The Spread of Infectious Disease with Household-Structure on the Complex Networks

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Abstract

In this paper we study the household-structure SIS epidemic spreading on general complex networks. The household structure gives us the way to distinguish inner and the outer infection rate. Unlike household-structure models on homogenous networks, such as regular and random networks, here we consider heterogeneous networks with arbitrary degree distribution p(k). First we introduce the epidemic model. Then rate equations under mean field appropriation and computer simulations are used here to analyze our model. Some unique phenomena only existing in divergent network with household structure is found, while we also get some similar conclusions that some simple geometrical quantities of networks have important impression on infection property of infectious disease. It seems that in our model even when local cure rate is greater than inner infection rate in every household, disease still can spread on scale-free network. It implies that no disease is spreading in every single household, but for the whole network, disease is spreading. Since our society network seems like this structure, maybe this conclusion remind us that during disease spreading we should pay more attention on network structure than local cure condition.

Key words: Infectious Disease; SIS Model; Networks
PACS: 89.75.-Hc; 05.70.Ln; 02.10.Yn; 87.23.Ge; 64.70.-p

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Preprint submitted to Elsevier Science 9 February 2008
1 Introduction

The spread of disease has been one of the focuses in the field of statistical physics for many years. The dynamical behavior of so-called susceptible-infected-susceptible (SIS) model and susceptible-infected-removed (SIR) model have been widely investigated on regular network and complex networks[1-12]. Within the studying, individuals are modeled as sites and possible contacts between individuals are linked by edges between the sites. It is easy to see that both the properties of disease and topological character of network determine the dynamics of the spread of disease. Studies have showed that there is an epidemic threshold $\lambda_c$ on regular networks. If the effective spreading rate $\lambda>\lambda_c$, the infection spreads and becomes endemic; otherwise the infection will die out. While the threshold disappears on scale-free networks[4].

Usually, infectious diseases, such as HIV and computer virus, have the similar spreading property. They not only can spread in one household, but also can spread from one household to another. To study this spreading character, there have been of considerable interests to epidemic models spreading among a community of households[12-17]. These studies were concerned with SIR model, which cannot appear endemic behavior. In 1999, Ball introduced the SIS household-structure model[18], in which the population is partitioned into $m$ households with $N$ members in each household. A threshold parameter $R_*$ was defined. It is shown that for the household with 2 members, if $R_* < 1$ then the epidemic die out; if $R_* > 1$ the epidemic will exist at an endemic equilibrium. This model has also been studied on homogeneous network by the mean of self-consistent field[19,20]. The similar results have been obtained. These previous studies about household-structure epidemic model were mainly on regular networks. However, studies have showed that a large number of systems, such as Internet, world-wide-web, physical, biological, and social networks, exhibit complex topological properties[21-23]. In particular, small-world properties[24] and scale-free degree distributions[25] appear in many real network systems. In this paper, we will analyze the SIS household-structure epidemic model on complex networks. The outline is as follows: 1) introduction; 2) description of the model; 3) mean-field equations; 4) steady-state solutions; 5) simulation; 6) summary.

2 Model

In complex networks with degree distribution $p(k)$, which is the probability that a given site has $k$ connections (links) that connect it with other $k$ sites (We say that the given site’ degree is $k$.), there are $N$ individuals that are grouped as a household on every site. We assume that these $N$ individuals
contact each other fully. A healthy individual may get infected from within the household and from outside its household. The parameters $\lambda$ and $\beta$ are the infection rates from outside and from within the household respectively. We give each site $x$ a number $i(i \in [0, N])$, which means that there are $i$ infected individuals in the household at site $x$. The number of infected individuals at a given site $x$ changes according to the following transition rates:

$$0 \rightarrow 1 \text{ at rate } \lambda \sum_{(x,y)} i_y$$

$$i \rightarrow i + 1 \text{ at rate } i\beta \text{ for } 1 \leq i \leq N - 1$$

$$i + 1 \rightarrow i \text{ at rate } \gamma \text{ for } 0 \leq i \leq N - 1$$

In the above expressions $\langle x, y \rangle$ means that site $x$ and site $y$ are nearest neighbors, and we suppose there is a (connection) link between them. Infected individuals may infect healthy individuals in their household with rate $\beta$, and also can infect healthy individuals in their nearest neighbors with rate $\lambda$. We assume that once a site is infected, infections within the site are much more likely than infections from outside, so we can neglect the latter. And also, an infected individual in a site can recover with rate $\gamma$. We suppose that all the individuals in a household have the same external connectivity and do not take the birth and death into account.

