Integrated Environment-Occupant-Pathogen Information Modeling to Assess and Communicate Room-Level Outbreak Risks of Infectious Diseases

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4 Abstract

5 Microbial pathogen transmission within built environments is a main public health concern. The 6 pandemic of coronavirus disease 2019 (COVID-19) adds to the urgency of developing effective 7 means to reduce the pathogen transmission in mass-gathering public buildings such as schools, 8 hospitals, and airports. To inform occupants and guide facility managers to prevent and respond 9 to infectious disease outbreaks, this study proposed a framework to assess the room-level 10 outbreak risks in buildings by modeling built environment characteristics, occupancy 11 information, and pathogen transmission. Building information modeling (BIM) is exploited to 12 automatically retrieve building parameters and possible occupant interactions that are relevant 13 to pathogen transmission. The extracted information is fed into an environment pathogen 14 transmission model to derive the basic reproduction numbers for different pathogens, which 15 serve as proxies of outbreak potentials in rooms. A web-based system is developed to provide 16 timely information regarding outbreak risks to occupants and facility managers. The efficacy of 17 the proposed method was demonstrated by a case study, in which the building characteristics, 18 occupancy schedules, pathogen parameters, as well as hygiene and cleaning practices are 19 considered for outbreak risk assessment. This study contributes to the body of knowledge by 20 computationally integrating building, occupant, and pathogen information modeling for infectious 21 disease outbreak assessment, and communicating actionable information for built environment 22 management. 23

24 Keywords

Building Information Modeling; Pathogen Transmission; Outbreak Risk; COVID-19; Health
 26

27 **1. Introduction**

28 People spend most of their time in buildings, including homes, offices, schools, stores,

29 restaurants, theaters, and many others. The buildings become hotspots for pathogen

30 transmission and exposure, decimating populations through epidemics and everyday infections.

31 The disastrous impacts of infectious diseases highlight the urgent need to reduce the

- transmission of pathogens, and their exposure to occupants in buildings. Humans can be
- infected by microbial pathogens via contacting contaminated objects, referred to as fomites.
- Fomite-based transmission is an important route in built environments for transferring diseasecausing microbiomes to a new human host [1]. The mechanism of fomite-mediated transmission
- causing microbiomes to a new human host [1]. The mechanism of fomite-mediated transmission
 involves three steps. First, a surface is contaminated by infectious pathogens. The
- 37 contamination can occur when an infected person touches the surface or bioaerosols containing
- 38 pathogens settle down on the surface. Second, a person touches a contaminated surface with
- 39 his or her hand, transferring the pathogens to the hand. Third, the person touches susceptible
- 40 sites (mucous membranes) on his or her body with the contaminated hand, which inoculates the
- site with pathogens, resulting in potential infection. A recent study [2] found that contamination
- 42 of a single doorknob or tabletop can spread the infectious pathogens to other commonly
- touched objects, exposing 40-60% of people in the buildings.
- 44
- 45 Many pathogens can be transmitted via fomites. For example, during flu seasons, measurable
- 46 levels of influenza virus can be found on all common building surfaces [1,3], underlining the
- 47 importance of fomite in influenza transmission. The pandemic of coronavirus disease 2019
- 48 (COVID-19) has swept the entire world with more than 29.6 million infections and 935,898
- 49 deaths as of September 16, 2020 [4]. During the pandemic of COVID-19, viable severe acute
- 50 respiratory syndrome coronavirus 2 (SARS-CoV-2) can be detected on various surfaces. High

51 concentration of SARS-CoV-2 are found on surfaces in healthcare facilities where COVID-19

52 patients are treated [5,6]. Norovirus can also be transmitted via fomite [7,8], causing 93% of

53 nonbacterial gastroenteritis outbreaks in the U.S. In addition, pathogens including

54 staphylococcus aureus, Clostridium difficile, Staphylococcus aureus, Pseudomonas aeruginosa,

- 55 Pseudomonas putida, and Enterococcus faecalis can also be transmitted by surface contact [9].
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57 Models have been developed for environmental risk assessment and environmental infection 58 transmission [10]. Fomite-mediated transmission has received increased attention [11,12]. To 59 assess pathogen transmission to susceptible hosts, the models such as the environmental 60 infection transmission system modeling framework consider the dynamics of contact and pathogen transfer between individuals via their hands and fomites, pathogen persistence in the 61 62 environment, pathogen shedding, and recovery of infected individuals. Studies [13-15] also 63 exploited experimentation approaches to measure the transfer of microbiomes between fomites 64 and humans. The measured microbiological and epidemiological data can be used to assess 65 the transmissibility of the pathogens and used in the models for risk assessment. Despite research efforts made in epidemiology, the modeling of building, occupant, and pathogen has 66

67 not been well linked to predict the microbial burdens and outbreak risks.

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69 Predicting outbreak risks in buildings and communicating actionable information to occupants 70 and facility managers are challenging. First, pathogen burdens could differ considerably in 71 rooms even in the same building. Building design and operation can influence indoor microbial 72 communities [16,17]. The microbial communities in different rooms with different functionalities 73 and spatial configurations are found to exhibit very different patterns [18,19]. Occupancy also 74 significantly affects the microbial communities in buildings. For example, bacteria taxa in spaces 75 with a high occupant diversity and a high degree of physical connectedness are different from 76 that in spaces with low levels of connectedness and occupant diversity [16]. Humans can 77 transfer microbiomes including pathogens to the environment via skin-to-surface contact and 78 direct shedding of large biological particles [20.21]. The microbial exchange between occupants 79 and surfaces can occur in both directions [12]. With different uses and occupancy levels, 80 outbreak risks could vary depending on the locations in a building, underlining the need for a 81 spatially-adapted modeling approach. However, there lacks a computational modeling approach 82 to link the coupled physical-biological processes of buildings, occupants, and pathogens to 83 automatically assess the spatially-varying infection and outbreak risks at unprecedented scales. Therefore, it is imperative to establish the computational framework to guickly compute the risk 84 85 in buildings to inform end-users and guide adaptive operations.

