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Long-range correlations improve understanding the influence of network structure on contact dynamics

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Abstract

Models of infectious diseases are characterized by a phase transition between extinction and persistence. A challenge in contemporary epidemiology is to understand how the geometry of a host's interaction network influences disease dynamics close to the critical point of such a transition. Here we address this challenge with the help of moment closures. Traditional moment closures, however, do not provide satisfactory predictions close to such critical points. We therefore introduce a new method for incorporating longer-range correlations into existing closures. Our method is technically simple, remains computationally tractable, and significantly improves the approximation's performance. Our extended closures thus provide an innovative tool for quantifying the influence of interaction networks on spatially or socially structured disease dynamics. In particular, we examine the effects of a network's clustering coefficient, as well as of new geometric measures, such as a network's square clustering coefficients. We compare the relative performance of different closures from the literature, with or without our long-range extension. In this way, we demonstrate that the normalized version of the Bethe approximation - extended to incorporate long-range correlations according to our method - is an especially good candidate for studying influences of network structure. Our numerical results highlight the importance of the clustering coefficient and the square clustering coefficient for predicting disease dynamics at low and intermediate values of transmission rate, and demonstrate the significance of path redundancy for disease persistence.

Keywords: Contact process; Interaction-network structure; Long-range correlation; Moment closure; Phase transition.

1 Introduction

Infectious diseases without permanent immunity are classically represented as SIS (susceptible–infected–susceptible) systems. Such models can be implemented on lattices, or more generally on graphs. The resultant models are characterized by a dichotomy between two possible dynamics: the disease either undergoes rapid extinction or it persists over a long time. This change of behavior can be understood as a bifurcation or phase transition governed by properties such as a disease's transmission rate. The value of the control parameter at which such a transition occurs is known as the critical value. Critical values have received considerable attention in statistical physics because (*i*) they delimit ranges of parameters values within which the qualitative behavior of a system remains unchanged and (*ii*) classical approximations – such as the mean-field approximation – are known to yield accurate predictions only far way from critical values. Studying the critical dynamics of an infectious disease is important for understanding the outbreaks and prevalence of the disease close to the transition. A better knowledge of factors influencing critical values is therefore important for improving disease control.

In many spatially or socially structured disease models, the transmission is restricted to nearest neighbors. The geometric pattern of transmission-enabling contacts between hosts can be described by a graph. The resultant interaction networks play an important role in the understanding and prediction of disease dynamics (Keeling, 1999; Eguiluz and Klemm, 2002; Keeling, 2005; Colizza et al., 2006). In particular, the geometric features of such networks can influence the critical values of a disease's phase transition. However, while the existence of this influence is well known, the important geometric characteristics of arbitrary networks, favoring or limiting disease propagation and persistence, are still not understood in general. For a long time, the study of interaction networks has been restricted to random graphs (resulting in mean-field models when fully connected, or in island models otherwise) or to regular lattices (resulting, e.g., in stepping-stone models). More realistic families of graphs have received mounting attention only recently: these graphs are characterized by statistical features such as degree distributions and clustering coefficients (Albert and Barabási, 2002; Colizza et al., 2006). In this study, we propose new geometric characteristics of graphs that are particularly useful for characterizing disease dynamics.

Individual-based spatially explicit stochastic systems are often called interacting particle systems (IPSs; Liggett, 1985; Durrett and Levin, 1994). IPSs are useful for capturing the propagation of infectious diseases (e.g. Keeling, 2003). They are also often used to describe ecological dynamics, especially when individual contacts occur among nearest neighbors (Van Baalen and Rand, 1998). However, very few exact results have been demonstrated for IPS models, which, therefore, are mostly studied through simulations. By applying mean-field approximations, IPS can be reduced to ODE (ordinary differential equation) models, with suitable state variables, such as the mean densities of susceptible and infected hosts. Critical values can then be approximated by the bifurcation points of these ODE models. However, examples of the failure of such mean-field models close to critical values are numerous (Durrett and Levin, 1994; Filipe and Gibson, 1998; Dieckmann et al., 2000 Part B; Keeling, 2005). A common reason for these failures is the inability of mean-field models to capture correlations in the state of neighboring individuals. Therefore, considerable attention has been given to the design of approximations that behave better than mean-field approximations in the vicinity of the critical values of an IPS model. Such methods are often referred to as moment closure methods (Bolker and Pacala, 1997). Prominent examples include the pair approximation (Matsuda et al., 1992) and cluster variation methods (Kikuchi, 1951; Krishnarajah et al., 2005) such as the Bethe approximation. While these methods have become popular in theoretical epidemiology and ecology (e.g., Dieckmann et al., 2000), they account for a network's structure only in limited ways, and thus cannot be used to study the influence of such structure more in general.

