

`<div style="margin: 0 auto; padding: 20px 15px;"><section style="margin-bottom: 40px;">
<h2 style="color: #000; border-bottom: 2px solid #000; padding-bottom: 8px;">Introduction</h2>`

Effective modular cleanroom layout design plays a crucial role in manufacturing efficiency, product quality, and regulatory compliance across industries like pharmaceuticals, microelectronics, and biotechnology. A well-planned modular cleanroom layout design can reduce contamination risks by up to 85% compared to traditional facilities, while offering significant advantages in flexibility, rapid deployment, and scalability.

This comprehensive guide provides a complete framework for modular cleanroom layout design, from initial concept through final implementation. We'll explore the fundamental principles, systematic design steps, and optimization strategies that ensure your modular cleanroom layout design meets both current operational

needs and future expansion requirements.

</section><!-- Part 2: Understanding Modular Cleanroom Basics -->

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<h2 style="color: #000; border-bottom: 2px solid #000; padding-bottom: 8px;">Understanding Modular Cleanroom Basics</h2>

Modular cleanrooms consist of prefabricated components that offer significant advantages over traditional construction. The key elements include insulated wall panels (typically 50mm-100mm thickness), fan filter units (FFUs) with HEPA/ULPA filtration, suspended ceiling grids, anti-static vinyl flooring, and specialized doors/windows with airtight seals.

These standardized components enable rapid assembly and reconfiguration. For example, a typical Deiang TM modular cleanroom can be installed 40-60% faster than conventional

construction. The modular approach allows for precise control over environmental parameters, with temperature maintained at $21 \pm 2^{\circ} \text{C}$, humidity at $45 \pm 5\% \text{ RH}$, and pressure differentials between zones typically at 10-15 Pascals.

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<p style="font-size: 14px; color: #666; margin-top: 8px;">Figure: Key components of a modular cleanroom system</p>

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</section><!-- Part 3: Fundamental Principles of

Cleanroom Layout Design -->

<section style="margin-bottom: 40px;">

<h2 style="color: #000; border-bottom: 2px solid #000; padding-bottom: 8px;">Fundamental Principles of Cleanroom Layout Design</h2>

Successful modular cleanroom layout design requires adherence to several core principles that ensure contamination control and operational efficiency. These principles form the foundation of any effective cleanroom environment regardless of industry application.

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<h3 style="color: #000; margin-top: 0;">3.1 Unidirectional Flow Principle</h3>

Personnel and materials should follow a predetermined path from lower to higher cleanliness areas, preventing cross-contamination. For example, in pharmaceutical applications, personnel typically follow this sequence: entrance

→ gowning area (ISO 8) → airlock → main clean area (ISO 7) → core process area (ISO 5).

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<h3 style="color: #000; margin-top: 0;">3.2 Cleanliness Zoning/Grading</h3>

Areas should be classified according to required cleanliness levels (ISO 5 to ISO 8), with proper isolation between zones. A typical electronics cleanroom might have ISO 7 for assembly areas and ISO 5 for critical processes, separated by airlocks maintaining 15 Pa pressure differential.

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<h3 style="color: #000; margin-top: 0;">3.3

Pressure Differential

Higher cleanliness areas must maintain positive pressure relative to adjacent less-clean areas. The standard pressure cascade is typically: ISO 5 > ISO 6 > ISO 7 > ISO 8 > non-clean area, with minimum pressure differentials of 10-15 Pa between adjacent zones.

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3.4 Maintainability & Cleanability

Design must facilitate easy maintenance and cleaning. This includes providing adequate space for filter replacement (minimum 600mm clearance above ceiling), smooth and non-porous surfaces, and rounded corners (minimum 25mm radius) to prevent particle accumulation.

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<h3 style="color: #000; margin-top: 0;">3.5 Scalability & Flexibility</h3>
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Modular cleanrooms should accommodate future expansion or reconfiguration. Deiiang™ systems typically allow for 20-30% expansion without structural modifications, with panel connections designed for easy disassembly and reassembly.

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<h3 style="color: #000; margin-top: 0;">3.6 Safety & Emergency</h3>
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Layout must incorporate safety features including emergency exits, eyewash stations, fire detection systems, and clear evacuation routes. Emergency exits should be located within 30m of any workstation and open outward without special

knowledge or effort.

