



Oklahoma Association of Healthcare Engineers

August 23, 2019







Session 1: FGI 2018 Overview

Oklahoma Association of Healthcare Engineers < 2019 Summer Regional Event

August 23, 2019



Session 1 Learning Objectives

- 1. RECOGNIZE KEY ORGANIZATIONAL CHANGES AND REQUIREMENTS TO SUCCESSFULLY APPLY GUIDELINES FOR DESIGN AND CONSTRUCTION OF HOSPITALS, 2018 EDITION
- 2. IDENTIFY AND EXPLAIN NEW INITIATIVES FOUND IN GUIDELINES FOR DESIGN AND CONSTRUCTION OF HOSPITALS, 2018 EDITION
- 3. DESCRIBE UPDATED VENTILATION REQUIREMENTS (ASHRAE 170 2017) FOR WHICH HOSPITALS NEED TO BE PREPARED AS COMPARED TO PREVIOUS REQUIREMENTS (ASHRAE 170 – 2008, NFPA 99 -2012 AND ASHRAE 170 - 2013, FGI 2014) IN GUIDELINES FOR DESIGN AND CONSTRUCTION OF HOSPITALS, 2018 EDITION
- 4. DESCRIBE UPDATED REQUIREMENTS FOR NURSE CALL, RECEPTACLES, AND MEDICAL GAS OUTLETS



AGENDA

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- Managing Director and Market Leader for HFG Architecture's Oklahoma Region.
- NCARB and LEED Accredited.
- Member of the American Institute of Architects, the American Society of Healthcare Engineers, and the Oklahoma Association of Healthcare Engineers.

Jessica Zvonek, PE

- Mechanical Associate at Professional Engineering Consultants
- EDAC and LEED Accredited.
- Responsibilities include the design of plumbing, heating, ventilation, and air conditioning (HVAC) systems from the initial concept stage to the final bid documents with extensive experience in healthcare.



INTRODUCTION





Session 1: FGI 2018 Overview

- 1. FGI 2018 ORGANIZATIONAL STRUCTURE
- 2. HOSPITAL ITEMS OF SIGNIFICANCE
 - **KEY FRONT-END CONCEPTS**
 - ACOUSTIC DESIGN
 - PATIENT CARE UNITS / PATIENT ROOMS ٠
 - TELEMEDICINE
 - OPERATING AND PROCEDURE ROOMS
- OUTPATIENT ITEMS OF SIGNIFICANCE З.
- VENTILATION AND SYSTEMS OVERVIEW 4.
- FUTURE OF FGI 5.
- QUESTIONS AND ANSWERS 6.



AGENDA













"...the most significant change to the 2018 edition of the Guidelines is that these important design standards are now presented as three independent documents."

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Guidelines for Design and Construction of Hospitals

Provides standards for designing and constructing hospital facilities.

<u>Guidelines for Design and Construction of Outpatient Facilities</u>

Provides standards for designing and constructing outpatient facilities.

Guidelines for Design and Construction of Residential Health, Care, and Support Facilities

Provides standards for designing and constructing residential facilities.











Basic Organization of the Guidelines

Main body

Part 1 contains chapters that address considerations applicable to all hospitals/outpatient facilities, except as modified in specific facility chapters in Part 2.

Part 2 addresses facilities where inpatient/outpatient care is provided, with chapters devoted to common elements.

Hospital Guidelines includes general hospitals, critical access hospitals, psychiatric hospitals, rehabilitation hospitals, and children's hospitals. Chapters on freestanding emergency departments and mobile/transportable medical units are also included.

Outpatient Guidelines includes specific requirements for outpatient facility types.

Part 3 contains the full text of the American Society of Heating, Refrigerating and Air-Conditioning Engineers (ASHRAE) Standard 170-2017: Ventilation of Health Care Facilities.

Appendix items are incorporated within the main body of the text.

An asterisk (*) preceding a section or paragraph number indicates that explanatory or educational material can be found in an appendix item located at the bottom of the page.

Appendix items are identified by the letter "A" preceding the section or paragraph number in the main text to which they relate.

Cross-references. Cross-references are used throughout the Guidelines: example: See Section 2.2-2.1.3 (Accommodations for Care of Patients of Size).

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Major Additions and Revisions of the 2018 Guidelines

Functional Programs

Changes include additional requirements for space programs.

Acoustics

Many new requirements and guidance for acoustics

Patients of Size

Many new requirements and guidance for patients of size

Sustainable Design

Sustainable design becomes a part of the *Guidelines*, though much of it is Appendix content.

Telemedicine

Telemedicine is now addressed in the *Guidelines*

Imaging Facilities

Now classified according to use in Hospitals Guidelines and included separately in Outpatient Guidelines.

Pre- and Post-Procedure Patient Care

Separate or combined areas. Minimums established for required ratios based on imaging rooms or procedure/Ors

Sterile processing

- One room processing facility no longer permitted except as an exception for small countertop, limited workflow SP facilities. Two-room minimum, including decontam. and clean workroom.
- Requirements and guidance for storage of clean instruments are also provided in Hospital and OP Guidelines









Major Additions and Revisions of the 2018 Guidelines

Technology distribution room

TDR space requirements revised to provide a minimum three-foot clearance on all sides of equipment racks vs. 2014 minimum of 12 feet by 14 feet for the TDR.

Critical care unit patient rooms

In new construction, all patient rooms in critical care units except NICUs will be single-patient rooms with potential ٠ exceptions for renovations.

Procedure and operating rooms

- Better alignment of requirements in room types where procedures occur based on levels of invasiveness and risk to patient.
- Addresses support spaces for patients. ٠
- Table provided to quickly identify which procedures are to occur in which spaces.



Guidelines for Design and Construction of Hospitals

Provides standards for designing and constructing hospital facilities.

List of Tables 2018 Health Guidelines Revisions Major Additions and Revisions Glossary List of Acronyms Part 1: General Chapter 1.1: Introduction Chapter 1.2: Planning, Design, Construction, and Commissioning Chapter 1.3: Site Chapter 1.4: Equipment Part 2: Hospital Facility Types Chapter 2.1: Common Elements for Hospitals Chapter 2.2: Specific Requirements for General Hospitals Chapter 2.3: Specific Requirements for Freestanding Emergency Care Facilities Chapter 2.4: Specific Requirements for Critical Access Hospitals Chapter 2.5: Specific Requirements for Psychiatric Hospitals Chapter 2.6: Specific Requirements for Rehabilitation Hospitals Chapter 2.7: Specific Requirements for Children's Hospitals Chapter 2.8: Specific Requirements for Mobile/Transportable Medical Units Part 3: Ventilation of Hospitals ANSI/ASHRAE/ASHE Standard 170-2017 Ventilation of Health Care

Facilities

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Guidelines for Design and Construction of Hospitals

Provides standards for designing and constructing hospital facilities.

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The Facility Guidelines Institute

2018 edition





Includes ANSI/ASHRAE/ASHE Standard 170-2017: Ventilation of Health Care Facilities



* 1.2-1.2 Multidisciplinary Project Team

1.2-1.2.1 Multidisciplinary groups/persons (stakeholders) affected by and integral to the design shall be included in the project planning and implementation process.

A1.2-1.2 Project team

a. The multidisciplinary project team should be assembled as early as possible in the design process.

b. The multidisciplinary team should include administrators, clinicians, infection preventionists, architects and other design professionals, facility managers, safety officers, security managers, users of equipment, and support staff relevant to the areas affected by the project as well as those with knowledge of the organization's functional goal for the project. Inclusion of patient advocates/consumers, A/E consultants, and construction specialists should be considered.

1.2-1.2.2 The scope and nature of the project shall dictate others to be involved on the project team.



* 1.2-3 Space Program

A space program, organized by department, now must be provided **separately** from the functional program.

Must include:

- Each room in the project
- Size of each room, including gross square footage and clear square footage.
- Must cite relevant paragraphs from the FGI Document

A1.2-3 Project gross floor area

a. Gross floor area for the project should be aggregated by department, and multiplying factors should be applied to reflect circulation and wall thicknesses within the department or functional area. This result is referred to as department gross square footage (DGSF).

b. DGSF for the project should be aggregated, and multiplying factors should be applied to reflect interdepartment circulation patterns, exterior wall thicknesses, engineering spaces, general storage spaces, vertical circulation, and any other areas not included within the intra-department calculations. This result is referred to as building gross square footage (BGSF) and reflects the overall size of the project.



HOSPTIALS - PART 1

* 1.2-6 Planning and Design Considerations and Requirements

Acoustic design plays a large role in the newest guidelines.

Must include :

- Accommodations for site exterior noise
- Considerations for facility noises that may reach nearby residences •
- Surfaces
- Maximum room noise levels
- Speech privacy
- Building vibration



Acoustic Design

The appendix in this section provides useful information that will help direct acoustical design efforts, including:

- Definitions
- Separate limits for daytime or nighttime
- Reference codes for acoustic design
- Guidance on how to address facility site noise when the hospital cannot operationally control the noise

Means of measurement may include:

- Exterior site observations
- Sound-level monitoring surveys

Strategies may include:

Distance, Orientation, Shielding





Acoustic Design – A1.2-6.1.2 Site Exterior Noise

This section provides design guidance on how to address environmental noise at a facility site

- May include on-site noise or off-site noise
- May be under hospital's operational control (plant, generators)
- May be under hospital's control, but with limited ability to control (helipad or heliport)
- May not be under hospital's control (roads, rail, airports, power plants, etc.)

This section is meant to provide a means for screening sites to help determine which exterior wall/window assemblies are suitable to address site noise; it is not intended to be used as a means to qualify the suitability of a site with respect to environmental noise exposure.









