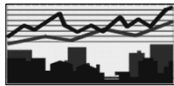


Using Surface Dust Samples to Assess Occupant Exposures to Mold

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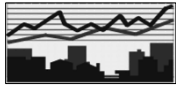
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Topics

- **Collecting Dust Samples**
- **The ERMI Method**
- **Analyzing Dust Samples and Reporting Results**
- **Assessing Occupant Exposure Potentials**
- **HERTSMI (Briefly)**

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Collecting the Sample

A Critical Step

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Inspection Objectives

- **HOW WILL THE SAMPLES SUPPORT THE OBJECTIVE?**
- Possible Objectives
 - Assessing Building-Related Contamination
 - Assessing Occupant Exposure Potential
 - Or Both

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The Sampling “Rationale”

- **HOW WILL THIS SAMPLE SUPPORT THE OBJECTIVE?**
- 1st: Why am I collecting this sample, in this location, using this sampler?
- 2nd: What do I expect to learn from the sample result?
 - What will it mean if the result is positive? If negative?
- 3rd: How should the sample be analyzed to support the objective?
- 4th: Which interpretation method should I use to support the objective?

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Assessing HVAC Samples

- **Air Return Filter**
 - Central collection point for airborne contaminants
 - Composite of what has been circulating in the air
 - Indicator of chronic occupant exposure *since changed*
- **Air Supply Grill**
 - Distribution point for contaminants in HVAC system
 - Contaminants collected from the grill/boot were probably reaching the breathing zones of the occupants

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Different Methods for Different Objectives

- | BUILDING
CONTAMINATION | OCCUPANT
EXPOSURE |
|--|---|
| • Airborne: 5-minute SIC samples, Total spore counts | • Airborne: 60-90 minute filter cassette samples with qPCR analysis |
| • HVAC: Inspect the cooling coils, return plenum | • HVAC: Sample the air supply grill |
| | • Dust: Air return filter |
| | • qPCR analysis |

The Sample Describes the Environment

- **Koppen:** “One of the primary challenges of assessing exposures (to environmental contaminants) is the collection of representative samples, with the sampling step typically contributing the largest variability.”

[Koppen, R., et. al.; Determination of mycotoxins in foods: current state of analytical methods and limitations; Applied Microbiology and Biotechnology (May, 2010)]

- A sample that is not representative of the fungal loading in an indoor space will not be representative of the occupant exposure potential in that space

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Sampling and Interpretation Methods

- Should consider meeting the “SOCS” criteria
 - **Significance** of the sample result
 - Numerical guidelines for interpreting the lab report
 - **Objective** decision criteria for interpreting the sample result (High, Low, Average)
 - Independent of training, experience, or bias
 - **Consistent** assessments of condition
 - across projects and between inspectors
 - **Stable** basis for assessment
 - Does not vary with local conditions

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“Professional” Sampling Methods



Problems exist even with “standard” methods

My opinion:
Only one of these four sampling methods can meet the “SOCS” criteria and actually describe the exposure potential of a carpet dust sample

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Occupant-Collected Samples

Goal: A sample representative of the indoor environment and occupant exposure potential

Sampling Method	Swiffer, combined locations (Bsmt+LR)
Sample Size	One sample collected on one day
Sample Matrix	Mixed surface types (Typical)

- Collecting a **representative** sample is actually not a simple task.
- Occupants may need as much guidance as possible when being asked to collect samples.
- Requires a knowledgeable and informed mentor and guide.

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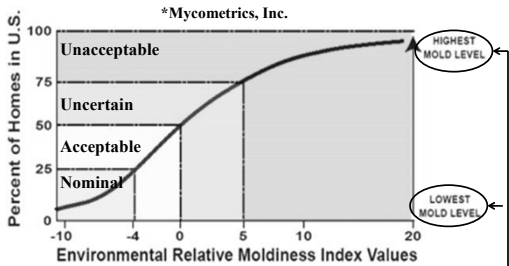
The ERM Method

- Unless the collection method looked like the photo, and it was collected from a carpet, it was not an ERM sample, and should not be assessed using an ERM score.



