© ASHRAE (www.ashrae.org). For personal use only. Additional reproduction, distribution, or transmission in either print or digital form is not permitted without ASHRAE's prior written permission.

ASHRAE Standing Standard Project Committee 170 Cognizant TC: TC 9.6, Healthcare Facilities SPLS Liaison 2011-2013: Douglass S. Abramson SPLS Liaison 2009-2011: Byron W. Jones SPLS Liaison 2008-2009: William F. Walter

Paul T. Ninomura, Chair Hamid Habibi Richard D. Moeller Chris P. Rousseau, Co-Vice Chair and Secretary Jeffery M. Hardin Tyler Ninomura Michael P. Sheerin, Co-Vice Chair Richard D. Hermans Russell N. Olmsted Judene M. Bartley* Michael R. Keen Heather L. Platt Theodore Cohen Anand K. Seth Marvin L. Kloostra Rajendra N. Shah John M. Dombrowski Peter H. Langowski Douglas S. Erickson Michael F. Mamayek Gordon P. Sharp James (Skip) Gregory Farhad Memarzadeh Andrew J. Streifel Michael E. Woolsev

ASHRAE STANDARDS COMMITTEE 2013–2014

William F. Walter, Chair John F. Dunlap Rick A. Larson Richard L. Hall, Vice-Chair James W. Earley, Jr. Mark P. Modera Karim Amrane Steven J. Emmerich Cyrus H. Nasseri Joseph R. Anderson Julie M. Ferguson Janice C. Peterson James Dale Aswegan Krishnan Gowri Heather L. Platt Charles S. Barnaby Cecily M. Grzywacz Douglas T. Reindl Steven F. Bruning Rita M. Harrold Julia A. Keen, BOD ExO Thomas E. Werkema, Jr., CO John A. Clark Adam W. Hinge

Waller S. Clements Debra H. Kennoy
David R. Conover Malcolm D. Knight

Stephanie C. Reiniche, Manager of Standards

SPECIAL NOTE

This American National Standard (ANS) is a national voluntary consensus standard developed under the auspices of ASHRAE. Consensus is defined by the American National Standards Institute (ANSI), of which ASHRAE is a member and which has approved this standard as an ANS, as "substantial agreement reached by directly and materially affected interest categories. This signifies the concurrence of more than a simple majority, but not necessarily unanimity. Consensus requires that all views and objections be considered, and that an effort be made toward their resolution." Compliance with this standard is voluntary until and unless a legal jurisdiction makes compliance mandatory through legislation.

ASHRAE obtains consensus through participation of its national and international members, associated societies, and public review.

ASHRAE Standards are prepared by a Project Committee appointed specifically for the purpose of writing the Standard. The Project Committee Chair and Vice-Chair must be members of ASHRAE; while other committee members may or may not be ASHRAE members, all must be technically qualified in the subject area of the Standard. Every effort is made to balance the concerned interests on all Project Committees.

The Manager of Standards of ASHRAE should be contacted for:

- a. interpretation of the contents of this Standard,
- b. participation in the next review of the Standard,
- c. offering constructive criticism for improving the Standard, or
- d. permission to reprint portions of the Standard.

DISCLAIMER

ASHRAE uses its best efforts to promulgate Standards and Guidelines for the benefit of the public in light of available information and accepted industry practices. However, ASHRAE does not guarantee, certify, or assure the safety or performance of any products, components, or systems tested, installed, or operated in accordance with ASHRAE's Standards or Guidelines or that any tests conducted under its Standards or Guidelines will be nonhazardous or free from risk.

ASHRAE INDUSTRIAL ADVERTISING POLICY ON STANDARDS

ASHRAE Standards and Guidelines are established to assist industry and the public by offering a uniform method of testing for rating purposes, by suggesting safe practices in designing and installing equipment, by providing proper definitions of this equipment, and by providing other information that may serve to guide the industry. The creation of ASHRAE Standards and Guidelines is determined by the need for them, and conformance to them is completely voluntary.

In referring to this Standard or Guideline and in marking of equipment and in advertising, no claim shall be made, either stated or implied, that the product has been approved by ASHRAE.

^{*}This edition is dedicated to our friend and colleague, Judene Bartley. This Standard benefited tremendously from her insight and tireless contributions regarding healthcare infection prevention.

© ASHRAE (www.ashrae.org). For personal use only. Additional reproduction, distribution, or transmission in either print or digital form is not permitted without ASHRAE's prior written permission.

CONTENTS

ANSI/ASHRAE/ASHE Standard 170-2013, Ventilation of Health Care Facilities

SECTION -oreword		PAGE
		2
	1 Purpose	2
	2 Scope	
	3 Definitions	2
	4 Compliance	3
	5 Planning	4
	6 Systems and Equipment	4
	7 Space Ventilation	7
	8 Planning, Construction, and System Startup	14
	9 Normative References	15
	Informative Appendix A—Operations and Maintenance Procedures	16
	Informative Appendix B—Informative References and Bibliography	
		18

NOTE

Approved addenda, errata, or interpretations for this standard can be downloaded free of charge from the ASHRAE Web site at www.ashrae.org/technology.