Acoustic Design – 1.2-6.1.2 Site Exterior Noise

Various means are suggested to evaluate noise in order to better design for the local acoustical environment

- Four categories in which to classify environmental noise (A-D) •
- Table 1.2-3 and appendix table A1.2-b offer descriptions of sound categories

Table 1.2-3: Categorization of Hospital Sites by Exterior Ambient Sound with Design Criteria
for Sound Isolation of Exterior Shell in New Construction

Exterior Site Noise Exposure Category							
	Α	В	С	D			
General description	Minimal	Moderate	Significant	Extreme			
Outdoor day-night average		(5. (0)	70.74	≥ 75			
sound level during (L_{dn}) $(dBA)^{i}$	< 65	65–69	70–74				
Outdoor average hourly nominal							
maximum sound level $(L_{01})^2$	< 75	75–79	80-84	≥ 85			
(dBA)							
Design Criteria for Sound Isolation of Exterior Shell in New Construction ³							
Minimum exterior shell	OITCe: 25	OITCc: 30	OITCe: 35	OITCc: 40			
composite sound transmission	or	or	or	or			
rating ^{4, 5, 6}	STCe: 35	STCc: 40	STCc: 45	STCc: 50			

OITC - Outdoor-Indoor Transmission Class

STC – Sound Transmission Class





STC RATING

VERSUS

COMPARING THE 2 TYPES OF SOUND RATINGS



Acoustic Design – Interiors (1.2-6.4-7)

Design Criteria for Acoustic Surfaces

Normally occupied spaces in a hospital shall now require surfaces that help achieve minimum sound absorption coefficients. There are several important considerations

Other Design Criteria:

- Room Noise Levels (HVAC, etc.) (Table 1.2-5)
- Interior wall and floor/ceiling construction (Table 1.2-6)
- Speech privacy (Table 1.2-7)
- Vibration control
- Alarm fatigue top priority in hospitals; leads to patient stress
- OR noise levels may be difficult to achieve due to HVAC systems noise
- Vibration produced by equipment & activities (i.e. footfall)









<u>Guidelines for Design and Construction of Hospitals</u>

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Part 2: Hospital Facility Types Chapter 2.1: Common Elements for Hospitals Section 2.1-1: General Section 2.1-2: Patient Care Units and Other Patient Care Areas Section 2.1-3: Diagnostic and Treatment Areas Section 2.1-4: Patient Support Facilities Section 2.1-5: General Support Facilities Section 2.1-6: Public and Administrative Areas Section 2.1-7: Design and Construction Requirements Section 2.1-8: Building Systems Chapter 2.2: Specific Requirements for General Hospitals Chapter 2.3: Specific Requirements for Freestanding Emergency Care Facilities Chapter 2.4: Specific Requirements for Critical Access Hospitals Chapter 2.5: Specific Requirements for Psychiatric Hospitals Chapter 2.6: Specific Requirements for Rehabilitation Hospitals Chapter 2.7: Specific Requirements for Children's Hospitals Chapter 2.8: Specific Requirements for Mobile/Transportable Medical Units



Hospitals

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Includes ANSI/ASHRAE/ASHE Standard 170-2017: Ventilation of Health Care Facilities



2.1-2 – Patient Care Units and Other Patient Care Areas Accommodations for Patients of Size

- Not only "bariatric" but all patients of size
- The number of POS spaces is to be determined by hospital based on projected need.
- Special design requirements for POS include:
 - A.I.I. rooms
 - Space provisions
 - Exam / treatment rooms
 - Toilet rooms and bathing facilities, including patient lifts to assist caregivers







PART 2

HOSPTIALS –



2.1-3 – Diagnostic and Treatment Areas 2.2-2 – Patient Care Units

Single Patient Exam Room: 120 SF clear floor area, 10 ft. min. dim.

Minimum 3 feet at sides, foot of bed (see notes in • chapter about placement)

Multiple Patient Exam Room: 80 SF clear *per station*

- 5 feet between beds
- 4 feet between beds/walls

SAFE Room: same as single patient exam room

- Private shower •
- A room for consultation, family, support services, and law enforcement shall be readily accessible





*2.1-3.3 Accommodations for Telemedicine Services

A2.1-3.3 Patient experience. Remote communications via electronic equipment, although not a replacement for in-person care, may be offered as a supplement where in-person care is not available or medically necessary. To assist in the adoption of telemedicine and maximize its benefits for elderly patients, those unaccustomed to electronic communication, and those with vision, hearing, or cognitive impairments, care should be given to remove technological barriers and provide telemedicine endpoints that facilitate natural communication for the widest range of participants. Facilities and systems used for telemedicine communications should strive to maintain the level of safety, privacy, quality of care, and patient experience that would be expected for in-person communication.





A2.1-3.3.1 Telemedicine service types

a. Services may include one-on-one interactions, consultations with a patient and family members (e.g., pediatric or elderly patients), examinations supported by a telemedicine presenter located with the patient, or specialty services such as dermatology or orthopedics. ...to achieve a functional design, it is important to know what services will be provided.





A2.1-3.3.2 Design considerations for telemedicine

- Be careful with camera placement. Eye-to-eye camera angle important.
- Provide adequate HVAC and electrical support based on equipment
- b. Architectural details
- Doors should allow for maximum privacy
- Doors should not be placed behind the patient







- Equipment based on exam type
- Lighting should be both direct & indirect, from sides of patient
- Room color can affect quality • of exam





HOSPTIALS - PART 2

Patient Care Stations (PCS)

The number of required patient care stations is defined by the Guidelines.

- When combined into one area, provide at least 2 PCS per Class 2 and Class 3 imaging, procedure, or operating room.
- If separate:
 - Pre-Procedure Room or Area: 1 PCS / imaging, procedure, or OR.
 - Phase | PACU: 1 PCS / Class 3 imaging or OR (was 1.5 PACU / OR)
 - Phase II Recovery Room or Area: 1 Phase II PCS per Class 2 or Class 3 imaging, procedure, or OR





Patient Care Stations (PCS)

- Where cubicles are used, the following minimum clearances shall be provided:
 - 3 feet between the sides of patient beds/gurneys/lounge chairs and adjacent walls or partitions (was 4 feet)
 - 2 feet between the foot of patient beds/gurneys/lounge chairs and the cubicle curtain (was 3 feet)
 - Where bays or cubicles face each other, an aisle with a minimum clearance of 8 feet independent of the foot clearance between patient stations or other fixed objects shall be provided







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Patient Care Stations (PCS)

- Where bays are used, the following minimum clearances shall be provided:
 - 5 feet between the sides of patient beds/gurneys/lounge chairs
 - 3 feet between the sides of patient beds/gurneys/lounge chairs and adjacent walls or partitions (was 4 feet)
 - 2 feet between the foot of patient beds/gurneys/lounge chairs and the cubicle curtain (was 3 feet)



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Patient Care Stations (PCS)

Where single-patient rooms are used, 3 feet shall be provided • between the sides and foot of beds/gurneys/lounge chairs and adjacent walls or partitions.





Patient Care Stations (PCS)

Where single-patient rooms are used, 3 feet shall be provided between the sides and foot of beds/gurneys/lounge chairs and adjacent walls or partitions.





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Part 2: Hospital Facility Types

Chapter 2.1: Common Elements for Hospitals

Chapter 2.2: Specific Requirements for General Hospitals

Section 2.2-1: General

Section 2.2-2: Patient Care Units

Section 2.2-3: Diagnostic and Treatment Facilities

Section 2.2-4: Patient Support Facilities

Section 2.2-5: General Support Facilities

Section 2.2-6: Public and Administrative Areas

Section 2.2-7: Design and Construction Requirements

Section 2.2-8: Building Systems

Chapter 2.3: Specific Requirements for Freestanding Emergency Care Facilities

Chapter 2.4: Specific Requirements for Critical Access Hospitals

Chapter 2.5: Specific Requirements for Psychiatric Hospitals

Chapter 2.6: Specific Requirements for Rehabilitation Hospitals

Chapter 2.7: Specific Requirements for Children's Hospitals

Chapter 2.8: Specific Requirements for Mobile/Transportable Medical Units

Guidelines

FOR DESIGN AND CONSTRUCTION OF Hospitals

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2018 ORGANIAZATION

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Includes ANSI/ASHRAE/ASHE Standard 170-2017: Ventilation of Health Care Facilities



Procedures 2.2-3.3.3 Operating Rooms

*Operating rooms are where invasive procedures are to be performed

- Minimum OR size is 400 SF clear
- Image guided surgery (or other procedures requiring large equipment or more personnel) requires 600 SF clear (renovations: 500 SF w/ 20 ft clear dim possible)
- No longer are "clear dimensions" required, however minimum clearances are now established as:
 - 8.5 ft on each side of the operating table, gurney, or procedural chair
 - 6 ft at the head. This dimension shall result in an anesthesia work zone with a clear floor area of 6 ft x 8 ft





Procedures 2.2-3.3.2 Procedure Room

Governing body to complete a clinical assessment of procedures to determine the appropriate room type and location and document this in the functional program.

A2.2-3.3.2.1 (1) (c) Procedures that require different pressure relationships cannot be provided in the same procedure room; if these rooms are also used for other procedures, the other procedures must be able to be performed in the same pressure environment.

Table 2.2-1: Examination/Treatment, Procedure, and Operating Room Classification¹

Room	الدم			Design Requirements ²
Room	Use	Room Type	Location	Surfaces
Exam or treatment room	Patient care that may require high- level disinfected or sterile instruments but does not require the environmental controls of a procedure room	Unrestricted area	Accessed from an unrestricted area	<i>Flooring:</i> cleanable and wear-resistant for the location; stable, firm, and slip-resistant <i>Wall finishes:</i> washable <i>Ceiling:</i> cleanable with routine housekeeping equipment; la in ceiling permitted
Procedure room	Patient care that requires high-level disinfection of the room, sterile instruments, and some environmental controls but does not require the environmental controls of an operating room Endoscopic procedures	Semi- restricted area	Accessed from an unrestricted or a semi- restricted area	 <i>Flooring:</i> cleanable and wear-resistant for the location; stable, firm, and slip-resistant <i>Floor and wall base assemblies in cystoscopy, urology, and endoscopy procedure rooms and endoscope processing room:</i> monolithic floor with integral coved wall base carried up the wall a minimum of 6 inches <i>Wall finishes:</i> washable <i>Wall finishes in endoscopy procedure room and endoscope processing room:</i> washable; free of fissures, open joints, or crevices <i>Ceiling:</i> smooth and without crevices, scrubbable, non-absorptive, non-perforated; capable of withstanding cleaning chemicals; lay-in ceiling permitted if gasketed or each ceiling; tile weighs at least one pound per square foot and no perforated, tegular, serrated, or highly textured tiles
Operating room	Invasive procedures ³ Any procedure during which the patient will require physiological monitoring and is anticipated to require active life support	Restricted area	Accessed from a semi- restricted area	<i>Flooring:</i> cleanable and wear-resistant for the location; stable, firm, and slip-resistant <i>Floor and wall base assemblies:</i> monolithic floor with integral coved wall base carried up the wall a minimum of 6 inches <i>Wall finishes:</i> washable; free of fissures, open joints, or crevices <i>Ceiling:</i> monolithic, scrubbable, capable of withstanding cleaning and/or disinfecting chemicals, gasketed access openings



HOSPTIALS - PART 2

Procedures 2.2-3.3.2 Procedure Room

Procedure rooms shall be a minimum clear floor area of 130 SF, unless anesthesia is administered using anesthesia machine and supply carts which shall have a minimum clear floor area of 160 SF

Clearance requirements are

- 3.5 ft. on each side of gurney/chair
- 3 ft at the head and foot
- *Where an anesthesia machine and associated supply cart are used, the clearance at the head shall be 6 ft

A2.2-3.3.2.2 (2) (b) Anesthesia work zone. On the outside edge of the anesthesia work zone, 2 ft x 8 ft may serve as part of the circulation pathway.