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ERMI Score v Fungal Loading



Implies the ERM Score is a measure of total fungal loading, but it is a difference (Gr 1 – Gr 2), not a total

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ERMI Score v Fungal Loading

Fungal ID \ Unit	Spore E./mg	Spore E./mg
Aspergillus flavus/oryzae	ND	ND
Aspergillus fumigatus	ND	1
Aspergillus niger	ND	2
Aspergillus ochraceus	5	9
Aspergillus penicilloides	4	730
Aspergillus restrictus*	ND	ND
Aspergillus sclerotiorum	ND	ND
Aspergillus sydowii	ND	<1
Aspergillus umbratus	ND	8
Aspergillus versicolor	ND	550
Aureobasidium pullulans	690	395
Chaetomium globosum	ND	ND
Cladosporium sphaerosperrum	7	26
Eurotium (Asp.) amstelredamum*	1	150
Phaeoannaea variabilis	ND	ND
Penicillium brevicompactum	ND	170
Penicillium corylophilum	ND	74
Penicillium crustosum*	ND	28
Penicillium parvum/penum	ND	ND
Penicillium spinulosum*	ND	1
Penicillium variable	ND	3
Scopulariopsis brevicaulis/fusca	ND	ND
Scopulariopsis chartarum	ND	4
Stachybotrys chartarum	ND	140
Trichoderma viride*	ND	<1
Wallemia zeevi	ND	460
Sum of Logs (Group 1)	4.98	25.36

ERMI Score not related to fungal loading

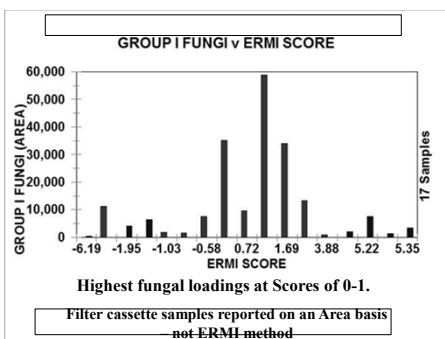
GR 1	GR 2	ERMI
100,000	1,000	+ 1
100,000	10,000	+ 1

ERMI can actually decrease as the fungal loading increases

GR 1	GR 2	ERMI
100,000	10,000	+ 1
100,000	50,000	+ 0.3

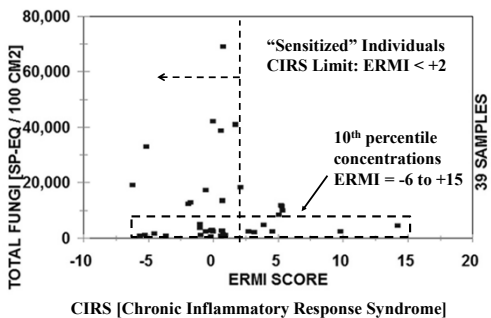
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Group 1 Fungi v ERMi Scores



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CARPET DUST SAMPLES ERMi: TOTAL FUNGI v ERMi SCORE

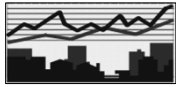


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Characteristics of ERMi Scores

- ERMi scores only apply to carpet dust sampled using a specific sampling method
 - Not to mixed surfaces sampled using a Swiffer
- Results are reported on a Weight-Analyzed Basis rather than an Area Basis
 - Sample results cannot be compared
- Scores are calculated based on the difference between Group 1 and Group 2 fungi
 - Scores do not reflect total fungal loading (OEP)
- A score does not represent a unique fungal loading for assessing OEP

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Analyzing Dust Samples and Reporting the Results

You Have Options

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Reporting Surface Dust Samples

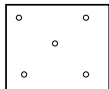
- How the sample results are reported by the laboratory affects the ability to assess BRC and OEP
- HOW SAMPLE RESULTS CAN BE REPORTED
 - Standard; Weight-Analyzed Basis: Sp-Eq/mg
 - Sp-Eq/mg of dust analyzed
 - Option 1; Total Weight Basis: Sp-Eq/mg x (total mg)
 - Sp-Eq/Sample
 - Option 2; Area Basis: [Sp-Eq/mg x (total mg)] / Area
 - Sp-Eq/in² Sp-Eq/100 cm²

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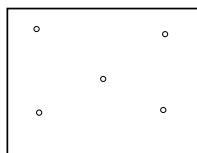
Carpet Dust Example

