Modeling COVID-19 infection in a confined space

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Abstract In this paper, we construct a stochastic model of the 2019-nCoV transmission in a confined space, which gives a detailed account of the interaction between the spreading virus and mobile individuals. Different aspects of the interaction at mesoscopic level, such as the human motion, the shedding and spreading of the virus, its contamination and invasion of the human body and the response of the human immune system, are touched upon in the model, their relative importance during the course of infection being evaluated. The model provides a bridge linking the epidemic statistics to the physiological parameters of individuals and may serve a theoretical guidance for epidemic prevention and control.

Keywords 2019-nCoV · Stochastic Model · Epidemic prevention and control · Confined space

1 Introduction

The novel coronavirus (2019-nCoV) was first widely known after the detection of unexplained pneumonia in the early 2020. Due to the lack of knowledge on this newly emerging virus, the 2019-nCoV began to spread rapidly and a large outbreak of the disease was globally reported. As the research goes on, a serious yet common scenario is the possibility of the aerial spreading in a

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relatively closed environment. Today's world is characterized with large traffic flows at all scales. As a result, a large number of people stay in a closed space like cars, trains and airplanes. Even if there is no contact between people, the airborne virus may still cause infection. For example, the Diamond Princess, the world's most luxurious ocean liner, which had on board more than 3700 people. A passenger was confirmed the 2019-nCoV infection on February 1st, which was followed with a total of 691 infections by the date of February 24th, with an infection rate of 18.6% [15].

With the pandemic inflicting the globe, a large number of passengers return home by air or other means of transportation. Due to the strict prevention and control measures of many governments, local traffic flow of people is restricted and residential areas become relatively isolated. According to statistics [1], humans spend 87% of their time indoors and 6% in confined microenvironments, i.e. vehicles. Therefore, living and working in a closed environment constitute the main scene in our everyday life. Studying the dynamics of infection in a closed or semi-enclosed environment is of great significance for the ongoing and future epidemic prevention and control.

After the outbreak of the epidemic, much research has been done to describe its development and containment, many of which are based on the susceptible-exposed-infected-removed (SEIR) [2, 19] compartment model, incorporating data from different temporal and spatial regions [18]. Through large-scale data analysis and extensive numerical simulation, the basic regeneration numbers R_0 of 2019-nCoV [3, 4, 16, 17] is estimated, according to which appropriate measures may be planned and implemented for an efficient prevention and control of the disease. As a typical example, according to the latest data, Zifeng Yang [5] plotted the future development curve of the pandemic, based on an artificial intelligence (AI) approach, to estimate the parameters for large-scale simulations of phenomenological models such as the SEIR, similar to what has been done in many other researches.

However, little has been done to model the viral spreading in a confined space and check how physiological specifics of individuals interact with ambient viruses. Here, we build a fully stochastic model, trying to answer some of these questions. In our model, individual's immunity to 2019-nCoV is randomly drawn from a probability distribution. The contamination, shedding and spreading of the virus are modeled in detail for each agent. All the stochastic simulation is done based on the Gillespie algorithm [12]. The results obtained are highly consistent with the statistics procured from the real data [13, 14].

In this study, a new model is built to simulate the the dynamics of the 2019-nCoV infection in a confined space, which includes different characteristics of individuals during the transmission of 2019-nCoV and in the immune response. We estimate various parameters in our model in view of the data in news reports and published articles. Finally, by comparing results in different conditions, reasonable suggestions are made for the actual epidemic prevention and control.