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87 Second, due to the absence of an effective means for information communication, end-users 88 have limited access to easy-to-understand information regarding the outbreak risks to make 89 necessary interventions. Building information modeling (BIM) uses standardized machine-90 readable information created or gathered about a facility throughout its lifecycle for all stakeholders involved [22]. Information can be extracted from building information models, as 91 92 they are the shared digital representations of physical and functional characteristics of any built 93 objects [22]. In addition, BIM has also been used as a powerful tool to visualize the parametric 94 building model with computed rich information [23]. However, to the authors' best knowledge, 95 existing studies have not explored the capability of BIM in environmental pathogenic infection assessment, and leverage BIM as a platform to visualize and communicate outbreak risk 96 97 information to end-users for facility management. 98 99 This study aims to develop a framework for room-level outbreak risk assessment based on

100 integrated building-occupancy-pathogen modeling to mitigate the spread of infectious disease in

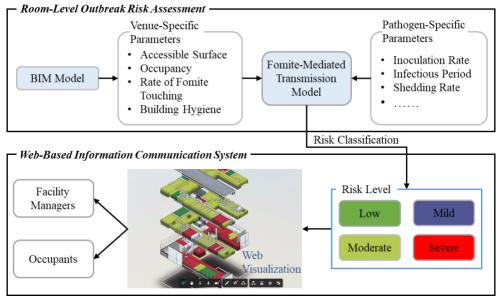
101 buildings. The rationale is twofold. First, buildings are highly heterogeneous with a variety of

102 compartments of distinctive functionalities and characteristics, providing diverse habitats for 103 humans and various pathogens [17,18]. Modeling the pathogen transmission and exposure 104 within a building at the room level will provide useful information at an unprecedented resolution 105 to implement appropriate disease control strategies. Second, the spread of infectious diseases 106 can be mitigated if occupants and facility managers have adequate and timely information regarding the outbreak risks within their buildings. Communicating actionable information to 107 108 occupants and facility managers through an easily accessible interface will help occupants to 109 follow hygiene and social distancing practice, and help facility managers to schedule disinfection 110 for rooms with high outbreak risks.

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112 2. Method

113 To address the knowledge gaps, a novel environment-occupant-pathogen modeling framework 114 and a web-based information visualization system are developed to assess the outbreak risks 115 and mitigate the spread of infectious diseases in buildings (Fig. 1). First, to assess the outbreak 116 risks, the fomite-based pathogen transmission model proposed in [24] is adopted in this study. 117 The limitation of the model is that the environmental parameters and occupant characteristics 118 are not automatically extracted and incorporated in the model, hindering the computation of the 119 spatially-varying environmental infection risks in buildings. To overcome this limitation, BIM is 120 exploited to automatically retrieve venue-specific parameters including building characteristics 121 and occupancy information that are relevant to pathogen transmission and exposure. Then, the 122 extracted building and occupant parameters are used with pathogen-specific parameters in a 123 human-building-pathogen transmission model to compute the basic reproduction number R_0 for 124 each room in a building. R_{θ} is used as a proxy to assess the outbreak risks of different infectious 125 diseases. Second, a web-based system is developed to enable information visualization and communication in an interactive manner to provide guidance for occupants and facility 126 127 managers. This study innovatively establishes the computational links among building, 128 occupant, and pathogen modeling to predict outbreak risks. The risk prediction for spatially and 129 functionally distributed rooms in a building provides useful information for end-users to combat 130 and respond to the spread of infectious diseases, including the seasonal flu and COVID-19. The 131 developed method and system add a health dimension to transform the current building 132 management to a user-centric and bio-informed paradigm.



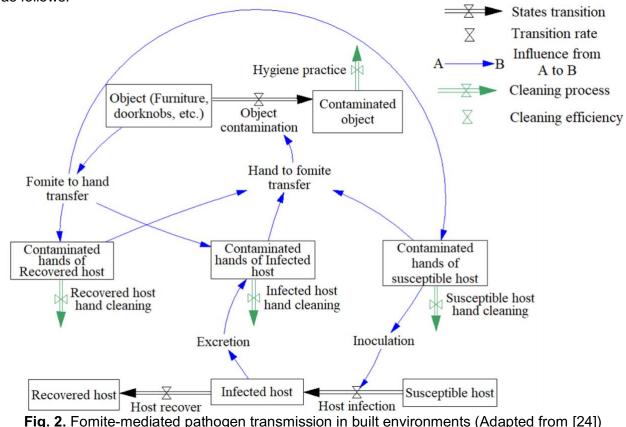
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Fig. 1. Research Framework

136 2.1. Room-Level Outbreak Risk Assessment

137 Employing the model proposed in [24], individuals are divided into three categories, i.e., 138 susceptible, infectious, and recovered. Pathogens may survive outside the host and 139 contaminate either object surfaces or human hands. The pathogen exchange in built 140 environments can occur through hand-surface contacts. Contaminated hands of hosts can 141 contaminate surfaces of accessible objects, while susceptible people can get infected by 142 touching the contaminated surfaces and self-inoculation. Fig. 2 shows the fomite-mediated 143 pathogen transmission process in built environments. Building characteristics, occupant 144 behavior, and pathogen parameters collectively determine the transmission ability through the 145 dynamic processes of pathogen inoculation, fomite touching and transfer, pathogen excretion, 146 pathogen decay, individual recovery, and building disinfection and individual hygiene. 147 Characteristics of the built environment (e.g., contaminated objects and building hygiene) and occupant behavior (e.g., fomite touching and hand cleaning) are critical in the process of fomite-148 149 mediated pathogen transmission in the built environment and are considered as venue-specific 150 parameters. In addition, the transmission efficiency of different diseases also depends on 151 pathogen-specific parameters, such as recovery rates and pathogen excretion. The

152 determination and acquisition of venue-specific and pathogen-specific parameters are detailed 153 as follows.



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157 2.1.1. Venue-specific parameters

158 Because venue-specific parameters vary across rooms with different functions and occupancy 159 levels, it is important to develop an effective means to accurately and automatically extract the

- 160 venue-specific parameters to assess the outbreak risks at the resolution of room level. A
- building information model captures the relationships among different elements in a building,
- and allows the storage and extraction of detailed geometric and non-geometric information in a

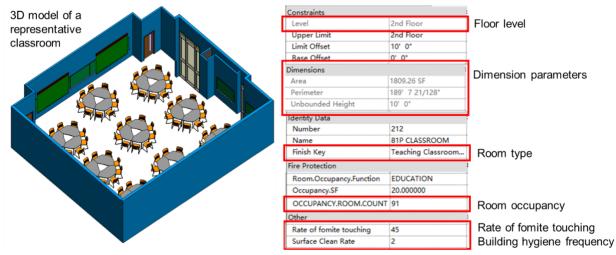
163 3D virtual representation. The non-geometric information includes semantic and topological 164 information, describing the attributes of elements and the relationship between components,

165 respectively [25]. Hence, it is feasible and efficient to extract venue-specific parameters from a 166 building information model.