Table 2.2-1: Examination/Treatment, Procedure, and Operating Room Classification¹

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Procedure room	Patient care that requires high-level disinfection of the room, sterile instruments, and some environmental controls but does not require the environmental controls of an operating room Endoscopic procedures	Semi- restricted area	Accessed from an unrestricted or a semi- restricted area	 <i>Flooring:</i> cleanable and wear-resistant for the location; stable, firm, and slip-resistant <i>Floor and wall base assemblies in cystoscopy, urology, and endoscopy procedure rooms and endoscope processing room:</i> monolithic floor with integral coved wall base carried up the wall a minimum of 6 inches <i>Wall finishes:</i> washable <i>Wall finishes in endoscopy procedure room and endoscope processing room:</i> washable; free of fissures, open joints, or crevices <i>Ceiling:</i> smooth and without crevices, scrubbable, non-absorptive, non-perforated; capable of withstanding cleaning chemicals; lay-in ceiling permitted if gasketed or each ceiling tile weighs at least one pound per square foot and no perforated, tegular, serrated, or highly textured tiles
Operating room	Invasive procedures ³ Any procedure during which the patient will require physiological monitoring and is anticipated to require active life support	Restricted area	Accessed from a semi- restricted area	<i>Flooring:</i> cleanable and wear-resistant for the location; stable, firm, and slip-resistant <i>Floor and wall base assemblies:</i> monolithic floor with integral coved wall base carried up the wall a minimum of 6 inches <i>Wall finishes:</i> washable; free of fissures, open joints, or crevices <i>Ceiling:</i> monolithic, scrubbable, capable of withstanding cleaning and/or disinfecting chemicals, gasketed access openings



Procedures 2.2-3.4 Imaging Services

Now classified according to use in Hospitals Guidelines and included separately in Outpatient Guidelines. Per Guidelines, this is intended to aid in easier adaptation of new technologies and equipment.

- Class 1 (unrestricted area) for services that use a natural orifice entry (noninvasive)
- Class 2 (semi-restricted area) for diagnostic and therapeutic procedures
- Class 3 (restricted area) for **invasive** procedures and any Class 2 procedure during which the patient will require physiological monitoring and is anticipated to require life support.
- Class 2 and 3 rooms should have a physically separate control room

Table 2.2-2: Classification of Room Types for Imaging Services

Deem	llas	Use Design Requirements ²		
Room	- Ose	Room Type	Location	Surfaces
Class 1 imaging room	Diagnostic radiography, fluoroscopy, mammography, computed tomography (CT), ultrasound, magnetic resonance imaging (MRI), and other imaging modalities Services that use natural orifice entry and do not pierce or penetrate natural protective membranes	Unrestricted area	Accessed from an unrestricted area	<i>Flooring:</i> cleanable and wear-resistant for the location; stable, firm, and slip-resistant <i>Wall finishes:</i> washable <i>Ceiling:</i> cleanable with routine housekeeping equipment; lay-in ceiling permitted
Class 2 imaging room	Diagnostic and therapeutic procedures such as coronary, neurological, or peripheral angiography Electrophysiology procedures	Semi- restricted area	Accessed from an unrestricted or a semi- restricted area	 <i>Flooring:</i> cleanable and wear-resistant for the location; stable, firm, and slip-resistant <i>Floor and wall base assemblies:</i> monolithic floor with integral coved wall base carried up the wall a minimum of 6 inches <i>Wall finishes:</i> washable; free of fissures, open joints, or crevices <i>Ceiling:</i> smooth and without crevices, scrubbable, non-absorptive, non-perforated; capable of withstanding cleaning chemicals; lay-in ceiling permitted if gasketed or each ceiling tile weighs at least one pound per square foot and no perforated, tegular, serrated, or highly textured tiles
Class 3 imaging room	Invasive procedures ³ Any Class 2 procedure during which the patient will require physiological monitoring and is anticipated to require active life support	Restricted area	Accessed from a semi- restricted area	<i>Flooring:</i> cleanable and wear-resistant for the location; stable, firm, and slip-resistant <i>Floor and wall base assemblies:</i> monolithic floor with integral coved wall base carried up the wall a minimum of 6 inches <i>Wall finishes:</i> washable; free of fissures, open joints, or crevices <i>Ceiling:</i> monolithic, scrubbable, capable of withstanding cleaning and/or disinfecting chemicals, gasketed access openings



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Guidelines FOR DESIGN AND CONSTRUCTION OF **Outpatient Facilities** The Facility Guidelines Institute 2018 edition FGI FGI

Guidelines for Design and Construction of Outpatient Facilities

Provides standards for designing and constructing outpatient facilities.

Major Additions and Revisions

Glossary List of Acronyms Part 1: General Chapter 1.1: Introduction Chapter 1.2: Planning, Design, Construction, and Commissioning Chapter 1.3: Site Chapter 1.4: Equipment Part 2: Outpatient Facility Types Chapter 2.1: Common Elements for Outpatient Facilities Chapter 2.2: Specific Requirements for General and Specialty Medical Services Facilities Chapter 2.3: Specific Requirements for Outpatient Imaging Facilities Chapter 2.4: Specific Requirements for Birth Centers Chapter 2.5: Specific Requirements for Urgent Care Centers Chapter 2.6: Specific Requirements for Infusion Centers Chapter 2.7: Specific Requirements for Outpatient Surgery Facilities Chapter 2.8: Specific Requirements for Freestanding Emerg. Care Facilities Chapter 2.9: Specific Requirements for Endoscopy Facilities Chapter 2.10: Specific Requirements for Renal Dialysis Centers Chapter 2.11: Specific Requirements for Outpatient Psychiatric Facilities Chapter 2.12: Specific Requirements for Rehabilitation Therapy Facilities Chapter 2.13: Specific Requirements for Mobile/Transportable Medical Units Chapter 2.14: Specific Requirements for Dental Facilities

Part 3: Ventilation of Hospitals Index



ANSI/ASHRAE/ASHE Standard 170-2017 Ventilation of Outpatient Facilities

Guidelines for Design and Construction of Outpatient Facilities

Outpatient *Guidelines* are applied as:

- Specific outpatient project types in facility type chapters in Part 2
 - Part 1 applies to all facility types
 - Requirements in Chapter 2.1, Common Elements for Outpatient Facilities, apply only when crossreferenced in the facility chapter.
- Note, per *Guidelines*, cross-references to the common elements often begin with "where provided," which allows for **flexible application** in facility design to fit particular services provided.









Major Additions and Revisions of the 2018 *Guidelines* Facilities Chapters

Freestanding Diagnostic and Treatment Facilities (Outpatient Imaging Facility)

- In Guidelines since 1990
- Major overhaul in 2018 to become outpatient imaging center with reference to common elements and rad. therapy requirements.

Urgent Care Centers

Revised to allow more flexible design.

Outpatient Surgery Centers

- Procedure and operating room requirements were moved into the common elements chapter, where they can be easily cross reference from other chapters
- Descriptions of unrestricted, semi restricted, and restricted areas were updated to correlate with AORN requirements,
- Support areas were reorganized to clarify their location in the outpatient surgery facility: in the semi-restricted area, directly accessible to the semi-restricted area, or elsewhere in the facility.
- Operating Rooms of 255 SF require the following clearances:
 - 6 ft on each side, 5 ft at head and foot
- Operating Rooms of 270 SF require the following clearances:
 - 6 ft on each side. 5 ft at the foot
 - Anesthesia work zones require 6 ft by 8 ft clear at head of operating table.
- Operating Rooms of 400 SF requires a sterile field of:
 - 8 ft each side, 6 ft at head, 7 ft at foot of operating table
 - Anesthesia work zone requires the same 6 ft x 8 ft clear at head of operating table.

Endoscopy

- Procedure room reduced from 200 SF to 180 SF.
- Processing rooms updated to match sterile processing requirements (dirty to clean)



FGI Guidelines Resources

Fgiguidelines.org

- Design Guide for the Built Environment of Behavioral Health Facilities lacksquare
- FGI Acoustics Working Group (AWG) 2011 white paper •
- Articles by FGI HGRC members in peer-reviewed journals lacksquare
- Articles elaborating on changes to the Guidelines ullet

Madcad.com (electronic version)

FGI Errata

Presentations on the 2018 Guidelines from the 2017 HCD Conference + Expo https://www.fgiguidelines.org/resource/presentations-2018-guidelines-2017-hcd-conference-expo/#



1.1-8 Codes, Standards, and Other Documents Referenced in the Guidelines

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Requirements:	ASHRAE 170 (2008) NFPA 99 (2012) / Current CMS Requirement	ASHRAE 170 (2013) FGI 2014 / Current OSDH Requirement		
	Ventilation shall be required for the following spaces: <> All rooms <> PE rooms <> Class B & C Operating rooms, including Caesarean	Ventilation shall be required for the following spaces: <> All rooms <> PE rooms <> Class B & C Operating rooms, including Caesarean	Ventilat <> All r <> PE r <> Clas	
Reserve Heating	Capacity of back up/reserve sources shall be sufficient to provide for sterilization and diatary purposes and to provide heat for operating, delivery, birthing, labor, recovery, emergency, intensive care, nursery, and inpaitient rooms.	Capacity of back up/reserve sources shall be sufficient to provide for domestic hot water, sterilization and diatary purposes and to provide heat for operating, delivery, birthing, labor, recovery, emergency, intensive care, nursery, and inpaitient rooms.	Capacity domest heat for intensiv	
Reserve Cooling	Greater than 400 tons of cooling there shall be more than one chiller for redundnacy to maintain facility operation plan.	for Greater than 400 tons of cooling there shall be more than one chiller to redundnacy to maintain facility operation plan.		
Outdoor Air Intakes	Outside air intakes shall be: <> 25' away from exhaust and vents <> 6' above grade <> 3' above roof	Outside air intakes shall be: <> 25' away from exhaust and vents <> 6' above grade <> 3' above roof	Outside <> 25' a <> 6' ab <> 3' a Exception distance	
Filters:	<> MERV 12 filters and above shall have differential pressure device. <> Filter 1 shall be upstream of coil. <> Filter 2 shall be downstream of coil. *Refer to Table 6-1 Minimum Filter Efficiencies	<> MERV 12 filters and above shall have differential pressure device. <> Filter 1 shall be upstream of coil. <> Filter 2 shall be downstream of coil. *Refer to Table 6.4 Minimum Filter Efficiencies	<> MER <> Filte <> Filte *Refer t	
Humidifiers:	Locate humidity sensor a suitable distance downstream from steam injection source. Controls shall limit duct humidity to a max of 90% RH. Steam valve shall remain off when AHU is not in operation.	Steam humidifiers shall be used. Locate humidity sensor a suitable distance downstream from steam injection source. Controls shall limit duct humidity to a max of 90% RH. Steam valve shall remain off when AHU is not in operation. Duct takeoffs shall not be within the absorption distance.	Steam h distance duct hu AHU is r distance <u>Humidi</u>	



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ion shall be required for the following spaces: ooms ooms

B & C Operating rooms, including Caesarean

of back up/reserve sources shall be sufficient to provide for ic hot water, sterilization and diatary purposes and to provide operating, delivery, birthing, labor, recovery, emergency, e care, nursery, and inpaitient rooms.

than 400 tons of cooling there shall be more than one chiller for nacy to maintain facility operation plan.

air intakes shall be: away from exhaust and vents oove grade bove roof

<u>on: Gas fired package air handling does not have to meet the 25'</u> e between OSA intake and unit flue.

