belongia, E.A., B.A. Kieke, J.G. Donahue, R.T. Greenlee, A. Balish, A. Foust, S. Lindstrom, and D.K. Shay. 2009. Effectiveness of inactivated influenza vaccines varied substantially with antigenic match from the 2004–2005 season to the 2006–2007 season. *Journal of Infectious Diseases* 199(2):159–67. DOI: 10.1086/595861.
- BOMA. 2012. Emergency Preparedness Guidebook: The Property Professional's Resource for Developing Emergency Plans for Natural and Human-Based Threats. Washington, DC: Building Owners and Managers Association International.
- Brankston, G., L. Gitterman, Z. Hirji, C. Lemieux, and M. Gardam. 2007. Transmission of influenza A in human beings. *Lancet Infectious Disease* 7:257–65.
- Bucher, S.J., 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 Reports* 123:52–60.
- Catanzaro, A. 1982. Nosocomial tuberculosis. *American Review of Respiratory Diseases* 125:559–62.
- CDC. 2001. Recognition of illness associated with the intentional release of a biologic agent. Journal of the American Medical Association 286:2088–90.
- CDC. 2003. *Guidelines for Environmental Infection Control in Health-Care Facilities*. Atlanta: Center for Disease Control and Prevention.
- CDC. 2014. NIOSH-approved N95 particulate filtering facepiece respirators. Atlanta: Center for Disease Control and Prevention. www.cdc.gov/niosh/npptl/topics/respirators/disp_part/n95list1.html.
- Chu, C.M., V.C. Cheng, I.F. Hung, K.S. Chan, B.S. Tang, T.H. Tsang, K.H. Chan, and K.Y. Yuen. 2005. Viral load distribution in SARS outbreak. *Emerging Infectious Diseases* 11(12):1882–86.
- Cole, E.C., and C.E. Cook. 1998. Characterization of infectious aerosols in health care facilities: An aid to effective engineering controls and preventive strategies. *American Journal of Infection Control* 26(4):453–64.
- D'Alessio, D.J., C.K. Meschievitz, J.A. Peterson, C.R. Dick, and E.C. Dick. 1984. Short-duration exposure and the transmission of rhinoviral colds. *Journal of Infectious Diseases* 150(2):189–94.
- Dick, E.C., C.R. Blumer, and A.S. Evans. 1967. Epidemiology of infections with rhinovirus types 43 and 55 in a group of University of Wisconsin student families. *American Journal of Epidemiology* 86(2):386–400.
- Dick, E.C., L.C. Jennings, K.A. Mink, C.D. Wartgow, and S.L. Inhorn. 1987. Aerosol transmission of rhinovirus colds. *Journal of Infectious Diseases* 156:442–8.
- Duguid, J.P. 1946. The size and duration of air-carriage of respiratory droplets and droplet nucleii. *The Journal of Hygiene* (London) 44:471–79.

- Fennelly, K.P., J.W. Martyny, K.E. Fulton, I.M. Orme, D.M. Cave, and L.B. Heifets. 2004. Coughgenerated aerosols of Mycobacterium tuberculosis: A new method to study infectiousness. *American Journal of Respiratory and Critical Care Medicine* 169:604–609.
- Gao, N.P., and J.L. Niu. 2004. CFD study on micro-environment around human body and personalized ventilation. *Building and Environment* 39:795–805.
- Gao, X., Y. Li, P. Xu, and B.J. Cowling. 2012. Evaluation of intervention strategies in schools including ventilation for influenza transmission control. *Building Simulation* 5(1):29, 37.
- Gwaltney, J., and J.O. Hendley. 1978. Rhinovirus transmission: One if by air, two if by hand. *American Journal of Epidemiology* 107(5):357–61.
- Han, K., X. Zhu, F. He, L. Liu, L. Zhang, H. Ma, X. Tang, T. Huang, G. Zeng, and B.P. Zhu. 2009. Lack of airborne transmission during outbreak of pandemic (H1N1) 2009 among tour group members, China, June 2009. *Emerging Infectious Diseases* 15(10):1578–81.
- Harriman, L., G. Brundrett, and R. Kittler. 2006. *Humidity Control Design Guide for Commercial and Institutional Buildings*. Atlanta: ASHRAE.