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168 The BIM model can be divided into six Levels of Development (LOD) [26] that are suitable for 169 conceptual design (LOD 100), schematic design (LOD 200), design development (LOD 300), 170 construction documentation (LOD 350), fabrication and assembly (LOD 400), and maintenance 171 and operation (LOD 500). To effectively capture the characteristics of buildings and occupants, this study uses LOD 500 BIM model that reflects the as-built conditions regarding the geometry 172 173 information and non-graphical building attributes, as well as occupancy information. Fig. 3 174 shows an example of a representative classroom in the BIM model. For most public buildings 175 such as schools and hospitals, and particularly during the pandemic, the occupancy can be 176 predetermined and incorporated in the BIM model as attributes.

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178 179 180

Fig. 3. Building and Occupancy Information Modeling

181 The following venue-specific parameters will be extracted from the model. 182

183 1) Accessible surface. The surfaces of objects, including doorknobs, stair railings, tables, and chairs, which people frequently interact with are considered as accessible surfaces. The 184 accessible surface is computed as the summation of surface area of all touchable objects in 185 186 a room. The proportion of accessible surface λ is defined as the ratio of accessible surface 187 to the total area of surfaces within a room that includes both accessible surface and interior 188 surface. The calculation is shown in Eq. 1.

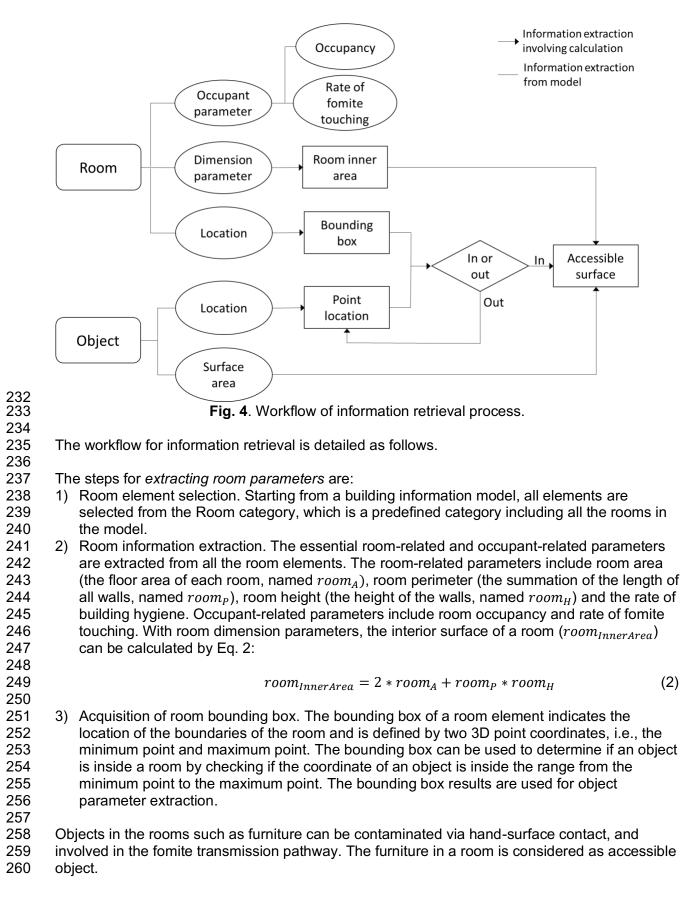
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$$\lambda = \frac{\sum Accessible \ surface \ area}{\sum Accessible \ surface \ area + Room_{InnerArea}}$$
(1)

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191 2) Occupancy. The occupancy is the number of individuals present in a room per day. In this 192 study, it is assumed that the occupancy of each room is predefined based on room capacity. 193 During the pandemic of COVID-19, many buildings such as university campus buildings and 194 office buildings have developed detailed occupancy schedules, which can be updated in the 195 BIM model and then extracted for analysis. Consistent with the prior study [24], it is 196 assumed that all individuals are identical within each room regarding susceptibility, contact 197 rates, and infectiousness as well as other individual characteristics. This assumption

- simplifies the model to capture the complex nature of pathogen transmission process. The
 only difference among the individuals is the state associated with them: Susceptible *S*,
 Infected *I*, or Recovered *R* during the pathogen transmission process.
- 201 202 3) Rate of fomite touching. The rate of fomite touching is the frequency that occupants interact 203 with the objects inside a room on an hourly basis. A higher frequency of interaction indicates 204 a higher possibility of pathogen transmission between objects and hosts. In this study, the 205 rate of fomite touching is determined based on different functionalities of the rooms 206 considering the primary age group present in the rooms. For example, classrooms and 207 offices in a school building are two main types of rooms considered in this study. It is 208 assumed that the rate of fomite touching in classrooms is higher than that in offices because 209 the occupants in classrooms are younger people who are more likely to interact with the 210 built environment. According to the observations in [27], an average rate of touching 211 common areas (e.g., chairs, desks, facilities) in a school office is 12 times per hour. 212 Therefore, in this study, the rate of fomite touching is set as 12 times per hour for offices, 213 and that for classrooms is set as 45 times per hour based on [28]. Furthermore, to incorporate the possible variation in different scenarios, a range of (0, 30) and (30, 60) is 214 215 considered for offices and classrooms, which also aligns with the setting in [28]. Analyses 216 will be conducted to examine the influence of the rate of fomite touching on outbreak risk. 217
- 4) Building Cleaning and Hand Hygiene. Building cleaning plays an important role in object decontamination. For fomite-mediated transmission, surface cleaning can significantly decrease the pathogen reproductive process. The frequency of building cleaning is determined by the adopted sanitation schedule of the building. Hand hygiene removes pathogens picked up from contaminated objects. For infected individuals, hand cleaning also removes pathogens excreted to hand, and thus, preventing contaminating objects through hand touching.
- In this study, a computational tool is developed based on Dynamo [29] to extract the geometry
 and properties of each room in a building, and to compute the corresponding venue-specific
 parameters. Fig. 4 shows the workflow of the information retrieval process. Lines in Fig. 4
 indicate direct information retrieval from the models and arrows indicate the information retrieval
 involving calculations.
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- 262 The steps for extracting the object parameters are:
- Furniture element selection. All the elements under the category "Furniture" are selected
 from the model. This category contains information of all the furniture in the model.
- 2) Furniture information extraction. The essential furniture parameters are extracted from all
 the furniture elements. The parameters include area (the surface area of furniture) and
 location (the point location of each furniture element). The location of furniture is
 transformed to a 3D point (a point with x, y, z coordinates) using a default function in
 Dynamo. The coordinates represent the location of the furniture.
- 272 3) Location relationship between room and furniture. For each room element, the coordinates
 273 of furniture in the model are compared with the coordinates of the room bounding box. This
 274 process checks the 3D location relationship between each room and furniture.
- Thereafter, the total furniture area in each room (Named $furniture_A$) is calculated by summing up the surface area of all furniture inside the room. The proportion of accessible surface (λ) of each room is calculated using Eq. 3.
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$$\lambda = \frac{furniture_A}{furniture_A + room_{innerArea}}$$
(3)

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2.1.2. Pathogen-specific parameters

Pathogen characteristics affect the transmission process through inoculation, excretion,
 inactivation (decay), and recovery. According to the study [24], Table 1 lists the pathogen specific parameters used in the fomite-mediated transmission model.

