

Properties of Hantavirus Infection Model

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ABSTRACT

Hantavirus is a group of viruses that can cause *Hantavirus Pulmonary Syndrome (HPS)* and *Hemorrhagic Fever with Renal Syndrome (HFRS)*. Human contract this virus through their rodent hosts. Since the discovery of *Hantavirus*, there is still no vaccination for the diseases caused by the viruses. This situation has inspired researchers to explore the ecology and epidemiology of *Hantaviruses* in order to come out with a better solution in preventing this infection. In this paper, we investigate the mathematical model of *Hantavirus* infection known as *Abramson-Kenkre (AK)* model. The result shows that by changing the values of certain data such as the transmission rate responsible for infection, the birth rate and the natural death rate, different results can be obtained.

Keywords: *Hantavirus* infection, *Abramson-Kenkre (AK)*, disease

Introduction

Numerous studies have shown a strong correlation between generalist rodent species and infectious diseases carried by them such as *Hantavirus Pulmonary Syndrome (HPS)*. *HPS* is caused by a group of viruses that were first isolated in the laboratory from striped field mice captured near Korea's Hantaan River called as *Hantavirus*. Their potential for causing severe human illness makes these viruses an important public health concern.

The *Hantavirus* infection is carried by mice that move from location to location, and is transmitted to other mice through fight. The mice do not die or are impaired, from contraction of the virus. Vertical transmission (infected mother transfer the disease to offspring), appears to be negligible or nonexistent. Humans get the virus from the mice but have no feedback effects on the mice in the infection process (Kenkre, 2005).

In this research, we investigate the mathematical model in modelling and analyzing *Hantavirus* infection developed by Abramson and Kenkre (2002) or known as *AK* model. The *AK* model analyzes spatio-temporal patterns in the spread of *Hantavirus* infection based on differential equation system. Figure 1 shows the result of analysis by *AK* model.

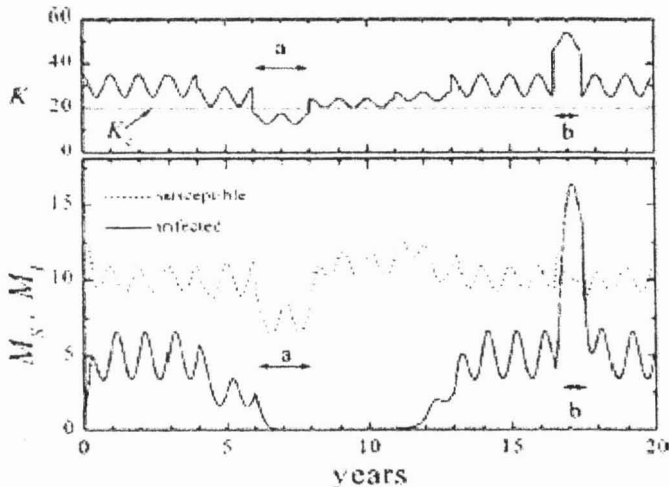


Figure 1: Temporal evolution of the population of mice (bottom) in a caricature time-dependent carrying capacity (top). Two special events are marked: (a) The carrying capacity is below the K_c threshold (shown as a horizontal line). (b) An extraordinary one-year event of greater carrying capacity. Model parameters are $a = 0.1$, $b = 1$, and $c = 0.5$. (Abramson & Kenkre, 2002)

AK model is able to reproduce several observed features in Hantavirus epidemics such as refugia which is foci of infection in the landscape. It also has qualitative and semi-quantitative success in explaining observations such as spatio-temporal patterns in the epidemics. Besides, AK model can successfully explain several field observations as environmentally controlled phase transitions, and thus provide an analytical support to biological hypotheses such as the trophic cascade discussed in Yates et. al, 2002(Abramson, 2007).

However, based on the AK result (as shown in Figure 1), there is no relation between K (carrying capacity) and M_s (susceptible mouse) or M_i (infected mouse). In this paper we investigate in detail the relationship between K and M_s or M_i due to the changes of data. This relation is important as the changes in K , will affect the value of susceptible and infected mice. Thus, by changing the values of K , the relation between them can be obtained.

Preliminary

The AK model for mouse population with two classes represented by M_s for susceptible mouse and M_i for infected mouse is shown by Equation (1):

$$\begin{aligned} \frac{dM_s}{dt} &= bM - cM_s - \frac{M_s M}{K} - aM_s M_i \\ \frac{dM_i}{dt} &= -cM_i - \frac{M_i M}{K} + aM_s M_i \end{aligned} \quad (1)$$

where a , b , c and K are constant with

M is the total population,

a represent the rate of transformation from susceptible to infected, due to fight when the population is over crowded

b signify the births of mice, given that all mice are born susceptible,

c characterizes the death rate for natural reason such that the rodent not infected by the virus carried by them,

K is carrying capacity that is the resource used by mice for the survival such as food and shelter and,

$-(M_s M \text{ or } M_i M)/K$ show the limitation process of population growth.

The range of the deer mice comprises a diverse landscape with a variety of habitats. The inhomogeneous way in which this spatial extent affects local populations can be included in a spatially extended version of the model, where M_s , M_i , and K become functions of a space variable x (Abramson, 2007). By using a simple diffusion model for the transport mechanism of the population, the extended model is shown by Equation (2):

$$\begin{aligned} \frac{dM_s}{dt} &= f(M_s, M_i) + D_s \nabla^2 M_s \\ \frac{dM_i}{dt} &= g(M_s, M_i) + D_i \nabla^2 M_i \end{aligned} \quad (2)$$

where f and g are the right-hand sides of Equation (1) and the included separate diffusion coefficients D_s and D_i for the two classes of mice (Abramson & Kenkre, 2002). The solution of Equation (2) and even its stationary solution may be impossible to find, analytically, for an arbitrary function $K(x)$ (Abramson, 2007).

