

MOLD, BACTERIA AND BIRD FECES STUDY

US Army Corps of Engineers South Pacific Division Laboratory 25 Liberty Ship Way Sausalito, California 94965

December 1, 2004

Project No.: U04-2732



MOLD, BACTERIA AND BIRD FECES INVESTIGATION REPORT

US ARMY CORPS OF ENGINEERS SOUTH PACIFIC DIVISION LABORATORY 25 LIBERTY SHIP WAY SAUSALITO, CALIFORNIA 94965

PROJECT NO.: U04-2732

DECEMBER 1, 2004

Prepared for:

Veterans Administration Hospital San Francisco, California

Prepared by:

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TABLE OF CONTENTS

1.0 EXECUTIVE	E SUMMARY	2
2.0 INTRODUC	TION	5
3.0 MOLD INVI	ESTIGATION	6
4.0 BACTERIA	& BIRD FECES INVESTIGATION	6
5.0 CONCLUSIO	ONS AND RECOMMENDATIONS	8
6.0 OPINIONS	OF COST	11
7.0 LIMITATIO	NS	14
APPENDICES		
Appendix A:	Sampling Methods	
Appendix B:	Laboratory Results & Chain of Custody	
Appendix C:	Sample Location Drawings	
Appendix D:	Photographic Documentation	
Appendix E.	Credentials	
Appendix F:	Microbiological Species Information	

1.0 EXECUTIVE SUMMARY

On October 26, 2004, an indoor air quality (IAQ) investigation was conducted by Advanced Solutions Group (ASG) at the United States Army Corps of Engineers (USACOE) South Pacific Division Laboratory located at 25 Liberty Ship Way in Sausalito, Marin County, California. The purpose of the investigation was to evaluate mold, bacteria, and bird feces, which have previously been visually identified within the building.

The subject building is a two-story unoccupied building, which was constructed in the late-1940s. The building has been vacant since 1997 and contains a total of approximately 37,800 square feet. Previous episodes of water intrusion were observed in various locations inside the building. In addition, numerous pigeons and other fowl, have been occupying the building.

Our visual and laboratory findings found the following:

- Mold/Fungi Stachybotrys was found on the interior air samples where the visible black growth was observed to be most concentrated and visual evidence of significant water damage was apparent. This species, as well as Aspergillus/Penicillium and Fusarium, which were also found, has been known to produce mycotoxins. Also, relatively high concentrations Yeast were found on the water damaged ceiling tiles and wood floors. In addition, Stachybotrys and Aspergillus, which may also produce mycotoxins, were found on the surfaces in the southwest areas on levels 1 and 2, where the heaviest visual growth was observed.
- Bacteria Various bacteria species were identified from sampling of the bird droppings and these species are believed to be opportunistic pathogens, which may cause illnesses. In addition, *Brevibacterium* was found on the concrete floor below the bird droppings. This species is believed to be an opportunistic pathogen, which may also cause illnesses.
- Bird Feces Excessive bird droppings were observed on the concrete floor, namely on level 1 north side and central area. Also, dead birds were seen in these areas.

Based upon these findings, we recommend the following:

- Repair the roof and possible plumbing leaks, including broken windows, namely at the southwestern side of the building.
- Replace the building's HVAC system and disinfect the duct work and grills/vents.
- Remove all carpeting and disinfect the flooring below if concrete.

- Remove all porous surfaces including water damaged wooden floors, sheet rock walls and ceiling, and ceiling tiles.
- Remove all bird droppings and dead bird from the building. Disinfect the concrete surfaces
 where bird droppings were located. Seal the building as to not allow further intrusion of
 fowl.
- Disinfect all surfaces and building products remaining in the building.
- Retest the air and building surfaces, once these items have been conducted.

Opinions of Cost

Based upon the mold, fungi, bacteria, and bird droppings identified at the site, we are providing these opinions of cost:

Removal and disinfection \$ 90,000
 Industrial hygiene monitoring \$ 18,875
 Total Estimate \$108,875

2.0 INTRODUCTION

On October 26, 2004, Mr. Terry Bleckner, Certified Industrial Hygienist (CIH), conducted an indoor air quality (IAQ) investigation at the USACOE South Pacific Division Laboratory located at 25 Liberty Ship Way in Sausalito, Marin County, California. The purpose of the investigation was to evaluate water damaged building materials, observe general conditions as they relate to fungal growth, identify possible bacteria, and determine the extent and health risks associated with the bird feces located in the building.

The subject building is a two-story unoccupied building, which was constructed in the late-1940s. The building was originally occupied by the US Army Corps of Engineers South Pacific Division Laboratory. The building has been vacant since 1997 and contains a total of approximately 37,800 square feet. Visual evidence of water damage was observed in the following locations:

- Level 1 drafting west wall and the engineers office at the southwest corner
- Level 2 –south hallway by men's restroom, south office area (room 220), west offices by special studies, the central area by chemical analysis room (Room 205), and the general area of the southwest corner offices

In addition, the majority of the north side, level 1 concrete floor, is covered with bird feces and bird carcasses.

The roof contains numerous package units, which heat and cool the building; however, during our investigation, the HVAC system was inoperable.

3.0 MOLD INVESTIGATION

There are currently no established regulatory guidelines for concentrations of fungi. Guidelines for determining qualitative growth of various fungi have been provided for use as part of this report. The guidelines used do not represent threshold values having medical or health significance with respect to exposure, nor are they necessarily representative of an unacceptable indoor environment. Rather, the guidelines are intended as a reactionary threshold to prompt further investigation as to the cause of what is considered to be an above average concentration for indoor fungi. Occupational laboratories generally use qualitative guidelines when evaluating wipe, bulk and vacuum samples for fungi concentrations. The following guidelines have been used for determining qualitative growth of various fungi in wipe, bulk and vacuum samples.

<1000 CFU Light Growth 1,000 – 10,000 CFU Moderate Growth >10,000 CFU Heavy Growth

Research by independent laboratories has suggested that typical levels for indoor fungal prompt in air samples should be below a total of 300 CFU/ m³. With the exception of *Cladosporium*, no individual type of fungi should contribute more than 50 CFU/ m³ to the total. In general, indoor populations should be consistent with outdoor populations. The following text and tables outline the results of our bioaerosol testing.

The following fungal samples were collected during this investigation:

- Air samples via spore trap (6 samples plus 1 blank)
- Bulk samples (3 samples)
- Tape-lift samples (2 samples plus 1 blank)

On October 26, 2004, ASG conducted sampling for fungi at the USACOE South Pacific Division Laboratory. During this sampling, we collected samples inside and outside of the building. The exterior samples are used as control samples. Interior samples should show the same type and similar concentrations to exterior samples. Interior samples showing higher concentrations compared to outside samples, different genera make up as compared to exterior samples, or differing order of domination may indicate an interior environmental condition favorable to microorganism growth. The samples were analyzed by NCG Laboratories in Muncie, Indiana.