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Table 1. Description of pathogen parameters

Pathogen parameters	Symbol	Unit	Parameter description
Infectious period	$1/\gamma$	days	The period that an infectious individual can excrete and transmit pathogens
Shedding rate	α	pathogens/ (hours × people)	Infectious individual releases pathogens at rate α
Pathogen inactivation rate on surfaces	μ_F	1/hours	Pathogens decay at rate μ_F on surfaces
Pathogen inactivation rate on hands	μ_H	1/hours	Pathogens decay at rate μ_H on hands
Transfer efficiency from fomite to hand	$ au_{FH}$	1/touch	Pathogens transfer from fomite to hand at rate $ au_{FH}$
Transfer efficiency from hand to fomite	$ au_{HF}$	1/touch	Pathogens transfer from hand to fomite at rate $ au_{HF}$
Pathogen excreted to hand	$arphi_{H}$	unitless	The proportion that pathogens are shed on hands
Dose response of pathogens on mucosa	π	unitless	The infectivity of a pathogen
Inoculation rate	ρ	1/hours	Rate of touching mouth or other routes of infection

In this study, three pathogens, i.e., influenza, norovirus, and SARS-CoV-2 are considered.
Table 2 shows the parameter values used in the model. The pathogen-specific parameters of
the first two viruses are determined based on [24]. The parameters of SARS-CoV-2 were
determined based on a number of studies up to date. For the parameters that are still under
research, the values are set based on surrogate viruses and assumptions, which are described
as follows.

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Table 2. Values of pathogen-specific parameters of four viruses					
Pathogen-specific parameter	Influenza	Norovirus	SARS-CoV-2		
1/γ	6	15	8 [30]		
α	1E4	2.88E3	1.99E4 (1.8E3, 2.39E4)		
μ_F	0.121	0.288	0.059		
μ_H	88.2	1.07	0.8		
$ au_{FH}$	0.1	0.07	0.37		
$ au_{HF}$	0.025	0.13	0.14		
$arphi_{H}$	0.15	0.9	0.15		
π	6.93E-05	4.78E-04	6.58E-06 [31]		
ρ	15.8	15.8	15.8		

 Table 2. Values of pathogen-specific parameters of four viruses

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297 1) The inactivation rates on surfaces (μ_F) and hands (μ_H). The inactivation rate on surfaces is 298 determined based on the study [32], which provides the half-life of infectivity $(t_{0.5})$ on 299 surfaces under common temperature and relative humidity. The inactivation process of the 300 virus is assumed as a first-order kinetic model in this paper, and the inactivation rate is 301 calculated as $ln2/t_{0.5}$. Under the circumstance of 74°F and 40 of relative humidity, the estimated half-life of infectivity on surfaces is 11.78 hours, and the approximate inactivation 302 303 rate is 0.059 per hour. Due to the lack of exact data of μ_H , the parameter inactivation rate on skin of Middle East Respiratory Syndrome (MERS-CoV) is used in the paper, which is 0.8 304 305 per hour [33]. 306

- 307 2) Transfer efficiency from fomite to hand (τ_{FH}) and transfer efficiency from hand to fomite 308 (τ_{HF}) . The transfer efficiency coefficients are estimated using parameters of MERS-CoV in 309 [33] due to the absence of data. The transfer efficiency varies with surface materials. 310 Compared with porous surfaces (e.g. fabrics, clothes, and sponges), non-porous surfaces 311 such as desks, chairs, and door handles are more appropriate to represent the material of furniture surfaces considered in this paper. Thus, the transfer rates between hands and non-312 313 porous surfaces are used to indicate the transfer efficiency between hands and fomites. 314 According to the results in [33], τ_{FH} is set as 0.37, and τ_{HF} is set as 0.14.
- 3) Pathogen excreted to hand (φ_H) . Because the virus excretion behavior of SARS-CoV-2 such as coughing, sneezing, and exhaling is similar to the excretion behavior of influenza, φ_H of SARS-CoV-2 is estimated using the same parameter of influenza.
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4) Shedding rate (α). In the paper, coughing is considered as the primary way for virus shedding. The shedding rate is determined by the number of viruses in the respiratory tract that is shed via coughing per hour per infectious individual. The equation for shedding rate calculation is shown in Eq. 4.

$$\alpha = V_{droplet} \times F_{cough} \times N_{droplet} \times L \tag{4}$$

 $V_{droplet}$ indicates the volume per infectious droplet in cm^3 , F_{cough} is the coughing frequency 324 per hour, N_{droplet} is the number of droplets excreted per cough, L is the viral load in the 325 respiratory tract in copies/mL. According to [34], the viral load of SARS-CoV-2 for children 326 327 aging 0-22 is 6.2 log₁₀ RNA copies/ml, which is adopted in this study as the occupants are 328 primarily children in school buildings. Due to the lack of data, other parameters are 329 estimated using parameters of MERS-CoV in [33]. V_{droplet} is calculated considering the 330 largest diameter for infectious droplets that best fits the scenario of fomite transmission. The diameter is set as 100 μm . F_{cough} is set as 12 times per hour. $N_{droplet}$ is set to be 2000 per 331 332 cough. Based on the calculation above, α is set to be 1.99E4. Besides, as the accurate 333 shedding rate is still not well understood, it is assumed within the range of (1.8E3, 2.39E4), 334 where the lower bound is set according to [28], and the higher bound is set as 1.2 times of 335 the estimated value to allow potential higher shedding rate value.

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5) Dose response of pathogens on mucosa (π). The infectivity is determined based on study [31]. [31] found that the exponential model $p = 1 - \exp(-d/k)$ can well demonstrate the dose-response function of SARS-CoV-2, where the constant k ranges from 6.19E4 to 7.28E5. In the paper, k is set as 1.52E5, representing 50% of contribution from airborne particles to the total dose. π is set as the inverse of k, which is 6.58E6.