Analytically tractable models of statistical physics, such as the Ising model, reveal that approximations often break down because of the persistence, or even amplification, of long-range correlations close to critical values (Marro and Dickman, 1999; Snyder and Nisbet, 2000; Dickman and Martins de Oliveira, 2005). Hence, one way to improve upon classical approximations is to take long-range correlations into account, to the extent possible, thereby going beyond the correlations between nearest neighbors, which are the basis of the classical pair approximations and Bethe approximations on graphs. Here we take as our point of departure several published approximations relying on correlations between nearest neighbors only, and propose for each of them new approximations by incorporating correlations at distances larger than 1. We show that, in some cases, this method leads to more accurate predictions of critical parameter values, and also better describes the prevalence of a disease. In addition, our new

approximations allow us to explicitly take into account geometric features of graphs not captured by the classical approximations, which makes it possible to study their influence on disease prevalence.

The new method introduced here is but one of several propositions for improving classical approximations (Sato et al., 1994; Snyder and Nisbet, 2000; Filipe and Gibson, 2001; Petermann and De Los Rios, 2004). None of these earlier methods, however, was dedicated to exploring and quantifying the influence of the distinctive geometric features graphs possess. To overcome this limitation, Petermann and De Los Rios (2004) suggested an alternative method for improvements, through the careful examination of basic clusters defined by local patterns larger than pairs. While the resultant approximations can improve the description of disease dynamics for a given interaction network, the systematic comparison of disease dynamics across different types of graph is not straightforward using their method, since different basic clusters would then have to be identified and used as the basis of (hence essentially different) approximation schemes. This is why we have developed an alternative approach.

To illustrate our approach, we focus on a simple model of disease propagation on graphs by analyzing the well-studied contact process (Harris, 1974; Marro and Dickman, 1999), which serves as a standard for the spatially explicit definition of SIS models. This model has been used for modeling phenomena involving excitable media, like forest-fire models (Drossel and Schwabl, 1992), or for modeling the ecological dynamics of metapopulations (Franc, 2004). The contact process has also been widely used for studying the spread of diseases (Filipe and Gibson, 1998; Pastor-Satorras and Vespignani, 2001; Eguiluz and Klemm, 2002).

This article is organized as follows. Section 2 introduces the contact process, and Section 3 reviews classical moment closures. In Section 4, we explain in detail, using the Bethe approximation as an example, the main methodological contribution of this study: how to extend moment closure approximations based on nearest neighbors to longer-range correlations. Exploiting the insights thus obtained, we propose new geometric characteristics of graphs that have the potential to influence the dynamics of contact processes. In Section 5, we present a comparison of different approximation schemes, by critically evaluating approximations that have already been published against approximations based on our new method. We thus select the approximation best suited for studying the influence of the identified graph characteristics. We conclude with a discussion of possible improvements of our method and with guidelines for selecting suitable moment closures.

2 The contact process

The contact process is a simple model for describing the spread of an epidemic in continuous time (Harris, 1974; Marro and Dickman, 1999). We consider a graph, whose nodes are individual hosts and whose edges connect hosts that can infect one another. We will focus on graphs with homogeneous degree, in which each node has the same number h of neighbors. At any time t > 0, a host i can be in either of two states z_i : susceptible $(z_i = 0)$ or infected $(z_i = 1)$. If i is infected, it becomes susceptible with probabilistic rate μ . As units are arbitrary, we choose $\mu = 1$ without loss of generality, so that the unit of time is the expected duration between a host's infection and recovery. If i is susceptible and has a_i infected neighbors, it becomes infected with probabilistic rate βa_i , with β denoting the probabilistic transmission rate between neighboring hosts,

$$P(z_i^{t+dt} = 0 \mid z_i^t = 1) = dt , \qquad (1a)$$

$$P(z_i^{t+dt} = 1 | z_i^t = 0, a_i = a) = a\beta dt .$$
(1b)

This process implies a critical value of β that depends on the considered graph: for a transmission rate below the critical value the infection dies out, while above this value the infection spreads.

There is no known analytical solution to the contact process, and several approximations have been derived. Approximations based on moment closure have been shown to behave fairly well far from the critical point; these are described in the next section. Our goal in this study is to propose an improvement of existing moment closure approximations close to the critical point by taking into account correlations between hosts at distances larger than that between nearest neighbors.