© 2013 ASHRAE

(This foreword is not part of this standard. It is merely informative and does not contain requirements necessary for conformance to the standard. It has not been processed according to the ANSI requirements for a standard and may contain material that has not been subject to public review or a consensus process. Unresolved objectors on informative material are not offered the right to appeal at ASHRAE or ANSI.)

FOREWORD

ANSI/ASHRAE/ASHE Standard 170, Ventilation of Health Care Facilities, is one of a family of documents that offers guidance, regulation, and mandates to designers of health care facilities. It is first and foremost a mandatory minimum requirement and, as such, may not always offer the state-of-the-art best practice for health care ventilation design. Other publications, such as the ASHRAE HVAC Design Manual for Hospitals and Clinics, 2nd Edition, complement the standard, providing additional depth and detail for the designer. In addition, the health care designer must refer to any design requirements from the appropriate jurisdiction that has authority. Many jurisdictions use or refer to Guidelines for Design and Construction of Health Care Facilities, published by the Facility Guidelines Institute (FGI). Where practical, the committee was cognizant of these other documents in the development of this standard.

Ventilation design for health care spaces is a combination of tasks that leads to a set of documents used in construction. One such task requires medical planners to develop departmental programs of spaces. These programs include space names that suggest the use for which the space is intended, and health care ventilation designers depend upon these names to determine the ventilation parameters for their designs. This standard provides these ventilation parameters.

Ventilation systems and designs for health care facilities are intended to provide a comfortable environment for patients, health care workers, and visitors while diluting, capturing and exhausting airborne contaminants including potentially infectious airborne agents such as M. tuberculosis. Without high-quality ventilation in health care facilities, patients, health care workers, and visitors can become exposed to contaminants through normal respiration of particles in the air. Poorly ventilated health care facilities may increase the concentration of airborne contaminants including fungi or mold, which may cause allergic responses in even healthy workers and occupants. Some patients are profoundly immunosuppressed for prolonged periods and, if exposed, are highly susceptible to infection from fungi. For such patients, fungal spores become invasive pathogens and lead to high rates of severe morbidity and mortality. For all these reasons, and considering the various occupancies and patient populations, great care must be taken in the design of health care ventilation systems.

1. PURPOSE

The purpose of this standard is to define ventilation system design requirements that provide environmental control for comfort, asepsis, and odor in health care facilities.

2. SCOPE

- **2.1** The requirements in this standard apply to patient-care areas and related support areas within health care facilities, including hospitals, nursing facilities, and outpatient facilities.
- **2.2** This standard applies to new buildings, additions to existing buildings, and those alterations to existing buildings that are identified within this standard.
- **2.3** This standard considers chemical, physical, and biological contaminants that can affect the delivery of medical care to patients; the convalescence of patients; and the safety of patients, health care workers, and visitors.

3. DEFINITIONS

absorption distance: the distance downstream of a humidifier required for all moisture to be absorbed into the airstream.

addition: an extension or increase in floor area or height of a *building*, building system, or equipment.

airborne infection isolation (AII): the isolation of patients infected with organisms spread by airborne droplet nuclei less than 5 µm in diameter (see FGI [2010] in Informative Appendix B). For the purposes of this standard, the abbreviation "AII" refers to the room that provides isolation.

airborne infection isolation room: a room that is designed according to the requirements of this standard and that is intended to provide airborne infection isolation.

alteration: a significant change in the function or size of a space, in the use of its systems, or in the use of its equipment, either through rearrangement, replacement, or addition. Routine maintenance and service shall not constitute an alteration.

authority having jurisdiction: the agent or agency responsible for enforcing this standard.

average velocity: the volumetric flow rate obtained by dividing the air quantity issuing from an air distribution device by the nominal face area of the device.

building: a structure that is wholly or partially enclosed within exterior walls and a roof, or within exterior and party walls and a roof, and that affords shelter to persons, animals, or property. In this standard, a building is a structure intended for use as a hospital or health care facility.

classification of surgeries:

procedure room (Class A surgery): provides minor surgical procedures performed under topical, local, or regional anesthesia without preoperative sedation. Excluded are intravenous, spinal, and epidural procedures, which are Class B or C surgeries.

operating room (Class B surgery): provides minor or major surgical procedures performed in conjunction with oral, parenteral, or intravenous sedation or performed with the patient under analgesic or dissociative drugs.

operating room (Class C surgery): provides major surgical procedures that require general or regional block anesthesia and/or support of vital bodily functions.

(For more information on this method of classifying surgeries, see ACS (2000) in Informative Appendix B.)