The following tables outline the fungal results obtained:

3.1 Air Samples (filter method) for Fungi

Stachybotrys was found on the interior samples (Samples VA-AC-05 and VA-AC-06) where the visible black growth was observed to be most concentrated. Also, visual evidence of significant water damage was noted in these areas. This species, as well as *Aspergillus/Penicillium* and *Fusarium* has been known to produce mycotoxins. Note that although *Aspergillus/Penicillium* a was found on the interior, they were also found on the exterior samples

	Table 1				
	Air Samples (spore trap method) for Fungi				
Sample Number	Location	CFU/M ³	Analyte		
VA-AC-01	Exterior – east side	157	Ascospore		
		10,365	Aspergillus/Penicillium		
		628	Cladosporium		
		11,151	TOTAL		
VA-AC-02	Level 1 – east side	314	Ascospore		
		6,753	Aspergillus/Penicillium		
		1,099	Cladosporium		
		157	Zygomycetes		
		8,324	TOTAL		
VA-AC-03	Level 1 – north side	1,728	Ascospore		
		6,596	Aspergillus/Penicillium		
		2,356	Cladosporium		
		10,689	TOTAL		
VA-AC-04	Level 2 – west side –	157	Ascospore		
	hallway by men's room	6,282	Aspergillus/Penicillium		
		785	Cladosporium		
		7,224	TOTAL		
VA-AC-05	Level 1 – engineer's office	3,769	Aspergillus/Penicillium		
	at southwest corner	1,256	Chaetomium		
		942	Cladosporium		
		157	Fusarium		
		3,141	Stachybotrys		
		9,266	TOTAL		
VA-AC-06	Level 2 – southwest corner	3,612	Aspergillus/Penicillium		
		628	Chaetomium		
		785	Cladosporium		
		157	Stachybotrys		
		5,183	TOTAL		
VA-AC-07	Blank	N/A	No spores seen		

 CFU/M^3 = colony forming units per cubic meter

3.2 Bulk Samples for Fungi

Relatively high concentrations of *Yeast* were found on the water damaged ceiling tiles and wood floors. In addition, *Penicillium* was found in relatively high concentrations on the water damaged ceiling tiles. These species have been known to produce mycotoxins.

Table 2				
	Bulk Samples fo	r Fungi		
Sample Number	Location / Material	CFU/gram	Analyte	
VA-B-01	Level 1 – west side	25,000	Penicillium	
	Ceiling tile	15,000	Yeast	
		27,000	Unidentified Mildew	
		67,000	TOTAL	
VA-B-02	Level 2 – southwest	10,000	Yeast	
	Ceiling tile	164,000	Unidentified Conidia	
		174,000	TOTAL	
VA-B-03	Level 2 – Room 203 – hall	428,571	Yeast	
	Wood floor	428,571	TOTAL	

CFU/gram = colony forming units per gram

3.3 Tape Lift Samples for Fungi

Stachybotrys and Aspergillus have been known to produce mycotoxins and were found in the southwest areas on levels 1 and 2 where the heaviest visual growth was observed.

Table 3					
Tape Lift Samples for Fungi					
Sample Number	Location (surface)	Analyte (loading)			
VA-T-01	Level 1 – southwest corner –	Aspergillus/Penicillium			
	engineer's office	Chaetomium			
	(ceiling tile)	Cladosporium			
		Stachybotrys			
		(light)			
VA-T-02	Level 2 – southwest corner office	Stachybotrys			
	(office 203)	(very heavy)			
	(sheet rock wall)	·			
VA-T-03	Blank	None seen			

4.0 BACTERIA & BIRD FECES INVESTIGATION

There are currently no identified acceptable levels for indoor bacterial bioaerosols. Independent laboratory research suggests that Bacillus, Staphylococcus, Micrococcus, Corynebacterium and other gram positive bacteria typically dominate indoor environments. Bacillus species are widely distributed in nature, particularly in soil, dust, water, and on materials of plant or animal origin. The majorities of *Bacillus* species apparently have little pathogenic potential and are rarely associated with disease in humans. Most Staphylococcal species are common inhabitants of the human skin and mucous membranes. Micrococcus are common on skin, in soil, dust, water, and elsewhere and are not considered pathogenic to humans, plants, or animals. Corynebacterium species is a non sporulating bacilli which are found in diverse habitats ranging from soil to skin and mucous membranes of mammals. Gram negative bacilli include such organisms as Escherichia coli, Salmonella, Shingella, Pseudomonas, Klebsiella, Proteus, Enterobacter, Legionella and a number of genera to numerous to discuss. Most of these bacteria are associated with moisture sources and tend to be subject to desiccation more than other bacteria. Their natural habitats range from the intestinal tract of mammals to stagnant fresh water and seawater. An important aspect of these bacteria is their ability to produce endotoxin, a component of the cell structure. Endotoxins are the lipopolysaccharide complexes of the cell wall which are released into the environment when the cell breaks up. These endotoxins can be relatively toxic to mammals. Exposure can result in fever, malaise, respiratory distress, headaches, alterations of white blood cell counts, and death. When air samples are collected with an Andersen air sampler, total indoor concentrations are typically in the range of 250 colony forming units per cubic meter (CFU/m³) in air, however, greater concentrations can occur relative to activity level.

The following bacteria samples were collected during this investigation:

- Bulk samples (1 sample)
- Wipe samples (1 sample plus 1 blank)

On October 26, 2004, ASG conducted sampling for bacteria at the USACOE South Pacific Division Laboratory. During this sampling, we collected samples of various surfaces within the building, namely where visible bird feces was observed. The samples were analyzed by NCG Laboratories in Muncie, Indiana.

The following tables outline the bacterial results obtained:

4.1 Bulk Samples for Bacteria

Various bacteria species were from sampling the bird droppings and these species are believed to be opportunistic pathogens which may cause illnesses.

Table 4							
Bulk Samples for Bacteria							
Sample Number	Location / Material	CFU/gram	Analyte				
VA-BA-01	Level 1 – north side	3,300	Curtobacterium				
	Bird droppings	100	Rhodococcus				
	3,400 TOTAL						

CFU/gram = colony forming units per gram

4.2 Wipe Samples for Bacteria

A bacterium species, *Brevibacterium*, was found on the concrete floor below the bird droppings. This species is believed to be an opportunistic pathogen, which may cause illnesses.

Table 5					
Wipe Samples for Bacteria					
Sample Number	Location (Surface)	CFU/ft ²	Analyte		
VA-W-01	Level 1 – north side	900	Brevibacterium		
	(concrete floor)				
VA-w-02	Blank	N/A	No Growth		

 CFU/ft^2 = colony forming units per square foot

4.3 Bird Feces

Some heath risks associated with birds and feces in buildings include Urban pest birds promote the transmission of several hundred types of disease infections and illnesses. Buildings infested with pest birds place at risk the building occupants, customers and maintenance workers. These nuisance birds: pigeons, starlings, crows, house sparrows and nuisance gulls, to mention only a few, transmit disease in four ways: food and water contaminated with feces; inhalation of contaminated dust; transference by parasites associated with nuisance birds, i.e., fleas, ticks, mites and other ectoparasites; and direct contact with feces.

The following visual observations were made on October 26, 2004:

- Excessive bird droppings were observed on the concrete floor, namely on level 1 north side and central area
- Dead birds were also seen in these areas.

5.0 CONCLUSIONS AND RECOMMENDATIONS

Through laboratory analysis and visual inspection, we recommend the following activities take place to alleviate potential indoor air quality problems.