3 Moment dynamics and moment closures

Instead of tracing through time the states of all nodes of a graph, the essential features of the contact process dynamics can be captured by salient summary statistics. ODEs can thus be constructed to describe the expected dynamics of configuration probabilities

over realizations of the stochastic process. For infinitely extended graphs, the configuration probabilities in individual realizations are identical to their expected values across realizations.

3.1 Singlet dynamics

We denote the probability that a host is infected at time t by $\rho(t)$. The dynamics of ρ are then given by

$$\frac{d\rho}{dt} = h\beta P^{1}(01) - \rho , \qquad (2)$$

(Filipe and Gibson, 2005; Peyrard and Franc, 2005), where $P^{1}(01)$ is the probability that a pair of nodes linked by an edge is in state 01. Such nodes are also said to be (nearest) neighbors, adjacent, or directly connected; their distance is defined as 1. More generally, here and in the following $P^{d_{ij}}(z_i, z_j)$ denotes the probability for two arbitrary nodes *i* and *j* at distance d_{ij} to be in states z_i and z_j . The symmetry relation $P^{d_{ij}}(z_i, z_j) = P^{d_{ij}}(z_j, z_i)$ holds by definition. These probabilities do not refer to specific nodes and must be interpreted as expectations over all pairs in the graph that possess the specified distance.

3.2 Pair dynamics

Closing Eq. (2) requires information about the dynamics of $P^{1}(01)$,

$$\frac{dP^{1}(01)}{dt} = P^{1}(11) + \beta(h-1)P^{1,1}(100)$$
$$-P^{1}(01)\left[1 + \beta + \beta(h-1)\frac{P^{1,1}(101)}{P^{1}(01)}\right]$$
(3)

(Filipe and Gibson, 2005; Peyrard and Franc, 2005), where $P^{1,1}(100)$ is the probability to observe a node in state 0 and two of its neighbors in states 0 and 1, respectively. The definition of $P^{1,1}(101)$ is analogous. Denoting the set of all neighboring pairs of nodes by *E*, we thus have $P^{1,1}(100) = P(z_i = 1, z_j = 0, z_k = 0 | (i, j), (j, k) \in E)$ and $P^{1,1}(101) =$ $P(z_i = 1, z_j = 0, z_k = 1 | (i, j), (j, k) \in E)$. Eq. (3) may seem complicated at first sight, but is nothing more than a straightforward consequence of accounting for infection and recovery events starting from, or leading to, a pair of neighbors in state 01 (Appendix 1 provides details on this derivation). Note that the roles of nodes *i* and *k* in the triplets discussed here are equivalent, while that of the central node *j* is different: it is for this reason that the ordering of states in $P^{1,1}(100) = P^{1,1}(001)$ and $P^{1,1}(101)$ is important. It also needs to be appreciated that the triplets (*i*, *j*, *k*) can either be open or closed (resulting in a triangular configuration of edges), depending on whether or not nodes *i* and *k* are neighbors.

Just as the dynamics of singlets in Eq. (2) depends on pair states, the dynamics of pairs in Eq. (3) depends on triplet states. In general, whenever interactions occur only along edges, the dynamics of configurations of order n depends on the state of configurations of order n+1. To close a dynamical system like the one jointly given by Eqs (2) and (3), one therefore needs to truncate the resultant hierarchy of equations at a certain order by approximating the required higher-order joint probabilities as functions of lower-order ones. Such relations are referred to as moment closures. Moment closure approximations go back to the cluster variation method developed in solid-state physics during the 1930s (Kikuchi, 1951; Lavis and Bell, 1999).

3.3 Order-1 closure

The simplest closure is a closure at order 1, also known as the mean-field approximation (*MF*): $P^{1}(01) = \rho(1-\rho)$. This yields a closed equation in ρ (Marro and Dickman, 1999),

$$\frac{d\rho}{dt} = h\beta\rho(1-\rho) - \rho \quad . \tag{4}$$

The SIS model often used in non-spatial epidemiology is thus recovered (with the probability ρ being interpreted as the proportion, or frequency, of infected hosts). Eq. (4) is easy to solve and study, but except in particular cases (for contact graphs with a high value of h, for contact graphs in high dimensions, or for diseases with high prevalence) the mean-field assumption, implying a well-mixed population of hosts, is too restrictive, so that this approximation's performance typically is quantitatively poor (Durrett and Levin, 1994; Filipe and Gibson, 1998; Dieckmann et al. 2000; Keeling, 2005).

