

Blacktrail Environmental, Inc.

ENVIRONMENTAL LIABILITY MANAGEMENT

August 5, 2022

Mr. Rolf Eggers
Cape Boats
2170 Wallingford Loop
Mount Dora, Florida 32757

RE: Mold Assessment and Air Sampling for Toxic Mold Concentrations

The Capital Gallery, 109 North 4th Street, Bismarck, North Dakota
Blacktrail Project No. MLD2243

Dear Mr. Eggers,

Concern about indoor exposure to mold has been increasing as the public becomes aware that exposure to mold can cause a variety of health effects and symptoms, including allergic reactions. Molds are part of the natural environment. Outdoors, molds play a part in nature by breaking down dead organic matter such as fallen leaves and dead trees, but indoors, mold growth should be avoided. Molds reproduce by means of tiny spores; the spores are invisible to the naked eye and float through outdoor and indoor air. Mold may begin growing indoors when spores land on surfaces that are wet. There are many types of mold, and none of them will grow without water or moisture. Molds are usually not a problem indoors, unless mold spores land on wet or damp spots and begin growing. Molds have the potential to cause health problems. Molds produce allergens (substances that can cause allergic reactions), irritants, and in some cases, potentially toxic substances (mycotoxins).

A wide variety of symptoms have been attributed to the toxic effects of fungi. Inhaling or touching mold or mold spores may cause allergic reactions in sensitive individuals. Allergic responses include hay fever-type symptoms, such as sneezing, runny nose, red eyes, and skin rash (dermatitis). According to documents published by the U.S. Environmental Protection Agency (USEPA), allergic reactions to mold are common (approximately 1 in 5 individuals). They can be immediate or delayed. Some of the symptoms related to fungal exposures are non-specific, such as discomfort, inability to concentrate, and fatigue. Severe illnesses such as Organic Dust Toxic Syndrome (ODTS), Hypersensitivity Pneumonitis (HP), and pulmonary hemosiderosis have also been attributed to fungal exposures. Molds can also cause asthma attacks in people with asthma who are allergic to mold. In addition, mold exposure can irritate the eyes, skin, nose, throat, and lungs of both mold-allergic and non-allergic people. Symptoms other than the allergic and irritant types are not commonly reported as a result of inhaling mold.

Symptoms of ODTS are describe as the abrupt onset of fever, flu-like symptoms, and respiratory symptoms in the hours following a exposure to dust containing organic material including fungi. It differs from HP in that it is not an immune-mediated disease and does not require repeated exposures to the same causative agent. ODTS may be caused by a variety of biological agents including common species of fungi (e.g., species of *Aspergillus* and *Penicillium*). ODTS has been documented in farm workers handling contaminated material but is also of concern to workers performing renovation work on building materials contaminated with fungi.

Some studies have suggested an association between *Stachybotrys* and pulmonary hemorrhage/hemosiderosis in infants, generally those less than six months old. Pulmonary hemosiderosis is an uncommon condition that results from bleeding in the lungs. The cause of this condition is unknown, but may result from a combination of environmental contaminants and conditions (e.g., smoking, fungal contaminants and other bioaerosols, and water-damaged homes), and currently its association with *Stachybotrys* is unproven, but not dismissed (USEPA).

Temperature and Relative Humidity Measurements

Air sampling for concentrations of airborne toxic molds was performed in the commercial space on August 1, 2022. Temperature and relative humidity measurements were recorded to assist in determining conditions that may be favorable for mold growth. A digital MannixTM digital sling psychrometer/thermo-hygrometer Model Sam 990 DW was used to measure temperature and relative humidity. The temperature and relative humidity sensors were calibrated prior to the sampling event.

The measured temperature in the commercial space at the time of the sampling was 72°F. The American Society of Heating, Refrigerating and Air Conditioning Engineers, Inc. (ASHRAE) recommends an acceptable temperature range during the heating season between 69°F and 74.5°F, and between 72°F and 78°F during the cooling (air-conditioning) season. Based on ASHRAE recommendations, the temperature inside of the commercial space at the time of the assessment was within the acceptable range.

The measured relative humidity level in the commercial space was 36%. ASHRAE recommends an acceptable relative humidity range below 70% to prevent bacteria, yeast, and mold growth from occurring in air ducts, heat exchangers and on other surfaces, but suggests that relative humidity levels be kept between 30% and 50% for optimum prevention of mold growth and individual comfort. Commercial buildings with central air conditioning generally range from 20% relative humidity in the winter season to 50% in the summer season. The measured reading was below 70% and within the ASHRAE recommendations for prevention of mold growth; therefore, *there is not* a potential for molds (or bacteria or yeast) to flourish and sustain indoor atmosphere concentration levels that would cause adverse health effects *without a direct source of water*.

