





Within the first 24-48 hours there are probably just wet materials. After 2-3 days, there are probably wet, moldy materials.



These options are listed in the general order of preference.



Repairs are typically not an effective method for removing mold from the indoor environment, for addressing the source/s of water intrusion, nor for preventing occupant exposure.



A 1 to 10 dilution of household bleach (never use straight bleach) has been reported to kill about 95% of viable (living) mold in actual field trials. However, the remaining 5% of viable mold left on the contaminated surface is still a lot of mold, and it can start growing again as soon as the repair is completed.



Restoration dries the wet materials and will probably stop any further mold growth but leaves any mold that has grown on the wet materials in-place. This may not necessarily be an issue, but mold-sensitive individuals should be aware of this potential issue.

Room	Kitchen	Hall Bath	
Cavity	Range Base	Vanity Base	
Asp/Pen	67,749	70,233	
Total (Sp/m <sup>3</sup> )	68,257	100,268	
~~~	Dryin a pat expo	ng holes can prov thway for occupai sure	

Background concentrations of indoor airborne Asp/Pen spores are typically several hundred spores/m<sup>3</sup>. Here we have concentrations of about 70,000. In addition, the drying holes that were made to allow the spaces below the cabinets to dry also allow the contaminant mold spores to enter the occupied space.





## **Remediation Containments**

**Configuration and Criteria for Assessing Airborne Spores** 







### Dehumidifier

#### • Dehumidifier

- Removes water vapor, not spores
- Dessicant Units
- Condensing Units
  - Warm, wet environment
  - Can become contaminated and a potential source of airborne spores inside containment

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An isolated AFU tends to re-clean the same localized mass of air repeatedly, doing a poor job of cleaning the air in the containment. Placing two AFU in containment with ducts in opposing corners of containment for example, creates a circulation pattern that minimizes short-circuiting.



Phase 1 is a pre-inspection by the Contractor to make sure the Containment is ready for inspection by a third-party IEP. It is also intended to pre-clear any cavities that may be sealed prior to the Phase 2 inspection by an IEP. Phase 2 would typically be performed by an IEP.



ATP testing is a rapid and inexpensive method for assessing the condition of surfaces. A table or graph of ATP values for general conditions can be established for reference.



The Contractor can then determine RLU acceptance criteria based their instrument.



Collecting an airborne sample inside containment is obvious. Collecting a sample immediately outside containment is not as obvious, but three reasons for doing so are given.





This is an opinion at this point in the presentation. The objective is to present a logical argument to support this opinion.



The discussions of the following three example containments are intended to support these opinions, as well as illustrate how criteria can be established.



These are common containment configurations that illustrate how acceptance criteria can be modified to accommodate varying conditions. With a little thought, criteria can be established for any configuration or set of conditions.



Spores	Outside Contain	Percent	Spores	Inside Contain	Percent
Clad	133	47 %	Clad	45	39 %
Asp/Pen	57	20 %	Asp/Pen	31	27 %
Altern	34	12 %	Altern	17	15 %
Epicoc	35	13 %	Epicoc	7	6%
Curvul	24	8 %	Curvul	2	2 %

Compare the sample inside containment to the sample outside containment (two samples in each location is even better). First, are the rank orders for Asp/Pen similar? Yes. Second, was Asp/Pen as a percentage of total spores about the same in both samples? Yes, similar. Conclusion: condition in containment was Acceptable.

Outside Containment	PERCENT	Inside Containment	PERCENT
<b>Cladosporium</b>	50%	Cladosporium	40%
Alternaria	25%	Asp/Pen	30%
Epicoccum	20%	Alternaria	20%
Asp/Pen	5%	Epicoccum	10%
<i>Asp/Pen</i> spores w containment, but Not an acceptabl	vere 5% of tot t 30% inside o le result	al spores in the ro f containment	om outside of

Compare the sample inside containment to the sample outside containment. First, are the rank orders for Asp/Pen similar? No. 4<sup>th</sup> in the outside sample and 2<sup>nd</sup> in the inside sample, indicating a possible source in containment. Second, was Asp/Pen as a percentage of total spores about the same in both samples? No, 5% outside and 30% inside, again indicating a possible source inside containment. Conclusion: condition in containment was Not Acceptable.



If Asp/Pen was present in the make-up air, then it should be at a similar or lower concentration in containment.





An 8 ft by 10 ft chamber with an 8-ft ceiling height. Settling time of 300 minutes or 5 hours. Starting concentrations were 100% and decreased to just a little more than 0.1% of the initial concentration during that time. All the spore types essentially "completely" settled during that time period in an undisturbed, quiescent environment.

# Criteria 2

- Containment sealed & "quiescent" for some time
  - No negative pressure
  - Quiescent conditions, so shorter settling time
    - Five hours or longer prior to sampling
- Expect "low" total spore counts with "nominal" concentrations of *Asp/Pen* detected



This configuration is isolated from the adjacent room, so rank order analysis or indoor/outdoor comparisons are not appropriate. The "criterion" has to come from within the containment or from prior experience.



This is the distribution of Asp/Pen concentrations from sampling about 90 mold remediation containments while in scrub mode. The median concentration was 70 spores/m<sup>3</sup> and the average concentration was 200 spores/m<sup>3</sup>, with 300 spores/m<sup>3</sup> one standard deviation above the median concentration. Based on these 90 remediations, which were performed by different Contractors, the Expected Range of Asp/Pen spore concentrations would be 300 spores/m<sup>3</sup> or less.



78% of 109 Stachybotrys samples were nondetects, 12% were 10 spores/m<sup>3</sup>, and the remaining 10% of the samples ranged from 30-80 spores/m<sup>3</sup>. The expected concentration of Stachybotrys was Nondetect.

# Criteria 3 • Containment sealed and AFU in scrub mode • No negative pressure • Aggressive conditions • Expect "moderate" total spore counts with concentrations of *Asp/Pen* in an "expected range" - 200 Sp/m³ to 300 Sp/m³ or less

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#### Post-Remediation Acceptance Criteria for Airborne Mold Spores

- We may not want to think in terms of a single "clearance criterion" for airborne mold samples
  - The "clearance criterion" may depend on the configuration of the containment
- The "clearance criterion" can, and should, be adjusted to accommodate the
  - Configuration of the containment
  - Aggressive or quiescent conditions in the containment

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#### Containment Configurations and Criteria for Assessing Airborne Samples



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