Methodology

In this research, the Matlab programming has been used to generate the result when the data is changed. The equation is solved by using Runge-Kutta 4th Order Method where this method has the advantage of high order local truncation error. Based on the AK result, the value of K is divided into few values depending on the interval of years as shown in Table 1. The value at the end of one partition becomes the initial value for next partition. This is to make sure that the figure will be continuous through 20 year

Table 1. Value of K for different intervals

Value of K	Interval of years
40	[0, 2]
30	[2, 4]
20	[4, 6]
5	[6, 8]
40	[8, 10]
30	[10, 12]
20	[12, 14]
60	[14, 16]
40	[16, 18]
30	[18, 20]

Result and Discussion

The numerical experiments on AK equation have been conducted and the result of the simulation has been studied with the initial condition: $M_s = 15$, $M_i = 1$, $a=0.1$, $b = 1$, $c = 0.5$.

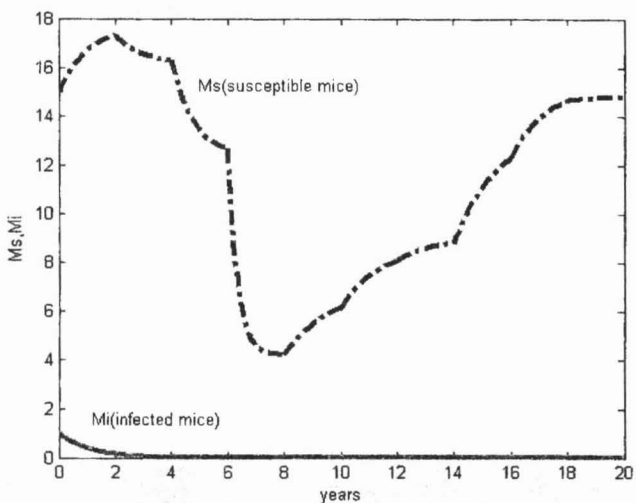


Figure 2: Changes of the number of infected and susceptible mice when the transmission responsible for infection decreased

From Figure 2, the rate of transformation is reduced from 0.1 to 0, which means that there is no transmission from susceptible mice to infected mice. The number of infected mice will drop to zero as time approaches to infinity. As mentioned earlier, mice do not die because of the virus, so it is probably because of competition to obtain the space. The susceptible mice survive due to no transformation of the virus from the infected mice.

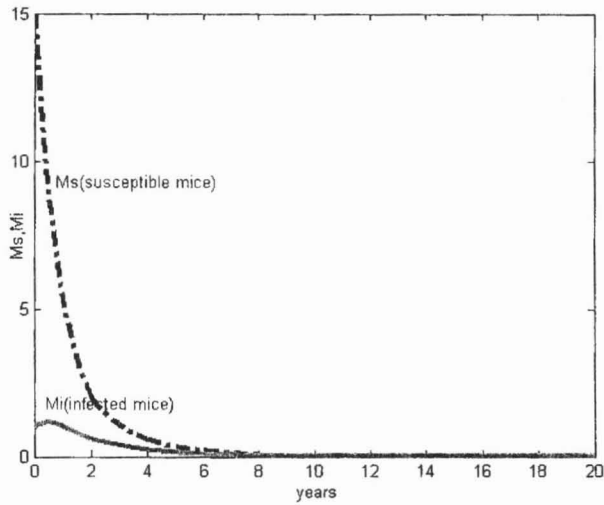


Figure 3: Changes of the number of infected and susceptible mice when the birth rate decreased

Furthermore, graph in Figure 3 shows that when the birth rate decreased from 0.1 to 0, both of the graph M_s and M_i will be dropped. This situation happened because when there is no increasing of mice so there will be no new infected and susceptible mice.

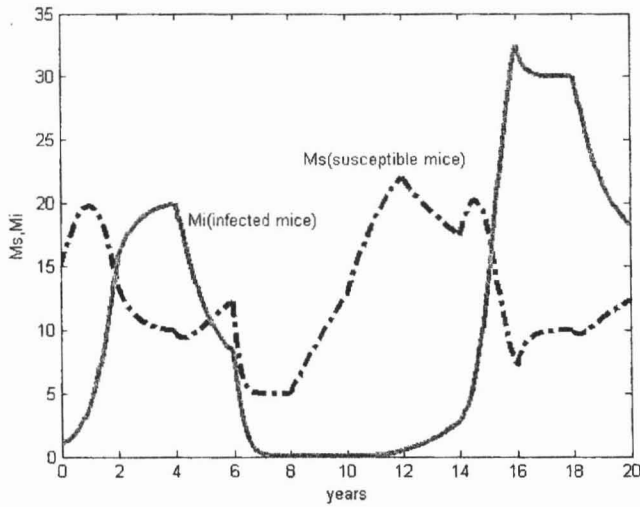


Figure 4: Changes of the number of infected and susceptible mice when the natural death rate decreased

Last but not least, graph in Figure 4 shows the situation that will occur if the natural death rate is decreased. The graph clearly shows that the number of infected mice changed inversely with the number of susceptible mice. Even when the rate of death is decreasing during that time, the rate of birth of the mice is still increasing as usual. This shows that when the number of the infected mice increased, the number of susceptible mice decreased. Conversely, when the number of the infected mice decreased, the number of susceptible mice increased.

Conclusion

This study is conducted to investigate on the properties of AK model and the relationship between K and M_s or M_i due to the changes of data. The result shows that changing in some of the data such as the transmission rate responsible for infection, the birth rate and the natural death rate will affect the values of infected and susceptible mice. As a conclusion, AK model is suitable to demonstrate the Hantavirus infection.

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