2 Model

According to the characteristics of human infection, in the susceptibility-exposure-infection-recover (SEIR) compartment model, each individual can be in one of the five states (see Fig. 1A) : susceptible (S), exposed (E, infected without symptoms), hidden (H, infected without symptoms for a very long incubation period compared to the exposed), infected (I, with symptoms and infectiousness) and recover (R, never susceptible anymore). In our new model, we do not explicitly classify people into these five categories and the state of each individual is determined by two variables (n_v, t_v) , being respectively the concentration of the 2019-nCoV in his body and the body tolerance to this virus. A person with $n_v = 0$ can be regarded as a susceptible person (S). For $0 < n_v < N$, where N is a given threshold which satisfies a uniform distribution in the interval $[0, 8]$, the virus carrier can be regarded as being in an incubation period (E). As the concentration of virus increases, in general symptoms such as cough, fever, or slow movement, become more and more obvious. However, a high tolerance will exempt a person from these symptoms. As different people have different tolerances to the COVID-2019, for convenience, we assume that the tolerance t_v to the 2019-nCoV satisfies a uniform distribution in $[0, 10]$. However, it is hard to directly measure t_v at this time and hence we assume that the moving speed of a person $s_v = s_{speed} * \frac{1}{1+2*10^{-4}*n_v^{11-t_v}}$, where $s_{speed} = 4000m/day$, such that s_v may be used to map the tolerance. For $n_v > N$ but $s_v < T$ where $T (= 2000m/day)$ is another given threshold, the person is deemed to be infected (I) with symptoms (slow movement). However, for $n_v > N$ and $s_v > T$, the person would be identified as asymptomatic (H) with no obvious symptoms. In our model, states of individuals with diverse physiological conditions are subject to dynamic changes according to their interaction with viruses in the closed environment, where spatial factors play an essential role in that the spreading of the virus and the infection of human are tightly correlated in both space and time.

Even if a susceptible individual is exposed to the virus, he may not become infected, since a small number of viruses invading the human body can be completely degraded and eliminated by our immune system. Only when the virus concentration accumulates to a point that defies the immune system, the virus starts to proliferate exponentially. In the model, the rate of virus reproduction in the body is assumed to be sigmoidal $P_T = \alpha * [1 + \tanh[z * (n_v - N)]]$, where α is a basic production rate and z controls the steepness of the switch function. The parameter N is the previously mentioned threshold with an average value of 4. As shown in Fig.1B, when the virus concentration in the body exceeds $4 \times 10^4 units/m^3$ [6], the rate of virus reproduction will be greatly increased.

In the simulation, each individual will be infected with certain probability after contacting with the virus in the environment. Here we distinguish the virus in the body surface of concentration n_v from that inside the body, characterized by the surface contamination degree b_s . Each grid point (m, n) can only be occupied by at most one person and the change of the virus concentration

Fig. 1 Different states of individuals. A. Boxes with different colors indicate different states, arrows for transitions between states. B. The rate of virus reproduction P_T vs the virus concentration n_v . $\alpha = 3, z = 1, N = 4$

$p_t(m, n)$ could be described by the following reactions

$$p_t(m, n) \xrightarrow{\gamma_b} b_s, b_s \xrightarrow{r_b} p_t(m, n), b_s \xrightarrow{\gamma_c} n_v,$$

where $\gamma_b = 2.3 \times 10^4 \text{ units/day}$ is the contamination rate, $r_b = 0.1 \times 10^4 \text{ units/day}$ is the rate at which the virus peels off into the air from the body surface and $\gamma_c = 1.8 \times 10^4 \text{ units/day}$ is the rate of viral invasion into the human body.

Since individuals in the asymptomatic incubation period are already infectious, they can release the virus to the environment at a rate r_f , so does the individual in state I. The changes of the virus concentration in the body and the antibody concentration secreted by immune cells are described by the reactions below

$$\begin{aligned} n_v &\xrightarrow{r_f} p_t(m, n), n_v \xrightarrow{P_T} 2n_v, \\ n_v &\xrightarrow{r_c * n_{Ab}} \phi, Ag \xrightarrow{r_{Ab}} 2Ag, \end{aligned}$$

where $r_f = \frac{1}{\tau} \times (1 + \frac{2 \times 10^{-4} * n_v^3}{1 + 2 \times 10^{-4} * n_v^3})$ with $\frac{1}{\tau}$ the basic rate, and P_T is the rate of virus reproduction in the body. r_c is the rate of virus being killed by antibody, and n_{Ab} is the antibody concentration in the body, which is produced at the rate r_{Ab} being assumed to have a uniform distribution in the interval $[0 \sim 0.01 \times 10^4]$. With the increase of virus concentration in the body, the epidemic symptoms like coughing is turning obvious, so the releasing of the virus to the environment is getting fast. However, we assume that when each

infected individual releases the virus to the environmental in this way the body virus concentration does not change.