3.4 Order-2 closures

In order-2 closures, triplet probabilities are approximated in terms of singlet and pair probabilities. Correlations between two nodes linked by an edge are thus no longer ignored. However, two nodes at a distance larger than 1 are still assumed to be independent. Let us introduce here the notation for triplet probabilities: $P^{d_{ij},d_{jk},d_k}(z_i,z_j,z_k)$ and $\hat{P}^{d_{ij},d_{jk},d_k}(z_i,z_j,z_k)$ denote the exact and approximated probabilities, respectively, for three arbitrary nodes i, j, and k at distances d_{ij} , d_{jk} , and d_{ik} to be in states z_i , z_j , and z_k . As for pair probabilities, these triplet probabilities must be interpreted as expectations over all triplets in the graph that possess the specified distances. Corresponding probabilities that are applicable when the third distance d_{ik} is unknown are denoted analogously, by $P^{d_{ij},d_{jk}}(z_i,z_j,z_k)$ and $\hat{P}^{d_{ij},d_{jk}}(z_i,z_j,z_k)$; for an example see Eq. (6b).

The most classical order-2 closure is known as pair approximation (PA). This approximation (Matsuda et al, 1992) has become a standard for simplifying spatiotemporal complexity in epidemiological and ecological models in which space is discrete (Iwasa, 2000; Sato and Iwasa, 2000; van Baalen, 2000),

$$\widehat{P}_{PA}^{1,1,1}(z_i, z_j, z_k) = \widehat{P}_{PA}^{1,1,2}(z_i, z_j, z_k) = \widehat{P}_{PA}^{1,1}(z_i, z_j, z_k) = \frac{P^1(z_i, z_j)P^1(z_j, z_k)}{P(z_j)} .$$
(5).

Note that $\hat{P}_{PA}^{1,1,d_{ik}}(z_i, z_j, z_k) = 0$ for $d_{ik} > 2$, since in a triplet with two pairs at distance 1 the third distance is always less or equal to 2. The pair approximation implies the assumption that the states of nodes *i* and *k* are independent given the state of node *j*.

This limitation can be overcome by the so-called Bethe approximation (*BA*), which distinguishes between closed triplets ($d_{ik} = 1$) and open triplets ($d_{ik} = 2$). Denoting by θ the graph's clustering coefficient (i.e., the probability that two neighbors of a node are neighbors of each other; Albert and Barabási, 2002; Newman, 2003), $P^{1,1}$ can, in general, be expressed as $P^{1,1} = \theta P^{1,1,1} + (1-\theta)P^{1,1,2}$. The Bethe approximation (Morita, 1994; Yedidia et al., 2005) for closed triplets is then given by the so-called Kirkwood approximation (Singer, 2004),

$$\hat{P}_{BA}^{1,1,1}(z_i, z_j, z_k) = \frac{P^1(z_i, z_j)P^1(z_j, z_k)P^1(z_i, z_k)}{P(z_i)P(z_j)P(z_k)} , \qquad (6a)$$

while the Bethe approximation for open triplets is given by the pair approximation, $\hat{P}_{BA}^{1,1,2} = \hat{P}_{PA}^{1,1,2}$. Accordingly, the Bethe approximation for $P^{1,1}$ is defined by

$$\hat{P}_{BA}^{1,1}(z_i, z_j, z_k) = \theta \hat{P}_{BA}^{1,1,1}(z_i, z_j, z_k) + (1 - \theta) \hat{P}_{BA}^{1,1,2}(z_i, z_j, z_k) .$$
(6b)

However, as opposed to pair approximation, the Bethe approximation of closedtriplet probabilities according to Eq. (6a) is not normalized, so that the probabilities $\hat{P}_{BA}^{1,1,1}(z_i, z_j, z_k)$ when summed over all combinations of z_i , z_j , and z_k do not necessarily add up to 1. We will therefore also consider the following, appropriately normalized version of the Bethe approximation (*NBA*),

$$\hat{P}_{NBA}^{1,1,1}(z_i, z_j, z_k) = \frac{\hat{P}_{BA}^{1,1,1}(z_i, z_j, z_k)}{\sum_{z_i, z_j, z_k} \hat{P}_{BA}^{1,1,1}(z_i, z_j, z_k)} .$$
(7)