Air Sampling Results

Air sampling for the project included collection of three air samples inside the commercial space; an air sample was not collected from outside the subject building for comparison purposes (the “control sample”) because the subject area was a commercial space with no windows that open to allow outside air to enter the sampling area(s). The air sampling was conducted with Air-O-Cell[®] cassettes based on the recommendations of the manufacturer, Zefon InternationalTM. Air sampling for mold with Air-O-Cell[®] cassettes is a widely accepted practice among the industrial hygiene community and is recognized by the American Industrial Hygiene Association (AIHA). Air sampling with Air-O-Cell[®] cassettes identifies and quantifies

(with normal statistical parameters) concentrations of airborne mold spores. The Air-O-Cell[®] cassette consists of a treated microscope slide that is contained within a cassette. The slide is treated with a sticky substance onto which mold spores and allergen particulate will adhere upon impact onto its surface. The cassette is then analyzed with the use of a Phase Contrast Microscope (PCM) for the identification of the materials on the treated microscope slide. During the analysis, the number of countable structures per spore/particle type is documented and those numbers are entered along with the scope field of view, number of traverses of the trace (if different than 6) and the air volume.

The analysis of the Air-O-Cell[®] cassettes was performed by EMSL Analytical, Inc. laboratory (EMSL) in Indianapolis, Indiana. The laboratory is an AIHA accredited environmental microbiology laboratory. The analyses of the samples was supervised and reviewed by Nathan Husted, a mycologist with experience in identifying indoor air molds and bacteria. The analyses of the air samples were performed to identify which contaminants in the commercial space are of a possible health concern. If *Stachybotrys* spores were observed on any of the sampling mediums, a second analyst verified its presence.

The air samples were analyzed for different types of common mold (fungi) spores that are of a potential concern because of their toxicity, commonly found in the environment, and routinely cause adverse health effects or irritation to select individuals. The samples were also analyzed for skin and insect fragments, fibrous particulate, and pollen. No areas of concern were identified from the analyses of the air samples for skin and insect fragments, fibrous particulate, and pollen. The toxic mold spores of potential concern are listed in Table 1. Results of the individual types of mold spores identified in the air samples are present in an attachment to this letter report. Guidelines for relative exposures to mold are presented in Table 2.

Table 1 - Mold Spores of Potential Concern		
<i>Alternaria (Ulocladium)</i> <i>Ascospores</i> <i>Aspergillus/Penicillium</i> <i>Basidiospores</i> <i>Bipolaris</i> <i>Chaetomium</i>	<i>Cladosporium</i> <i>Curvularia</i> <i>Epicoccum</i> <i>Fusarium</i> <i>Ganoderma</i> <i>Myxomycete</i>	<i>Pithomyces</i> <i>Rust</i> <i>Scopulariopsis</i> <i>Stachybotrys</i> <i>Zygomycetes</i> <i>Unidentifiable Spores</i>

Although there are no indoor air quality standards for interpreting mold spore counts, the National Allergy Bureau has set some guidelines based on ecological measurements for outdoor air. The National Allergy Bureau is a section of the American Academy of Allergy, Asthma and Immunology (AAAAI) Aeroallergen Network that is responsible for reporting current pollen and mold spore levels to the media. As health effects are dependent on individual susceptibility, the relative exposure index is not based on health effects. They are relative numbers and limits.

Table 2 National Allergy Bureau Guideline for Relative Exposures to Outdoor Mold Spores (Spores Per Cubic Meter of Air [Spores/m³])				
Very Low	Low	Medium	High	Very High
<500	500-1,000	1,000-5,000	5,000-20,000	>20,000

In North Dakota, generally normal outside air counts during summer conditions range from 3,000 to 50,000 total toxic mold spores per cubic meter of air (spores/m³), and winter conditions range from 50 to 5,000 spores/m³. *Preferred* air counts for indoors (statistically 95% of “clean homes and commercial buildings”) should be below 2,000 spores/m³. Additionally, airborne counts of *Stachybotrys* (the “black mold”) should be kept totally absent, but at least less than or equal to 20 spores/m³. The following table presents the results of the laboratory analyses of the air samples collected for toxic mold concentrations.

