Influence of indoor conditions on microbial diversity and quantity in schools

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Abstract. HVAC systems consumes 37 % of the electricity in educational buildings. Energy consumption varies depending on the ventilation strategy. School buildings have a responsibility of ensuring energy performance and maintenance of indoor environmental quality for its occupants. This study assesses the efficiency of the ventilation system in terms of the microbes present in the spaces. Microorganisms and particulate matters were sampled in classrooms, locker rooms, libraries in two high schools and an elementary school. Six bacterial species and seven fungal species were identified. The most abundant microorganisms were *Staphylococcus* sp., *Bacillus* sp., and *Micrococcus* sp. Elementary school, especially kindergarten, showed higher microbial concentration and particulate matter as a result of higher human activity. Microbial concentration was influenced by the types of room in the same building with the same ventilation system. Canonical correspondence plot (CCA) determined that gender and school type have significant effects. The presence of *Bacillus* sp., *Aerococcus* sp., *Corynebacterium* sp., and *Penicillium* sp. was significantly related to gender, while *Staphylococcus* sp. and *Alternaria* sp. were related to type of school. The presence of *Aerococcus* sp. and *Micrococcus* sp. depended on the distance from the vent location. Relatively lower number of *Aerococcus* sp. was detected when the distance between sampling site and vent was less than 2 m. *Micrococcus* sp. was generally detected when the distance was < 3.5 m. The distance from the door was not significant.

1 Introduction

Poor air quality causes health problem [1, 2], discomfort [3, 4], and deterioration of learning performance [5]. More and more buildings are being designed as air-tight structures to save energy, however, this could lead to degradation of indoor air quality through containment of contaminated air in dead zones in the room, or introduce outdoor pollutants indoors depending on the ventilation condition [6, 7]. Even though ventilation systems have evolved over decades, poor air quality persists. Therefore, appropriate ventilation strategy is required to maintain air quality and prevent waste of energy.

According to the data from EIA (Energy Information Administration, US), HVAC systems consume most of the electricity following lighting, refrigeration, and computing in educational buildings in USA [8]. Schools spend approximately \$130 per student for electricity [8]. School's median energy costs based on EPA data are \$1.30/ft² in elementary school, \$1.38/ft² in middle school, and \$1.35/ft² in high school, respectively [9]. Educational buildings which include classrooms, preschool/day-care, elementary/middle/high school, and college/university consume 37 % of total electricity consumption for HVAC systems [8]. Energy usage in schools is not only related to the schools' operational

expenditure but is also connected to reducing energy consumption and CO₂ emission. School as a public building has a responsibility of energy performance and indoor environmental quality [10]. A recent study has predicted that the total world energy usage will increase by 56 % from 2010 to 2040 [11]. Buildings consume 50 % of total building energy usage and 20 % of total energy consumption in USA to operate HVAC systems [12]. According to a previous study comparing energy usage of offices in U.S. cities, energy consumption reduced 1.5 % ~ 29.0 % when the ventilation strategy changed from mechanical to natural ventilation [13]. HVAC systems have become essential for thermal comfort and desired indoor air quality. The increase of building's energy consumption is inevitable. Since different school levels have different human density and occupational time, consequently, different energy consumption [10], schools' requirements need to be considered before determining an energy efficient ventilation strategy which will also ensure health and productivity.

Several studies have investigated the effect of ventilation system using carbon dioxide (CO₂) or particulate matter as tracers/indicators [14, 15], but few studies have assessed the impact on indoor microbes. Microorganisms in an indoor space reflect the combined

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effects of the presence of occupants, activities, environmental parameters and specific characteristics of the space. For instance, human-related bacteria such as *Staphylococcus* sp. and *Micrococcus* sp. are frequently found in building environment [16]. Different age groups have different microbial flora [17]. Since *Lactobacillus* sp. is a vaginal microorganism, it has been frequently detected in female related spaces while *Corynebacterium* sp. has been found in male-related rooms [18]. Meadow and his colleagues confirmed that individuals have their own biological cloud and release their distinctive microbes in the spaces [19]. Human activity changes the indoor microbial communities and concentrations [20].

Although numerous studies have found that the indoor microorganisms differ depending on the indoor conditions, current ventilation standards do not use microbial presence as a way of assessing, 'adequate ventilation' [21]. Indoor microbial communities were highly influenced by the ventilation strategy [22]. In this study, the relationship between indoor microbial characteristics and engineering controls have been investigated.