343 2.1.3. Risk Assessment

344 In epidemic dynamics, the basic reproductive number (R_0) is an estimation of a pathogen's 345 transmission ability of an infectious disease. R_0 is the expected number of cases generated by 346 one single infected person, supposing all other individuals are susceptible to the epidemic [35]. In this study, R_0 is used to represent the outbreak potential of each pathogen across different 347 348 rooms in the building. Given the fomite-mediated transmission model described in the previous section, R_0 is computed using the next generation matrix method [36], which consists of two 349 matrices, i.e., the matrix of disease transmission and matrix of host state transition. R_0 is 350 351 identified as the dominant eigenvalue of the product of the two matrices, computed using Eq. 5 352 proposed in [24].

353

$$\begin{cases}
R_{0} = R_{0,F} + R_{0,H} \\
R_{0,F} = \frac{a_{F}}{\gamma} P_{inoculation} P_{pickup} P'(0) \\
R_{0,H} = \frac{a_{H}}{\gamma} P_{inoculation} P_{pickup} P_{deposit} P'(0) \\
P_{inoculation} = \frac{\rho \chi}{\mu_{H} + \rho_{HF} + \rho \chi + \theta_{H}} \\
P_{pickup} = \frac{\frac{N\rho_{FH}}{1 - \frac{N\rho_{FH}}{(N\rho_{FH} + \mu_{F} + \theta_{F})}(\frac{\rho_{HF}}{(\mu_{H} + \rho_{HF} + \rho \chi + \theta_{H})}} \\
P_{deposit} = \frac{\rho_{HF}}{\mu_{H} + \rho_{HF} + \rho \chi + \theta_{H}}
\end{cases}$$
(5)

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354

356 $R_{0,F}$ represents direct fomite contamination route, $R_{0,H}$ is hand-fomite contamination route,

357 $P_{inoculation}$ is the proportion of pathogens that are self-inoculated to susceptible hosts; P_{pickup} is 358 the proportion of pathogens picked up by hands from fomites; $P_{deposit}$ is the proportion of 359 pathogens excreted to hands that are deposited to the fomites. P'(0) is the slope of the dose 360 function, indicating the infectivity of a dose of the pathogen.

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362 In the above equations, $a_F = \alpha (1 - \varphi_H) \lambda$, representing the rate pathogens excreted to surfaces, 363 where α is the shedding rate, φ_H is the proportion that pathogens are shed on hands, both

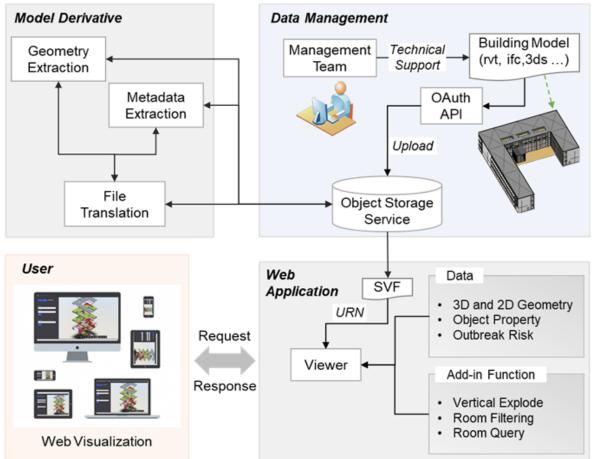
- 364 defined in Table 1. λ is the proportion of accessible surfaces, calculated by parameters 365 extracted from the BIM model. $a_H = \alpha \varphi_H$, representing the rate pathogens excreted to hands. 366 Infectious period $1/\gamma$, inoculation rate ρ , and pathogen inactivation rate in hands μ_H and in fomites μ_F , are all pathogen-specific parameters that are defined in Table 1. χ is the proportion 367 368 of pathogens self-inoculated by susceptible hosts, set as 1 in this study. $\rho_{HF} = \rho_T \tau_{HF}$, indicating 369 the rate of pathogen deposited from hand to fomite, where ρ_T is the rate of fomite touching 370 extracted from the BIM model, τ_{HF} is the transmission efficiency defined in Table 1. θ_{H} is the effective hand cleaning rate, which is set as the rate of hand washing. N is the occupancy of 371 each room, extracted from the BIM model. $\rho_{FH} = N \rho_T \tau_{FH} \kappa$, representing the rate of pathogen 372 picked up by hands, where τ_{FH} is the transmission efficiency from fomites to hands, κ is the fingertip to surface ratio, set as $\frac{6E-06}{\lambda}$ according to study [24]. θ_F is the effective fomite cleaning rate, which is set as the rate of building cleaning and can be extracted from BIM model. 373 374 375
- 376

377 In epidemiology literature, R_0 is one of the most widely used indicators of transmission intensity

- to demonstrate the outbreak potential of an infectious disease in a population. Commonly, $R_0 > 1$ means the epidemic begins to spread in the population, $R_0 < 1$ means the disease will
- 380 gradually disappear, and $R_0 = 1$ means the disease will stay alive and reach a balance in the 381 population. With the increase of R_0 , the outbreak risk will increase, and more severe control 382 measures and policies will be needed [37]. In this study, we categorize the level of outbreak risk 383 into low, mild, moderate, and severe based on the range of R_0 . Specifically, the risk is low when 384 $R_0 < 1$; the risk is mild when $1 \le R_0 < 1.5$ because there is a fair chance that the transmission 385 will fade out as R_0 is not much larger than 1 [38]; the risk is moderate when $1.5 \le R_0 < 2$, indicating an epidemic can occur and is likely to do so [39,40]; and the risk is severe when R_0 > 386 387 2 and immediate actions should be taken by facility managers, such as cleaning the surfaces, to 388 reduce the risk.
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390 **2.2.** Web-Based Information Communication System

391 To better communicate the infection risk to occupants and facility managers, a web-based 392 system was developed to visualize the outbreak risk of different pathogens in each room within 393 a building. Fig. 5 illustrates the architecture of the web-based system, which consists of four 394 modules, i.e., data management, model derivative, web application, and user. The data 395 management module is maintained by the management team and allows them to upload 396 building models. In the model derivative module, the uploaded model is translated into the SVF 397 format which is the format used by the web application. The web application module displays 398 the building model and provides customized functionalities to facilitate visualization of pathogen 399 risk within the building. Finally, the user can access the web-based system and visualize the 400 room-level risk of pathogens. The web-based system is developed using Autodesk Forge that is 401 a collection of APIs to develop cloud-based platforms to access, manage, and visualize design 402 and engineering data. Each module is detailed below.