- Mold/Fungi Stachybotrys was found on the interior air samples where visible black growth was observed to be most concentrated. In addition, visual evidence of significant water damage was located in these areas. This species, as well as Aspergillus/Penicillium and Fusarium, which were also found, has been known to produce mycotoxins. Also, relatively high concentrations Yeast were found on the water damaged ceiling tiles and wood floors. In addition, Stachybotrys and Aspergillus, which may produce mycotoxins, were found on the surfaces in the southwest areas on levels 1 and 2, where the heaviest visual growth was observed.
- Bacteria Various bacteria species were identified from sampling the bird droppings, and
 these species are believed to be opportunistic pathogens which may cause illnesses. In
 addition, *Brevibacterium*, was found on the concrete floor below the bird droppings. This
 species is believed to be an opportunistic pathogen, which may also cause illnesses.
- Bird Feces Excessive bird droppings were observed on the concrete floor, namely on level 1 north side and central area. Also, dead birds were also seen in these areas.

Recommendations

- Repair the roof and possible plumbing leaks, including broken windows, namely at the southwestern side of the building.
- Replace the building's HVAC system and disinfect the duct work and grills/vents.
- Remove all carpeting and disinfect the flooring below if concrete.
- Remove all porous surfaces including water damaged wooden floors, sheet rock walls and ceiling, and ceiling tiles.
- Remove all bird droppings and dead bird from the building. Disinfect the concrete surfaces where bird droppings were located. Seal the building as to not allow further intrusion of fowl.
- Disinfect all surfaces and building products remaining in the building.
- Retest the air and building surfaces, once these items have been conducted.
- The HVAC system throughout the facility, including the duct system, if it remains, should be disinfected and cleaned using HEPA vacuuming protocols.

6.0 OPINIONS OF COST

Based upon the various mold, fungi, bacteria and bird droppings identified at the site, we are providing these opinions of cost:

Table 6					
Removal / Cleaning — Opinions of Cost					
Task	Comments		Cost for Removal		
Mold and fungal removal	Removal of ceiling tile,	sheet rock, wood floors	\$50,000		
Disinfecting	Floors, duct work, wood	len structures	\$20,000		
Bird dropping removal	Removal of visible mate	erial	\$10,000		
Carpet removal	Removal and disposal o	f carpet	\$10,000		
SUBTOTAL (removal /disinfecting)			\$90,000		
IND	USTRIAL HYGIENE	E CONSULTING			
On-Site Management and	30 Days – 75 samples	\$ 500 per day +	\$16,375		
samples		\$40/sample			
Abatement Specifications and	LS	\$ 2,500			
Bidding Documents					
SUB-TOTAL (monitoring)	\$ 18,875				
GRAND TOTAL (Est.)			\$108,875		

These opinions of costs are estimates only, based upon our experience of having contractors perform similar projects in scope and size. They do not, however, include the fees for contractors bonding or insurance or replacement costs, such as re-flooring or HVAC replacement or roof inspection and repair. The most accurate way to determine an estimated cost of removal and disinfecting is to have several qualified contractors provide actual bids to perform the work.

7.0 LIMITATIONS

Information contained herein was obtained by means of on-site observations and analytical data. Conclusions of this survey are based on reasonably accessible information pertaining specifically to this survey. However, this is not to suggest that the information obtained is a complete compilation of all existing information, which may be pertinent to this site. The intent of this survey is to sample the indoor atmospheric conditions as they relate to the intent of the building's structure and content to ensure that conditions remain parallel to comfort levels established by regulatory agencies which govern indoor atmospheric conditions. This survey is not intended to represent an exhaustive research of all-potential hazards or conditions, which may exist.

This report does not purport to represent future indoor conditions or events. Situations or activities, which transpire subsequent to this report, which result in adverse environmental, construction and/or engineering impacts, are not to be construed as relevant to this study.

This report is intended for the sole use of ASG's client. This report may not be used or relied upon by any other party without the written consent of ASG. The scope of services performed in execution of the evaluation may not be appropriate to satisfy the needs of other users, and the use or re-use of this document or the finding, conclusions, or recommendations is at the risk of said user.

We appreciate the opportunity to be of service to you on this project. If you have any questions or comments, please contact us at (801) 261-4616.

Prepared By:

Terry Bleckner, CIH Industrial Hygienist/Utah Branch Manager Paul E. Johnson Indoor Air Quality Manager

C:report/U04-2732-IAQ

APPENDIX A SAMPLING METHODS

AIR SAMPLING FOR MOLDS AND FUNGI

Air Sampling: Agar Impaction Method

The single stage multi-hole impactors were used for sampling for molds and spores.

Sampling procedures:

Air flow: 28.3 1/m

Calibrated with a primary standard

Sampling time: Sampling time of 2-5 minutes

Media: Malt extract agar

Filter cassettes were shipped to the laboratory by overnight courier for morning delivery. It was necessary that the samples arrive at the laboratory and be incubated at appropriate temperatures before visible growth develops. One blank media dish was submitted with each set of samples.

SURFACE DUST SAMPLING FOR MOLDS AND FUNGI

Surface samples are collected by removing material with a suction device or by washing a prescribed area with a wetted swab, cheesecloth, gauze, or filter.

Suction Device Method

Cassette: 37 mm, MCE filter, 0.8 mm pore size

Equipment: High volume personal sampling pump

Calibration: 20-40 liters per minute (L/m) calibrated with a primary standard

Sample area: 1 meter x 1 meter

Sample volume: Carpet was vacuumed for a minimum of 5 minutes.

Method: Open face cassette

Wipe Sample Method

A sterile gauze pad was used to wipe a 10 cm x 10 cm area.

Aseptic techniques were used to avoid contamination.

Samples were shipped to the laboratory within 24 hours.

AIR SAMPLING FOR BACTERIA

Air Sampling: Agar Impaction Method

The single stage multi-hole impactors were used for sampling for bacteria.

Sampling procedures:

Air flow: 28.3 1/m

Calibrated with a primary standard

Sampling time: Sampling time of 2-5 minutes

Media: Tryptic soy agar

Filter cassettes were shipped to the laboratory by overnight courier for morning delivery. It was necessary that the samples arrive at the laboratory and be incubated at appropriate temperatures before visible growth develops. One blank media dish was submitted with each set of samples.

SURFACE SAMPLING FOR BACTERIA

Surface samples are collected by washing a prescribed area with a sterile swab, cheesecloth, gauze, or filter.

Wipe Sample Method

A sterile gauze pad was used to wipe a 10 cm x 10 cm area.

Aseptic techniques were used to avoid contamination.

Samples were shipped to the laboratory within 24 hours.

APPENDIX B

LABORATORY RESULTS AND CHAIN OF CUSTODY FORMS





Phone: (765) 286-5142 Fax: (765) 286-5152

Tape Report

Client:		ARG	
Company:			
Address:			
Phone Number:			
Fax Number:			
E-Mail Address:			
Client Number:		ARG001	
Project Number:		U04-2732	
Project Name:		Sausilito	
Lab ID Number:	355		
Analyst:	AJK		
Date:	10/28/2004		

Tape Definitions:

Concentrations of greater than 100% are due to overlap of spores, pollen, hyphae, debris, etc.

Concentrations of less than 100% are due to clean sections on the sample where nothing is present.

Spores identified in bold may produce fungi that are known mycotoxin producing organisms.

** Pathogenic versus non-pathogenic fungi cannot be differentiated using this method. Viable versus nonviable fungi cannot be differentiated using this method. Samples cannot be identified beyond the genera level. Some fungi must be grouped together due to their similar spore structure. If these fungi can be identified separately they will be. This is dependent on observing hyphae along with the spores.