RV 12 filters and above shall have differential pressure device. r 1 shall be upstream of coil. r 2 shall be downstream of coil.

o Table 6.4 Minimum Filter Efficiencies

umidifiers shall be used. Locate humidity sensor a suitable downstream from steam injection source. Controls shall limit midity to a max of 90% RH. Steam valve shall remain off when not in operation. Duct takeoffs shall not be within the absorption Requirements for Steam humidifiers vs Adiabatic Atomizing fiers provided.

		ACHDAE 170 (2012)	
Requirements:	ASHRAE 170 (2008) NFPA 99 (2012) / Current CMS Requirement	ASHRAE 170 (2013) FGI 2014 / Current OSDH Requirement	
	Discharge air from All rooms, bronchoscopy rooms, emergency department waiting rooms, nuclear medicine laboratories, radiology waiting, and laboratory chemical fume hoods shall: have ductwork under negative pressure; discharge 10' above roof level; be located to minimize recirculation of air back into building	<> Discharge air from All rooms, bronchoscopy rooms, emergency department waiting rooms, nuclear medicine laboratories, radiology waiting, and laboratory chemical fume hoods shall: have ductwork under	
Ductwork:	Spaces listed in Table 7-1 Design Parameters that have required pressure relationshiops shall be served by fully ducted returns.	Smoke and Fire Dampers shall be located on design drawings and provided with access to all dampers. Spaces listed in Table 7.1 Design Parameters that have required pressure relationships shall be served by fully ducted returns and exhaust as well as the following areas: <> Surgery and critical care recovery rooms, critical and intensive care areas, intermediate care areas, and wound intensive care units (burn units). <> Inpatient facilities patient care areas.	Smoke and F provided wit Spaces listed relationships as the follow <> Surgery a areas, intern units). <> Inpatient
Supply Diffusers:	<> Operating room diffusers shall allow for internal cleaning. <> Psychiatric, seclusion, and holding-patient rooms shall be designed with security diffusers, grilles, and registers. *Refer to Table 6-2 Supply Air Outlets	<> Operating room diffusers shall allow for internal cleaning. <> Psychiatric, seclusion, and holding-patient rooms shall be designed with security diffusers, grilles, and registers. *Refer to Table 6.7.2 Supply Air Outlets	<> Operating <> Psychiatri with security *Refer to Ta



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e air from All rooms, bronchoscopy rooms<u>, **sputum collectio**</u> tamidine administration rooms, emergency department ing rooms, nuclear medicine laboratories, radiology waiting awaiting chest x-rays for respiratory disease), pharmacy drug, and laboratory chemical fume hoods shall: Have nder negative pressure; -Discharge 10' above roof level; be minimize recirculation of air back into building Γowers drifts shall be located away from air intakes and meet uirements. discharge of All rooms, bronchoscopy, sputum collection, azardous-drug, and laboratory chemical fume hoods shall e the roof and discharge in a vertical direction. bry chemical fume hoods shall discharge with a stack velocity um of 2500 fpm. s, bronchoscopy, sputum collection, laboratory chemical

s shall be lcoated a minimum of 25 ft horizontally form intakes, openable windows/doors, and areas accessible to

Fire Dampers shall be located on design drawings and ith access to all dampers.

ed in Table 8.1 Design Parameters that have required pressure os shall be served by fully ducted returns and exhaust as well wing areas:

and critical care recovery rooms, critical and intensive care mediate care areas, and wound intensive care units (burn

nt facilities patient care areas.

ng room diffusers shall allow for internal cleaning. ric, seclusion, and holding-patient rooms shall be designed ty diffusers, grilles, and registers.

able 6.7.2 Supply Air Outlets

Requirements:	ASHRAE 170 (2008) NFPA 99 (2012) / Current CMS Requirement	ASHRAE 170 (2013) FGI 2014 / Current OSDH Requirement	
Energy Recovery Systems:	N/A	<> Energy Recovery systems shall be located upstream of Filter Bank No. 2. <> Energy Recovery systems with air leakage potential shall be designed to have no more than 5% of the total supply airsteram consisting of exhaust air. <> All rooms or combination All/PE rooms shall not be utilized for energy recovery.	<> Energenergenergenergenergenergenergenerg
Space Ventilation:	<> Movement of air shall be designed from gernerally clean to less clean areas. <> Ventilation rates are intended for comfort, asepsis, and odoer control. *Refer to Table 7-1 Design Parameters	<> Movement of air shall be designed from gernerally clean to less clean areas. <> Ventilation rates are intended for comfort, asepsis, and odoer control. <> Spaces permited to be reciculated by room units shall: not receive nonfiltered, non conditioned outside air; serve only a single space; provide a minumum of MERV 6 filter. <> Outdoor air quantity of AHU's serving multiple spaces can be calculated by the sum of individual space requirements or per the ASHRAE 62.1 method. *Refer to Table7.1 Design Parameters	<> Move areas. <> Venti control. <> Space nonfilter provide a <> Outd calculate ASHRAE *Refer to
Airborne Infection Isolation (AII) Rooms:	<> A pressure monitor device shall be installed locally to indicate (-) 0.01" wc negative differential is maintained when occupied by patients with airborne diseases. <> Air is exhausted directly outdoors without mixing with with non-All rooms. <> Exhaust grilles shall be located directly over patient beds or on the wall near the patient head.	<> A pressure monitor device shall be installed locally to indicate (-) 0.01" wc negative differential is maintained when occupied by patients with airborne diseases. <> Air is exhausted directly outdoors without mixing with with non-AII rooms. <> Exhaust grilles shall be located directly over patient beds or on the wall near the patient head. <> When Ante rooms accompany an AII room, the AII room shall be negative to the Ante room, and the Ante room shall be negative to the corridor. <> When an AII is not utilized for an airborne disease patient, it must still remain negative. Reversable airflow controls are not allowed.	A preserved of the second s



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rgy Recovery systems shall be located upstream of Filter Bank No.

rgy Recovery systems with air leakage potential shall be designed no more than 5% of the total supply airsteram consisting of air.

ooms or combination AII/PE rooms shall not be utilized for recovery.

vement of air shall be designed from gernerally clean to less clean

tilation rates are intended for comfort, asepsis, and odoer

ces permited to be reciculated by room units shall: not receive red, non conditioned outside air; serve only a single space; a minumum of MERV 6 filter.

door air quantity of AHU's serving multiple spaces can be ed by the sum of individual space requirements or per the 62.1 method.

to Table8.1 Design Parameters

essure monitor device shall be installed locally to indicate (-) 0.01" ative differential is maintained when occupied by patients with e diseases.

exhausted directly outdoors without mixing with with non-All

ust grilles shall be located directly over patient beds or on the ar the patient head.

en Ante rooms accompany an All room, the All room shall be e to the Ante room, and the Ante room shall be negative to the r.

n an All is not utilized for an airborne disease patient, it must still negative. Reversable airflow controls are not allowed.

Requirements:	ASHRAE 170 (2008) NFPA 99 (2012) / Current CMS Requirement	ASHRAE 170 (2013) FGI 2014 / Current OSDH Requirement	
	<> A pressure monitor device shall be installed locally to indicate (+) 0.01" wc positive differential is maintained when occupied by patients requireing a protective environment. <> Supply diffusers shall be located directly over patient beds, and return/exhaust grilles lcoated near the patient room door.	<> A pressure monitor device shall be installed locally to indicate (+) 0.01" wc positive differential is maintained when occupied by patients requireing a protective environment. <> Supply diffusers shall be located directly over patient beds, and return/exhaust grilles lcoated near the patient room door. <> When Ante rooms accompany a PE room, the PE room shall be positive to the Ante room, and the Ante room shall be poitive to the corridor. <> When a PE room is not utilized for a patient needing a protective environiment, it must still remain positive and shall be constant volume. Reversable airflow controls are not allowed.	<> A pre 0.01" we require (<> Supp return/e <> Whe positive corridor <> Whe environi Reversa
Combination Airborn Infectious Isolation/Protectiv e Environment (AII/PE) rooms:	N/A	<> Supply air diffusers shall be loated above patient bed with exhaust grilles located near patient room door. <> Two pressure monitor devices shall be installed locally to indicate the required pressure is maintained at both the AII/PE door to the Ante room and at the Ante room to the corridor. <> Ante room pressurization will either be positive to both the AII/PE room and corridor or negative to both the AII/PE room and corridor depending on what is required and defined by the hospital.	
Critical Care Units:	Wound Intensive Care Units (Burn Units) that require humidifiers to comply with Table 7-1 shall be provided with individual humidity control.	Wound Intensive Care Units (Burn Units) that require humidifiers to comply with Table 7.1 shall be provided with individual humidity control.	Wound I comply v



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ressure monitor device shall be installed locally to indicate (+) c positive differential is maintained when occupied by patients eing a protective environment.

ply diffusers shall be located directly over patient beds, and exhaust grilles lcoated near the patient room door.

en Ante rooms accompany a PE room, the PE room shall be e to the Ante room, and the Ante room shall be poitive to the or.

en a PE room is not utilized for a patient needing a protective iment, it must still remain positive and shall be constant volume. able airflow controls are not allowed.

ply air diffusers shall be lcoated above patient bed with exhaust located near patient room door.

o pressure monitor devices shall be installed locally to indicate the ed pressure is maintained at both the AII/PE door to the Ante room the Ante room to the corridor.

e room pressurization will either be positive to both the AII/PE and corridor or negative to both the AII/PE room and corridor ling on what is required and defined by the hospital.

Intensive Care Units (Burn Units) that require humidifiers to with Table 8.1 shall be provided with individual humidity control.