Sample Number	Sample Location	<i>Aspergillus/</i> <i>Penicillium</i> (Spores/m ³)	<i>Stachybotrys</i> (Spores/m ³)	Total Result (Spores/m ³)
MO	Marci's Office	0	0	140
G-O	Between Gallery and (Gift) Shop	80	0	240
BACK	Back Storage Area	0	0	320

The lab analyses results for the commercial space indicate acceptable levels well below the guideline of 2,000 spores/m³ (i.e. no current concern to occupants). The concentration level of *Stachybotrys* (the “black mold”) spores was within the guidelines for a “clean” commercial space (equal to or less than 20 spores/m³).

Aspergillus/Penicillium is an “indicator” of mold inside walls and ceilings, under cabinets and carpets because they germinate, grow, and multiply quickly. Very low concentrations of *Aspergillus/Penicillium* molds associated with a “clean” commercial space were identified in the air samples. A concentration of *Aspergillus/Penicillium* molds above 1,000 spores/m³ usually indicates a significant area(s) of hidden mold in a subject building (i.e. a “mold problem”).

Stachybotrys (the “black mold”) spores take 8 to 10 days to germinate while remaining moist or wet. No *Stachybotrys* spores were identified in the air samples collected inside the commercial space. Based on the air sampling results it can be surmised that no *Stachybotrys* mold is currently flourishing inside the commercial space.

Conclusions

The total concentration of airborne toxic mold spores in commercial space is within the statistical range of a “clean” commercial space and is acceptable for all individuals, even for infants and individuals with respiratory ailments such as asthma and chronic obstructive pulmonary lung disease (COPD), etc. In addition, a TramexTM moisture meter did not identify elevated levels of moisture in the building material(s) throughout the commercial space in the building.

To additionally assess the air quality conditions of the commercial space the assessment included measurements of carbon monoxide (CO), oxygen (O₂), and hydrogen sulfide (H₂S). A MSA™ multi-gas meter was used to measure these parameters in the commercial space. The CO, O₂, and H₂S sensors were calibrated in accordance with manufacturer's instructions prior to the assessment. No level of carbon monoxide (0 ppm) or hydrogen sulfide (0 ppm) was found in the commercial space, and oxygen was identified to be at the standard concentration (20.8%).

References

Information in the following references was considered for this project:

- *Mold Remediation in Schools and Commercial Buildings*, March 2001, U.S. Environmental Protection Agency.
- *Bioaerosols: Assessment and Controls*, 1999, American Conference of Governmental Industrial Hygienists.
- *Field Guide for the Determination of Biological Contaminants in Environmental Samples*, 1996, American Industrial Hygiene Association.
- *Guidelines on Assessment and Remediation of Fungi in Indoor Environments*, New York City Department of Health, 2000.
- *Fungal Contamination in Buildings: A Guide to Recognition and Management*, 1995, Health Canada.
- *Ventilation for Acceptable Indoor Air Quality*, ANSI/ASHRAE standard 62-2001.
- *Identifying Filamentous Fungi*, 1996, Guy St-Germain and Richard Summerbell.
- *Standard and Reference Guide for Professional Mold Remediation*, December 2005, Institute of Assessment, Cleaning and Restoration (IICRC).
- *Indoor Air Quality Sampling Methodologies*, 2002, Kathleen Hess-Kosa.
- *Indoor Air Quality Handbook*, 2000, John D. Spengler, Jonathan M. Samet, and John F. McCarthy.
- *The Mold Help Organization*, 1255 Broadway St. NE Ste. #410, Salem, Oregon.
- *Fundamentals of Industrial Hygiene* (Third Edition), 1988, National Safety Council.

Limitations

Currently, there are no federal standards for treatment of fungal contamination. This report has been prepared to assist the client only. Blacktrail provided services consistent with the level and skill ordinarily exercised by members of the profession currently practicing under similar conditions, and with similar budget constraints. This statement is in lieu of other statements either expressed or implied. This report is intended for the sole use of client. The intent of the project is to aid the client in locating fungi growth (mold) and identify concentrations of mold that may be impacting the occupants. This report is not intended to serve as a bidding document nor as a project specification document and actual site conditions and quantities should be field verified. The scope of services performed in execution of this evaluation may not be appropriate

to satisfy the needs of other users, and use or re-use of this document, the findings, conclusions, or recommendations is at the risk of said user. Although every attempt has been made to identify suspect fungal (mold) growth in the areas identified, the assessment techniques used are inherently limited in the sense that only full demolition procedures will reveal all building materials of a structure and therefore all areas of potential fungal growth. The size of the area impacted by fungal impact is based on professional judgment and practicality. Additionally, other possible building material hazards such as asbestos and lead-based paint were not included as part of this evaluation and may require proper sampling for identification prior to disturbance. Other unidentified microbiological impact may be located within walls, ceiling cavities, below flooring or grade, and other non-accessible areas. Precaution should be used during any remediation activities (if any).