Moment closures have been used to approximate population dynamics in continuous space. In this context, other order-2 closures have been proposed recently (Law and Dieckmann, 2000; Dieckmann and Law, 2000; Law et al., 2003; Murrel et al., 2004). They are referred to as power-1 closure (*P1*) and power-2 closure (*P2*) according to the number of pair probabilities that get multiplied in the terms of the closure,

$$\hat{P}_{P1}^{1,1,1}(z_i, z_j, z_k) = P(z_i)P^1(z_j, z_k) + P(z_j)P^1(z_i, z_k) + P(z_k)P^1(z_i, z_j) -2P(z_i)P(z_j)P(z_k),$$
(8)

$$\widehat{P}_{P2}^{1,1,1}(z_i, z_j, z_k) = \frac{1}{2} \left[\frac{P^1(z_i, z_j) P^1(z_j, z_k)}{P(z_j)} + \frac{P^1(z_i, z_j) P^1(z_i, z_k)}{P(z_i)} + \frac{P^1(z_i, z_k) P^1(z_j, z_k)}{P(z_k)} - P(z_i) P(z_j) P(z_k) \right].$$
(9)

The corresponding equations for $\hat{P}^{1,1,2}$ are obtained by replacing $P^1(z_j, z_k)$ with $P(z_i)P(z_k)$. For obvious reasons, the Bethe/Kirkwood approximation is referred to as a power-3 closure (Law and Dieckmann, 2000; Dieckmann and Law, 2000; Law et al., 2003; Murrel et al., 2004). A drawback of the power-1 and power-2 closures is that they may lead to negative values (which then have to be replaced with 0); for the contact process, this may happen when β , and thus ρ , are low.

4 Extensions to longer-range correlations

For each of the five closures described above (*PA*, *BA*, *NBA*, *P1*, and *P2*), it is implicitly assumed that any two nodes at distances larger than 1 are independent. We will now show how to surmount this restriction, by deriving extended versions of these closures that account for correlations at larger distances. A method for incorporating distance-2 and distance-3 correlations is explicitly detailed and illustrated for the Bethe closure. While the presented method readily extends to larger distances, only short correlation distances are computationally tractable.

4.1 Incorporating distance-2 correlations

We define the extended Bethe approximation as

$$\hat{P}^{d_{ij},d_{jk},d_{ik}}(z_i,z_j,z_k) = \frac{P^{d_{ij}}(z_i,z_j)P^{d_{jk}}(z_j,z_k)P^{d_{ik}}(z_i,z_k)}{P(z_i)P(z_j)P(z_k)} .$$
(10)

In the above expression, we approximate a pair probability, for instance $P^{d_{ik}}(z_i, z_k)$, with $P(z_i)P(z_k)$ if $d_{ik} > 2$ (instead of if $d_{ik} > 1$, as assumed in the classical closures).

To construct an extended Bethe approximation, we again use the relationship $P^{1,1} = \theta P^{1,1,1} + (1-\theta)P^{1,1,2}$. The approximation of $P^{1,1,1}(z_i, z_j, z_k)$ applies to closed triplets and remains given by Eq. (6a). For $P^{1,1,2}(z_i, z_j, z_k)$, i.e., for an open triplet, even if there is no edge between nodes *i* and *k*, many other paths of lengths larger than 1 may exist. The shortest of them is the path through the central node *j* of the triplet, which is a path of length 2. Taking into account this correlation (which of course is only part of the set of the long-range correlations existing along all possible paths), the distance-2 Bethe closure for $P^{1,1,2}(z_i, z_j, z_k)$ is

$$\hat{P}^{1,1,2}(z_i, z_j, z_k) = \frac{P^1(z_i, z_j)P^1(z_j, z_k)P^2(z_i, z_k)}{P(z_i)P(z_j)P(z_k)} .$$
(11)

The correction factor relative to the classical Bethe approximation is $\frac{P^2(z_i, z_k)}{P(z_i)P(z_k)}$ and accounts for the fact that the states z_i and z_k are not independent.