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<p style="font-size: 14px; color: #666; margin-top: 8px;">Figure: Unidirectional personnel flow from "dirty" to "clean" areas</p>

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</section><!-- Part 4: Key Steps in Modular Cleanroom Layout Design -->

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Key Steps in Modular Cleanroom Layout Design</h2>

The systematic approach to modular cleanroom layout design ensures all critical factors are considered from the beginning. Following these steps methodically reduces the risk of costly redesigns and ensures regulatory compliance while optimizing operational efficiency.

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4.1 Phase 1: Requirement Analysis & Goal Setting</h3>

Thorough analysis of operational requirements forms the foundation of successful modular cleanroom layout design. Key considerations include:

<ul style="padding-left: 20px;">

Application Pharmaceutical	Purpose: (GMP compliance),
---	--

microelectronics (particle control), biotechnology (microbial control)

Cleanliness

Requirements: ISO Class 5-8 based on process sensitivity

Process Flow: Sequential operations, batch vs continuous processes

Equipment

Specifications: Dimensions, heat load, utility requirements

Personnel Requirements:

Number of operators, gowning protocols

Material Flow: Raw

material entry, in-process handling, finished goods exit

Space Constraints:

Building dimensions, ceiling height, column locations

Budget & Timeline:

Capital investment, installation schedule

Future Expansion: 2-5

year growth projections

For example, a pharmaceutical filling line might require ISO 5 for critical zones, ISO 7 for background, and specific temperature control at 21 ± 1 ° C with $45 \pm 5\%$ RH.

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<h3 style="color: #000; margin-top: 0;">4.2 Phase 2: Preliminary Conceptual Layout</h3>

Based on requirement analysis, create initial sketches dividing the space into functional zones. A typical 100m² cleanroom might allocate:

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15% for entrance/gowning area (15m²)

10% for material airlock (10m²)

60% for main process area (60m²)

10% for equipment/service area (10m²)

5% for exit/waste handling (5m²)

Initial FFU placement follows the rule of thumb: ISO 5 requires 25-40% ceiling coverage, ISO 7 requires 15-25%, and ISO 8 requires 5-15%.

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<p style="font-size: 14px; color: #666; margin-top: 8px;">Figure: Preliminary conceptual layout with functional zones</p>

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<h3 style="color: #000; margin-top: 0;">4.3 Phase

3: Detailed Layout Design</h3>

This phase transforms the conceptual layout into a detailed, operational design addressing all critical aspects:

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Detailed planning of personnel movement: entrance → locker area → pre-gowning → gowning room → airlock → clean corridor → workstations → exit. Each transition includes appropriate contamination control measures like sticky mats, air showers, or hand sanitation stations.

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Planning of material movement: receiving →

quarantine → decartoning area → pass-through
→ process area → packaging → finished
goods. Critical calculation: pass-through
dimensions = largest component + 150mm
clearance (minimum 600 × 600mm).

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<h4 style="color: #000;">Equipment
Placement</h4>

Precise positioning of all equipment with adequate
clearances: operational space (minimum 900mm),
maintenance access (minimum 600mm), and
safety zones. Utility connections planned with
10-15% spare capacity for future modifications.

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<h4 style="color: #000;">Cleanliness Zoning
& Pressure Control</h4>

Defining cleanroom classifications and pressure cascades: ISO 8 (15Pa) → ISO 7 (25Pa) → ISO 5 (40Pa). Airlocks sized for 2-3 personnel (minimum 2m²) with interlocking doors and pressure monitoring.

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<h4 style="color: #000;">Airflow Pattern Design</h4>

Determining airflow direction, velocity, and distribution. Unidirectional flow for ISO 5 (0.45m/s ± 20%), non-unidirectional for ISO 6-8 (20-40 air changes/hour). Return air grilles strategically placed low on walls for optimal particle removal.