404 405

Fig. 5. Web-based alert system

406 407 The data management module supports a variety of 3D model formats such as rvt, ifc, and 3ds, 408 where rvt is the file format used by the Autodesk Revit; ifc is an open international standard data 409 schema for BIM data that are supported by various software products such as AutoCAD, Revit, 410 and Tekla Structures; 3ds is the file format used by the Autodesk 3ds Max 3D modeling, 411 animation and rendering software. The management team needs to log into their account to 412 obtain authorization from the Forge OAuth API to access the Object Storage Service (OSS). 413 Model files are uploaded to the OSS and stored as objects in buckets. In the second module, 414 the model derivative translates the uploaded model into SVF format and extracts design 415 metadata such as geometric data and object properties (e.g. room area and occupancy). The 416 translated model and extracted data are also stored in the OSS. The model derivative 417 component generates a unique identifier called URN for each translated model. The URN is 418 then fed into the web application for the building model visualization. 419 420 The web application is built on the Forge Viewer API with customized functions. The Viewer API 421 is a WebGL-based JavaScript library to render both 2D and 3D models. It is developed to 422 display translated models generated by the model derivative component. ExpressJS was 423 selected to develop the web application due to its flexibility and scalability. ExpressJS is a 424 prebuilt NodeJS framework that is designed to create server-side web applications [41], and it 425 allows the web application to handle multiple requests concurrently. As such, pathogen risk 426 information can be quickly communicated to facility users even at times of peak traffic of the

427 website. ExpressJS allows the developer to design customized functionalities in the web

- 429 (HTTP) request. The routing technique manages the way the web application responds to user
- requests. This technique is derived from the HTTP method [42] and attached to the ExpressJS
 router instance. POST and GET methods were used to send and retrieve data from the
- 431 router instance. POST and GET methods were used to send and retrieve data from th 432 webserver.
- 433

434 Three add-in functions were developed to help users visualize the interior layout of the building 435 and color-coded rooms with their corresponding risk levels, as well as search specific room-436 related disease outbreak risk information. The first add-in function is "vertical explode", which is 437 used to view each level of the building. This function can help the user visualize the interior and 438 room layout. The facility users can also use this function to visualize the outbreak risk of rooms 439 on each floor and take appropriate practices. For facility managers, the "vertical explode" function enables them to obtain a holistic view of risk distribution at each level and take 440 441 informed actions, such as limiting the number of occupants and implementing cleaning and 442 disinfection protocols, to control the spread of the disease. This function is integrated with the 443 web-based system, and clicking buttons were created to activate and deactivate it. The second 444 function is "room filtering", which is used to highlight rooms at different risk levels for a specific 445 pathogen. The user needs to first select one of the three pathogens from the dropdown menu: 446 SARS-CoV-2, Influenza, and norovirus. Thereafter, the user can set a risk threshold to highlight 447 rooms with R_0 greater than a specific value. In addition, different highlighting colors are used to 448 represent different infection risk levels. Low, mild, moderate, and severe risks are represented 449 by color green, blue, celery, and red, respectively. The third function is "room guery", which 450 enables the user to search for a specific room and retrieve infection risk for the three pathogens. 451 The "room query" function is displayed as a search box on the web-based system. The users can easily find the potential risk of a specific room using this function. Finally, end users can 452 453 access the web-based information communication system and obtain information about 454 outbreak risk in each room of the building through various channels, including laptops, 455 smartphones, and tablets.

456

457 3. Case Study

A hypothetical case study is used as an example to demonstrate the efficacy of the proposed
framework and the newly developed web-based system. The building information model of a
six-floor school building with 221,000 square feet is used. The building contains classrooms and
faculty and graduate assistant offices.

462

463 3.1. Disease Outbreak Risk in Different Rooms

The room types considered in the case study include offices and classrooms. Five offices and five classrooms were selected. The venue-specific parameters of the rooms are extracted and listed in Table 3, and the computed R_0 values of the three diseases are listed in Table 4.

Table 3. Venue-specific parameters in representative rooms					
Room Type	Room #	Accessible surface area (square feet)	Proportion of accessible surface	Occupancy (number of people)	Rate of fomite touching (times per hour)
	#1	45.5	0.018	36	45 (30, 60)
	#2	45.5	0.017	37	45 (30, 60)
Classroom	#3	176.3	0.138	19	45 (30, 60)
	#4	1328.9	0.194	91	45 (30, 60)
	#5	410.9	0.151	26	45 (30, 60)
Office	#1	36.6	0.052	2	12 (0, 30)

#2	106.8	0.115	13	12 (0, 30)
#3	52.1	0.062	10	12 (0, 30)
#4	1289.8	0.306	9	12 (0, 30)
#5	53.7	0.053	15	12 (0, 30)

Т	Table 4. R_0 values of the three diseases of representative rooms				
	Room #		<i>R</i> ₀ values		
Room Type		Influenza	Norovirus	COVID-19	
	#1	0.078	9.704 ²	0.962	
	#2	0.079	10.441 ²	0.970	
Classroom	#3	0.014	0.092	0.168	
	#4	0.237	2.603 ²	1.803 ¹	
	#5	0.020	0.117	0.224	
Office	#1	0.002	0.023	0.022	
	#2	0.010	0.073	0.118	
	#3	0.008	0.098	0.099	
	#4	0.007	0.023	0.078	
	#5	0.011	0.169	0.146	
			1 4		

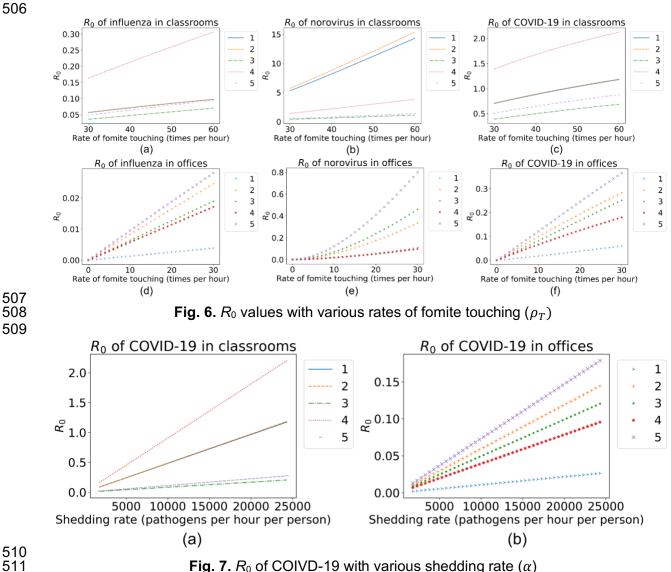
Note: The superscripts indicate the risk level of the diseases, where 1 represents a moderate
 risk level and 2 represents a severe risk level. Values without superscripts indicate the risk level

473 is low.

