Safe Building Practices for Laboratories and Pharmaceutical Facilities

What is the problem?

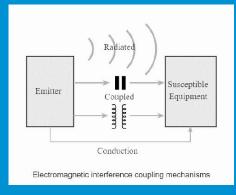
- Fire, water, gas, dust, EMI, sound, and pests can pose major threats.

What can happen?

- If the seals of the building, room or equipment are breached, there could be severe consequences, such as the loss of life or company assets.

SEALING IN ADVANCED FACILITIES

- Fire protection: 2 4 hours
- Equipment Protection: IP 55 69k; NEMA 4X
- Acoustics and Vibrations: STC Maintenance
- Hazardous/Explosive Areas: Zones 0 1 2
- Electromagnetic Interference & Grounding



CLEANROOMS:

Pressurized locations require effective sealing to prevent contaminants from entering or exiting.

> **CRITICAL AREAS:** ·Laboratory walls ·Glove boxes



Cleanroom and Contained Room Requirements

- Fire Protection
- Watertight Protection
- Gas/smoke Protection
- Acoustic Sealing
- Operating across a wide range of pressures
- Good material reaction to common fumigants
- Tamper resistance service retention
- Follows GMP Guidelines

Good Manufacturing Practices

ISO 14644: Specifies the classification of air cleanliness in terms of concentration of airborne particles in cleanrooms and clean zones

- For GMP compliance, all surfaces in a cleanroom should be "smooth and impervious" and:
- not generate their own contamination (does not create dust, or peel, flake, corrode or provide a place for microorganisms to proliferate)
- are easy to clean (all surfaces are easily accessible, there should not be any ledges or recesses)



Building envelope
 Mechanical and electrical rooms

 Interior equipment

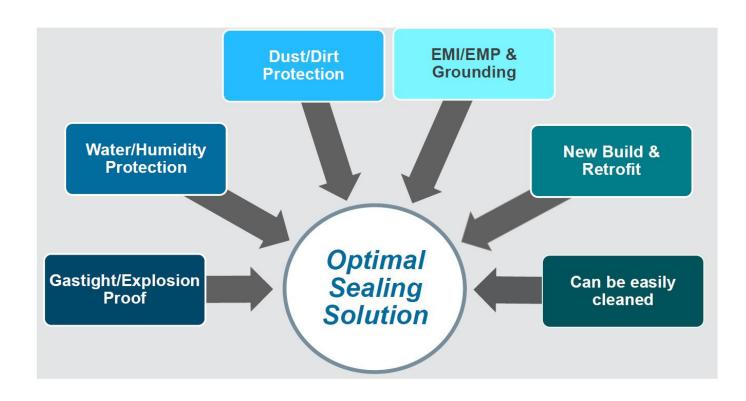


Fumigant Exposure Compliance Airtightness Report:

Formaldehyde VHP (Vaporized Hydrogen Peroxide) Chlorine Dioxide Acetate

Separate Sectors Secto	
Airtightness in Belgian high biocontainment facilities	
Overview of four BSL3 facilities and recommendations for optimal airtightne	155
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Does your current sealing method meet your needs?



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