Requirements:	ASHRAE 170 (2008) NFPA 99 (2012) / Current CMS Requirement	ASHRAE 170 (2013) FGI 2014 / Current OSDH Requirement		
Operating Rooms:	<> Surgry rooms shall maintain a (+) 0.01" wc positive differential to all adjacent spaces. <> Surgery supply difusers shall be laminar flow diffusers delivering air	adjacent spaces and each room shall have individual temperature control. nd <> Surgery supply difusers shall be laminar flow diffusers delivering air between 25-35 ft/min covering a minimum of 70% of the patient bed and		
Imaging Procedures Rooms:	If invasive procedures take place it shall be designed as Class A surgery, if anesthesia gas is used it shall be designed as Class B or C surgery.	If invasive procedures take place it shall be designed as Class A surgery, if anesthesia gas is used it shall be designed as Class B or C surgery.	lf invasi anesthe	
Morgue and Autopsy Rooms:	<> Low sidewall exhaust shall be provided or exhaust removed through an autopsy table and shall be exhausted directly outdoors without mixing with any other exhaust systems.	<> Low sidewall exhaust shall be provided or exhaust removed through an autopsy table and shall be exhausted directly outdoors without mixing with any other exhaust systems. <> A differential pressure between the morgue and autopsy and adjacent spaces shall maintain a (-) 0.01" wc negative pressure.	<> Low an auto with any <> A dif spaces s	
Bronchoscopy Rooms:	N/A	<> A differential pressure between the bronchoscopy procedure or sputum induction room and adjacent spaces shall maintain a (-) 0.01" wc negative pressure. <> Local exhaust shall be provided for sputum collection procedures.	<> A dif sputum negative <> Loca	



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rgry rooms shall maintain a (+) 0.01" wc positive differential to all ent spaces and each room shall have individual temperature ol.

rgery supply difusers shall be laminar flow diffusers delivering air en 25-35 ft/min covering a minimum of 70% of the patient bed and ound the patient bed.

oms shall have (2) low sidewall return or exhaust grilles at opposite rs approximately 8" above finished floor. In addition to the

ed low wall grilles, grilles may also be placed high on the walls. Iditonal supply diffusers shall be permiteed within the room le of the primary array to provide additional environmental ements.

sive procedures take place it shall be designed as Class A surgery, if nesia gas is used it shall be designed as Class B or C surgery.

w sidewall exhaust shall be provided or exhaust removed through topsy table and shall be exhausted directly outdoors without mixing iny other exhaust systems.

lifferential pressure between the morgue and autopsy and adjacent s shall maintain a (-) 0.01" wc negative pressure.

lifferential pressure between the bronchoscopy procedure or m induction room and adjacent spaces shall maintain a (-) 0.01" wc ve pressure.

cal exhaust shall be provided for sputum collection procedures.

Table 6.4 Minimum Filter Efficiencies

Space Designation (According to Function)	Filter Bank No. 1 (MERV) ^a	Filter Bank No. 2 (MERV) ^a
Operating rooms (ORs); inpatient and ambulatory diagnostic and therapeutic radiology; inpatient delivery and recovery spaces	7	14
Inpatient care, treatment, and diagnosis, and those spaces providing direct service or clean supplies and clean processing (except as noted below); AII (rooms)	7	14
Protective environment (PE) rooms	7	HEPA ^{c,d}
Laboratory work areas, procedure rooms, and associated semirestricted spaces	13 ^b	NR
Administrative; bulk storage; soiled holding spaces; food preparation spaces; and laundries	7	NR
All other outpatient spaces	7	NR
Nursing facilities	13	NR
Psychiatric hospitals	7	NR
Resident care, treatment, and support areas in inpatient hospice facilities	13	NR
Resident care, treatment, and support areas in assisted living facilities	7	NR

The new FGI 2018 requirements have more requirements than CMS.

NR = not required

ASHRAE 17C

a. Informative Note: The minimum efficiency reporting value (MERV) is based on the method of testing described in ANSI/ASHRAE Standard 52.2 (ASHRAE [2017a]).

b. Additional prefilters may be used to reduce maintenance for filters with efficiencies higher than MERV 7.

c. As an alternative, MERV-14 rated filters may be used in Filter Bank No. 2 if a tertiary terminal HEPA filter is provided for these spaces.

The new FGI 2018 requirements have

more requirements than CMS.

d. Informative Note: High-efficiency particulate air (HEPA) filters are those filters that remove at least 99.97% of 0.3 micron-sized particles at the rated flow in accordance with the testing methods of IEST RP-CC001.6 (IEST [2016]).

Table 6.7.2 Supply Air Outlets

Space Designation (According to Function)	Supp
Operating rooms (ORs) ^b , procedure rooms	Supp nona
	Addi
Protective environment (PE) rooms	Grou
Wound intensive care units (burn units)	Grou
Trauma rooms (crisis or shock)	Grou
AII rooms	Grou
Single-bed patient or resident rooms e	Grou
All other patient care or resident care spaces	Grou
All other spaces	No re

RAE [2017c]), for definitions related to outlet classification and performance. b. Surgeons may require alternate air distribution systems for some specialized surgeries. Such systems shall be considered acceptable if they meet or exceed the requirements of this

standard.

c. Air distribution systems using Group D diffusers shall meet the following requirements:

1. The system shall be designed according to "Design Guidelines" in System Performance Evaluation and Design Guidelines for Displacement Ventilation, Chapter 74. 2. The supply diffuser shall be located where it cannot be permanently blocked (Informative Note: e.g., opposite the foot of the bed). 3. The room return/exhaust grille shall be located in the ceiling, approximately above the head of the patient bed.

4. The transfer grille to the toilet room shall be located above the occupied zone.



PEC – PROFESSIONAL ENGINEERING CONSULTANTS PA



ply Air Outlet Classification^a

ply diffusers within the primary supply diffuser array: Group E, aspirating

litional supply diffusers within the room: Group E

up E, nonaspirating

up E, nonaspirating

up E, nonaspirating

up A or Group E

up A, Group D, or Group E

up A or Group E

equirement

New Space Types – New Requirements

Table 7.1	Design	Para	eters-	-Hospital	Spa

Function of Space	Pressure Relationship to Adjacent Areas (n)	Minimum Outdoor ach	Minimum Total ach	All Room Air Exhausted Directly to Outdoors (j)	Air Recirculated by Means of Room Units (a)	Design Relative Humidity (k), %	Design Temperature (°F/°C
DIAGNOSTIC AND TREATMENT Laboratory work area, media transfer (f), (v)	Positive	2	4	NR	NR	NR	70-75/21-24
Laboratory work area, microbiology (f), (v)	Negative	2	6	Yes	NR	NR	70-75/21-24
Laboratory work area, nuclear medicine (f), (v)	Negative	2	6	Ves	NR	NR	70-75/21-24
Laboratory work area, pathology (f), (v)	Negative	2	6	Yes	NR	NR	70-75/21-24
Laboratory work area, serology (f), (v)	Negative	2	6	Yes	NR	NR	70-75/21-24
Laboratory work area, sterilizing (f)	Negative	2	10	Yes	NR	NR	70-75/21-24
Medication room	NR	2	4	NR	NR	Max 60	70-75/21-24
Nonrefrigerated body-holding room (h)	Negative	NR	10	Yes	No	NR	70-75/21-24
Nuclear medicine hot lab	Negative	NR	6	Yes	No	NR	70-75/21-24
Nuclear medicine treatment room	Negative	2	6	Yes	NR	NR	70-75/21-24
Pharmacy (b)	Positive	2	4	NR	NR	NR	NR
Physical therapy	Negative	2	6	NR	NR	Max 65	72-80/22-27
Special examination room (aa)	NR	2	6	NR	NR	Max 60	70-75/21-24
Treatment room	NR	2	6	NR	NR	Max 60	70-75/21-24
STERILIZING	NK	2	0	NK	INK	Max 00	70-75/21-24
	Negative	NR	10	Yes	N	NR	NR
Sterilizer equipment room STERILE PROCESSING DEPARTMENT ^z	Negative	NK	10	Yes	No	NK	NK
	D			10			(0. 53 /00. 00
Clean workroom	Positive	2	4	NR	No	Max 60	<u>68–73</u> /20–23
Decontamination room	Negative	2	6	Yes	No	NR	<u>60–73</u> /16–23
Sterile storage room	Positive	2	4	NR	NR	Max 60	Max 75/24
SERVICE							
Bathroom	Negative	NR	10	Yes	No	NR	72-78/22-26
Bedpan room	Negative	NR	10	Yes	No	NR	NR
Clean linen storage	Positive	NR	2	NR	NR	NR	72-78/22-26
Dietary storage	NR	NR	2	NR	No	NR	72-78/22-26
Food preparation center (i)	NR	2	10	NR	No	NR	72-78/22-26
Janitor's closet	Negative	NR	10	Yes	No	NR	NR
Laundry, general	Negative	2	10	Yes	No	NR	NR
Linen and trash chute room	Negative	NR	10	Yes	No	NR	NR
Soiled linen sorting and storage	Negative	NR	10	Yes	No	NR	NR
Warewashing	Negative	NR	10	Yes	No	NR	NR
SUPPORT SPACE							
Clean workroom or clean holding	Positive	2	4	NR	NR	NR	NR
Hazardous material storage	Negative	2	10	Yes	No	NR	NR
Soiled workroom or soiled holding	Negative	2	10	Yes	No	NR	NR
Note: NR = no requirement		-	10				
Normative Notes for Table 7.1: a. Except where indicated by a "No" in this column, recircul HVAC units (with heating or cooling coils) are acceptable ing that portion of the minimum total air changes per hour mitted by Section 7.1 (subparagraph [a][5]). Because of th difficulty and potential for buildup of contamination, re room units shall not be used in areas marked "No." Re devices with high-efficiency particulate air (HEPA) filte permitted in existing facilities as interim, supplemental env	for provid- that is per- e cleaning infiltra circulating restrict circulating or the rs shall be the nut	e proper makeup a ASHRAE Standar- tion to or from en ions of NFPA 904 maximum defined	ir to kitchen exh d 154 ⁶ . In some kit corridors con A ⁷ , the pressure in the table. Dur s to any extent re	h) shall be that required aust systems as specified cases, excess exfiltration apromises the exit corri- equirements of NFPA 9 ing operation, a reduction quired for odor control sh	in s. For intermed or delivery/reco dor ted when sup ing and coolin to t. The protective all patient from	very/postpartum rooms plemental heating and/o ng, baseboard heating, o e environment airflow common environment	ry/recovery rooms, and lat , four total ach shall be perr or cooling systems (radiant he