- Hoge, C.W., M.R. Reichler, E.A. Dominguez, J.C. Bremer, T.D. Mastro, K.A. Hendricks, D.M. Musher, J.A. Elliott, R.R. Facklam, and R.F. Breiman. 1994. An epidemic of pneumococcal disease in an overcrowded, inadequately ventilated jail. *New England Journal of Medicine* 331(10):643–8.
- Klontz, K.C., N.A. Hynes, R.A. Gunn, M.H. Wilder, M.W. Harmon, and A.P. Kendal. 1989. An outbreak of influenza A/Taiwan/1/86 (H1N1) infections at a naval base and its association with airplane travel. *American Journal of Epidemiology* 129:341–48.
- Ko, G., M.W. First, and H.A. Burge. 2002. The characterization of upper-room ultraviolet germicidal irradiation in inactivating airborne microorganisms. *Environmental Health Per*spectives 110:95–101.
- Kujundzic, E., M. Hernandez, and S.L. Miller. 2007. Ultraviolet germicidal irradiation inactivation of airborne fungal spores and bacteria in upper-room air and in-duct configurations. *Journal of Environmental Engineering and Science* 6:1–9.
- Li, Y., G.M. Leung, J.W. Tang, X. Yang, C.Y.H. Chao, J.Z. Lin, J.W. Lu, P.V. Nielsen, J. Niu, H. Qian, A.C. Sleigh, H-J. J. Su, J. Sundell, T.W. Wong, and P.L. Yuen. 2007. Role of ventilation in airborne transmission of infectious agents in the built environment—A multi-disciplinary systematic review. *Indoor Air* 17(1):2–18.
- Li, Y., H. Qian, I.T.S. Yu, and T.W. Wong. 2005. Probable roles of bio-aerosol dispersion in the SARS outbreak in Amoy Gardens, Hong Kong. Chapter 16. In *Population Dynamics and Infectious Disease in the Asia-Pacific*. Singapore: World Scientific Publishing.
- Li, Y., X. Huang, I.T.S. Yu, T.W. Wong, and H. Qian. 2005. Role of air distribution in SARS transmission during the largest nosocomial outbreak in Hong Kong. *Indoor* Air 15:83–95.
- Lowen, A.C., S. Mubareka, J. Steel, and P. Palese. 2007. Influenza virus transmission is dependent on relative humidity and temperature. *PLOS Pathogens* 3:1470–76.
- Mahida, N., N. Vaughan, and T. Boswell. 2013. First UK evaluation of an automated ultraviolet-C room decontamination device (Tru-D). *Journal of Hospital Infection* http://dx.doi.org/10.1016/j.jhin.2013.05.005.
- Mandell, G. 2010. *Mandell, Douglas, and Bennett's Principles and Practice of Infectious Diseases*, 7th ed. London: Churchill Livingstone.
- McLean, R.L. 1961. The effect of ultraviolet radiation upon the transmission of epidemic influenza in long-term hospital patients. *American Review of Respiratory Diseases* 83(2):36–8.

- MDH. 2013. Airborne Infectious Disease Management Manual: Methods for Temporary Negative Pressure Isolation. St Paul, MN: Minnesota Department of Health.
- Memarzadeh, F. 2011. Literature review of the effect of temperature and humidity on viruses. *ASHRAE Transactions* 117(2).
- Memarzadeh, F., R.M. Olmsted, and J.M. Bartley. 2010. Applications of ultraviolet germicidal irradiation disinfection in healthcare facilities: Effective adjunct, but not stand-alone technology. *American Journal of Infection Control* 38:S13–24.
- Miller, S.L., J. Linnes, and J. Luongo. 2013. Ultraviolet germicidal irradiation: Future directions for air disinfection and building applications. *Photochemistry and Photobiology* 89:777–81.
- Moser, M.R., T.R. Bender, H.S. Margolis, G.R. Noble, A.P. Kendal, and D.G. Ritter. 1979. An outbreak of influenza aboard a commercial airliner. *American Journal of Epidemiology* 110(1):1–6.
- Myatt, T.A., S.L. Johnston, Z. Zuo, M. Wand, T. Kebadze, S. Rudnick, and D.K. Milton. 2004. Detection of airborne rhinovirus and its relation to outdoor air supply in office environments. *American Journal of Respiratory and Critical Care Medicine* 169:1187–90.
- 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 Reports* 123:52–60.