4.2 Resultant closed dynamical system

We can now exploit the distance-2 order-2 closure defined above to close the open dynamical system given by Eqs (2) and (3). This will be achieved by approximating $\hat{P}^{1,1}(100)$ based on Eqs (6a) and (11),

$$\hat{P}^{1,1}(100) = \theta \hat{P}^{1,1,1}(100) + (1-\theta) \hat{P}^{1,1,2}(100)$$

$$= \theta \frac{P^{1}(01)^{2} P^{1}(00)}{\rho(1-\rho)^{2}} + (1-\theta) \frac{P^{1}(01) P^{1}(00) P^{2}(01)}{\rho(1-\rho)^{2}}$$

$$= \frac{P^{1}(01) P^{1}(00)}{\rho(1-\rho)^{2}} [\theta P^{1}(01) + (1-\theta) P^{2}(01)].$$
(12)

The approximation $\hat{P}^{1,1}(101)$ is defined analogously based on a decomposition with respect to the third distance. We thus see that an additional equation is required for describing the dynamics of $P^2(01)$. Denoting by $T^2(z_1, z_2 \rightarrow z_3, z_4)$ the transition rate of two nodes at distance 2 from states z_1 and z_2 to states z_3 and z_4 , respectively, and accounting for all four types of transition starting from or leading to state 01, we obtain

$$\frac{dP^{2}(01)}{dt} = -P^{2}(01)[T^{2}(01 \to 11) + T^{2}(01 \to 00)] + P^{2}(00)T^{2}(00 \to 01) + P^{2}(11)T^{2}(11 \to 01) .$$
(13)

The involved transition rates at distance 2 are then approximated by

$$T^{2}(01 \to 11) = \beta(h-1)\frac{\hat{P}^{1,2}(101)}{P^{2}(01)} + \beta \frac{\hat{P}^{1,1,2}(011)}{P^{2}(01)} , \qquad (14a)$$

$$T^2(01 \to 00) = 1$$
, (14b)

$$T^{2}(00 \to 01) = \beta(h-1)\frac{\hat{P}^{1,2}(100)}{P^{2}(00)} + \beta \frac{\hat{P}^{1,1,2}(010)}{P^{2}(00)} , \qquad (14c)$$

$$T^2(11 \to 01) = 1$$
 (14d)

(Appendix 2 provides details on this derivation). In $\hat{P}^{1,2}$, only two distances are known. Three values are possible for the third distance: 1, 2, or 3 (see top row of Fig. 1). Again using the extended Bethe approximation, while ignoring correlations at distance 3, we obtain

$$\hat{P}^{1,2}(z_i, z_j, z_k) = \frac{P^1(z_i, z_j)P^2(z_j, z_k)}{P(z_i)P(z_j)P(z_k)}$$
$$[\alpha_1 P^1(z_i, z_k) + \alpha_2 P^2(z_i, z_k) + \alpha_3 P(z_i)P(z_k)]$$
(15)

The coefficients α_1 , α_2 , and $\alpha_3 = 1 - \alpha_1 - \alpha_2$ represent the proportions of the three corresponding triplet configurations. The extended closure therefore enables us to capture the influence of new geometric characteristics of the graph that go beyond the clustering coefficient θ and which, according to the approximation derived here, must be expected to affect the dynamics systematically. We refer to α_1 , α_2 , and α_3 as a graph's square clustering coefficients. Their influence can be explored based on the distance-2 order-2 closure derived here.

Finally, pair probabilities for states 00 and 11 are readily recovered from the singlet probability for state 1 and the pair probability for state 01,

$$P^{1}(11) = \rho - P^{1}(01) , \qquad (16a)$$

$$P^{1}(00) = 1 - \rho - P^{1}(01) . \tag{16b}$$

The dynamical system given by Eqs (2), (3), and (13) has thus been closed by the approximations in Eqs (12) (and the corresponding equation for $\hat{P}^{1,1}(101)$), (14), (15) and (16).

4.3 Incorporating distance-3 correlations

The method detailed above can be used to refine moment closure approximations by including correlations between nodes separated by more than two edges. We have implemented one more step in this scheme for successive refinement, by accounting for distance-3 correlations. It turns out that this enhancement modifies only the term with weight α_{3+} in Eq. (15), as this is the only place where a distance larger than 2 needs to be considered between the three nodes involved in the triplet probabilities.

Applying the same logic as introduced above for distance-2 correlations, we derive this further extension through the following steps:

1. We add a fourth equation in the system, to describe the dynamics of $P^{3}(01)$.

- 2. We express the transition rates $T^3(01 \rightarrow 11)$ and $T^3(00 \rightarrow 01)$ in terms of the probabilities $P^{1,3}(z_i, z_j, z_k)$ that three nodes *i*, *j*, and *k* are in states z_i , z_j , and z_k , given that $d_{ij} = 1$, $d_{jk} = 3$, and d_{ik} is unknown (while *i* is not on the path between *j* and *k*).
- We decompose the approximation of P^{1,3}(z_i, z_j, z_k) considering three possibilities for d_{ik}: 2, 3, or 4 (see bottom row of Fig. 1). The case d_{ik} = 1 does not occur, since it would imply d_{jk} = 2. The proportions γ₂, γ₃, and γ₄₊ = 1-γ₂ γ₃ of these three triplet configurations can be determined for any graph.
- 4. We apply the extended Bethe approximation in Eq. (10) for $d_{ik} = 2$, 3, and 4while ignoring correlations at distance 4, $P^4(z_i, z_k) = P(z_i)P(z_k)$.