The diffusion and decay of virus concentration in the environment is modelled by the following reactions

$$p_t(m, n) \xrightarrow{r_d} \phi, p_t(m, n) \xrightarrow{D} \frac{1}{2} * p_t(m, n)$$

$$\sum_{(m_1, n_1)} p_t(m_1, n_1) = \frac{1}{2} * p_t(m, n) .$$

Since the coronavirus can survive in the air for almost 3 hours [20], its decay rate is $r_d = 8 \times 10^4 \text{ units/day}$. The diffusion coefficient of virus is very weak ($D_v = 2.9 \times 10^{-11} \text{ m}^2/\text{s}$) [5]. However, due to the movement of individuals in space and sneezing to the air, the small-scale air flow increases effectively the diffusion and an effective coefficient may be set as $D = 2.5 \times 10^{-5} \text{ m}^2/\text{s}$. In total, the concentration change in a time interval Δt is

$$\begin{aligned} \Delta p_t(m, n) &= p_{t+\Delta t}(m, n) - p_t(m, n) \\ &= D * \left[\sum_{(m_1, n_1)} p_t(m_1, n_1) - d * p_t(m, n) \Delta t \right] , \quad (1) \\ &\quad - r_d * p_t(m, n) * \Delta t + r_b * b_s * \Delta t + r_f * n_v * \Delta t \end{aligned}$$

where D is the virus diffusion coefficient and $d = 4$ is the coordination number of the simulation lattice. The summation is over the adjacent lattice points.

We use Gillespie algorithm to carry out the simulation of the stochastic dynamics. Information vectors are used for each individual and each grid point and get updated in each time step. The closed space is modeled as a square grid with a side length of $100m$, where the distance of adjacent grid points is $5m$. These individuals are homogeneously mixed in the space and no attention in the model is paid to his or her own social contact network. Due to the epidemic, in a typical situation, everyone's activities is restricted and most people spend 90% of their time in a more or less stationary state [1] and 10% in active motion. We treat each movement of an individual as a reaction. If the reaction occurs, the person can move to a randomly chosen adjacent grid point. The spreading of 2019-nCoV is a random motion in which not all viruses move together to one adjacent grid point. Only half the of the viruses move to the neighbouring points, such that concentration on the grid point (m, n) is reduced by half, and the other half is randomly distributed to the four adjacent grid points (Fig.2).

3 Results

At the initial moment, we have 74 susceptible (S) individuals in a closed environment among which one is infected. The initial concentration of the virus in the infected (I) individual is $50 \times 10^4 \text{ units/m}^3$. As shown in Fig.3, after 24

Fig. 2 Schematic representation of human movement and virus spreading: people (green circles), viruses (yellow blob), and the width of the blue arrow signals the probability of diffusion.

days of simulation, 14 people become infected in the confined space. The highest concentration at some grid point can reach almost $20 \times 10^4 \text{ units/m}^3$. The number of people infected has been increasing over time. There are two asymptomatic infections. After averaging through multiple simulations, as shown in Fig.3D, we display the evolution of the infection rate within 24 days, and compare it with the real data on the Diamond Princess. The curves seem to be in good agreement. The first day of the horizontal axis represents February 1, 2020, with the first confirmed case among tourists. On February 24, an average of 13.4 people were infected with the 2019-nCoV in model, and the infection rate was 18.1%, which was basically the same as the infection rate of 18.6% on the Diamond Princess.

The basic reproduction number R_0 depends on the exponential growth rate of the outbreak, as well as the serial interval (SI), which is defined as the time duration between a primary and a secondary case-patient (infector) having symptom onset [9] (Fig.4). In the model, because no epidemic prevention measures are taken, the number of infected people is growing exponentially, so we calculate the exponential growth rate as $r = \ln(I(t))/t$, where $I(t)$ is the number of infected people by time t [4]. Taking the average value over a few realizations, the exponential growth rate is estimated to be $0.1081/\text{day}$ (the corresponding doubling time was 6.19 days). Based on the SEIR model, Joseph T Wu [3] estimated the basic regeneration number as 2.68 and the doubling time as 6.4 days; Qun Li [7] analyzed data on the first 425 confirmed cases in the early stage and found that the epidemic is doubled in size every 7.4 days with a mean serial interval of 7.5 days. According to the definition of SI, we can calculate the mean and standard deviation of the SI (T_c, σ) = (6.22, 2.98) in our model which is considerably shorter than that of