	HVAC units (with heating or cooling coils) are acceptating that portion of the minimum total air changes per homitted by Section 7.1 (subparagraph [a][5]). Because o difficulty and potential for buildup of contamination,
	room units shall not be used in areas marked "No."
	devices with high-efficiency particulate air (HEPA) fi permitted in existing facilities as interim, supplemental e controls to meet requirements for the control of airbon
	agents. The design of either portable or fixed systems si stagnation and short circuiting of airflow. The design of
	shall also allow for easy access for scheduled prevent nance and cleaning.
b.	Pharmacy compounding areas may have additional air c ential pressure, and filtering requirements beyond the
	this table, depending on the type of pharmacy, the regula ments (which may include adoption of USP-797), the
	level of risk of the work, and the equipment used in the s mative Note: See USP (2017a) in Appendix B.
c.	The term trauma room as used herein is a first-aid room gency department room used for general initial treatment

gency department room used tor general initial treatment of accident victims. The OR within the trauma center that is routinely used for emergency surgery is considered to be an OR by this standard. Pressure relationships need not be maintained when the room is

- ments. Higher ventilation in. dictated by the labor iontial cor rentilation rates above the total ach listed shall be used when tes above me training and the hazard rever-atory program requirements and the hazard rever-minants in each laboratory work area. Lower tota shall be permitted when a Hazard Assessment per Managemet of the potential contaminants in e ach ventilation rates shall be perm formed as part of an effective Laboratory Ventilation Management Plan per ANSI/AIHA/ASSE Z9.5, American National Standard for Laboratory Ventilation13 determines that either (a) acceptable expo are concentrations in the laboratory work area can be achieved with a ower minimum total ach ventilation rate than is listed in Table 7.1 or p. lower minimum total ach ventilation rate than is listed in Table 7.1 or p.
 (b) a demad control approach with active sensing of contaminants or appropriate surrogates is used as described in *ASHRAE Handbook*-*HVAC Applications*, Chapter 16, "Laboratorics" (*Mnormative Note:* 9.
 Sea ASHRAE [2015] in Informative Appendix B).
 All air need not be exhausted if darkroome equipment has a seaveng-ing exhaust duct attached and meets ventilation standards regarding NIOSH³, OSHA, and local employee exposure limits.
 h. A nonrefrigerated body-holding room is applicable only to facilities that do not perform autopsies on-site and use the space for short periods while waiting for the body to be transferred.

		2017 A	SHRAE	170			Г
eters—Hospital Spaces	Pressure Relationship to Adjacent Areas (n)	Minimum Outdoor ach	Minimum Total ach	All Room Air Exhausted Directly to Outdoors (j)	Air Recirculated by Means of Room Units (a)	Design Relative Humidity (k), %	Design Temperature (l), °F/°C
AL CARE							
are	NR	2	6	NR	No	30-60	70-75/21-24
ean) (m), (o)	Positive	4	20	NR	No	20-60	68-75/20-24
decontamination	Negative	2	12	Yes	No	NR	NR
exam/treatment room (p)	NR	2	6	NR	NR	Max 60	70-75/21-24
public waiting area	Negative	2	12	Yes (q)	NR	Max 65	70-75/21-24
	NR	2	6	NR	NR	Max 60	70-75/21-24
	Positive	3	15	NR	No	20-60	70-75/21-24
storage (r)	Negative	NR	8	Yes	NR	NR	NR
	Positive	2	6	NR	No	30-60	72-78/22-26
)	Positive	4	20	NR	No	20-60	68-75/20-24
oscopic rooms (m), (o)	Positive	4	20	NR	No	20-60	68-75/20-24
	Positive	3	15	NR	No	20-60	70-75/21-24
ns	Negative	2	12	Yes (q), (w)	NR	Max 60	70-75/21-24
	NR	2	6	NR	No	20-60	70-75/21-24
	NR	2	6	NR	No	NR	NR
shock) (c)	Positive	3	15	NR	No	20-60	70-75/21-24
	NR	2	6	NR	NR	20-60	70-75/21-24
	Negat <u>i</u> ve	2	12	Yes (q)	NR	Max 60	70-75/21-24
burn unit)	NR	2	6	NR	No	40-60	70-75/21-24
uni unit)		-			110	10 00	10 10/21 21
	(c)	NR	10	Yes	No	NR	NR
	Negative	2	12	Yes	No	Max 60	70-75/21-24
teroom	(e)	- NR	10	Yes	No	NR	NR
om	Positive	2	12	Yes	No	Max 60	70-75/21-24
	N/R	2	6	N/R	No	30-60	72-78/22-26
(LDR) (s)	NR	2	6	NR	NR	Max 60	70-75/21-24
//postpartum (LDRP) (s)	NR	2	6	NR	NR	Max 60	70-75/21-24
(b) (b)	NR	2	6	NR	No	30-60	72-78/22-26
om	NR	2 NR	2	NR	NR	NR	NR
511	NR	NR	2	NR	NR	NR	NR
	NR	2	2 4 (y)	NR	NR	Max 60	70-75/21-24
						NR	
room (t)	(e) Regitive	NR 2	10	NR	No		NR
room (t)	Positive	2	12	NR	No	Max 60	70-75/21-24
	Negative	NR	10	Yes	No	NR	NR
	Nagativa	NR	10	Var	No	NR	70 75/21 24
	Negative NR	2	6	Yes NR	No NR	NR	70-75/21-24 70-75/21-24
		2				NR	
its (dining	Negative		6	NR	NR		70-75/21-24
/ity/dining	NR	4	4	NR	NR	NR	70-75/21-24
	NR	2	2	NR	NR	NR	70-75/21-24
	NR	NR	4	NR	NR	NR	NR
	Nagatina	2	10	Var	No	ND	NP
	Negative	2	10	Yes	No	NR	NR
reatment)	NR	2	6	NR	NR	Max 60	72-78/22-26
are and catheterization)	Positive	3	15	NR	No	Max 60	70-75/21-24
ATMENT							
	Negative	2	12	Yes	No	NR	68-75/20-24
collection, istration	Negative	2	12	Yes	No	NR	68-73/20-23
	NR	2	6	NR	NR	NR	72-78/22-26
oom	Negative	NR	10	Yes	No	NR	NR
	NR	2	4	NR	NR	Max 60	72-78/22-26
	Manufact	2	10	N.			

Yes

NR

NR

NR

Yes

Yes

Yes NR

Yes

Yes

10

6

2

No

No

NR

NR

NR

NR

NR

NR

NR

NR

No longer required to have pressure relationships, but CMS still requires that they do.

Humidity range is 20% to 60%; but
CMS still requires 30% to 60%.

Less ACH now required, but CMS still requires higher.

Laboratories no longer have a requirement to be circulated by room units, but CMS still requires that they cannot.



Table 7.1 Design Paramet

Emergency department of Emergency department p

Medical/anesthesia gas s Newborn intensive care Operating room (m), (o) Operating/surgical cysto Procedure room (o), (d) Radiology waiting rooms Recovery roon Substerile service area Trauma room (crisis or s Treatment room (p) Triag

Wound intensive care (b INPATIENT NURSING AII anteroom (u) AII room (u) Combination AII/PE ant

Combination AII/PE roo Continued care nursery Labor/delivery/recovery Labor/delivery/recovery/ Newborn nursery suite

Patient corrido Patient room PE anteroom (Protective environment Toilet room

NURSING FACILITY Bathing room

Occupational therap Physical therapy Resident gathering/activ Resident room Resident unit c RADIOLOGY

Darkroom (g)

X-ray (diagnostic and tre

X-ray (surgery/critical ca

Bronchoscopy, sputum c and pentamidine adminis

Dialysis treatment area Dialyzer reprocessing ro

ECT procedure room

Endoscope cleaning

Hydrotherapy

General examination room

Gastrointestinal endoscopy procedure room (x)

Laboratory work area, bacteriology (f), (v)

Laboratory work area, biochemistry (f), (v)

Laboratory work area, cytology (f), (v)

Laboratory work area, general (f), (v)

Laboratory work area, glasswashing (f)

Laboratory work area, histology (f), (v)

Negativ

NR

NR

Negative

Negative

Negative

Negative

Negative

Negative

DIAGNOSTIC AND TREA

Autopsy room

ASHRAE 1

Laser eye room

Function of Space SURGERY AND CRITICA Critical and intensive car Delivery room (Caesarea Emergency department

PEC – PROFESSIONAL ENGINEERING CONSULTANTS PA

NR

20-60

Max 60

NR

NR

NR

NR

NR

NR

NR

NR

68-73/20-23

70-75/21-24

72-80/22-27

70-75/21-24

70-75/21-24

70-75/21-24

70-75/21-24

70-75/21-24

NR

aces (Continued)

should preven f such system tative mainte

hange, differ minimum o atory require spaces. Infor

nt of acciden

exhaust air shall be discharged directly to the outdoors and not recirculated to other areas. Individual circumstances may require special consideration for air exhausted to the outdoors. To satisfy exhaust eeds, constant replacement air from the outdoors is necessary when

he system is in operation The RH ranges listed are the minimum and/or maximum allowable at

- The Kritanges instead are the minimum and/or maximum and/water and any point within the design temperature range required for that space. Systems shall be capable of maintaining the rooms within the range during normal operation. Lower or higher temperature shall be per-mitted when patients' comfort and/or medical conditions require National Institute for Occupational Safety and Health (NIOSH) crite-
- National institute for Occupational stately and relatin (NOSH) critic ria documents' regarding occupational exposure to waste anesthetic gases and vapors and control of occupational exposure to nitrous oxide indicate a need for both local exhaust (seavening) systems and general ventilation of the areas in which the respective gases are used. Refer to NFPA 90¹⁶ for other requirements. If pressure-monitoring device alarms, short-eterm excursions from required pressure relationships shall be allowed while doors are mov-
- ing or temporarily open. Simple visual methods such as smoke trail, ball-in-tube, or flutterstrip shall be permitted for verification of airflow direction.

- bull-in-tube, or fluttestrip shall be permitted for verification of aris, bunnity ranges, and/or air distribution methods, but exceed the minimum indicated ranges.
 Direatment rooms used for bronchoscopy shall be treated as bronchoscopy rooms. Treatment rooms used for bronchoscopy shall be treated as bronchoscopy shall contain provisions for cellusting aventifies to the outdoors instead of exhausting an estification the return air passes through the HEPA filters shall be permitted that the return air passes through the HEPA filters shall be permitted to a use of a for last of a for a for the segarage respiratory distance of a full standard ST29⁻¹¹ for additional information for these spaces. The entire minimum total air infinished of exclusion between the sealing area. Informative Nete: The intent here is to not requiring the volume calculated base to the sealing area chalandor to the sealing area.
 In a real-adulation to include a very large space (e.g., an artium) just because a waiting area opens on to it.