Background and Overall Particle Densities:

<u>Light</u> - Discriptor for very little observed on analysis of entire sample.

Moderate - Discriptor for a medium amount of coverage observed on analysis of entire sample.

Heavy - Discriptor for a high amount of coverage observed on analysis of entire sample.

Very Heavy - Discriptor for a very high amount of coverage or almost complete coverage observed on analysis of entire sample.

- ***Background Densities of heavy or very heavy may occlude some spores, pollen, and insect fragments causing concentrations to be lower than what they actually are.
- ***Occasionally surface layers or borders are only able to be analyzed due to high background densities. This will be noted in the Comment Section as it occurs.

Specific Genera Densities: 1=<10%, 2=10% to 25%, 3=25% to 50%, 4=50 to 75%, 5=75 to 90%, 6=>90% Specific Genera Densities are based on overall particle densities.

Overloaded means there was too much background debris &/or spores present to be able to accurately see individual particles. See the Comment Section for whether it is due to background debris or spores.

Individual concentration amounts may differ from the total concentration amount due to the rounding of significant figures.

We analyze a minimum of 4 traverses completely across the sample. Each traverse is done in a random place on the sample. Percents of identified spores only include the concentration of spores observed. Hyphae is not included in this percent concentration, but is noted if it is seen!



Phone: (765) 286-5142 Fax: (765) 286-5152

Project Number:	U04-2732
Lab ID Number:	355

ND

ND

ND

Tape Report

Sample Number:	VA-T-1		Sample Number:	VA-T-2	
Sample Description:	L1 - SWC - Eng.	. Office - DCT	Sample Description:	L2 - SWC office	e - SR wall
			<u> </u>		
Background Density:	Light		Background Density:	Very Heavy	
Overall Particle Density: Light			Overall Particle Density:		
	J			, ,	
Comments:			Comments:		
Hyphae Observed:	No		Hyphae Observed:	Yes	
Hyphal Fragments Observed:			Hyphal Fragments Observed:		
	1.1.4				
Particle Identification	Concentration	Comments	Particle Identification	Concentration	Comments
Acremonium	ND		Acremonium	ND	
Alternaria	ND		Alternaria	ND	
Ascospore	ND		Ascospore	ND	
Aspergillus	ND		Aspergillus	ND	
Aspergillus/Penicillium	1		Aspergillus/Penicillium	ND	
Bipolaris	ND		Bipolaris	ND	
Chaetomium	3		Chaetomium	ND	
Cladosporium	1		Cladosporium	ND	
Curvularia	ND		Curvularia	ND	
Epicoccum	ND		Epicoccum	ND	
Fusarium	ND		Fusarium	ND	
Myxomycetes	ND		Myxomycetes	ND	
Penicillium	ND		Penicillium	ND	
Peronospora	ND		Peronospora	ND	
Stachybotrys	3		Stachybotrys	6	spores & hyphae obs
Ulocladium	ND		Ulocladium	ND	
Zygomycetes	ND		Zygomycetes	ND	
Unidentified Mildew	ND		Unidentified Mildew	ND	

Unidentified Conidia

Insect Fragments

Pollen

and of Rock

Lab Coordinator Signature:

ND

ND

ND

Unidentified Conidia

Insect Fragments

Pollen



Lab Coordinator Signature:

NCG Laboratories 5826 West Kilgore Avenue Muncie, IN 47304

Phone: (765) 286-5142 Fax: (765) 286-5152

Project Number:	U04-2732
Lab ID Number:	355

Tape Report

			1000.		
Sample Number:	VA-T-3		Sample Number:		
Sample Description:	Blank		Sample Description:		
			<u> </u>		
Background Density:	Light		Background Density:		
Overall Particle Density:			Overall Particle Density:		
•	<u>I</u>				
Comments:			Comments:		
Hyphae Observed:	No		Hyphae Observed:		
Hyphal Fragments Observed:			Hyphal Fragments Observed:		
<i>y</i> ,	-		71.	<u></u>	
Particle Identification	Concentration	Comments	Particle Identification	Concentration	Comments
Acremonium	ND		Acremonium		
Alternaria	ND		Alternaria		
Ascospore	ND		Ascospore		
Aspergillus	ND		Aspergillus		
Aspergillus/Penicillium	ND		Aspergillus/Penicillium		
Bipolaris	ND		Bipolaris		
Chaetomium	ND		Chaetomium		
Cladosporium	ND		Cladosporium		
Curvularia	ND		Curvularia		
Epicoccum	ND		Epicoccum		
Fusarium	ND		Fusarium		
Myxomycetes	ND		Myxomycetes		
Penicillium	ND		Penicillium		
Peronospora	ND		Peronospora		
Stachybotrys	ND		Stachybotrys		
Ulocladium	ND		Ulocladium		
Zygomycetes	ND		Zygomycetes		
Unidentified Mildew	ND		Unidentified Mildew		
Unidentified Conidia	ND		Unidentified Conidia		
Pollen	ND		Pollen		
Insect Fragments	ND		Insect Fragments		
Lah Coordin	ator Signature	Ands	g Rock		



Phone: (765) 286-5142 Fax: (765) 286-5152

Micro5 Report

Client:		ARG		
Company:				
Address:				
Phone Number:				
Fax Number:				
E-Mail Address:				
Client Number:		ARG 001		
Project Number:		U04-2732		
Project Name:		Sausolito		
Lab ID Number:	355			
Analyst:	AJK			
Date:	10/28/2004			

Micro5 Definitions:

ND=None Detected.

TNTQ=Too Numerous To Quantify.

Spores identified in bold may produce fungi that are known mycotoxin producing organisms.

** Pathogenic versus non-pathogenic fungi cannot be differentiated using this method. Viable versus nonviable fungi cannot be differentiated using this method. Samples cannot be identified beyond the genera level. Some fungi must be grouped together due to their similar spore structure.

Debris & Skin Densities: 0=0%, 1=0-25%, 2=25% to 50%, 3=50% to 75%, 4=>75%

Densities of 3 or 4 may occlude some spores, pollen, and insect fragments causing counts to be lower than what they actually are.

Overloaded means that there was too much non-fungal debris/skin present to be able to accurately count particles.

Individual concentration amounts may differ from the total concentration amount due to the rounding of significant figures.

We analyze a minimum of 25% or 300 spores of one genera.

Micro5 Equations for Analysis, From Environmental Monitoring Systems Users Manual

Equation 1: Air Volume Calculation

Air volume (m^3)=(Sampling rate [liters per minute]/1000)XNo. of minutes

Equation 2: Raw Count Multiplier Calculation

Raw Count Multiplier=(Total Impact Area [4.154]**)/(# Fields AnalyzedXMicroscope field of view surface area [.0962]*)

Equation 3: Count per meter^3 Calculation

Cts/m^3=No. Particles CountedXRaw Count MultiplierX(1/Volume [m^3])

Equation 4: % of Trace Analyzed Calculation

% Analyzed=(No. Fields Analyzed)/(Total No. Fields [43]***)X100%

*Microscope field of view surface area was measured using a slide graticule, calculated, and found to be 0.0962mm^3.

- **Total Impact Area of 4.154mm^3 is provided in the Micro5 Users Manual.
- ***Total No. Fields of 43 is provided in the Micro5 Users Manual.