Additionally, the passage of time may result in a change in the environmental characteristics of the subject building. This report does not warrant against future operations or conditions that could affect the recommendations made. The results, findings, conclusions, and recommendations expressed in this report are based only on conditions that were observed during Blacktrail's assessment of the site.

Please call me at 701-527-0274, if you have questions or would like additional information.

Sincerely,



John Spilman, CSP, CHMM
Sr. Industrial Hygienist

Attachments: Laboratory Results and Chain-of-Custody
Certifications of the Sampling Professional



EMSL Analytical, Inc.

6340 CastlePlace Dr. Indianapolis, IN 46250

Tel/Fax: (317) 803-2997 / (317) 803-3047

<http://www.EMSL.com / indianapolislabs@emsl.com>

EMSL Order: 162217506

Customer ID: BTRL34

Customer PO:

Project ID:

Attention: John Spilman

Blacktrail Environmental, inc.

1112 Southport Loop

Bismarck, ND 58504

Phone: (701) 527-0274

Fax:

Collected Date: 08/01/2022

Received Date: 08/02/2022 09:57 AM

Analyzed Date: 08/04/2022

Project: THE CAPITAL GALLERY, 109 NORTH 4TH STREET, BISMARCK ND 58501

Test Report: Air-O-Cell(™) Analysis of Fungal Spores & Particulates by Optical Microscopy (Methods MICRO-SOP-201, ASTM D7391)

Lab Sample Number:	162217506-0001			162217506-0002			162217506-0003		
Client Sample ID:	MO			G-O			BACK		
Volume (L):	150			150			150		
Sample Location:	MARCI OFFICE			BETWEEN GALLERY AND SHOP			BACK STORAGE AREAS		
Spore Types	Raw Count	Count/M ³	% of Total	Raw Count	Count/M ³	% of Total	Raw Count	Count/M ³	% of Total
Alternaria (Ulocladium)	1	20	14.3	-	-	-	2	40	12.5
Ascospores	-	-	-	2	40	16.7	2	40	12.5
Aspergillus/Penicillium	-	-	-	4	80	33.3	-	-	-
Basidiospores	2	40	28.6	4	80	33.3	-	-	-
Bipolaris++	-	-	-	-	-	-	-	-	-
Chaetomium++	-	-	-	-	-	-	-	-	-
Cladosporium	4	80	57.1	1	20	8.3	9	200	62.5
Curvularia	-	-	-	-	-	-	-	-	-
Epicoccum	-	-	-	-	-	-	-	-	-
Fusarium++	-	-	-	-	-	-	-	-	-
Ganoderma	-	-	-	1	20	8.3	-	-	-
Myxomycetes++	-	-	-	-	-	-	1	20	6.3
Pithomyces++	-	-	-	-	-	-	1	20	6.3
Rust	-	-	-	-	-	-	-	-	-
Scopulariopsis/Microascus	-	-	-	-	-	-	-	-	-
Stachybotrys/Memnoniella	-	-	-	-	-	-	-	-	-
Unidentifiable Spores	-	-	-	-	-	-	-	-	-
Zygomycetes	-	-	-	-	-	-	-	-	-
Total Fungi	7	140	100	12	240	100	15	320	100
Hyphal Fragment	1	20	-	1	20	-	-	-	-
Insect Fragment	-	-	-	-	-	-	-	-	-
Pollen	-	-	-	-	-	-	-	-	-
Analyt. Sensitivity 600x	-	21	-	-	21	-	-	21	-
Analyt. Sensitivity 300x	-	7*	-	-	7*	-	-	7*	-
Skin Fragments (1-4)	-	1	-	-	1	-	-	1	-
Fibrous Particulate (1-4)	-	1	-	-	1	-	-	1	-
Background (1-5)	-	1	-	-	1	-	-	2	-

++ Includes other spores with similar morphology; see EMSL's fungal glossary for each specific category.