3 Mean-field equations

We now solve the above model with mean-field method. Let $u_{k,i}$ be the density of individuals whose household has $i$ infected individuals and the corresponding site’s degree is $k$, which means this site has $k$ nearest neighbors. It is worth noticing that $\sum_{i=0}^{N} u_{k,i} = 1$. According to the transitions rate described in the above section, the evolution equations of $u_{k,i}$ are written as below[4,6]:

$$\frac{\partial u_{k,0}(t)}{\partial t} = \gamma u_{k,1} - \lambda k \Theta_k(t) u_{k,0} \quad (1)$$

$$\frac{\partial u_{k,1}(t)}{\partial t} = \lambda k \Theta_k(t) u_{k,0} - \beta u_{k,1} + \gamma u_{k,2} - \gamma u_{k,1} \quad (2)$$

$$\frac{\partial u_{k,i}(t)}{\partial t} = (i - 1) \beta u_{k,i-1} - i \beta u_{k,i} + \gamma u_{k,i+1} - \gamma u_{k,i} \quad (i \in [2, N - 1]) \quad (3)$$

$$\frac{\partial u_{k,N}(t)}{\partial t} = (N - 1) \beta u_{k,N-1} - \gamma u_{k,N} \quad (4)$$
In equations (1)-(4), $\Theta_k(t)$ is the probability that a link from a site points to another site with at least one infected individual. And the expression of $\Theta_k(t)$ is:

$$\Theta_k(t) = \sum_{k'} p(k'/k) \sum_{j=1}^{N} j u_{k',j}(t)$$

(5)

where $p(k'/k)$ is the probability that a site with $k$ degrees points to another site with $k'$ degrees. For uncorrelated networks the expression of $p(k'/k)$ is [7]:

$$p(k'/k) = \frac{k' p(k')}{\sum_{k'} k' p(k')} = \frac{1}{\langle k \rangle} k' p(k')$$

(6)

Substituting (6) to (5), we get $\Theta_k(t) = \Theta(t)$ independent of $k$:  

$$\Theta_k(t) = \Theta(t) = \frac{1}{\langle k \rangle} \sum_{k'} k' p(k') \sum_{j=1}^{N} j u_{k',j}(t)$$

(7)

Now we are going to get steady solutions of Eqs. (1)-(4)

4 Steady-state solutions

4.1 For $N=2$

The evolution equations are simplified as:

$$\frac{\partial u_{k,1}(t)}{\partial t} = k \Theta(t) (1 - u_{k,1} - u_{k,2}) + \gamma u_{k,2} - \beta u_{k,1} - \gamma u_{k,1}$$

(8)

$$\frac{\partial u_{k,2}(t)}{\partial t} = \beta u_{k,1} - \gamma u_{k,2}$$

(9)

In Eq. (8), we have used the equality: $u_{k,0} + u_{k,1} + u_{k,2} = 1$. When $t \to \infty$, $\frac{\partial u_{k,1}}{\partial t} = 0$ and $\frac{\partial u_{k,2}}{\partial t} = 0$, then the steady-state solutions are:

$$u_{k,1} = \frac{\lambda \gamma k \Theta}{\gamma^2 + (\gamma + \beta) \lambda k \Theta}$$

(10)

$$u_{k,2} = \frac{\lambda \beta k \Theta}{\gamma^2 + (\gamma + \beta) \lambda k \Theta}$$

(11)
Substituting (10) and (11) to (7), we get the self-consistent equation of $\Theta$:

$$\Theta = \frac{1}{\langle k \rangle} \sum_{k} kp(k)(u_{k,1} + 2u_{k,2}) = \frac{1}{\langle k \rangle} \sum_{k} kp(k) \frac{(\gamma + 2\beta)\lambda k \Theta}{\gamma^2 + (\gamma + \beta)\lambda k \Theta}$$  \hspace{1cm} (12)

Clearly, $\Theta = 0$ is a solution of Eq.(12), which implies that $u_{k,0} = 1$, $u_{k,1} = 0$ and $u_{k,2} = 0$ is a steady-state solution of Eqs.(8) and (9). A nonzero steady-state solution $\Theta$ (That is: $u_{k,i} \neq 0$, for $i > 0$) is obtained when $\gamma$, $\beta$ and $\lambda$ satisfy the following inequality:

$$1 \frac{1}{\langle k \rangle} \sum_{k} kp(k) \frac{(\gamma + 2\beta)\lambda k}{\gamma^2} \geq 1$$

Then we can get the spreading threshold:

$$\lambda_c = \frac{\gamma^2}{\gamma + 2\beta} \langle k \rangle \langle k^2 \rangle \hspace{1cm} \text{(13)}$$

where $\langle k \rangle = \sum_{k} kp(k)$, $\langle k^2 \rangle = \sum_{k} k^2 p(k)$. In other words, the disease will die out when $\lambda < \lambda_c$; otherwise the disease will pervade the system. Clearly, the threshold $\lambda_c$ is the function of $\beta$, $\gamma$ and $\langle k \rangle \langle k^2 \rangle$. So the degree distribution of networks plays an important role on $\lambda_c$.