NCG Laboratories 5826 West Kilgore Avenue Muncie, IN 47304 Phone: (765) 286-5142 Fax: (765) 286-5152

Project Number:	U04-2732
Lab ID Number:	355

Micro5 Report

	oo ivebort
Sample Number: VA-AC-1	Sample Number: VA-AC-2
Sample Description: Exterior - East	Sample Description: L1 - East Side
Sample Volume (m^3): 0.0250	Sample Volume (m^3): 0.0250
Number of Traverses: 11	Number of Traverses: 11
Trace Length Counted (mm): 3.85	Trace Length Counted (mm): 3.85
% of Trace Analyzed: 26	% of Trace Analyzed: 26

Skin Fragment Density: 0	Skin Fragment Density: 1
Debris Density: 1	Debris Desity: 2

		Concentration	
Particle Identification	Raw	per m^3	%
Acremonium	ND	ND	ND
Alternaria	ND	ND	ND
Ascospore	1	157	1
Aspergillus/Penicillium	66	10365	93
Bipolaris	ND	ND	ND
Chaetomium	ND	ND	ND
Cladosporium	4	628	6
Curvularia	ND	ND	ND
Epicoccum	ND	ND	ND
Fusarium	ND	ND	ND
Myxomycetes	ND	ND	ND
Peronospora	ND	ND	ND
Stachybotrys	ND	ND	ND
Ulocladium	ND	ND	ND
Zygomycetes	ND	ND	ND
Unidentified Mildew	ND	ND	ND
Unidentified Conidia	ND	ND	ND
Hyphae	ND	ND	ND
Pollen	ND	ND	ND
Insect Fragments	ND	ND	ND
TOTAL	71	11151	100

		Concentration	
Particle Identification	Raw	per m^3	%
Acremonium	ND	ND	ND
Alternaria	ND	ND	ND
Ascospore	2	314	4
Aspergillus/Penicillium	43	6753	81
Bipolaris	ND	ND	ND
Chaetomium	ND	ND	ND
Cladosporium	7	1099	13
Curvularia	ND	ND	ND
Epicoccum	ND	ND	ND
Fusarium	ND	ND	ND
Myxomycetes	ND	ND	ND
Peronospora	ND	ND	ND
Stachybotrys	ND	ND	ND
Ulocladium	ND	ND	ND
Zygomycetes	1	157	2
Unidentified Mildew	ND	ND	ND
Unidentified Conidia	ND	ND	ND
Hyphae	ND	ND	ND
Pollen	ND	ND	ND
Insect Fragments	ND	ND	ND
TOTAL	. 53	8324	100

Lab Coordinator Signature:	Lab Coordinator Signature:	Andr	9	Koch,	
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Phone: (765) 286-5142 Fax: (765) 286-5152

Project Number:	U04-2732
Lab ID Number:	355

Micro5 Report

Sample Number: VA-AC	C-3	Sample Number:	VA-AC-4
Sample Description: L1 - N	orth Side	Sample Description:	L2 - West Side - Hall
Sample Volume (m^3): 0.0250)	Sample Volume (m^3):	0.0250
Number of Traverses: 11		Number of Traverses:	11
Trace Length Counted (mm): 3.85		Trace Length Counted (mm):	3.85
% of Trace Analyzed: 26		% of Trace Analyzed:	26

Skin Fragment Density: 1	Skin Fragment Density: 1
Debris Density: 3	Debris Desity: 1

		Concentration	
Particle Identification	Raw	per m^3	%
Acremonium	ND	ND	ND
Alternaria	ND	ND	ND
Ascospore	11	1728	16
Aspergillus/Penicillium	42	6596	62
Bipolaris	ND	ND	ND
Chaetomium	ND	ND	ND
Cladosporium	15	2356	22
Curvularia	ND	ND	ND
Epicoccum	ND	ND	ND
Fusarium	ND	ND	ND
Myxomycetes	ND	ND	ND
Peronospora	ND	ND	ND
Stachybotrys	ND	ND	ND
Ulocladium	ND	ND	ND
Zygomycetes	ND	ND	ND
Unidentified Mildew	ND	ND	ND
Unidentified Conidia	ND	ND	ND
Hyphae	ND	ND	ND
Pollen	ND	ND	ND
Insect Fragments	ND	ND	ND
TOTAL	68	10679	100

		Concentration	
Particle Identification	Raw	per m^3	%
Acremonium	ND	ND	ND
Alternaria	ND	ND	ND
Ascospore	1	157	2
Aspergillus/Penicillium	40	6282	87
Bipolaris	ND	ND	ND
Chaetomium	ND	ND	ND
Cladosporium	5	785	11
Curvularia	ND	ND	ND
Epicoccum	ND	ND	ND
Fusarium	ND	ND	ND
Myxomycetes	ND	ND	ND
Peronospora	ND	ND	ND
Stachybotrys	ND	ND	ND
Ulocladium	ND	ND	ND
Zygomycetes	ND	ND	ND
Unidentified Mildew	ND	ND	ND
Unidentified Conidia	ND	ND	ND
Hyphae	ND	ND	ND
Pollen	ND	ND	ND
Insect Fragments	ND	ND	ND
TOT	AL 46	7224	100

Lab Coordinator Signature:



Phone: (765) 286-5142 Fax: (765) 286-5152

Project Number:	U04-2732
Lab ID Number:	355

Micro5 Report

Sample Number: VA-AC-5	Sample Number: VA-AC-6
Sample Description: L1 - SWC Eng. office	Sample Description: L2 - SWC office
Sample Volume (m^3): 0.0250	Sample Volume (m^3): 0.0250
Number of Traverses: 11	Number of Traverses: 11
Trace Length Counted (mm): 3.85	Trace Length Counted (mm): 3.85
% of Trace Analyzed: 26	% of Trace Analyzed: 26

Skin Fragment Density: 1	Skin Fragment Density: 1
Debris Density: 1	Debris Desity: 1

		Concentration	
Particle Identification	Raw	per m^3	%
Acremonium	ND	ND	ND
Alternaria	ND	ND	ND
Ascospore	ND	ND	ND
Aspergillus/Penicillium	24	3769	41
Bipolaris	ND	ND	ND
Chaetomium	8	1256	14
Cladosporium	6	942	10
Curvularia	ND	ND	ND
Epicoccum	ND	ND	ND
Fusarium	1	157	2
Myxomycetes	ND	ND	ND
Peronospora	ND	ND	ND
Stachybotrys	20	3141	34
Ulocladium	ND	ND	ND
Zygomycetes	ND	ND	ND
Unidentified Mildew	ND	ND	ND
Unidentified Conidia	ND	ND	ND
Hyphae	ND	ND	ND
Pollen	ND	ND	ND
Insect Fragments	ND	ND	ND
TOTAL	59	9266	100

		Concentration	
Particle Identification	Raw	per m^3	%
Acremonium	ND	ND	ND
Alternaria	ND	ND	ND
Ascospore	ND	ND	ND
Aspergillus/Penicillium	23	3612	70
Bipolaris	ND	ND	ND
Chaetomium	4	628	12
Cladosporium	5	785	15
Curvularia	ND	ND	ND
Epicoccum	ND	ND	ND
Fusarium	ND	ND	ND
Myxomycetes	ND	ND	ND
Peronospora	ND	ND	ND
Stachybotrys	1	157	3
Ulocladium	ND	ND	ND
Zygomycetes	ND	ND	ND
Unidentified Mildew	ND	ND	ND
Unidentified Conidia	ND	ND	ND
Hyphae	ND	ND	ND
Pollen	ND	ND	ND
Insect Fragments	ND	ND	ND
TOTAL	. 33	5183	100