474

475 From Table 4, the values of R_0 vary across different rooms and different diseases. R_0 values in 476 offices are smaller than the values in classrooms, which stems from the small occupancy and 477 the low rate of fomite touching in offices compared to those in classrooms. For influenza, the R_0 478 values in all the rooms are less than 1, indicating that influenza is unlikely to outbreak in the 479 building through the fomite-mediated transmission. This could be partially explained by the 480 relatively short infectious period, high inactivation rate in hands, low hand-to-fomite pathogen 481 transmission efficiency, and relatively low infectiousness with the same amount of pathogens. 482 For COVID-19, the R_0 values in all rooms are higher than those of influenza, and the risk in 483 Classroom 4 reaches a moderate level, indicating that COVID-19 has the potential to outbreak 484 in the classroom. COVID-19 has a relatively high outbreak risk in most cases because it has a 485 high shedding rate, small surface inactivation rate, and high transfer efficiency from fomites to 486 hands. For norovirus, the R₀ values are high in most classrooms, which might be because of its 487 high infectivity, long infection period, and high hand-to-fomite transmission efficiency compared 488 to the other two diseases. This finding also aligns with the trend obtained in [24]. The above 489 results prove that the outbreak risk of an infectious disease is influenced by both venue-specific 490 and pathogen-specific parameters, which highlights the significance of integrating BIM and the 491 pathogen transmission model in assessing spatial-varying disease outbreak risk. 492

493 Sensitivity analysis was further conducted to evaluate the influence of the rate of fomite 494 touching (ρ_T) and the shedding rate (α) of SARS-COV-2 on R_0 based on the estimated ranges 495 of the two parameters (listed in Table 2). Fig. 6 illustrates the changes in R_0 with the increase of 496 ρ_T for all three diseases in both classrooms and offices. From Fig. 6, the disease outbreak risk 497 increases as the increase of ρ_T . The values of R_0 for norovirus and COVID-19 in Classroom 1, 498 2, and 4 may exceed 1 with the increase of ρ_T . On the other hand, the infection risk in offices and that for influenza in classrooms will remain low even occupants touch objects in the rooms more frequently. Therefore, it is particularly important to educate students in classrooms with relatively high occupancy to not touch the common areas frequently. Fig. 7 illustrates the changes in R_0 of COVID-19 with varying shedding rates. From the figure, α has a significant impact on the outbreak risk of COVID-19 in Classroom 1, 2, and 4. Therefore, for classrooms with relatively large occupancy, control strategies should be taken to reduce pathogen shedding from the occupants, such as using face masks, and covering the mouth when coughing.



512 513

3.2.

Influence of Cleaning Practice

514 Cleaning is an effective strategy to reduce fomite-mediated pathogen transmission in built 515 environments [43]. This study examined the impact of surface cleaning at different times per day 516 on reducing the disease outbreak risk. The timing of each cleaning practice is not included in 517 the disease transmission model and the average R_0 is estimated on an hourly basis. Fig. 8 518 illustrates the changes in R_0 with respect to various times of surface cleaning each day. 519

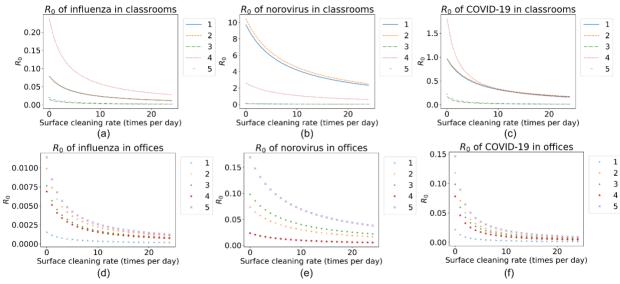




Fig. 8. R_0 values with various times of surface cleaning per day

523 From Fig. 8, surface cleaning can significantly reduce the outbreak risk of all three diseases in 524 both classrooms and offices. Based on the analysis, different surface cleaning practices can be 525 applied to different rooms to reduce the risks to an acceptable low level. Cleaning the surface 526 five times per day will decrease R_0 by over 50%, compared to no surface cleaning. Considering 527 the ongoing outbreak of COVID-19, classrooms with high occupancy (e.g., Classroom 4) should 528 be given particular attention on surface cleaning. Cleaning surfaces at least two times per day is 529 needed to achieve a low risk level. For norovirus, classrooms with relatively large occupancy 530 (e.g., Classroom 1, 2, and 4) will require more frequent surface cleaning to reduce the outbreak 531 risk to the low level. Other complementary strategies, such as increasing hand washing and 532 limiting occupancy, should be adopted to maintain a low level of outbreak risks.

533

534 **3.3.** Infection risk visualization via web-based system

Fig. 9 presents the user interface of the developed web-based system. The developed web application provides an intuitive and responsive user interface to visualize outbreak risk information in the building. The facility manager and user can navigate to the interior model to visualize the interior layout of the building using the "*Interior Model*" button. The user can select and visualize risk-related information for different diseases: COVID-19, influenza, and norovirus. Fig. 10 illustrates the developed web visualization tool.