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<p style="font-size: 14px; color: #666; margin-top: 8px;">Figure: Detailed layout with personnel flow (blue) and material flow (green)</p>

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<h3 style="color: #000; margin-top: 0;">4.4 Phase 4: 3D Modeling & Visualization</h3>

Using CAD/BIM software (AutoCAD, Revit, SolidWorks) to create precise 3D models that identify spatial conflicts, optimize ergonomics, and validate the design. Deiliang™ typically uses BIM for clash detection, reducing construction issues by 70-80%.

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<h3 style="color: #000; margin-top: 0;">4.5 Phase 5: Review, Revision & Final Approval</h3>
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Comprehensive review with all stakeholders including production, quality control, maintenance, and safety personnel. Typical review checklist includes:

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<ul style="padding-left: 20px;">
 Regulatory compliance verification
 Operational workflow efficiency
 Maintenance accessibility
 Safety and emergency protocols
 Utility distribution adequacy
 Future expansion capability

```

Following review, incorporate feedback and obtain formal approval before proceeding to fabrication and installation.

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</section><!-- Part 5: Specific Factors Influencing
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## Layout Design -->

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<h2 style="color: #000; border-bottom: 2px solid #000; padding-bottom: 8px;">Specific Factors Influencing Layout Design</h2>

Several industry-specific and operational factors significantly impact modular cleanroom layout design decisions. Understanding these variables ensures the final design meets both technical and regulatory requirements.

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<h4 style="color: #000; margin-top: 0;">5.1 Process Type & Particle Generation</h4>

Wet processes (pharmaceutical compounding) require chemical-resistant surfaces and drainage, while dry processes (electronics assembly) need enhanced ESD protection. High

particle-generating operations need increased air changes (up to 60 ACH for ISO 5).

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<h4 style="color: #000; margin-top: 0;">5.2  
Product Sensitivity</h4>

Products sensitive to temperature (  $\pm 0.5^{\circ}\text{C}$ ), humidity (  $\pm 3\%$  RH), or electrostatic discharge require specialized environmental controls and material selection. Pharmaceutical products may need  $21 \pm 1^{\circ}\text{C}$ , while electronics might need  $\leq 40\%$  RH for ESD control.

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#### 5.3 Equipment Maintenance Space

Adequate clearance around equipment: front access (minimum 900mm), rear access (minimum 600mm), overhead clearance for filter replacement (minimum 600mm). Large equipment may require removable wall panels for installation.

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#### 5.4 Regulatory & Standard Compliance

Industry-specific requirements: cGMP for pharmaceuticals, ISO 14644 for classification, FDA 21 CFR Part 11 for electronic records, and SEMI standards for microelectronics.

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<h4 style="color: #000; margin-top: 0;">5.5  
Energy Efficiency & Environmental  
Control</h4>

Optimizing layout to reduce energy consumption:  
variable air volume systems, heat recovery,  
efficient equipment placement. Proper design can  
reduce

<a href="https://www.deiiang.com/product-category/cleanroom-hvac/">HVAC</a> energy use by  
20-30% compared to conventional approaches.

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&nbsp;

[caption id="attachment\_4318" align="alignnone"

width="960"] Process type analysis for determining required cleanliness levels[/caption]

<p style="font-size: 14px; color: #666; margin-top: 8px;">Figure: Process type analysis for determining required cleanliness levels</p>

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</section><!-- Part 6: Common Modular Cleanroom Layout Configurations -->

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<h2 style="color: #000; border-bottom: 2px solid #000; padding-bottom: 8px;">Common Modular Cleanroom Layout Configurations</h2>

Several standard layout configurations have

proven effective for different applications and space constraints. Each offers distinct advantages for specific operational requirements.

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<h4 style="color: #000; margin-top: 0;">6.1
Linear/Straight-line Layout</h4>
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Process flow follows a straight path from raw material to finished product. Ideal for sequential operations with minimal backtracking.

**Advantages:** Simple workflow, easy supervision, minimal cross-traffic

**Disadvantages:** Requires longer space, limited flexibility

**Applications:** Assembly lines, packaging operations

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<h4 style="color: #000; margin-top: 0;">6.2  
U-shaped/L-shaped Layout</h4>