Lab Coordinator Signature:



Stachybotrys

Zygomycetes

Unidentified Mildew

Unidentified Conidia

Insect Fragments

Ulocladium

Hyphae

Pollen



Sample Number: VA-AC-7

ND

ND

ND

ND

ND

ND

ND

ND

0

TOTAL

ND

ND

ND

ND

ND

ND

ND

ND

0

Sample Description: Blank

NCG Laboratories 5826 West Kilgore Avenue Muncie, IN 47304

Phone: (765) 286-5142 Fax: (765) 286-5152

Project Number:	U04-2732
Lab ID Number:	355

Sample Number:

Sample Description:

Micro5 Report

Sample Volume (m^3):	NA			Sample Volume (m^3):			
Number of Traverses:	NA			Number of Traverses:			
Trace Length Counted (mm):	NA			Trace Length Counted (mm):			
% of Trace Analyzed:	100			% of Trace Analyzed:			
Skin Fragment Density:	0			Skin Fragment Density:			
Debris Density:	1			Debris Desity:			
		Concentration				Concentration	
Particle Identification	Raw	per m^3	%	Particle Identification	Raw	per m^3	%
Acremonium	ND	ND	ND	Acremonium			
Alternaria	ND	ND	ND	Alternaria			
Ascospore	ND	ND	ND	Ascospore			
Aspergillus/Penicillium	ND	ND	ND	Aspergillus/Penicillium			
Bipolaris	ND	ND	ND	Bipolaris			
Chaetomium	ND	ND	ND	Chaetomium			
Cladosporium	ND	ND	ND	Cladosporium			
Curvularia	ND	ND	ND	Curvularia			
Epicoccum	ND	ND	ND	Epicoccum			
Fusarium	ND	ND	ND	Fusarium			
Myxomycetes	ND	ND	ND	Myxomycetes			
Peronospora	ND	ND	ND	Peronospora			

ND

ND

ND

ND

ND

ND

ND

ND

0

Stachybotrys

Ulocladium

Hyphae

Pollen

Zygomycetes

Unidentified Mildew

Unidentified Conidia

Insect Fragments

TOTAL

	Andr	Q	Rock.	
Lab Coordinator Signature:		1	/	





Phone: (765) 286-5142 Fax: (765) 286-5152

Viable Fungi Report

Client:		ARG	
Company:		ARG	
Address:			
Phone Number:			
Fax Number:			
E-Mail Address:			
Client Number:		ARG001	
Project Number:		U04-2732	
Project Name:		V A Sausalite	
Lab ID Number:	356		
Analyst:	AJK		
Date:	11/11/2004		

Viable Fungi Definitions:

ND=None Detected.

TNTQ=Too Numerous To Quantify

Spores identified in bold may produce fungi that are known mycotoxin producing organisms.

**Holding times for samples are between 24-48 hours from collection to analysis.

Samples that have been identified to the species level are listed under the Comments heading.

Individual concentration amounts may differ from the total concentration amount due to the rounding of significant figures.

Detection Levels are:

Andersen Airs = $7/m^3$

Bulks = 100/gram, if 1gram of sample is submitted

Wipes/Swabs = 100/wipe or swab

Vacuum Samples = 25/foot^2

Water = 100/mL

Viable Equations for Analysis, Taken from Standard Methods 20th Edition

Equation 1: Dilution Factor Calculation

Dilution Factor={(Sample Amount)/(Amount of Sterile Buffer [99mL]*+Amount of Sample)}XAmount Plated[mL] Equation 2: Concentration Calculation

Concentration=No. Colonies CountedX(1/Dilution Factor)

Sterile Buffer is purchased in pre-aliquotted amounts of 99mL per bottle.





Phone: (765) 286-5142 Fax: (765) 286-5152

Project Number:	U04-2732
Lab ID Number:	356

Viable Fungi Report

	<u> </u>
Sample Number:	VA-B-1
Sample Description:	L1 - W. side - DCT

Sample Amount:	1 gram
Amount Plated:	0.1 mL
Dilution Factor:	0.001

***NOTE: Concentration=	per gra	am		
Identification	Raw	Co	ncentration	Comments
Acremonium species	ND	ND		
Alternaria species	ND	ND		
Aspergillus species	ND	ND		
Bipolaris species	ND	ND		
Chaetomium species	ND	ND		
Cladosporium species	ND	ND		
Curvularia species	ND	ND		
Epicoccum species	ND	ND		
Fusarium species	ND	ND		
Myxomycetes	ND	ND		
Penicillium species	25		25000	
Peronospora species	ND	ND		
Stachybotrys species	ND	ND		
Zygomycetes	ND	ND		
Yeast	15		15000	
Unidentified Mildew	27		27000	
Unidentified Conidia	ND	ND		
TOTAL	67		67000	

and of Rock



Phone: (765) 286-5142 Fax: (765) 286-5152

Project Number:	U04-2732
Lab ID Number:	356

Viable Fungi Report

Sample Number:	VA-B-2
Sample Description:	L2 - SW - DCT

Sample Amount:	1 gram
Amount Plated:	0.1 mL
Dilution Factor:	0.001

***NOTE: Concentration=	per gra	am	
Identification	Raw	Concentration	Comments
Acremonium species	ND	ND	
Alternaria species	ND	ND	
Aspergillus species	ND	ND	
Bipolaris species	ND	ND	
Chaetomium species	ND	ND	
Cladosporium species	ND	ND	
Curvularia species	ND	ND	
Epicoccum species	ND	ND	
Fusarium species	ND	ND	
Myxomycetes	ND	ND	
Penicillium species	ND	ND	
Peronospora species	ND	ND	
Stachybotrys species	ND	ND	
Zygomycetes	ND	ND	
Yeast	10	10000	
Unidentified Mildew	ND	ND	
Unidentified Conidia	164	164000	No Spores Detected
TOTAL	174	174000	



> Phone: (765) 286-5142 Fax: (765) 286-5152

Project Number: U04-2732
Lab ID Number: 356

Viable Fungi Report

Sample Number:	VA-B-3
Sample Description:	L2 - Rm 203 - Wood Floor

Sample Amount:	0.7 grams
Amount Plated:	0.1 mL
Dilution Factor:	0.001

***NOTE: Concentration=	per gra	am	
Identification	Raw	Concentration	on Comments
Acremonium species	ND	ND	
Alternaria species	ND	ND	
Aspergillus species	ND	ND	
Bipolaris species	ND	ND	
Chaetomium species	ND	ND	
Cladosporium species	ND	ND	
Curvularia species	ND	ND	
Epicoccum species	ND	ND	
Fusarium species	ND	ND	
Myxomycetes	ND	ND	
Penicillium species	ND	ND	
Peronospora species	ND	ND	
Stachybotrys species	ND	ND	
Zygomycetes	ND	ND	
Yeast	>300	>4285	71
Unidentified Mildew	ND	ND	
Unidentified Conidia	ND	ND	
TOTAL	>300	>428571	





Phone: (765) 286-5142

Fax: (765) 286-5152

Viable Bacteria Report

Client:		ARG	
Company:		ARG	
Address:			
Phone Number:			
Fax Number:			
E-Mail Address:			
Client Number:		ARG001	
Project Number:		U04-2732	
Project Name:		VA Sausalite	
Lab ID Number:	356		_
Analyst:	AJK		
Date:	11/11/2004		

Viable Bacteria Definitions:

ND=None Detected.