Failed Clearance: Selecting the Reporting Basis

Since the mold is in the dust and not in the fibers, results on a “weight” basis can be inaccurate & misleading



AREA = 100
MG of dust

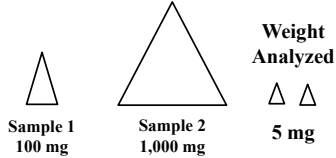


- Set Clearance Criteria (Mold/mg)
- Carpet cleaned
- 1st Clearance Failed
- Re-cleaned carpet
- 2nd Clearance Failed
- Problem
 - Larger area of carpet
 - But the same Mold/mg

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Standard Lab Procedure Weight-Analyzed Basis

Lab SOP: Not weight the sample, Analyze 5 mg of sample



1,000 Sp-Eq detected in both 5-mg samples
 200 Sp-Eq/mg reported for both 5-mg samples
 If Condition \approx fungal loading: Then 20,000 v 200,000 Sp-Eq
 Inspector: Same Exposure Potential reported for both samples

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Area Basis

1. Inspector measures the area that was sampled
2. Requests the result on a Total-Weight Basis as Sp-Eq/Sample
3. Divide Sp-Eq/Sample by the area sampled

Sample results are standardized & can be compared
 - by accounting for sample weights & areas sampled

Lab Report	Sample 1	Sample 2
Area Sampled	6" x 6"	12" x 12"
Weight Collected	400 milligrams	1,200 milligrams
Weight Analyzed	5 mg	5 mg
Fungi Detected	1,000 sp-eq/5 mg	1,000 sp-eq/5 mg
Sp-Eq/mg	200 sp-eq/mg	200 sp-eq/mg
Sp-Eq/Sample	80,000 sp-eq/Sample	240,000 sp-eq/Sample
REPORT	320,000 sp-eq/ft ²	240,000 sp-eq/ft ²

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Area Basis

- **1st: Health effects were better associated with results reported on an Area Basis rather than Weight Basis**

- Ulrike Gehring, et al. $\beta(1\rightarrow3)$ -Glucan in House Dust of German Homes: Housing Characteristics, Occupant Behavior, and Relations with Endotoxins, Allergens, and Molds; Environmental Health Perspectives; 109: (2), February 2001
- Paul J. Liroy, et al. Dust: A Metric for Use in Residential and Building Exposure Assessment and Source Characterization; Environmental Health Perspectives; 110: (10), October 2002

2nd: Results reported on an area basis allow Conditions and/or Occupant Exposure Potentials to be compared

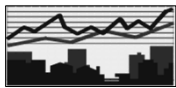
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Objective of Presentation

- Now that I've "rained on your parade", what now?
- What follows is knowledge
- You can use it to
 - collect more representative samples
 - Assist occupants as much as possible in characterizing their environments



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Assessing Occupant Exposure Potential based on total fungal loading

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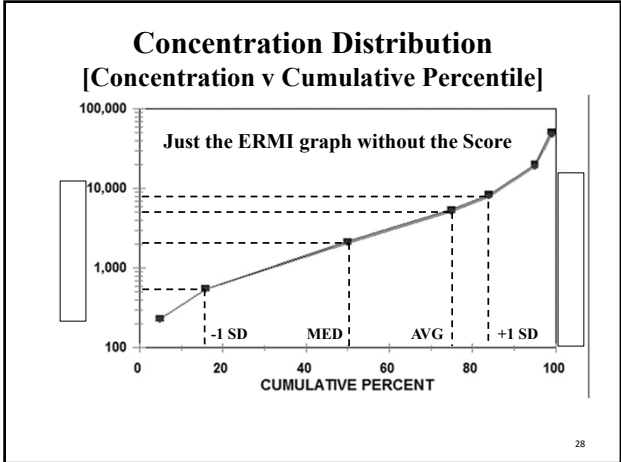
The Sampling Methods

- The data presented were collected using filter cassettes and surface swabs, so the data were not "ERMI" data
- Results were reported on an Area Basis rather than on a Weight-Analyzed Basis