Process flow follows a U or L pattern, suitable for space-constrained areas or processes requiring material return.

<strong>Advantages:</strong> Efficient use of space, compact footprint

<strong>Disadvantages:</strong> Potential congestion at corners, complex material handling

<strong>Applications:</strong> Laboratories, research facilities, small-scale production

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<h4 style="color: #000; margin-top: 0;">6.3  
Central Island Layout</h4>

Core process area in center with support areas  
surrounding it. Ideal for processes requiring  
multiple support functions.

<strong>Advantages:</strong>                      Excellent  
supervision, efficient material distribution

<strong>Disadvantages:</strong>                      Higher  
construction cost, potential airflow challenges

<strong>Applications:</strong>                      Biotechnology,  
pharmaceutical compounding, critical processes

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<th style="padding: 12px; text-align: left; border:
1px solid #ddd;">Space Efficiency</th>
<th style="padding: 12px; text-align: left; border:
1px solid #ddd;">Workflow Efficiency</th>
<th style="padding: 12px; text-align: left; border:
1px solid #ddd;">Expansion Potential</th>
<th style="padding: 12px; text-align: left; border:
1px solid #ddd;">Cost Impact</th>
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</thead>
<tbody>
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<td style="padding: 12px; border: 1px solid
#ddd;">Linear/Straight-line</td>
<td style="padding: 12px; border: 1px solid
#ddd;">Medium</td>
<td style="padding: 12px; border: 1px solid
#ddd;">High</td>
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<td style="padding: 12px; border: 1px solid
#ddd;">Low</td>
<td style="padding: 12px; border: 1px solid
#ddd;">Low</td>
</tr>
<tr style="background-color: #f8f9fa;">
<td style="padding: 12px; border: 1px solid
#ddd;">U-shaped/L-shaped</td>
<td style="padding: 12px; border: 1px solid
#ddd;">High</td>
<td style="padding: 12px; border: 1px solid
#ddd;">Medium</td>
<td style="padding: 12px; border: 1px solid
#ddd;">Medium</td>
<td style="padding: 12px; border: 1px solid
#ddd;">Medium</td>
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<tr>
<td style="padding: 12px; border: 1px solid
#ddd;">Central Island</td>
<td style="padding: 12px; border: 1px solid
#ddd;">Low</td>

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<td style="padding: 12px; border: 1px solid #ddd;">High</td>
<td style="padding: 12px; border: 1px solid #ddd;">High</td>
<td style="padding: 12px; border: 1px solid #ddd;">High</td>
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</tr>
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</table>
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<p style="font-size: 14px; color: #666; text-align: center; margin-top: 8px;">Comparison of different modular cleanroom layout configurations</p>
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</section><!-- Part 7: Design Tools & Best Practices -->
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<h2 style="color: #000; border-bottom: 2px solid #000; padding-bottom: 8px;">Design Tools & Best Practices</h2>
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Utilizing appropriate design tools and following

industry best practices significantly improves the quality and efficiency of modular cleanroom layout design projects.

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<h3 style="color: #000; margin-top: 0;">7.1
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Common Design Software</h3>
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<ul style="padding-left: 20px;">
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 AutoCAD: 2D drafting and detailed drawings
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 Revit (BIM): 3D modeling with building information
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```
 SolidWorks: Equipment modeling and integration
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 Specialized Cleanroom Software: Computational fluid dynamics for airflow simulation
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Deiiang™ typically uses BIM for complex projects, reducing design conflicts by up to 80% compared to traditional 2D approaches.

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<h3 style="color: #000; margin-top: 0;">7.2 Best Practices</h3>

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<li><strong>Early Multidisciplinary Integration:</strong> Involve production, quality, maintenance, and safety teams from concept stage</li>

<li><strong>Lifecycle Cost Consideration:</strong> Evaluate operational costs (energy, maintenance) alongside capital investment</li>

<li><strong>Airflow Simulation:</strong> Model particle distribution and airflow patterns before construction</li>

<li><strong>Detailed SOP Development:</strong> Create comprehensive standard operating procedures for all operations</li>

<li><strong>Continuous Validation:</strong>

Implement ongoing monitoring and periodic requalification</li>

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<h3 style="color: #000; margin-top: 0;">7.3  
Common Mistakes to Avoid</h3>