- Nicas, M., W.W. Nazaroff, and A. Hubbard. 2005. Toward understanding the risk of secondary airborne infection: Emission of respirable pathogens. *Journal of Occupational and Environmental Hygiene* 2:143–54.
- NIOSH. 2009. Environmental Control for Tuberculosis: Basic Upper-Room Ultraviolet Germicidal Irradiation Guidelines for Healthcare Settings. DHHS (NIOSH) Publication No. 2009-105. Cincinnati, OH: U.S. Department of Health and Human Services, Public Health Service, Centers for Disease Control and Prevention, National Institute for Occupational Safety and Health. www.cdc.gov/niosh/docs/2009-105.
- Noti, J.D., F.M. Blachere, C.M. McMillen, W.G. Lindsley, M.L. Kashon, D.R. Slaughter, and D.H. Beezhold. 2013. High humidity leads to loss of infectious influenza virus from simulated coughs. *PLOS ONE* 8(2):e57485.
- OSHA. 1999. OSHA Technical Manual. Washington, DC: Occupational Safety & Health Administration.
- Osterholm, M.T., N.S. Kelley, A. Sommer, and E.A. Belongia. 2012. Efficacy and effectiveness of influenza vaccines: A systematic review and meta-analysis. *Lancet Infectious Diseases*. 12(1):36–44. DOI: 10.1016/S1473-3099(11)70295-X.
- Peccia, J., H. Werth, S.L. Miller, and M. Hernandez. 2001. Effects of relative humidity on the ultraviolet-induced inactivation of airborne bacteria. *Aerosol Science & Technology* 35:728–40.
- Reed, N.G. 2010. The history of ultraviolet germicidal irradiation for air disinfection. *Public Health Reports* 125(1):15–27.
- Riley, R.L., and E.A. Nardell. 1989. Clearing the air: The theory and application of ultraviolet air disinfection. *American Review of Respiratory Diseases* 139(5):1286–94.
- Riley, R.L., C.C. Mills, F. O'Grady, L.U. Sultan, F. Wittestadt, and D.N. Shivpuri. 1962. Infectiousness of air from a tuberculosis ward—Ultraviolet irradiation of infected air: Com-

- parative infectiousness of different patients. *American Review of Respiratory Diseases* 85:511–25.
- Riley, E.C., G. Murphy, and R.L. Riley. 1978. Airborne spread of measles in a suburban elementary school. *American Journal of Epidemiology* 107:421–32.
- SA Health. 2013. Guidelines for Control of Legionella in Manufactured Water Systems in South Australia. Rundle Mall, South Australia: SA Health.
- Schaffer, F.L., M.E. Soergel, and D.C. Straube. 1976. Survival of airborne influenza virus: Effects of propagating host, relative humidity, and composition of spray fluids. *Archives of Virology* 51:263–73.
- Schoen, L.J. 2020. Guidance for building operations during the COVID-19 pandemic. *ASHRAE Journal Newsletter*, March 24, 2020. www.ashrae.org/news/ashraejournal/guidance-for-building-operations-during-the-covid-19-pandemic.
- Sun, Y., Z. Wang, Y. Zhang, and J. Sundell. 2011. In China, students in crowded dormitories with a low ventilation rate have more common colds: Evidence for airborne transmission. *PLOS ONE* 6(11):e27140.
- Sylvain, D., and L. Tapp. 2009. UV-C exposure and health effects in surgical suite personnel. Health hazard evaluation report: HETA-2007-0257-3082. Cincinnati, OH: U.S. Department of Health and Human Services, Public Health Service, Centers for Disease Control and Prevention, National Institute for Occupational Safety and Health. www.cdc.gov/niosh/nioshtic-2/20035372.html.
- Tang, J.W. 2009. The effect of environmental parameters on the survival of airborne infectious agents. *Journal of the Royal Society Interface* 6:S737–S746.
- Tang, J.W., Y. Li, I. Eames, P.K.S. Chan, and G.L. Ridgway. 2006. Factors involved in the aerosol transmission of infection and control of ventilation in healthcare premises. *Journal of Hospital Infection* 64(2):100–14.
- Tellier, R. 2006. Review of aerosol transmission of influenza a virus. *Emerging Infectious Disease* 12(11):1657–62.