For $p(k) = \delta_{k,k_c}$, the network is homogeneous and $\lambda_c = \frac{\gamma^2}{\gamma + 2\beta} \frac{1}{k_c}$. We can lift $\lambda_c$ to prevent infection in terms of increasing the recover rate $\gamma$ or decreasing the site degree $k_c$.

For $p(k) = Ck^{-\nu}$ ($\nu \in (2, 3)$), the networks are scale-free[19]. When $k \to \infty$, $\frac{\langle k \rangle}{\langle k^2 \rangle} \to 0$, the threshold is absent. This fact implies that for any positive value of $\lambda$ the infection can pervade the system, which is the same as the standard SIS model[4].

4.2 $N > 2$

Let $\frac{\partial u_{k,i}}{\partial t} = 0 (i = 1, 2, \cdots, N)$. Suppose $U_k = (u_{k,1}, u_{k,2}, \cdots, u_{k,N})^T$ and $V = (1, 0, \cdots, 0)^T$. Considering $\sum_{j=0}^{N} u_{k,j} = 1$, then Eqs.(1)-(4) can be written as:

$$SU_k = -\lambda k \Theta V \hspace{1cm} \text{(14)}$$

The matrix $S$ is:
\[
S = \begin{pmatrix}
-\gamma - \beta - \lambda k \Theta & \gamma & -\lambda k \Theta & \cdots & -\lambda k \Theta & -\lambda k \Theta \\
\beta & -2\beta - \gamma & \gamma & \cdots & 0 & 0 \\
0 & 2\beta & -3\beta - \gamma & \cdots & 0 & 0 \\
\vdots & \vdots & \vdots & \ddots & \vdots & \vdots \\
0 & 0 & 0 & \cdots & -(N - 1)\beta - \gamma & \gamma \\
0 & 0 & 0 & \cdots & (N - 1)\beta & -\gamma \\
\end{pmatrix}
\] (15)

Since \( \det(S) = (-\gamma)^N - \lambda k \Theta \sum_{j=1}^{N} (j - 1)!(-\gamma)^{N-j}(-\beta)^{j-1} \neq 0 \), so \( S^{-1} \) exists. Thus:

\[
U_k = -\lambda k \Theta S^{-1} V
\] (16)

\[
\sum_{j=1}^{N} j u_{k,j} = n \cdot U_k = (-\lambda k \Theta)nS^{-1}V = -\lambda k \Theta n(S^{-1}V) = -\lambda k \Theta \sum_{j=1}^{N} n_j S_{j1}^{-1}
\] (17)

where \( n = (1, 2, \ldots, N) \) and \( n_j = j \)

From (15), we get \( S_{j1}^{-1} \):

\[
S_{j1}^{-1} = \frac{(-\gamma)^{N-j}(-\beta)^{j-1}(j - 1)!}{(-\gamma)^N - \lambda k \Theta \sum_{j=1}^{N} (j - 1)!(-\gamma)^{N-j}(-\beta)^{j-1}}
\] (18)

Substituting (18) and (17) to (7), we get the self-consistent equation of \( \Theta \):

\[
\Theta = -\frac{1}{\langle k \rangle} \sum_{k} \sum_{j} \frac{\lambda k^2 p(k)(-\gamma)^{N-j}(-\beta)^{j-1} j! \Theta}{(-\gamma)^N - \lambda k \Theta \sum_{j=1}^{N} (j - 1)!(-\gamma)^{N-j}(-\beta)^{j-1}}
\] (19)

That is:

\[
\Theta = -\frac{1}{\langle k \rangle} \left\langle \sum_{j} \frac{\lambda k^2(-\gamma)^{N-j}(-\beta)^{j-1} j! \Theta}{(-\gamma)^N - \lambda k \Theta \sum_{j=1}^{N} (j - 1)!(-\gamma)^{N-j}(-\beta)^{j-1}} \right\rangle
\] (20)

Obviously, \( \Theta = 0 \) is a solution of Eq.(20). In addition, a non-zero solution with \( \Theta \neq 0 \) and \( u_{k,i} \neq 0 \) \((i = 1, 2, \ldots, N)\) is allowed if the following inequality
\[ -\frac{1}{\langle k \rangle} \left( \sum_j \frac{\lambda k^2 (-\gamma)^{N-j} (-\beta)^{j-1} j!}{(-\gamma)^N} \right) \geq 1 \]  

That is:

\[ \lambda \frac{\langle k^2 \rangle}{\langle k \rangle} \sum_{j=1}^{N} \frac{1}{\gamma} \left( \frac{\beta}{\gamma} \right)^{j-1} j! \geq 1 \]  