<ul style="padding-left: 20px;">

<li>Ignoring personnel and material flow intersections</li>

<li>Inadequate clearances for maintenance and filter replacement</li>

<li>Insufficient consideration of future expansion needs</li>

<li>Over-specifying or under-specifying cleanliness requirements</li>

<li>Poor placement of utilities and service connections</li>

<li>Inadequate emergency and safety provisions</li>

</ul>

According to industry data collected by Jason.peng, approximately 65% of cleanroom design issues stem from inadequate upfront planning and failure to involve all stakeholders early in the process.

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</section><!-- Part 8: FAQ -->

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<h2 style="color: #000; border-bottom: 2px solid #000; padding-bottom: 8px;">Frequently Asked Questions</h2>

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<h4 style="color: #000; margin-top: 0;">Can modular cleanroom layouts be easily modified?</h4>

Yes, one of the key advantages of modular cleanrooms is their reconfigurability. Deiiang™ systems typically allow for layout changes with minimal disruption, often completed in 30-50% of

the time required for traditional construction modifications. Wall panels can be repositioned, and additional modules can be integrated as needed.

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<h4 style="color: #000; margin-top: 0;">How is isolation maintained between different cleanliness zones?</h4>

Isolation is achieved through a combination of physical barriers (walls, doors), pressure differentials (typically 10-15 Pa between adjacent zones), and procedural controls (gowning protocols, material transfer procedures). Airlocks with interlocking doors prevent simultaneous opening between zones of different classifications.

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#### How to design a cleanroom layout for small spaces?

For limited spaces, consider vertical utilization with mezzanines, compact equipment layouts, and multi-functional areas. U-shaped or L-shaped configurations often work well. Optimize every square meter by carefully analyzing workflow and eliminating non-value-added movements. Deiiang<sup>TM</sup> has successfully implemented [ISO 5 cleanrooms](https://www.cleanroomequips.com/ISO-Class5-CleanRoom/) in spaces as small as 12m<sup>2</sup>.

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#### How is fire safety addressed in cleanroom design?

Fire safety incorporates early detection systems, appropriate fire-rated materials (typically 1-hour rating for walls), emergency lighting, clearly

marked exits, and automatic fire suppression systems. Cleanroom-specific considerations include compatibility of fire suppression agents with sensitive equipment and processes.

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<h4 style="color: #000; margin-top: 0;">How to determine the number and placement of FFUs?</h4>

<a href="https://www.deiiang.com/product-category/ffu/">FFU</a> quantity is calculated based on room volume, required air changes per hour, and filter efficiency. For ISO 5, typical coverage is 25-40% of ceiling area. Placement considers equipment layout, workflow, and return air locations. CFD analysis can optimize distribution for uniform airflow and particle control.

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</section><!-- Part 9: Conclusion & Call to Action  
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<h2 style="color: #000; margin-top: 0;">Conclusion</h2>

Effective modular cleanroom layout design requires careful consideration of multiple factors including workflow efficiency, contamination control, regulatory compliance, and future flexibility. By following the systematic approach outlined in this guide - from thorough requirement analysis through detailed design and validation - organizations can create cleanroom environments that support operational excellence while accommodating future growth and changes.

The modular approach offers significant advantages in speed of deployment, adaptability, and lifecycle cost, making it an increasingly

popular choice across industries requiring controlled environments. Proper planning and professional design consultation are essential investments that pay dividends throughout the cleanroom's operational life.

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<h3 style="color: #000;">Ready to Start Your Modular Cleanroom Project?</h3>

Contact our cleanroom specialists today for a free consultation and customized layout design.

<div style="display: flex; flex-wrap: wrap; justify-content: center; gap: 15px; margin: 10px 0;"><a style="display: inline-block; background-color: #000; color: white; padding: 12px 25px; text-decoration: none; border-radius: 4px; font-weight: bold;" href="#">Request Free Consultation</a>

<a style="display: inline-block; background-color: white; color: #000; padding: 12px 25px; text-decoration: none; border-radius: 4px; border: 2px solid #000; font-weight: bold;" href="#">Download Design Guide</a></div>

<p style="font-size: 14px; color: #666;">Design  
Specialist: Jason.peng | Brand: Deiiang™</p>

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