Kennedie Stansifer, Laboratory Manager
or other Approved Signatory

No discernable field blank was submitted with this group of samples.

EMSL maintains liability limited to cost of analysis. Interpretation and use of test results are the responsibility of the client. This report relates only to the samples reported above, and may not be reproduced, except in full, without written approval by EMSL. EMSL bears no responsibility for sample collection activities or analytical method limitations. The report reflects the samples as received. Results are generated from the field sampling data (sampling volumes and areas, locations, etc.) provided by the client on the Chain of Custody. Samples are within quality control criteria and met method specifications unless otherwise noted. High levels of background particulate can obscure spores and other particulates, leading to underestimation. Background levels of 5 indicate an overloading of background particulates, prohibiting accurate detection and quantification. Present = Spores detected on overloaded samples. Results are not blank corrected unless otherwise noted. The detection limit is equal to one fungal spore, structure, pollen, fiber particle or insect fragment. "" Denotes particles found at 300X. "-" Denotes not detected. Due to method stopping rules, raw counts in excess of 100 are extrapolated based on the percentage analyzed. Skin & Fibrous ratings: 1 (1-25%), 2 (26-50%), 3 (51-75%), 4 (76-100%) of the background particles.

Samples analyzed by EMSL Analytical, Inc. Indianapolis, IN AIHA-LAP, LLC-EMLAP Accredited #157245

Initial report from: 08/04/2022 01:05 PM

For information on the fungi listed in this report, please visit the Resources section at www.emsl.com

EMSL ANALYTICAL, INC.
LABORATORY • PRODUCTS • TRAINING162217506
EMSL Order Number (Lab Use Only):EMSL ANALYTICAL, INC.
6340 CASTLEPLACE DRIVE
INDIANAPOLIS, IN 46250
317-803-2997
317-803-3047

Company : Blacktrail Environmental, Inc.				EMSL-Bill to: <input checked="" type="checkbox"/> Same <input type="checkbox"/> Different If Bill to is Different please note in Comments**			
Street: 1112 Southport Loop				Third Party Billing requires written authorization from third party			
City: Bismarck		State/Province: ND		Zip/Postal Code: 58504		Country: USA	
Report To (Name): John Spilman				Fax #: NA			
Telephone #: 701-527-0274				E-mail Address: jspilman@blacktrail.co ("co" not ".com")			
Project Name/ Number: The Capital Gallery, 109 North 4 th Street, Bismarck ND 58501							
Please Provide Results: <input type="checkbox"/> Fax <input checked="" type="checkbox"/> E-mail		PO#		State Samples Taken: ND			
Turnaround Time (TAT) Options* - Please Check							
<input type="checkbox"/> 3 Hour <input type="checkbox"/> 6 Hour <input type="checkbox"/> 24 Hour <input checked="" type="checkbox"/> 48 Hour <input type="checkbox"/> 72 Hour <input type="checkbox"/> 96 Hour <input type="checkbox"/> 1 Week <input type="checkbox"/> 2 Week							
*Analysis completed in accordance with EMSL's Terms and Conditions located in the Analytical Price Guide. TATs are subject to methodology requirements							
Non Culturable Air Samples (Spore Traps)							
<ul style="list-style-type: none"> • M001 Air-O-Cell • M049 BioSIS • M030 Micro 5 		<ul style="list-style-type: none"> • M173 Allegro M2 • M003 Burkard • M174 MoldSnap 		<ul style="list-style-type: none"> • M004 Allergenco • M043 Cyclex • M176 Rella Smart 		<ul style="list-style-type: none"> • M032 Allergenco-D • M002 Cyclex-d • M130 Via-Cell 	
						<ul style="list-style-type: none"> • M172 Versa Trap 	
Other Microbiology Test Codes							
<ul style="list-style-type: none"> • M041 Fungal Direct Examination • M005 Viable Fungi ID and Count • M006 Viable Fungi ID and Count (Speciation) • M007 Culturable Fungi • M008 Culturable Fungi (Speciation) • M009 Gram Stain Culturable Bacteria • M010 Bacterial Count and ID – 3 Most Prominent • M011 Bacterial Count and ID – 5 Most Prominent • M013 Sewage Contamination in Buildings 		<ul style="list-style-type: none"> • M014 Endotoxin Analysis • M015 Heterotrophic Plate Count • M180 Real Time Q-PCR-ERMI 36 Panel • M018 Total Coliform (Membrane Filtration) • M020 Fecal <i>Streptococcus</i> (Membrane Filtration) • M210-215 <i>Legionella</i> Detection • M026 Recreational Water Screen • M027 Mycotoxin Analysis 		<ul style="list-style-type: none"> • M029 <i>Enterococci</i> • M019 Fecal Coliform • M133 MRSA Analysis • M028 <i>Cryptococcus neoformans</i> Detection • M120 <i>Histoplasma capsulatum</i> Detection • M033-39 Allergen Testing • M044 Group Allergen (Cat, Dog, Cockroach, Dustmites) • Other See Analytical Price Guide 			
Preservation Method (Water): None							
Name of Sampler: John Spilman				Signature of Sampler:			
Sample #	Sample Location	Sample Type	Test Code	Volume/Area	Temperature (°F)	Humidity	Date/Time Collected
MO	Marci Office	Air O Cell	M001	150 Liters	72	36	8/1/22
G-O	Between Gallery and Shop	↓	↓	↓	72	36	↓
BACK	Back Storage Areas	↓	↓	↓	72	36	↓
Client Sample # (s):		Total # of Samples:		3			
Relinquished (Client): John Spilman		Date: 8/1/22		Time: 5:55 pm			
Received (Client):		Date: 8/2/22		Time: 9:57 AM FAX			
Comments:							