Fig. 3 Plot of the simulation results. A. The spatial distribution of individuals in different states, and the virus concentration in space, with the infected individual (I, red X) and the susceptible (S, green circle). The time course of the number of I (B) and H (C). D. The simulated infection rate *vs* the real infection rate on the Princess Diamond.

SARS (8.4 days) or MERS (14.6 days), but close to the SI (= 7.5) estimated by Qun Li [7] and greater than the SI (= 3.96) from the 468 confirmed cases by Zhanwei Du [8] and Nishiura, H [9]. Now that we have the exponential growth rate ($r = 0.1081/day$) and the SI, we can calculate the basic reproductive number of the model [10],

$$R_0 = e^{r \cdot T_c - \frac{1}{2} \cdot r^2 \cdot \sigma^2} \quad (2)$$

and obtain $R_0 = 1.85$ lower than previous estimate [4, 11]. If the infection threshold N of the population is uniformly distributed with a mean value of 4, the average incubation time in our model would be 4.16 days, which is the same as the one estimated by Nanshan Zhong *et al* [5] based on extensive on-site data. If the distribution of the infection threshold adopts an exponential form with an average value of 4, the mean incubation time obtained would be 3.08 days. Therefore, the precise distribution of the population infection threshold N is a very important factor, which can determine how quickly the virus spreads in the population.

One effective way to contain the infection is to reduce the rate at which the infected individuals (I) spread the virus into the air, which cuts down the source of the virus. As shown in Fig.5, if we change the basic rate $\frac{1}{\tau}$ from $\frac{1}{2}$ to $\frac{1}{10}$, there would be only two infected person, which is hard to initiate an outbreak. It can be seen from the comparison between Fig.3A and Fig.5A that

Fig. 4 Schematic indication of the serial interval. There are two stages for an infected person: the incubation period (blue) and the infection period (red).

the highest virus concentration in the space is reduced from $20 \times 10^4 \text{ units}/m^3$ to only about $4 \times 10^4 \text{ units}/m^3$. In Fig.6A, if τ varies from 1 to 30, the infection rate will be greatly reduced, so limiting the amount of virus released to the environment is very effective. Another equally effective way is to reduce the contamination rate of the susceptible, as shown in Fig.6 when γ_b is reduced from $2.3 \times 10^4 \text{ units}/\text{day}$ to $0.6 \times 10^4 \text{ units}/\text{day}$. From Fig.6B, it can also be concluded that taking measures to make people stay in their dwelling position as much as possible also helps reduce the risk of infection.

With windows being open to ventilation in this confined space, the virus will slowly float out of the boundary. In the model, we add a directional ($v_f = 10m/\text{day}$) component to the virus velocity, aligned with the airflow. When the upper and lower boundaries of the space are open, as shown in Fig.8, compared with the enclosed space in Fig.3A, the virus concentration is much lower. However, in Fig.8A, the number of infected people is not much lower than that in the enclosed space, mainly because the infected individual is in the upstream of the airflow, which exposes a lot of susceptibles to the virus. As can be seen from Fig.8D, if instead the infected is put downstream of the airflow and next to the open boundary, the infection will decrease significantly. With all four boundaries being removed and the airflow blowing from the center, as shown in Fig.9A, the virus is hardly concentrated. In Fig.9D, a total of 30 simulations are conducted to find out how many people were infected 24 days later in a closed environment with the air blowing from the center. In the simulation, the initial location of the infected person is random. It is easy to see that in this condition, the number of infected people is considerably reduced when having airflow from center.

Fig. 5 The result of reducing the rate of virus release from the human body. A. The spatial distribution of individuals in different states, and the distribution of virus concentration in space, with the infected (I, red X) and the susceptible (S, green circle). The time course of the number of I (B) and H (C).