By going one step further in the refinement of the moment closure approximation we can thus introduce, and study the influence of, the two new graph characteristics γ_2 and γ_3 , which help characterize a graph's geometric features beyond its clustering coefficient θ and its square clustering coefficients α_1 and α_2 .

5 Numerical results

We derived the distance-2 and distance-3 extensions of the normalized Bethe approximation (*NBA*), power-1 closure (*P1*), and power-2 closure (*P2*) according to the general method detailed for the Bethe approximation in the previous section. On this basis, we systematically compared the performances of the following six closures: mean-field approximation (*MF*), pair approximation (*PA*), Bethe approximation (*BA*), *NBA*, *P1*, and *P2* without and, when applicable, with the extensions derived above for incorporating correlations at distances 2 and 3. The performance of closures was evaluated for the contact process, by estimating the transmission rate's critical value β_c and by examining the transients and equilibria of the singlet probability ρ . Accurate predictions of β_c , and of equilibrium densities ρ close to β_c , are known to be much more challenging than predictions of ρ far away from β_c .

5.1 Method of comparison

Three different graphs with homogeneous degree were studied (Fig. 2). The first graph is conventionally referred to as the triangular grid (h = 6, $\theta = 0.4$; Fig. 2a). The second graph is the 4-regular graph (h = 4, $\theta = 0.4$; Fig. 2b), which shares the clustering coefficient with the triangular grid, but differs with regard to the degree and the coefficients affecting distance-2 and distance-3 correlations. The third graph is the square grid (h = 4, $\theta = 0$; Fig. 2c), which shares a degree of 4 with the 4-regular graph, but is characterized by a vanishing clustering coefficient. All graph statistics necessary for implementing the closure-approximations are easily obtained by analytic combinatorial computation; the resultant values (assuming infinitely extended graphs) are reported in Table 1.

We used the Runge-Kutta integration method from the package odesolve of the scientific software environment R to solve the ODE systems resulting from the different moment closure approximations. Estimations of ρ during transients and at equilibrium were obtained from simulations of the continuous-time contact process by averaging over 100 different realizations, each of which proceeded for 200 000 iterations. An iteration was defined as the modification of the state of a single node in the graph, according to the rates given by Eq. (1). The time steps between successive iterations were drawn from an exponential distribution with mean $[N\rho + \frac{1}{2}h\beta NP^{1}(01)]^{-1}$. We ran simulations of the contact process on graphs with about N = 5000 nodes. Both the ODE models and the contact process simulations were initialized with a 20% proportion of infected nodes (allocated at random in the simulations).

5.2 Comparison of different closures

Prediction of critical transmission rate

Table 2 shows the simulation-based estimates of β_c , together with predictions resulting from the six moment closure approximations for the three studied graphs. Exact analytical results for β_c are available for *MF* and *PA* (1/*h* and 1/(*h*-1), respectively). When $\theta = 0$, classical *BA* and *NBA* are equal to *PA*.

Our results show that (1) with the exception of *P1*, predictions based on distance-1 (classical) closures are poor, following the ordering $MF < P2 < PA \le BA \le NBA$, where M1 < M2 means that method *M1* is less accurate than method *M2*; (2) *P1* at distance 1

is more accurate for the 4-regular graph than for the triangular grid; (3) accounting for correlations at distances 2 and 3 improves the predictions of *NBA*, whereas the improvement for *BA* is restricted to distance 2; and (4) the distance-2 and distance-3 extensions destabilize *P1*, because negative values for the triplet probabilities are produced. Fig. 3 illustrates results (2) and (3) around the values of β_c estimated from simulations.