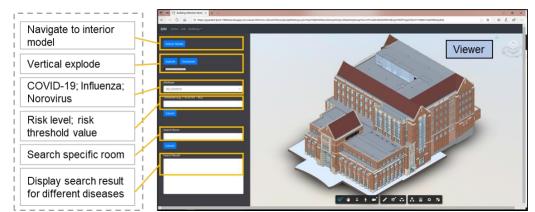
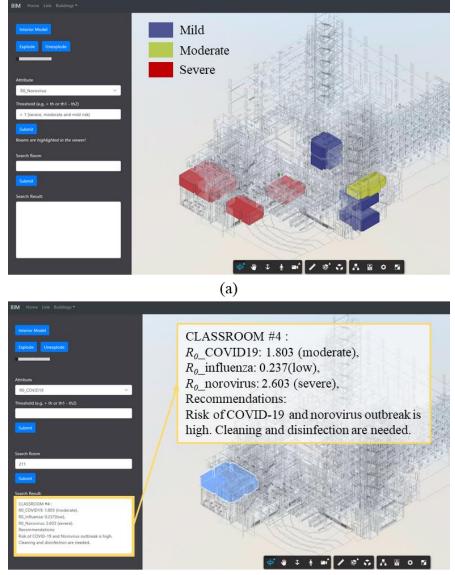




Fig. 9. The user interface of the developed web-based alert system



(b)

Fig. 10. Demonstration of pathogen risk visualization. (a) room filtering based on risk value threshold; (b) search specific room

549 As shown in Fig. 10, room filtering and room query functions can help the user easily locate rooms 550 with high risk and query risk information for a specific room. Specifically, Fig. 10 (a) shows an 551 exemplary output of the room filtering function that highlights the rooms with R_0 value greater than 552 1 for COVID-19. Fig. 10 (b) displays an example of the room query function in the web system. 553 The pathogen risk information for influenza, norovirus, and COVID-19 is retrieved with 554 corresponding recommendations. With the web-based information communication system, facility managers can take important measures to control the spread of diseases, such as designing 555 556 appropriate cleaning and disinfection strategies, promoting hand hygiene, reducing maximum 557 occupancy, and accommodating facility usage schedule based on risk distribution across rooms 558 within the building. For instance, deep cleaning and disinfection are required for rooms with 559 severe outbreak risk. In addition, facility managers can post signs at these high-risk areas to

remind occupants to take essential practices such as social distancing and hand hygiene. The web-based system will also keep facility users, including teachers, students, and other staff, aware of up-to-date outbreak risk information within the building, and thus taking informed actions to avoid further spread of diseases. For example, facility users can avoid entering rooms with high outbreak risk.

565

566 4. Discussion

567 The results and insights derived from the analysis have important implications on adaptive built 568 environment management to prevent infectious disease outbreak and respond to on-going 569 pandemic. Due to varying building characteristics, occupancy levels, and pathogen parameters, 570 the microbial burdens and outbreak risks differ significantly even in the same building, 571 highlighting the need for spatially-adaptive management of the built environment. The proposed 572 method automates the batch process for simulation and prediction of outbreak risks for different 573 pathogens at the room level, and visualizes the risks for adaptive management. The results on 574 outbreak risks at room level enables the paradigm for spatially-adaptive management of the 575 built environment. With the new streams of risk information, customizable interventions can be 576 designed. For instance, in consistent with the practice during the COVID-19 pandemic, reducing 577 the accessible surfaces in rooms and restricting the occupancy in the room are some of the effective strategies to reduce the outbreak risks. The spatially-varying risk information can also 578 579 guide the facility managers to pay close attention to high-risk areas by adopting more frequent 580 disinfection practices.

581

582 A BIM-based information system is developed to extract the necessary information for modeling 583 infection within buildings, and to visualize the derived information in an easy-to-understand and 584 convenient way through web pages. As such, the information-driven interventions could 585 alleviate the pathogenic burdens in the buildings to prevent the spread of infectious diseases. 586 Providing information to end-users is critically important for them to change behaviors. Human 587 behavior plays an important role in the transmission of pathogens such as the SARS-Cov-2. 588 Changing behaviors is critical to preventing transmission. Providing timely and contextual 589 information can be a promising option to motive the change of human behaviors. With the room-590 level outbreak risk information, the users could be motivated or persuaded by the visualized 591 risks to practice appropriate behaviors such as wearing a mask, social distancing, and hand-592 washing. The facility managers can use the information to conduct knowledge-based 593 management, such as limiting the occupancy in the room, managing crowd traffic, and 594 rearranging room layout.

595

596 This study has some limitations that deserve future research. First, the model does not consider 597 factors such as sunlight exposure, humidity, and airflow that may impact the persistence and 598 transmission of pathogens in built environments. This is mainly because the quantitative 599 impacts of these factors on pathogen persistence and transmission are largely ambiguous, if not 600 unknown. If these impacts can be quantified and the environmental parameters can be 601 monitored and modeled in BIM, our proposed framework can be extended to incorporate these 602 factors. Second, the computation of R₀ only considers the fomite-mediated transmission, and 603 does not consider the airborne and close contact transmission. Microbial pathogens may have 604 different transmission routes, including airborne, close-contact, and fomite-based transmission. 605 This study focused on fomite-based transmission to illustrate the modeling approach for 606 assessing the outbreak risks, and demonstrate the efficacy of the developed information system 607 to guide infection control practices and building operations. To fully assess the exposure risks 608 and outbreak potentials, all important routes need to be considered. In addition, the outbreak 609 potentials of a variety of pathogens can be considered together to develop an aggregate index,

610 which could be more intuitive for occupants and facility managers who are not public health

611 experts. Third, the system mainly relies on static models and does not make full use of dynamic

612 and real-time data regarding built environments and occupant behaviors such as presence and

613 interactions with objects. In future studies, the internet of things sensors can be installed in the 614 buildings and algorithms can be developed to retrieve dynamic data for integration with the

- 615 models for accurate and robust risk estimation. Fourth, the web-based system can be further
- 616 improved by connecting it with smart devices such as robots for automated cleaning and
- 617 disinfection and smartphones for precision notifications.
- 618

619 5. Conclusions

620 This study creates and tests a computational framework and tools to explore the connections 621 among built environment, occupant behavior, and pathogen transmission. Using BIM-based 622 simulations, building-occupant characteristics, such as occupancy and accessible surface, are 623 extracted as venue-specific parameters. The fomite-mediated transmission model is used to 624 predict the contamination risks in the built environment by calculating a room-by-room basic 625 reproductive number R₀, based on which the level of infection risk at each room is characterized 626 into low, mild, moderate, and severe. A web-based system is then created to communicate the 627 infection risk and outbreak potential information within buildings to occupants and facility 628 managers. The case study demonstrated the efficacy of the proposed methods and developed 629 systems. Practically, the method and system can be used in a variety of built environments, 630 especially, schools, hospitals, and airports, where transmission of infectious pathogens is of 631 particular concern. The outbreak risks predicted at room resolutions can inform the facility 632 managers to determine room disinfection and cleaning frequency, schedule, and standard. In 633 addition, appropriate operational interventions including access control, occupancy limits, social 634 distancing, and room arrangement (e.g. reducing the number of tables and chairs) can be 635 designed based on the derived information. The occupants can access the useful information 636 via webpage to plan their visit and staying time in the facilities, and practice appropriate

- 637 personal hygiene and cleaning practice based on the information. - Shuai Li
- 638

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