2 Materials and methods

Microorganism and particulate matter were obtained from sampling in classrooms, locker rooms, and libraries in two high schools and an elementary school. The spatial distribution of the microbial quantity and diversity was assessed in relation to the interior orientation of engineering controls (such as vents) and select interior 'décor' such as placement of desks, book shelves, sinks etc. All the schools were always mechanically ventilated and had windows (closed) except for the locker rooms which had no windows. Indoor microorganisms were collected through passive air sampling for 60 minutes using open Petri dishes filled with tryptic soy agar (TSA). The passive air sampling method follows the standard 1/1/1 scheme (1 hr exposure, 1 m above the floor, and 1 m far from the wall) [23]. The locations of the Petri dishes in this study were varied in relation to the HVAC system vents in order to investigate the influence of the ventilation system. Optical Particle Sizer 3330 (TSI Inc., MN, USA) was used to measure particle size distribution. Indoor environmental conditions were monitored using ABM-200 (Airflow and Environmental Meter, CPS Products, Inc., FL, USA). Temperature and humidity were recorded typically at ~ 26 °C and ~ 47 %, respectively. Samples were collected from October 2017 to February 2018. Occupancy and activity were recorded. The degree of activity was rated in three levels: -1 (no activity, unoccupied), 0 (normal activity: sitting, talking, or walking), and +1 (high activity: running or most people is moving around).

Sampled Petri dishes were incubated at the room temperature for 7 days. The shape, colour, and texture of the colony were evaluated for microbial identification. Colony forming units per m³ of indoor air (CFU/m³)

were calculated to quantify the concentration of indoor microbes applying equation (1) [24, 25]:

$$CFU / m^3 = \frac{5 \times n \times 10^4}{A \times t} \quad (1)$$

where n is the number of colonies on the Petri dish, A is the surface area of the Petri dish (cm²), and t is exposure time (min).

Overall 160 air samples were collected. For the data reliability, duplicate samples were taken at the same sampling location. R 3.5.1 and Minitab (Minitab Inc., PA, USA) were used for statistical analysis. The data significance was evaluated at p level of 0.05. Canonical correspondence analysis (CCA) was applied to determine the influence of sampling condition and distance from the vent. CCA represented the relationships between biological species and their environment [26]. One-way ANOVA test was applied to evaluate the differences among the samples.

3 Results and Discussion

3.1. Indoor air quality

Indoor air quality was evaluated with the concentrations and types of detected microorganism, and particulate matter. Overall 160 air samples were collected. Among the samples, six different bacteria and seven different fungi were identified. Detected bacterial species were *Bacillus* sp., *Staphylococcus* sp., *Curtobacterium* sp., *Aerococcus* sp., *Micrococcus* sp., and *Corynebacterium* sp. Detected fungi were *Rhodotorula* sp., *Cochliobolus* spp., *Pithomyces* sp., *Alternaria* sp., *Ascochyta* sp., *Aspergillus* sp., and *Penicillium* spp. The most abundant microorganisms were *Staphylococcus* sp., *Bacillus* sp., and *Micrococcus* sp. This result is in accordance with other studies in the literature. *Staphylococcus* sp., *Bacillus* sp., *Corynebacterium* sp., and *Micrococcus* sp. are frequently found in indoors such as residences, hospitals, offices, or schools since they are human-related bacteria [20, 27-29]. A study conducted in air-conditioned office buildings found *Micrococcus* sp., *Bacillus* sp., *Staphylococcus* sp., *Penicillium* sp., *Aspergillus* sp., and *Rhodotorula* sp. from the air and settled dust samples [28].

Indoor microbial concentration in each school and space is described in Fig. 1(A). Elementary school showed a higher microbial concentration, especially in kindergarten, than high school. The averaged concentrations were 135 CFU/m³ in high school and 293 CFU/m³ in elementary school, respectively. The ANOVA test confirmed that human activity ($p < 0.05$) had a significant influence on the microbial concentration. No activity was reported in the 5th grade classroom. Fig. 1(A) shows that the number of microbes detected is higher in male's locker room even when both locker rooms had similar activities and occupancy rates. Gender did not significantly impact the number of

detected microorganisms (CFU/m³) and microbial species ($p>0.05$). Fungal and bacterial concentration were highly influenced by the types of room ($p<0.05$). Relatively higher fungal level was measured in library (median: 23 %) than other rooms (median: 8 % in locker room and 11 % in classroom, respectively).

Fig. 1(B) shows the concentration of particulate matter (PM_{2.5}). In accordance with the indoor microbial concentration, Fig. 1(A), particulate matter was higher in kindergarten.

Fig. 1. (A) Microbial concentration (CFU/m³) and (B) PM_{2.5} (µg/m³) in each sampling location. H : High school, E : Elementary school, ML : Male's locker room, FL : Female's locker room, CL : Classroom, L : Library, K : Kindergarten, 2nd : 2nd grade classroom, and 5th : 5th grade classroom.

3.2 Influence of sampling condition

Canonical correspondence plot (Fig. 2) showed and confirmed the relation between sampling location and microbial community. The types of room are classified into locker room, classroom, and library. Adolescence and children represented two different age groups. The three schools are: two high schools and an elementary school. Gender indicates: female only, male only, and both genders present. CCA results show that gender and types of school had a significant effect ($p<0.05$). As shown in Fig. 2 the presence of *Bacillus* sp., *Aerococcus* sp., *Corynebacterium* sp., and *Penicillium* sp. is highly related to gender, while *Staphylococcus* sp. and *Alternaria* sp. are related to types of school. Numerous studies have found that *Corynebacterium* sp. is dominant in male-occupied rooms [17, 18, 30]. *Bacillus* sp. is often found in indoor air and dust samples [29],

Aerococcus sp. is known as an environmental species [31], and *Penicillium* sp. is one of the most common indoor fungi [31]. However, previous studies have not investigated the relation between *Bacillus* sp., *Aerococcus* sp., and *Penicillium* sp. and gender.

