



May 7, 2024

Dare County Schools  
Ian Adams  
3020 S. Wrightsville Avenue  
Nags Head, NC

**Re: Limited Indoor Air Quality Sampling with Airborne Fungal Sampling at Manteo Elementary School  
LRC Project – 24-2413**

At your request, on May 2, 2024, LRC Indoor Testing & Research, Inc. (LRC) performed a limited environmental fungal Indoor Air Quality (IAQ) sampling that included airborne fungal sampling at the property listed above.

LRC performs all water-damage and fungal investigations with sampling and recommendations in accordance with guidelines published in *Bioaerosols: Assessment and Control*, by the American Conference of Governmental Industrial Hygienists (ACGIH), in *Mold Remediation in Schools and Commercial Buildings* by the United States Environmental Protection Agency (USEPA), and in the currently recognized and accepted industry standards including the ANSI/IICRC S500 *Standard and Reference Guide for Professional Water Damage Restoration*, Fourth Edition (S500) and the ANSI/IICRC S520 *Standard and Reference Guide for Professional Mold Remediation*, Third Edition (S520).

Our inspection included the following:

1. Measure temperature and relative humidity indoors and outdoors.
2. Collect representative non-viable spore trap air samples indoors and one outdoors for comparison.
3. Collect representative Particle Counts, Carbon Dioxide and Carbon Monoxide counts at locations where air samples are collected.
4. Provide a written report describing the survey results and comparing those results to accepted guidelines and directives. This report includes a summary of data and Certificates of Laboratory Analysis.

## BACKGROUND

This inspection was limited to the collection of non-viable spore trap air samples, particle counts, CO and CO<sup>2</sup> measurements that were collected from locations selected at random throughout the building. Descriptions in this report are based on looking at the structure from the front main entrance door. Moisture measurements and visual inspections were not conducted on this day.

## SAMPLING METHODOLOGY

### Air Samples:

Currently there are no regulations regarding acceptable airborne fungal levels. Airborne fungal spores are ubiquitous in the outdoor and indoor environment. The guidelines followed in this report for the assessment and/or remediation of airborne and surface fungi are published in *Bioaerosols: Assessment and Control*, by the American Conference of Governmental Industrial Hygienists (ACGIH), in *Mold Remediation in Schools and Commercial Buildings* by the United States Environmental Protection Agency (USEPA), in *Recognition, Evaluation, and Control of Indoor Mold* by the American Industrial Hygiene Association (AIHA), and in the ANSI/IICRC S520 *Standard and Reference Guide for Professional Mold Remediation*, Third Edition (S520). Airborne fungal assessments are performed by comparing results from volumetric samples taken indoors to samples taken outdoors. Airborne fungi levels in non-problem indoor environments generally are less than or approximately the same as that outdoors and also show a similar composition and/or taxonomic predominance. Problems are usually implicated in the indoor air when one or more fungal genera or species are present in a much greater concentration indoors compared to outdoors. Sampling results are shown in the Certificates of Laboratory Analysis attached to this report. Results are discussed below.

### Temperature and Relative Humidity

Temperature and relative humidity readings were recorded by the Lighthouse Handheld 3016 Particle Counter. ASHRAE Standard 55 for thermal comfort suggested that the indoor temperature should be between 73°F to 79°F in the summer and 68°F to 75°F in the winter and that the indoor relative humidity should be between 20% to 65%.

The temperature and relative humidity are summarized in Table A below. The relative humidity met the current ASHRAE Standard to maintain indoor relative humidity below 65%.

## SAMPLING RESULTS

### Total Non-Viable Spore Air Sample Results:

Representative samples were taken for total airborne fungal spores with a calibrated Buck spore trap. Total airborne fungal spore sample volumes were 75-liters. The outdoor total fungal spore level (Sample 21) was measured at Spores/m<sup>3</sup> and was comprised of *Cladosporium* (%), Smuts (%), Ascospores (%), Basidiospores (%), *Penicillium/Aspergillus* group (%), Hyphal Elements (%), and 1% or less of various other fungal spores. The air sample results are summarized below in Table A.

**Table A – Air Sampling Results**

Sample #	Location	Total Airborne Spore Count (Spores/m <sup>3</sup> )	*Non-Fungal Background Particulate Level	Temperature °F	Relative Humidity %
01	Main Hall at Entrance	213	Moderate	80	41
02	Hall at B100	160	Moderate	76	49
03	Hall at B103	133	Low-moderate	74	51
04	CR B105	533	Heavy	74	47
05	CR B111	173	Low	75	49
06	Hall at B116	253	Moderate-heavy	74	52
07	Hall at B119	267	Moderate-heavy	74	52
08	CR B123	107	Low-moderate	73	52
09	CR B116	213	Low-moderate	75	49
10	Hall at 217	160	Low-moderate	74	51
11	Hall at 209	120	Low-moderate	76	47
12	Hall at 206	107	Low-moderate	75	48
13	CR 205	67	Low	75	50
14	Hall at B219	53	Low-moderate	74	48
15	CR B220	27	Low	74	54
16	Hall at CR C108	200	Moderate	75	51
17	CR D128	27	Low	75	47
18	CR D131	120	Low	73	47
19	Media	147	Low-moderate	72	42
20	Gym	173	Low-moderate	71	51
21	Outdoor Air	10,347	Low-moderate	86	60

\*The Background Particulate Level refers to non-fungal debris seen in the air sample; such as skin cells, hair, fibers, dust, dirt, etc.