- door air changes are still required. Constant-volume airflow is required for consistent ventilation for the protected environment. The pressure relationship to adjacent areas shall remain unchanged if the protective environment (PE) room is used as a normal patient roo Rooms with reversible airflow provisions for the purpose of switch between protective environment and AII functions shall not be pe
- The AII room described in this standard shall be used for isolatin the airborne spread of infectious discusses, such as measles, varicellä, or tuberculosis. Supplemental recirculating devices using HEPA fil-ters shall be permitted in the AII room to increase the equivalent room air exchanges; however, the minimum outdoor air changes of Table 7.1 are still required. AII rooms that are retrofitted from stan-dard patient rooms from which it is impractical to exhaust directly outdoors may be recirculated with air from the AII room, provided that air first passes through a HEPA filter. When the AII room is not used for airborne infection isolation, the pressure relationship to adjacent areas, when measured with the door closed, shall remain unchanged, and the minimum total air change rate shall be 6 ach. Room temperature ranges that exceed the minimum indicated range shall be permitted if required by the laboratory program or laboratory equipment. the airborne spread of infectious diseases, such as measles, var
- The requirement that all room air is exhausted directly to outdoor

Table 2.1-2 Locations for Nurse Call Devices in Hospitals'

Ontional KEV-Required

Section	Location	Patient Station	Bath Station	Staff Assistance Station	Emergency Call Station		Duty Station	Notes
NURSING UNIT	s	armino m	Printi o II	Dimitoli	CHILD CHILDE	Distort		
2.1-2.2.6	Patient toilet room		•					2
2.2-2.2.2	Medical/surgical unit patient bed	٠		•	•			1, 2, 3, 4
2.2-2.6.2	Critical care unit patient bed	•		•	•			1, 2, 4, 5
2.2-2.8.2	NICU							
2.2-2.9.3	LDR/LDRP room	٠		•	•			1, 2, 3, 4
2.2-2.10.3.1	Newborn nursery			•	•			
2.2-2.10.3.2	Continuing care nursery							
2.5-2.2.2	Psychiatric patient bedroom	•		•				2
2.5-2.4.2	Alzheimer's and other dementia unit	•						
	patient bedroom							
SUPPORT AREA								
2.1-2.8.2	Nurse/control station					•	_	
2.1-2.8.5	Multipurpose room							
2.1-2.8.8	Medication safety zone						•	
2.1-2.8.9	Nourishment area or room							
2.1-2.8.11.2	Clean workroom						•	
2.1-2.8.11.3	Clean supply room							
2.1-2.8.12.2	Soiled workroom						•	
2.1-2.8.12.3	Soiled holding room							
2.1-2.8.13.1	Clean linen storage							
2.1-2.8.13.2	Equipment storage room							
2.1-2.9.1	Staff lounge							
DIAGNOSTIC &	TREATMENT AREAS							
2.1-2.4.3	Seclusion room			•	•			
2.1-3.2	Examination room			•	•			
Table 2.2-2	Class 1 imaging room							
2.1-3.4.3	Pre-procedure patient care room or area	٠		•	•			1, 2
2.1-3.4.4	Phase I post-anesthetic (PACU)			•				
	nationt care station	-		-	•			2, 4
2.1-3.4.5	patient care station Phase II recovery patient care station			•	•			2, 4
2.1-3.4.5 2.2-2.9.11	P			_	-			
2.2-2.9.11	Phase II recovery patient care station Cesarean delivery room Emergency treatment room, triage			•	•			1, 2
	Phase II recovery patient care station Cesarean delivery room	•		•	•			1, 2
2.2-2.9.11 2.2-3.1.3.6 2.2-3.2.2	Phase II recovery patient care station Cesarean delivery room Emergency treatment room, triage area	•		•	•			1, 2
2.2-2.9.11 2.2-3.1.3.6	Phase II recovery patient care station Cesarean delivery room Emergency treatment room, triage area Observation unit patient care station Procedure room (including endoscopy)	•		•	•			1, 2 2 1, 2, 4
2.2-2.9.11 2.2-3.1.3.6 2.2-3.2.2 2.2-3.3.2	Phase II recovery patient care station Cesarean delivery room Emergency treatment room, triage area Observation unit patient care station Procedure room (including	•		•	•			1, 2 2 1, 2, 4
2.2-2.9.11 2.2-3.1.3.6 2.2-3.2.2 2.2-3.3.2 Table 2.2-2	Phase II recovery patient care station Cesarean delivery room Emergency treatment room, triage area Observation unit patient care station Procedure room (including endoscopy) Class 2 imaging room	•		•	•			1, 2 2 1, 2, 4 2, 4
2.2-2.9.11 2.2-3.1.3.6 2.2-3.2.2 2.2-3.3.2 Table 2.2-2 2.2-3.3.3 Table 2.2-2	Phase II recovery patient care station Cesarean delivery room Emergency treatment room, triage area Observation unit patient care station Procedure room (including endoscopy) Class 2 imaging room Operating room Class 3 imaging room Imaging waiting and changing area,	•		•	•			1, 2 2 1, 2, 4 2, 4
2.2-2.9.11 2.2-3.1.3.6 2.2-3.2.2 2.2-3.3.2 Table 2.2-2 2.2-3.3.3	Phase II recovery patient care station Cesarean delivery room Emergency treatment room, triage area Observation unit patient care station Procedure room (including endoscopy) Class 2 imaging room Operating room Class 3 imaging room Imaging waiting and changing area, including toilet room Electroconvulsive therapy (ECT) treatment room	•		•	•			1, 2 2 1, 2, 4 2, 4 2
2.2-2.9.11 2.2-3.1.3.6 2.2-3.2.2 2.2-3.3.2 Table 2.2-2 2.2-3.3.3 Table 2.2-2 2.2-3.4.10	Phase II recovery patient care station Cesarean delivery room Emergency treatment room, triage area Observation unit patient care station Procedure room (including endoscopy) Class 2 imaging room Operating room Class 3 imaging room Imaging waiting and changing area, including toilet room Electroconvulsive therapy (ECT)	•		•	•			1, 2 2 1, 2, 4 2, 4 2 2







Changed from no recommendation to required

Changed from required to no recommendation

Hard copy shows bath stations and staff call instead of staff call and emergency call station in these columns. The electronic version has been updated. 2014 matches more closely with the electronic version.

Table 2.1-3: Locations for Nurse Call Devices in Outpatient Facilities^{1,2}

Section	Location	Patient Station	Staff Assistance Station	Emergenc <u>y</u> Call Station	Notes
PATIENT CARE AND	DIAGNOSTIC AREAS				
2.1-3.2.2 Table 2.1-4	Procedure room (including endoscopy) Class 2 imaging room		•	•	2, 3
2.1-3.2.3 Table 2.1-4	Operating room Class 3 imaging room		•	•	2
2.1-3.7.3 2.1-3.7.5	Pre-procedure patient care station Phase II recovery patient care station	•	•	•	1, 2
2.1-3.7.4	Phase I post-anesthesia recovery (PACU) patient care station	•	٠	•	2, 3
2.8-3.4.2 2.8-6.2.2	Emergency treatment room Emergency triage area	•	٠		1, 2, 3
2.10-3.10.2	Dialysis facility patient toilet room		•	•	1
2.11-3.2.9.2 (2) 2.11-3.2.9.3 (2)	Electroconvulsive therapy (ECT) room ECT recovery patient care station		•	•	2



Nurse Cal

FGI Updates



Added areas

Table 2.1-1 Electrical Receptacles for Patient Care Areas in Hospitals

Section Location		Number of Single Receptacles ¹	Receptacle Locations
PATIENT BED LOO	CATIONS		
2.1-2.4.2	AII room ²	12	2 at each side of the head of the bed
2.2-2.2.2	Medical/surgical unit patient room2		2 on all other walls
2.2-2.2.4.4	Protective environment room ²		1 for a television, if used
2.2-2.5.2	Intermediate care unit patient room		1 for each motorized bed
2.2-2.9.2.2	Postpartum unit patient room ²		
2.2-2.11.2	Pediatric and adolescent unit patient room ²		
2.6-2.2.2	Rehabilitation unit patient room		
2.2-2.6.2	Critical care unit (CCU) patient room	16	Convenient? to head of bed with one on each
2.2-2.7.2	Pediatric critical care unit patient room		wall
2.2-2.8.2	Neonatal intensive care unit (NICU) patient care station		
2.2-2.9.3	LDR/LDRP room	16	8 convenient3 to head of mother's bed
			4 convenient3 to each bassinet with one on
			each wall
2.2-2.10.3.1	Newborn nursery patient care station	4	Convenient' to each bassinet
2.2-2.10.3.2	.2-2.10.3.2 Continuing care nursery patient care station		Convenient ² to head of each bed, crib, or bassinet (At least 50% of these outlets shall be connected to emergency system power and be so labeled.)
2.5-2.2.2	Psychiatric nursing unit	No minimum	
DIAGNOSTIC ANI	D TREATMENT AREAS		
2.1-3.2	Examination room	8	4 convenient3 to head of gurney or bed or on
Table 2.2-2	Class 1 imaging room		each lateral side of the imaging gantry
2,2-2,9,11	Cesarean delivery room	304	16 convenient3 to table placement
			2 on each wall
			6 in the infant care area
2.2-3.1.2.6	Treatment room for basic emergency services	12	Convenient ¹ to head of gurney or bed
2.2-3.1.3.3 Triage room or area in the emergency department		6	Convenient? to head of gurney or bed (At least 50% of these outlets shall be connected to emergency system power and be so labeled.)
2.2-3.1.3.6 (2) and (3)	Emergency department treatment room	12	Convenient ³ to head of gurney or bed
2.2-3.1.3.6 (4)	Trauma/resuscitation emergency room	16	Convenient) to head of gurney or bed
2.2-3.2.2	Observation unit patient care station	8	4 convenients to head of gurney or bed
2.2-3.3.2	Procedure room (including endoscopy)	124	8 convenients to table placement with at least
Table 2.2-2	Class 2 imaging room		one on each wall
2.2-3.3.3	Operating room	364	16 convenient? to table placement
Table 2.2-2	Class 3 imaging room		2 on each wall
2.2-3.10.2	Hemodialysis patient care stations	8	4 on each side of a patient bed or lounge chair. (Two on each side of the bed shall be connected to emergency power.)
POST-ANESTHESI	A CARE LOCATIONS		
2.1-3.4.4	Phase I post-anesthetic care (PACU) patient care station	8	Convenient ³ to head of gurney or bed
2.1-3.4.5	Phase II recovery patient care station	4	Convenient9 to gurney, lounge chair, or bed

Added areas that previously had minimums per NEC. Category 1 and 2 patient bed location minimums are still per NEC.