- VanOsdell, D., and K. Foarde. 2002. *Defining the Effectiveness of UV Lamps Installed in Circulating Air Ductwork—Final Report*. Arlington, VA: Air-Conditioning and Refrigeration Technology Institute.
- Wainwright, C.E., M.W. Frances, P. O'Rourke, S. Anuj, T.J. Kidd, M.D. Nissen, T.P. Sloots, C. Coulter, Z. Ristovski, M. Hargreaves, B.R. Rose, C. Harbour, S.C, Bell, and K.P. Fennelly. 2009. Cough-generated aerosols of Pseudomonas aeruginosa and other gram-negative bacteria from patients with cystic fibrosis. *Thorax* 64:926–31.
- Wang, B., A. Zhang, J.L. Sun, H. Liu, J. Hu, and L.X. Xu. 2005. Study of SARS transmission via liquid droplets in air. *Journal of Biomechanical Engineering* 127:32–8.
- Wang, Y., C. Sekhar, W.P. Bahnfleth, K. W. Cheong, and J. Firrantello. 2016. Effectiveness of an ultraviolet germicidal irradiation system in enhancing cooling coil energy performance in a hot and humid climate. *Energy and Buildings* 130, pp. 321–29. DOI: 10.1016/j.enbuild.2016.08.063.
- Wat, D. 2004. The common cold: A review of the literature. *European Journal of Internal Medicine* 15:79–88.
- Wells, W.F. 1948. On the mechanics of droplet nuclei infection; Apparatus for the quantitative study of droplet nuclei infection of animals. *Am J Hyg.* 47(1):1–10. DOI: 10.1093/oxfordjournals.aje.a119176.

- Wells, W.F. 1955. *Airborne Contagion and Air Hygiene*. Cambridge: Harvard University Press, pp. 191.
- WHO. 2007. Legionella and the prevention of Legionellosis. Geneva: World Health Organization.
- WHO. 2009. *Natural Ventilation for Infection Control in Health-Care Settings*. Geneva: World Health Organization.
- WHO. 2014. Influenza: Public health preparedness. Geneva: World Health Organization. www.who.int/influenza/preparedness/en.
- Wong, B.X., N. Lee, Y. Li, P.X. Chan, H. Qiu, Z. Luo, R.X. Lai, K.X. Ngai, D.X. Hui, K.X. Choi, and I.X. Yu. 2010. Possible role of aerosol transmission in a hospital outbreak of influenza. *Clinical Infectious Diseases* 51(10):1176–83.
- Xie, X.J., Y.G. Li, H.Q. Sun, and L. Liu. 2009. Exhaled droplets due to talking and coughing. *Journal of The Royal Society Interface* 6:S703–S714.
- Xie, X., Y. Li, A.T.Y. Chwang, P.L. Ho, and H. Seto. 2007. How far droplets can move in indoor environments—Revisiting the Wells evaporation-falling curve. *Indoor Air* 17:211–25.
- Xu, P., E. Kujundzic, J. Peccia, M.P. Schafer, G. Moss, M. Hernandez, and S.L. Miller. 2005. Impact of environmental factors on efficacy of upper-room air ultraviolet germicidal irradiation for inactivating airborne mycobacteria. *Environmental Science & Technol*ogy 39:9656–64.
- Xu, P., J. Peccia, P. Fabian, J.W. Martyny, K. Fennelly, M. Hernandez, and S.L. Miller. 2003. Efficacy of ultraviolet germicidal irradiation of upper-room air in inactivating bacterial spores and mycobacteria in full-scale studies. *Atmospheric Environment* 37:405–19.
- 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):792–8.
- Yang, W., and L. Marr. 2012b. Mechanisms by which ambient humidity may affect viruses in aerosols. *Applied and Environmental Microbiology* 78(19):6781. DOI: 10.1128/AEM.01658–12.
- Yang, W., S. Elankumaran, and L.C. Marr. 2012. Relationship between humidity and influenza A viability in droplets and implications for influenza's seasonality. *PLOS ONE* 7(10):e46789. DOI:10.1371/journal.pone.0046789.
- Yu, I.T., Y. Li, T.W. Wong, W. Tam, A.T. Chan, J.H. Lee, D.Y. Leung, and T. Ho. 2004. Evidence of airborne transmission of the severe acute respiratory syndrome virus. *N Engl J Med* 350:1731–39.