TNTQ=Too Numerous To Quantify

Samples that have been identified to the species level are listed under the Comments heading.

Individual concentration amounts may differ from the total concentration amount due to the rounding of significant figures.

Detection Levels are:

Andersen Airs = $7/m^3$

Bulks = 100/gram, if 1gram of sample is submitted

Wipes/Swabs = 100/wipe or swab

Vacuum Samples = 25/foot^2

Water = 100/mL

Viable Equations for Analysis, Taken from Standard Methods 20th Edition

Equation 1: Dilution Factor Calculation

Dilution Factor={(Sample Amount)/(Amount of Sterile Buffer [99mL]*+Amount of Sample)}XAmount Plated[mL] **Equation 2: Concentration Calculation**

Concentration=No. Colonies CountedX(1/Dilution Factor)

*Sterile Buffer is purchased in pre-aliquotted amounts of 99mL per bottle.

^{**}Holding times for samples are between 24-48 hours from collection to analysis.



Phone: (765) 286-5142 Fax: (765) 286-5152

Project Number:	U04-2732
Lab ID Number:	356

Viable Bacteria Report

Sample Number:	VA-W-1
Sample Description:	L1 - N side - conc. Floor

Sample Amount:	1 wipe
Amount Plated:	1 mL
Dilution Factor:	0.01

***NOTE: Concentration=	per wi	pe	
Identification	Raw	Concentration	Comments
Actinomycetes	ND	ND	
Brevibacterium	9	900	
Citrobacter species	ND	ND	
Corynebacterium sp	ND	ND	
Escherichia species	ND	ND	
Klebsiella species	ND	ND	
Lactococcus species	ND	ND	
Leclercia species	ND	ND	
Micrococcus species	ND	ND	
Oerskovia species	ND	ND	
Proteus species	ND	ND	
Pseudomonas species	ND	ND	
Serratia species	ND	ND	
Staphylococcus species	ND	ND	
Streptococcus species	ND	ND	
Gram Negatives	ND	ND	
Gram Positives	ND	ND	
TOTAL	9	900	



Phone: (765) 286-5142 Fax: (765) 286-5152

Project Number:	U04-2732
Lab ID Number:	356

Viable Bacteria Report

Sample Number:	VA-W-2
Sample Description:	Blank

Sample Amount:	1 wipe
Amount Plated:	1 mL
Dilution Factor:	0.01

***NOTE: Concentration=	per wi	ре	
Identification	Raw	Concentration	Comments
Actinomycetes	ND	ND	
Bacillus species	ND	ND	
Citrobacter species	ND	ND	
Corynebacterium sp	ND	ND	
Escherichia species	ND	ND	
Klebsiella species	ND	ND	
Lactococcus species	ND	ND	
Leclercia species	ND	ND	
Micrococcus species	ND	ND	
Oerskovia species	ND	ND	
Proteus species	ND	ND	
Pseudomonas species	ND	ND	
Serratia species	ND	ND	
Staphylococcus species	ND	ND	
Streptococcus species	ND	ND	
Gram Negatives	ND	ND	
Gram Positives	ND	ND	
TOTAL	0	0	

and of Rock





Phone: (765) 286-5142 Fax: (765) 286-5152

Project Number:	U04-2732
Lab ID Number:	356

Viable Bacteria Report

Sample Number:	VA-BA-1
Sample Description:	L1 - N side - Bird Droppings

Sample Amount:	1.0 grams
Amount Plated:	1 mL
Dilution Factor:	0.01

***NOTE: Concentration=	per gra	am	
Identification	Raw	Concentration	Comments
Actinomycetes	ND	ND	
Bacillus species	ND	ND	
Citrobacter species	ND	ND	
Curtobacterium	33	3300	
Escherichia species	ND	ND	
Klebsiella species	ND	ND	
Lactococcus species	ND	ND	
Leclercia species	ND	ND	
Micrococcus species	ND	ND	
Oerskovia species	ND	ND	
Proteus species	ND	ND	
Rhodococcus species	1	100	
Serratia species	ND	ND	
Staphylococcus species	ND	ND	
Streptococcus species	ND	ND	
Gram Negatives	ND	ND	
Gram Positives	ND	ND	
TOTAL	34	3400	

and g Rocky



1107 Hazeltine Blvd., Suite 400

Chaska, MN 55318

Phone: 952-448-9393 Fax: 952-448-1658

5826 West Kilgore

Phone: 765-286-5142 Fax: 765-286-5152

SAMPLE CHAIN OF CUSTODY

::: ::::::::::::::::::::::::::::::::::	C. tes 3 pm			3/2/2	FA=Filter Air
Client Information Project Name: Phone Number: Fax Number: E-mail Address:	Area/N		N/W	7.4%	AA=Anderson Air
Client: A LCs. Company: Address:	Analyses Requested**			711	ST=Spore Trap
Client No: A RC- oc 1. Project No: UC4-2732 Log-In No: 356 Date: 10/21/04	Sample Description Matrix* Ast SIDE Ast SIDE	Noard cine 1811	OFF: ce V.	21- SWC- Eng. office. DRT T LZ- SWC office- SR WALL T Blank	T=Tape Lift W=Water mation will not be accepted.
Building Number: NA Building Name: VA - SAUS OLITO Building Address: 3 LIBEATHIND SAUSCUITO CA	Extended 1	15 - 17 12- 12- 12- 12- 12- 12- 12- 12- 12- 12-	VA-AC-7 BIANE.	VA-T-1 61-5WC-E VA-T-2 62-5WC of VA-T-3 Blank	*Sample Matrix: B=Bulk W=Wipe T=Tape Lift **Samples missing Analyses Requested information will not be accepted

Date: Date:

Received By: $\subset \mathcal{U}$ Relinquished By: Turnaround Time***

Total # of Samples:

Date: Date:

***Turnaround times found on rate sheet.

Special Instructions:

Type of Analysis: Sampled By: Analyzed By:

Due:



5826 West Kilgore

Phone: 765-286-5142 Muncie, IN 47304

1107 Hazeltine Blvd., Suite 400 Chaska, MN 55318

Phone: 952-448-9393

Fax: 952-448-1658

SAMPLE CHAIN OF CUSTODY

Fax: 765-286-5152 Phone Number: E-mail Address. Project Name: Fax Number: Client Information Company: Address: Client: 2732 NCG Information Project No.:/ LO 4-Log-In No.: Date: Client No: **Building Information** Building Name: 🗸 A Building Number: Building Address:

E-mail Address:		M= 1++- 5pm	1.05	20,7	1.09	AA=Anderson Air FA=Filter Air
Watriv*	Floor W 2 DG	3	8 108	13	129 3 206	Lift W=Water ST=Spore Trap
Sample Number Sample Description	LI-N s. de - conc.	V4-W.2 Black		UA-13-3 LZ-1lm 203- WOOD Flood	VA-BA-1 L1- N 5. de - Bird D.	**Sample Matrix: B=Bulk W=Wipe T=Tape Lift **Samples missing Analyses Requested information will not be accented