The total fungal spore counts in the areas sampled indoors were lower than that found in the outdoor air. The types of fungal spores found in the indoor air samples were all common outdoor-type fungi present in low concentrations with no spikes in water loss fungi. Therefore, the results suggested a normal indoor fungal ecology in the areas sampled. Counts are somewhat higher in this school probably due to the school being occupied and doors being opened to the outdoors during testing.

The particulate in the indoor air samples was in the low to moderate-heavy range. The particulate that we see in the microscope at the magnification used is usually called ‘course particulate’ and consist of many things and can include the following: dirt, dust, mold, pollen, fiber, hair, skin cells, dust mites and other insects. Fine particulates (to include VOC’s – volatile organic compounds) are not seen with the magnification used for these samples.

#### Particle Count

A Lighthouse Handheld 3016 Particle Counter was used to measure the levels of airborne particulates. Airborne particles are solids suspended in the air. In a commercial setting, particle

levels should be significantly less than outside readings due to better filtration and better dilution with outside air.

Particle Count:

Table B below summarizes the minimum, maximum, and average particle count in the building in each size category during the sampling period. Each time one liter of air was drawn into the Particle Counter, and the Particle Counter calculated all particles that are greater than or equal to the particle size indicated in the size categories.

**Table B – Indoor Particle Count Sampling Results**

	<b>0.3 micron</b>	<b>0.5 micron</b>	<b>1.0 micron</b>	<b>2.5 micron</b>	<b>5.0 micron</b>	<b>10.0 micron</b>
Minimum	2,807	50	29	10	0	0
Maximum	11,206	1,693	1,296	1,038	587	144
Average	7,416	502	226	147	94	19

Table C below summarizes the average particle count in the outdoor air at the time of sampling. Each time one liter of air was drawn into the Particle Counter, and the Particle Counter calculated all particles that are greater than or equal to the particle size indicated in the size categories. In addition, temperature and relative humidity information were included.

**Table C – Outdoor Particle Count Sampling Results**

	<b>0.3 micron</b>	<b>0.5 micron</b>	<b>1.0 micron</b>	<b>2.5 micron</b>	<b>5.0 micron</b>	<b>10.0 micron</b>
Outdoor Air Average	12,918	1,930	745	178	8	1

Carbon Monoxide and Carbon Dioxide

A TSI IAQ-Calc Indoor Air Quality Meter (Model 7545) was used to measure the concentrations of CO and CO<sub>2</sub>. CO is a dangerous gas caused by incomplete combustion. The level of CO in an indoor environment should be low (none detected to 4 parts per million [ppm] depending on fuel sources used indoors) or same as outdoors. CO<sub>2</sub> is commonly used as an indicator of ventilation adequacy. American Society of Heating, Refrigeration, and Air Conditioning Engineers (ASHRAE) Standard 62.1 stated that the indoor concentration of CO<sub>2</sub> should be less than 700 ppm over the outdoor ambient air, which typically is around 400 ppm.

The average CO concentration indoors was similar to the CO concentration outdoors. In addition, the indoor CO concentrations were measured at low ppm in all locations. Therefore, the results suggested that the indoor CO concentrations were within normal ranges. Individual results area available upon request.

Therefore, the results suggested that the ventilation in the areas sampled was adequate.

## CONCLUSIONS

Results as reported by LRC apply only to the day of this inspection. LRC cannot and does not warrant that other parts of the structure were completely free or that the structure will remain free in the future from hidden sources of moisture or fungal contamination.

This inspection was limited to indoor air quality parameters. The indoor air samples did not show a fungal amplification or an altered indoor fungal ecology on this day. Relative humidity was in the recommended range. Particle counts were generally lower than that found in the outdoor air. Carbon Dioxide and Carbon Monoxide levels were within the normal recommended range. Total fungal spore counts in some air samples were higher than found in the other buildings and particle counts in some tests were higher than in other buildings. This is most likely due to the interior of this building being exposed to the outdoors during the collection of samples. The total numbers were still in the low range and the spores found indoors were similar to those found outdoors.

If you have any questions or concerns, please do not hesitate to contact us.

Sincerely,



Cathy Richmond, B.S.  
LRC Indoor Testing & Research



Tony Richmond, BBA, CAI, WRT  
LRC Indoor Testing & Research