THIS CERTIFIES THAT

JOHN S. SPILMAN

HAS SUCCESSFULLY MET ALL THE REQUIREMENTS OF EDUCATION, EXPERIENCE AND EXAMINATION, AND IS HEREBY DESIGNATED A

**CERTIFIED HAZARDOUS MATERIALS MANAGER®
CHMM®**



August 1, 1997
DATE OF CERTIFICATION

08452
CREDENTIAL NUMBER

August 31, 2023
CERTIFICATION EXPIRES

M. Patricia Bulley
DIRECTOR OF CERTIFICATION AND ACCREDITATION

VALID SO LONG AS THIS CREDENTIAL IS RENEWED ACCORDING TO SCHEDULE AND IS NOT OTHERWISE REVOKED.



Accredited by the American National Standards Institute and the Council of Engineering and Scientific Specialty Boards



BOARD OF CERTIFIED SAFETY PROFESSIONALS

AFFIRMS THAT

John S. Spilman

HAVING MADE APPLICATION FOR AND GIVEN SATISFACTORY EVIDENCE OF QUALIFICATION AS REQUIRED IN THE BY-LAWS; IS QUALIFIED TO RECEIVE AND IS HEREBY AUTHORIZED TO USE THE DESIGNATION

CERTIFIED SAFETY PROFESSIONAL
IN
Comprehensive Practice

SO LONG AS THIS CERTIFICATE OF QUALIFICATION IS RENEWED ANNUALLY AND NOT REVOKED

BOARD OF EXAMINERS IN WITNESS WHEREOF
WE HAVE HEREUNTO SET OUR HANDS AND
AFFIXED THE SEAL OF THE BOARD THIS
1ST DAY OF AUGUST, 1995

PRESIDENT

SECRETARY

NUMBER: 13474



Indoor Air Quality Project
sponsored by the U.S. Environmental Protection Agency
through the
UNIVERSITY OF MINNESOTA



Certificate of Course Completion

This certifies that

John Spilman

was enrolled in and completed the
Indoor Air Quality Orientation for Public Officials

June 8-10, 1993
Bismarck, North Dakota

Course Sponsors:
U.S. Environmental Protection Agency
University of Minnesota
American Lung Association of North Dakota
North Dakota Department of Health
and Consolidated Laboratories
Minnesota Building Research Center/Minnesota Extension Service

William J. Angell
William J. Angell
Director, IAQP
University of Minnesota

UNIVERSITY
OF MINNESOTA
**School of
Public Health**
Centers for Public Health
Education and Outreach

Midwest Center for Occupational Health and Safety
2221 University Avenue SE, Suite 350
Minneapolis, MN 55414
612.626.4515 cphed@uwm.edu

This certifies that

John Spilman

attended this continuing education course

Mold Identification

January 11, 2005 - January 13, 2005

Midwest Center for Occupational Health and Safety - 1.9 CEU's or 19 60-minute contact hours.
American Board of Industrial Hygiene for 3.0 Certification Maintenance Points. CM Approval # 04-2330.
Please see reverse side for additional accreditation.

Debra K. Olson

Debra K. Olson
Associate Dean for Public Health Practice Education
Retain this certificate for your records.