The presence of *Aspergillus* sp., *Rhodotorula* sp., *Ascochyta* sp., *Cochliobolus* sp., *Micrococcus* sp., and *Curtobacterium* sp. was not associated with the types of room, school, age, and gender. *Aspergillus* sp. is frequently detected indoors, while *Cochliobolus* sp. is an outdoor-related fungus [28, 31, 32]. *Rhodotorula* sp. is abundant in indoor and outdoor air samples [33].

Fig. 2. CCA plot showing overall detected microorganisms among different sampling conditions. B1 : *Aerococcus* sp., B2 : *Bacillus* sp., B3 : *Corynebacterium* sp., B4 : *Curtobacterium* sp., B5 : *Micrococcus* sp., B6 : *Staphylococcus* sp., F1 : *Alternaria* sp., F2 : *Ascochyta* sp., F3 : *Aspergillus* sp., F4 : *Cochliobolus* sp., F5 : *Penicillium* sp., F6 : *Pithomyces* sp., and F7 : *Rhodotorula* sp.

3.3 Influence of vent location

The spatial distribution of microbial diversity and quantity was investigated to determine the influence of vent location. Fig. 3 shows the location of sampling sites, vents, doors, and windows. The ratio of detected bacteria and fungi was not significantly influenced by the distance from vents ($p>0.05$). However, the presence of *Aerococcus* sp. and *Micrococcus* sp. were significantly influenced by the vent location ($p<0.05$). *Aerococcus* sp. is outdoor-related species and *Micrococcus* sp. is associated with human presence [31]. Relatively lower number of *Aerococcus* sp. was detected when the distance between sampling site and vent was less than 2 m. *Micrococcus* sp. was generally detected when the distance was lower than 3.5 m. Higher number of *Micrococcus* sp. was observed over 4 m away from the closest vent. This result demonstrates a possibility of the influence of the zones in the room with different range of velocity magnitudes influencing the presence of microbes. The number of detected bacteria (CFU/m³) was highly impacted by the distance from the closest window ($p<0.05$). However, each microbial species was

Fig. 3. Schematic diagram of the location of vents, doors, windows, and sampling sites.

not significantly influenced by the window location ($p > 0.05$). The distance between sampling site and the door was not significant among all the microbial species ($p > 0.05$).

4 Conclusion

The study investigated the microbial concentration and particulate matter in different sampling conditions. Samples were collected from two high schools and an elementary school. Air sampling from locker rooms, classrooms, and library was investigated to identify the influence of age, gender, space occupancy, and activity. Six different bacteria and seven different fungi were detected. Bacterial species were *Bacillus* sp., *Staphylococcus* sp., *Curtobacterium* sp., *Aerococcus* sp., *Micrococcus* sp., and *Corynebacterium* sp. Fungi were *Rhodotorula* sp., *Cochliobolus* spp., *Pithomyces* sp., *Alternaria* sp., *Ascochyta* sp., *Aspergillus* sp., and *Penicillium* spp. Higher microbial concentration was observed in kindergarten where the highest activity rate was measured. In agreement with previous studies, male's locker room showed relatively higher concentration of microorganisms and particulate matter [18, 19]. Our CCA result confirmed that the microbiome at different schools (high school and elementary school) and gender were significantly different. *Staphylococcus* sp. and *Alternaria* sp. are highly related to types of

school, while the presence of *Bacillus* sp., *Aerococcus* sp., *Corynebacterium* sp., and *Penicillium* sp. is mainly related to gender. The analysis determined that the ratio of bacteria and fungi was not significantly influenced by the vent location. However, several bacterial species showed differences depending on the distance from the closest vent, *Aerococcus* sp. and *Micrococcus* sp. The distance from the door was not significant among all the microbial species. *Aerococcus* sp. and *Micrococcus* sp. were detected in spaces with natural and mechanical ventilation system [34, 35].

The results of the analysis indicate that while the spaces were continually mechanically ventilated maintaining acceptable humidity and indoor temperature conditions, the effects on indoor microbial diversity and quantity needs further evaluation. Outdoor related microorganisms such as *Aerococcus* sp. and *Bacillus* sp. were continuously detected in the mechanically ventilated spaces. There was no impact of window or door locations to the presence or absence of the detected species. Understanding how other ventilation approaches such as natural, hybrid or personalized ventilation systems impact the spatial distribution of the microbiome can lead to more energy efficient designs and healthy indoor conditions. The characteristics of the microbiome can also have an influence, which remains to be investigated.

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