Fig. 6 Influence of two different measures on the infection rate. A. The number of the infected *vs* τ . B. The distribution of the number of the infected with people dwelling for 90% of the time (black) and for 50% of the time (green), in 30 repeated simulations.

Fig. 7 Simulation results with $\gamma_b = 0.6 \times 10^4 \text{ units/day}$. A. the spatial distribution of individual in different states, and the distribution of virus concentration in space, with the infected individual (I, red X) and the susceptible (S, green circle). The time course of the number of I (B) and H (C).

Fig. 8 Simulation for different virus source locations. The infected individual(I) is in the center of the space (A), and at the bottom of the space (D). The airflow is blowing down. The time course of the number of I (B, E) and H (C, F).

Fig. 9 The impact of the airflow. A. The distribution of people (circles) and viruses (encoded with bar colors). The air is blowing from the center. The time course of the number of I (B) and H (C). D. The distribution of the number of the infected in a confined space *vs* that in an open space from 30 realizations.

4 Discussion

Through the newly constructed virus infection model, we provide useful guidance for effective prevention and control of the COVID-19 spreading, such as restricting the movement of people and placing the ventilation opening in the right position, reducing the releasing of the virus and the contact of susceptibles with unfriendly environment. All the measures will reduce the infection rate and prevent the virus from spreading to various extents. Even very simple practice such as wearing a mask could be a very effective prevention measure, especially in confined spaces. Even in the presence of ventilation, if the airflow is slow and only in one direction, it does not necessarily reduce the infection rate. On the contrary, it helps spread the virus and makes people downstream in the airflow more accessible to viruses. To reduce the infection rate, proper arrangement of the ventilation is needed, for example, open the boundaries of the space as much as possible and send the fresh air from its center.

In the future, we should start an extensive investigation of the interaction between the virus and the human immune system to microscopically compute various simulation parameters for different people to see how they are related to human genetic features. From a physicist's point of view, it is also very interesting to construct a more realistic model to more accurately simulate different airflows from different ventilation facilities, and check the passive

motion of the virus under different air conditions. Expectedly, human social networks also play a very important role in the spreading of the disease, which should be carefully considered in the study of the large-scale infection.

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Conflict of Interest

The authors declare that they have no conflict of interest.

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Figures

Figure 1

Différent states of individuals. A. Boxes with différent colors indicate différent states, arrows for transitions between states. B. The rate of virus reproduction P_T vs the virus concentration n_v . $\alpha=3, z=1, N=4$

Figure 2

Schematic representation of human movement and virus spreading: people (green circles), viruses (yellow blob), and the width of the blue arrow signals the probability of diffusion.

Figure 3

Plot of the simulation results. A. The spatial distribution of individuals in different states, and the virus concentration in space, with the infected individual (I, red X) and the susceptible (S, green circle). The time course of the number of I (B) and H (C). D. The simulated infection rate vs the real infection rate on the Princess Diamond.

Figure 4

Schematic indication of the serial interval. There are two stages for an infected person: the incubation period (blue) and the infection period (red).

Figure 5

The result of reducing the rate of virus release from the human body. A. The spatial distribution of individuals in different states, and the distribution of virus concentration in space, with the infected (I, red X) and the susceptible (S, green circle). The time course of the number of I (B) and H (C).

Figure 6

Influence of two different measures on the infection rate. A. The number of the infected vs τ . B. The distribution of the number of the infected with people dwelling for 90% of the time (black) and for 50% of the time (green), in 30 repeated simulations.

Figure 7

Simulation results with $\gamma_b = 0.6 \times 10^4$ units/day. A. the spatial distribution of individual in different states, and the distribution of virus concentration in space, with the infected individual (I, red X) and the susceptible (S, green circle). The time course of the number of I (B) and H (C).

Figure 8

Simulation for different virus source locations. The infected individual(I) is in the center of the space (A), and at the bottom of the space (D). The airflow is blowing down. The time course of the number of I (B, E) and H (C, F).

Figure 9

The impact of the airflow. A. The distribution of people (circles) and viruses (en-coded with bar colors). The air is blowing from the center. The time course of the number of I (B) and H (C). D. The distribution of the number of the infected in a confined space vs that in an open space from 30 realizations.