Section	Room Type	Number of Single Receptacles ¹	Receptacle Locations ²	
PATIENT CARE	AND DIAGNOSTIC AREAS			
2.1-3.2.1 Table 2.1-4	Examination room/observation room Class 1 imaging room	8	4 convenient to head of exam table or gurney or on each lateral side of the imaging gantry	
2.1-3.2.2 Table 2.1-4	Procedure room (including endoscopy) Class 2 imaging room	123	8 convenient to table placement At least 1 on each wall	
2.1-3.2.3 Table 2.1-4	Operating room Class 3 imaging room	363	12 convenient to table placement 2 on each wall	Total count not aligns with NE
2.1-3.7.3 2.1-3.7.5	Pre-procedure patient care station Phase II recovery patient care station	4	Convenient to gurney, lounge chair, or bed	
2.1-3.7.4	Phase I post-anesthesia recovery (PACU) patient care station	8	Convenient to head of gurney or bed	
2.4-2.2	Birthing room	8	4 convenient to head of the mother's bed	
2.8-3.4.2	Treatment room (emergency facility)	12	4 convenient to head of exam table or gurney	
2.8-3.4.4	Trauma/resuscitation room (emergency facility)	16	Convenient to head of gurney or bed	
2.8-6.2.2	Triage area (emergency facility)	6	Convenient to head of gurney or bed (at least 3 outlets connected to emergency system power and so labeled)	
2.10-3.2.2	Hemodialysis patient care station	8	4 on each side of a patient bed or lounge chair (2 on each side of the bed connected to emergency power)	

Split out ED

treatment room

Total count now aligns with NEC



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Table 2.1-3 Station Outlets for Oxygen, Vacuum (Suction), Medical Air, and Instrument Air Systems in Hospitals'

Section	Location	Oxygen	Vacuum	Medical Air	WAGD ²	Instrument	Was requi	red in 2014							
PATIENT CARE UNIT	TS					Air									
2.1-2.4.2	Airborne infection isolation room	1/bed	1/bed	_	—	<u> </u>									
2.2-2.2.2	Patient room (medical/surgical)	1/bed	1/bed		_	- 1	3/bed in 2	014							
2.2-2.2.4.4	Protective environment room	1/bed	1/bed	_	_	- 1									م ادام ا
2.2-2.5.2	Intermediate care room	2/bed	2/bed	1/bed	_	- 1	GENERAL SUPPORT FACILITIES								Added — instrum
2.2-2.6.2	Critical care patient room	3/bed	3/bed	1/bed	_	- 1	2.1-5.1.2.2 (2) Two-room st	erile processing:	_	[_		1	9, 10, 11	require
2.2-2.6.4.2	Airborne infection isolation (critical care)	1					Decontamina							_	roquiro
2.2-2.7.2	Pediatric critical care room	1					2.1-5.1.2.2 (3) Two-room st workroom	erile processing: Clean	-	-	_	-	-	9, 10, 11	
2.2-2.8.2	Neonatal intensive care unit (NICU) infant	3/infant care bed	3/infant care bed	3/infant care bed	- 1	- 1		erile processing:			_	- 1		9,10,11	
	care bed							nination area							
2.2-2.9.2	Antepartum and postpartum unit	1/bed	1/bed	—	—	-	2.1-5.1.2.3 (3) Clean w 2.1-5.7.2.2 Autopsy root			1 mm moderation			_	-	
2.2-2.9.3	Labor/delivery/recovery (LDR)]					1.7	n rocessing room decontamination		1 per workstation	_	<u> </u>	<u> </u>	8, 9, 11	
2.2-2.9.3	Labor/delivery/recovery/postpartum (LDRP)						area	cocessing room decontainination							
2.2-2.9.3.9	Infant resuscitation space4 (LDR/LDRP)	3/bassinet	3/bassinet	3/bassinet	—	-	2.2-3.11.4.3 Endoscope p	rocessing room clean work area	—	—	8	—	-	8, 9, 11	
2.2-2.9.11	Cesarean delivery room	2/room	4/room	1/room	—			-	in a state of the	added					
2.2-2.9.11.1	Infant resuscitation space4 (cesarean delivery)	3/bassinet	3/bassinet	3/bassinet	—			St	pport facilities	added					
2.2-2.9.11.11	Recovery space for cesarean delivery	1/bed	3/bed	1/bed	-										
2.2-2.10.3.1	Newborn nursery	1/bassinets	1/bassinet5	1/bassinets	—	-				Added areas					
2.2-2.10.3.2	Continuing care nursery	1/bassinet	1/bassinet	1/bassinet	—	-		Table 2.1-2: Station		ygen, Vacuum, N	ledical Air	, and In	strume	nt Air Sy	stems in
2.2-2.11.2	Pediatric and adolescent patient room	1/bed	1/bed	1/bed	—	- 1		Outpatient Facilities Section	8	Location		Oversen	Vacuum	Modical Air	Instrument Air
2.2-2.12.2	Psychiatric patient room	—	—	_	—	-		PATIENT CARE AND DIAGN	DSTIC AREAS	Location		Oxygen	vacuum	Medical Air	Instrument Air
2.2-2.12.4.3	Seclusion treatment room (psychiatric	1						2.1-3.2.2	Procedure room			11	11	_	_
	unit[HL1])							Table 2.1-4	Class 2 imaging	room		2	2	11	_
	TREATMENT LOCATIONS							2.1-3.2.3.2 (1)(a)	Operating room	(255-square-foot O	R)	11	11		_
2.1-3.2	Examination room or emergency department	1/room	1/room	-	—	-		2.1-3.2.3.2 (1)(b)-(c)	Operating rooms	2		2	3	11	_
21244	treatment room	21.1.1	21.1.1	11.1.1				Table 2.1-4	Class 3 imaging						
2.1-3.4.4	Phase I post-anesthesia (PACU) patient care station	2/station	3/station	1/station	—	-		2.1-3.3.2	Airborne infectio			03	03	_	_
2.1-3.4.5	Phase II recovery patient care station	1/station	1/station*		_			2.1-3.7.4	•	sthesia recovery (PA	ACU)	1	1	_	—
2.2-3.1.2.6	Treatment room for basic emergency services	1/gurney	1/gurney					21275	patient care station			03	03		
2.2-3.1.3.3	Triage area (emergency department)	1/station	1/station					2.1-3.7.5	Cast room	patient care statio	n	0 ³	03		
2.2-3.1.3.6	Emergency department treatment room or	1/gurney	1/gurney	1/gurney				2.4-2.2	Birthing room			11	11		
2.2-5.1.5.0	area	nguney	i/guiney	1/guilley	_			2.8-3.4.2	÷	(emergency facility	л)	1	1		
2.2-3.1.3.6 (4)	Trauma/resuscitation room	2/gurney	3/gurney	1/gurney	_	- 1		2.8-6.2.2		rgency facility)-p	-	1	1		
	Plaster and cast room	1/room	1/room	_	_	- 1		2.8-3.4.4		tion room (emerge		2	2	1	
2.2-3.2.2	Observation unit patient care station	1/station	1/station		_	- 1			-per gurney		,,,	-	-		
Table 2.2-2	Class 1 imaging room	1/room	1/room	_	_	- 1		2.9-3.2.2	Endoscopy proce	edure room		1	3	_	_
2.2-3.3.2	Procedure room	2/room	2/room	1/room	_	- 1		2.11-3.2.9.2 (2)	Electroconvulsiv	e therapy treatment	t room	11	11	_	_
Table 2.2-2	Class 2 imaging room						Added separate		1						
2.2-3.3.3	Operating room	2/room	5/room	1/room	1/room	1/room 👍	— instrument air	2.1-4.3.2.2 (2)		g decontamination	room		-	—	11,4,5
Table 2.2-2	Class 3 imaging room						requirement	2.1-4.3.2.2 (3)	Sterile processin	g clean workroom					1, 4, 5
2.2-3.11.2	Endoscopy procedure room	1	3	—	_	-		2.1-4.3.2.3	One-room sterile	processing room		_		_	1, 4, 5
2.2-3.11.3	Endoscopy pre- and post-procedure patient	07	07	_	_	- 1		2.9-4.3.2		ssing room-decor	ntamination		3		1, 3, 5
	care area								area	sonig room decor	mation				
2.2-3.13.4	Hyperbaric suite pre-procedure patient care area	2	2	—	—	-		2.9-4.3.3		ssing room—clean	work area		3		1, 3, 5



—	—	—	19,10,11
_		_	9, 10, 11
	-		9,10,11
1 per workstation		-	—
_	Ļ	_	R, 9, 11
	-*	-	8, 9, 11

separate nent air ment

The Future of FGI

- Implementation of FGI in Oklahoma, Texas and Kansas 1.
- 2022 REVISION PROCESS 2.
- 2022 REVISION TOPICS З.

FGI Guidelines Adoption Map

Oklahoma will begin applying the 2018 FGI Guidelines to hospitals and outpatient surgery facilities on September 13th, 2019

KEY

2018	
2014	
2010	
2006	
2001	
1996–97	
Equivalency*	
HVAC only	

*Guidelines may be applied as an equivalency to state rules.





Health Guidelines Revision Committee (HGRC)



"Select multidisciplinary consensus body of about 100 clinicians, administrators, architects, engineers, and representatives from authorities having jurisdiction" (FGI).





Benefit-Cost Committee

"Evaluates the impact of changes to requirements in the *Guidelines*" (FGI)







2022 Public Feedback

- > 1st Opportunity for Feedback Public Proposal Period ended July 1, 2019
 - Feedback to suggestions for changes to the current baseline text and development of materials on operational and best-practice recommendations. November 1, 2018 – June 30, 2019
- > 2nd Opportunity for Feedback on the new language is open to the public following the release of the draft of the 2022 Guidelines in the summer of 2020.



2022 Topic Groups

Based on industry trends and user feedback and inquiries

- Acoustics and vibration
- Behavioral health
- Behavioral health in the ED
- Emergency preparedness, resiliency and business continuity
- > Inclusive environments (formerly, Geriatrics)
- Infection prevention and control



- ➤ Lighting
- Nurse call devices, electrical receptacles and med/gas outlets
- Palliative design
- > Pediatrics
- ➢ Rural health
- > Technology



2022 REVISION TOPICS

Publication of 2022 *Guidelines*

Every 4 Years there is a Revision Cycle of Guidelines

- ➤ The 2022 *Guideline* draft will be released to the public in summer 2020
- ➢ Final publication is estimated for the end of 2021



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The Facility G

2018 edition



Session 1

Contact Information

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