From (22), we get the epidemic threshold:

\[ \lambda_c = \frac{1}{f(N, \beta, \gamma) \langle k \rangle \langle k^2 \rangle} \]  

where \( f(N, \beta, \gamma) = \sum_{j=1}^{N} \frac{1}{\gamma} \left( \frac{\beta}{\gamma} \right)^{j-1} j! \), and \( f(N, \beta, \gamma) \) is an increasing function of \( N \) and \( \gamma \), but a decreasing function of the recover rate \( \gamma \). So the epidemic threshold is determined by three parameters \((N, \beta, \gamma)\) and the networks degree distribution \( p(k) \). We notice that the expression (23) involves multiplication of the well-known term \( \frac{\langle k \rangle}{\langle k^2 \rangle} \)\(^2\), which is closely related to the "average" number of secondary infections\(^7,8\). Not surprising, this result is the same as that of the standard SIS model\(^4\).

For \( p(k) = \delta_{k,k_c} \), the network is homogeneous. Then \( \lambda_c = \frac{1}{f(N, \beta, \gamma) k_c} \), we can increase the recover rate \( \gamma \) or decrease the site degree \( k_c \) and the size of the household \( N \) to lift \( \lambda_c \) to prevent the infectious disease from spreading. For large \( N \) the threshold is very small.

For \( p(k) = Ck^{-\nu} (\nu \in (2,3)) \), the network is scale-free\(^21\). When \( k \to \infty \), \( \frac{\langle k \rangle}{\langle k^2 \rangle} \to 0 \), then \( \lambda_c = \frac{1}{f(N, \beta, \gamma) \langle k \rangle / \langle k^2 \rangle} \to 0 \). So the threshold is absent for scale-free network. This implies that for any positive value of \( \lambda \), the infection can pervade the system even with high recover rate.

5 Simulation result

In above section, we have given the analytical result of the SIS model with household structure. We find that for regular network there is an epidemic threshold \( \lambda_c \); while for scale-free network the threshold disappears. For comparison, we simulate the model on regular network(see Fig1) and on scale free network(see Fig2) respectively. For simplicity(without lack of generality), we set \( \gamma = 1 \), \( N = 4 \). In Fig.1, we plot the fraction of infected individuals in the stationary state, \( \rho \), for different values of \( \beta \) on regular network with \( k_c = 4 \). Obviously, there is a threshold \( \lambda_c \) for each \( \beta \). For \( \beta = 0.6 \), \( \lambda_c \) is 0.026, in agreement with the corresponding analytical result, \( \lambda_c = 0.026 \), which can be obtained from(23). Only when \( \lambda \) is increased above \( \lambda_c \) is a significant prevalence
observed. In Fig.2, we plot the fraction of infected individuals in the stationary state, $\rho$, for different values of $\beta$ on scale-free network with $\langle k \rangle = 6$. We observe that $\lambda_c$ is absent. In contrast with the standard SIS model, of which the prevalence, $\rho$, increases slowly when increasing $\lambda$[24], our current epidemic model exhibits that $\rho$ increases rapidly with $\lambda$.

6 Summary

In this work, we analyze the SIS model that incorporates social household. We have focused on the impaction of geometrical property of complex networks and on the role of several parameters in the spreading threshold. Results show that the large household size $N$ and the high within household infection rate are more likely to cause the spread of disease. But it’s worth noticing that, even when local recovery rate is greater than effective infection rate, in divergent networks such as scale-free network, disease still can spread! This results tell us that even the local recover condition is good enough to give local protection, there are still some probability for a wide range disease spreading. It seems that this phenomenon can only exist in divergent networks with household structure. Maybe this imply that we have to care about the network structure much more than recover condition during disease spreading.

Of course, the model we have studied seems more ideal. For example, we have supposed that the existence of the $N$-member households do not affect the property of the complex networks, and also we do not take the move of the individuals into account. However, the result tells us that the properties of the complex networks play the most important role in the epidemic spreading.

ACKNOWLEDGMENT

This work was supported by the National Science Foundation of China under Grant No. 10175008. We thank research professor Yifa Tang for helpful discussion. We also acknowledge the support from The State Key Laboratory of Scientific and Engineering Computing (LSEC), Chinese Academic of Science.

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Captions of Figures

Fig1 The fraction of the infected individuals, $\rho$, as a function of the spreading rate $\lambda$ for household structure SIS model on regular networks with $k_c = 4$, $N = 4$. The simulations have been averaged over 200 different realizations.

Fig2 The fraction of the infected individuals, $\rho$, as a function of the spreading $\lambda$ for household structure SIS model on scale-free networks with $\langle k \rangle = 6$, $N = 4$. The simulations have been run in networks with $10^5$ nodes.
Fig. 1.

Fig. 2.