Due: Date:

Turnaround Time***

Total # of Samples:

Date: 10/ Date:

***Turnaround times found on rate sheet.

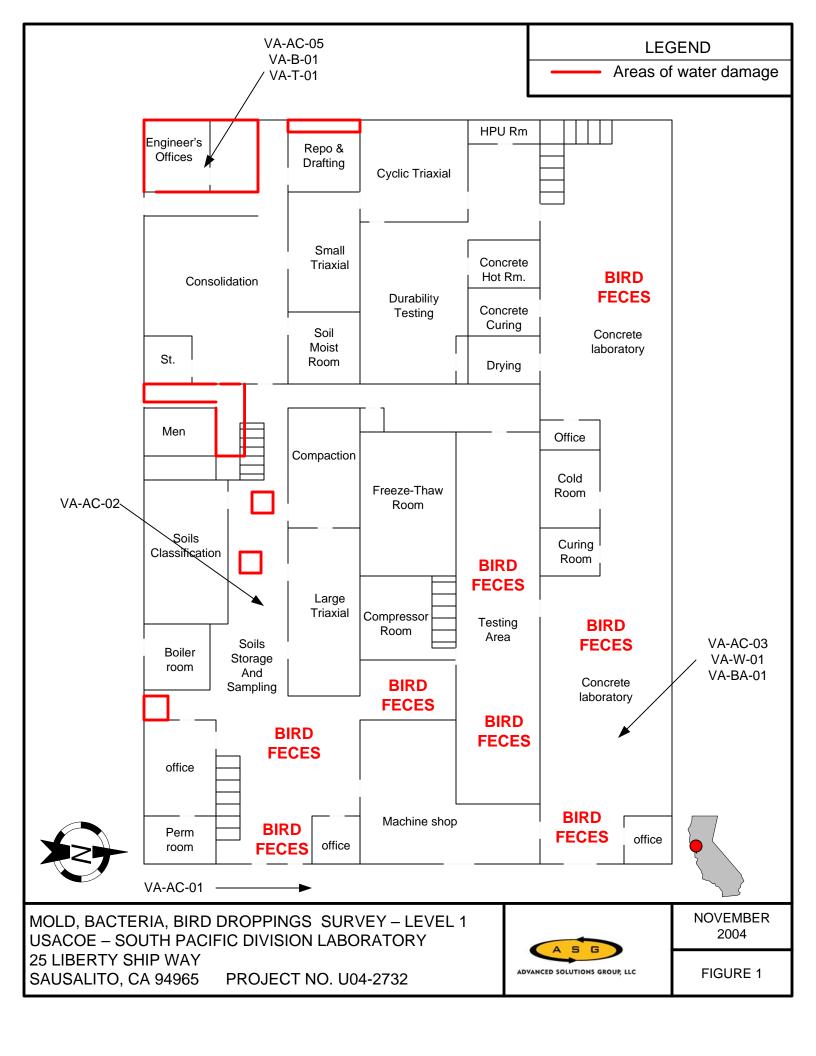
Special Instructions:

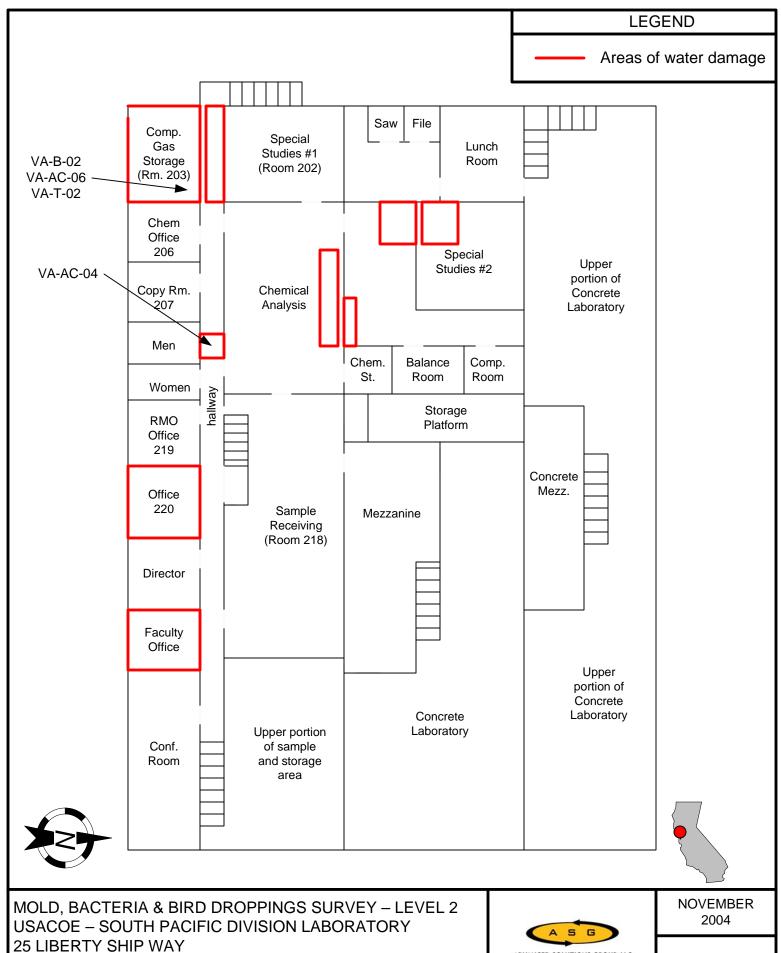
Analyzed By: Sampled By:

Type of Analysis:

Received By: Relinquished By:

APPENDIX C SAMPLE LOCATION DRAWINGS





SAUSALITO, CA 94965 PROJECT NO. U04-2732



FIGURE 2

APPENDIX D

PHOTOGRAPHIC DOCUMENTATION



Photo 1 – Subject Site



Photo 2 – bird droppings – level 1 – north side



Photo 3 – dead bird – level 1 – north side



Photo 4 – water damage – level 1 – wood doorway



Photo 5 – water damage – level 1 – SE sheet rock ceiling



Photo 6 – water damage – level 1 – south sheet rock ceiling



Photo 7 – water damage & significant growth – level 1 – engineer's office



Photo 8 – level 1 – broken windows – water intrusion



Photo 9 – water damage – level 2 – west ceiling tile



Photo 10 – water damage – level 2 – carpet – south side



Photo 11 – water damage – L2 – Chemical Analysis Rm.



Photo 12 – water damage – level 2 – s. hallway



Photo 13 – water damage & significant growth – level 2 – southwest offices and hallway



Photo 14 – water damage – level 2 – copy room



Photo 15 – water damage – level 2 – room 220



Photo 16 – water damage – level 2 – SEC room

APPENDIX E

CREDENTIALS

The

American Board of Industrial Hygiene^{*} ABIH ^{*}



organized to improve the practice of Industrial Hygiene proclaims that

Terence 1. Bleckner

having met all requirements through education, experience, and examination, is hereby certified in the

COMPREHENSIVE PRACTICE of INDUSTRIAL HYGIENE

and has the right to use the designations

CERTIFIED INDUSTRIAL HYGIENIST

GENING VI

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	-		ı

June 29, 2000

Chairman ABIH

7973 CP

Secretary ABI

APPENDIX F MICROBIOLOGICAL SPECIES INFORMATION