Just consider the implications of the concepts

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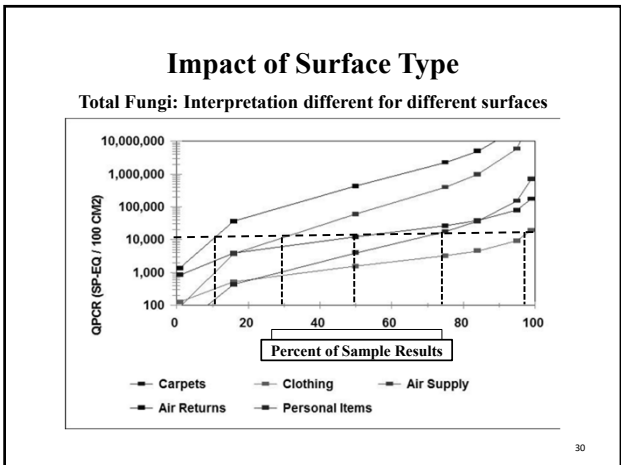
Interpreting Percentile Concentrations

Percentile Concentration	Significance
84 th %-tile	+1 "Std" Dev (Med)
73 rd %-tile	[Average Concentration]
50 th %-tile	Median Concentration
16 th %-tile	-1 "Std" Dev (Med)

Assessing BRC: Use the Median (50th) Percentile

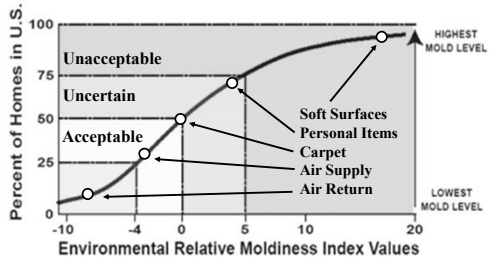
Assessing OEP: Use the Average (75th) Percentile
DOSE = [Average Conc X Time]

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Sample Collection and Assessment

qPCR: 10,000 sp-eq/100 cm²

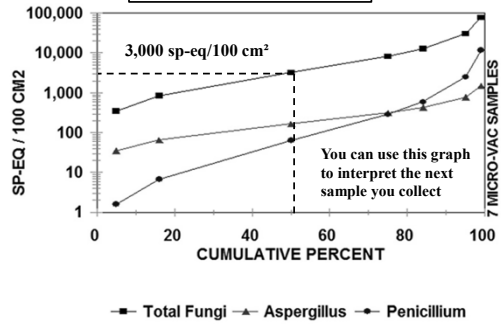


Conclusion: The surface sampled should be considered when assessing the sample result

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SOFT SURFACES

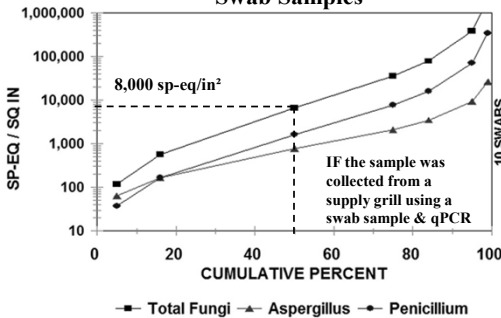
Micro-vacuum Cassette Samples



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AIR SUPPLY GRILLS

Swab Samples



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The HERTSMI-2 Method

- “Health Effects Roster of Type Specific Formers of Mycotoxins and Inflammagens”
- Derivative of the 36 ERMI fungi
- The concentration ranges of five ERMI fungi are used to calculate the HERTSMI score

<i>Aspergillus Penicillioides</i>
<i>Aspergillus versicolor</i>
<i>Chaetomium globosum</i>
<i>Stachybotrys chartarum</i>
<i>Wallemia sebi</i>

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HERTSMI: Percent of Total Fungi

Prevalence of HERTSMI-2 Fungi by Surface Type

93 Samples	<i>A pen</i>	<i>A ver</i>	<i>Chaet</i>	<i>Stachy</i>	<i>Wall</i>	SUM
Carpets	0.4%	0.4%	1.4%	0.6%	0.2%	3.0 %
Soft Surfaces	0.6%				0.5%	1.1 %
Clothing	0.7%				0.9%	1.6 %
Hard Surfaces		0.2%		0.1%		0.3 %
Air Return		0.2%			0.1%	0.3 %
Air Supply			0.1%			0.1 %

1. HERTSMI scores can be based on less than 3% of the fungal loading, typically on 1% or less.
2. The sampling locations can obviously influence the results.

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