Prediction of singlet probability during transients and at equilibrium

Two approximations are retained for testing predictions of ρ during transients and at equilibrium: *NBA* with extension to distance-3 correlations and *P1* without extension. Fig. 4 shows the dynamics of the singlet probability ρ for the 4-regular graph, based on simulations and on these two approximations. No other approximation, without or with extension when applicable, resulted in better predictions (results not shown). Three cases can be distinguished:

- 1. For $\beta < \beta_c$ (Fig. 4a), the transients towards extinction are well predicted by *NBA* with extension and by *P1* without extension (and with lesser accuracy by the other approximations).
- 2. For $\beta \approx \beta_c$ with $\beta > \beta_c$ (Fig. 4b), all approximations overestimate the probability ρ , both at equilibrium and during the transients; however, this overestimation is smaller for *NBA* with extension and for *P1* without extension than it is for the other approximations.
- 3. For $\beta \gg \beta_c$ (Fig. 4c), all methods correctly predict the probability ρ at equilibrium, while the transients are predicted less accurately.

5.3 Influence of network structure

The extended closures we have introduced above can also be used to understand and predict the influence of those geometric features of a graph that the closures have been shown to depend on. Owing to their key role in the resultant moment-based approximations, we propose these features as good candidates for characterizing a graph's dynamically relevant geometric properties.

We tested the influence of the clustering coefficient θ and of the square clustering

coefficients α_1 , α_2 , and α_3 on the equilibrium probability ρ using *NBA* with extension to distance-3 correlations. Whereas this closure and the classical *P1* provided the best approximations, the latter does not enable taking into account the square clustering coefficients. To test the influence of a given feature of the geometry (including θ and the vector (α_1 , α_2 , α_3), with $\alpha_1 + \alpha_2 + \alpha_3 = 1$), we solved the ODE system of the momentbased approximation for a range of values of the corresponding parameter, while leaving the other parameters unchanged.

Fig. 5 shows the influence of the clustering coefficient θ for different values of the transmission rate β . We see that the role of θ is highly dependent on β : for high values of β (implying a high probability ρ of infected nodes at equilibrium) there is little effect of θ . This is consistent with the acceptable predictive accuracy of the mean-field approximation (which completely ignores the graph's geometry) for large β . The effect of θ increases as β decreases towards its critical value. We recover the fact that a high clustering coefficient θ impedes disease propagation (Keeling, 1999; Proulx et al., 2005). This is because infected nodes in a highly clustered region of the interaction network share many nearest neighbors; these links are then wasted by becoming redundant as paths for disease propagation.

Fig. 6 shows the influence of the square clustering coefficients α_1 , α_2 , and α_3 . We see that the partitioning of triplet configurations as measured by these coefficients can significantly modify the equilibrium probability ρ . Their effect is especially strong when the transmission rate $\beta > \beta_c$ is relatively low. Specifically, when α_1 is dominant among the three square clustering coefficients, ρ is lowest, whereas when α_3 is dominant, ρ is highest. The interpretation of this result is analogous to that for θ : a high value of α_1 implies a redundancy of paths of length 2 between pairs of nodes, so that infected nodes have more overlapping contacts, which consequently are wasted from the perspective of disease propagation. Again, the sensitivity to these geometric features decreases as the transmission rate β increases and vanishes altogether for very high values of β .

6 Discussion

Stochastic dynamical systems on graphs provide powerful tools for understanding

and quantifying the role of spatial or social structures, and of the resultant contact networks, in the emergence and spread of infectious diseases. Simulations of such dynamics are computationally intensive and analytical results are rarely available. A standard method of investigation is thus the derivation of approximations based on order-2 moment closures. Here we introduced a new method for improving classical order-2 moment closure approximations by accounting for longer-range correlations, thus revealing the importance of previously unrecognized general geometric features of contact networks.

Longer-range correlations are particularly important in models of population dynamics in continuous space, defined through spatial kernels for local processes such as competition and movement, rather than through nearest-neighbor interaction. It is therefore not surprising that longer-range correlations have more naturally been taken into account in those models (Bolker and Pacala, 1997; Murrel et al., 2004). However, to our knowledge, the importance of longer-range correlations for the quality of moment closures has been little studied in discrete space up until now. While methods have been developed for improving classical order-2 closures on graphs, in order to compensate for neglected correlations (Sato et al., 1994; Snyder and Nisbet, 2000; Filipe and Gibson, 2001), these methods rely on the introduction of new parameters estimated through simulations or heuristic choices. By contrast, the approach we propose here is selfcontained and the new parameters appearing in the ODEs are clearly identified as geometric characteristic of the considered graph.

In analyzing the performance of our new method for moment closures, we have focused on the contact process. This simple model of disease dynamics involves as a single essential parameter a disease's transmission rate β between neighboring hosts. And yet this model's behavior is difficult to predict accurately, especially in the vicinity of the critical value β_c below which the disease dies out. For these reasons the contact process serves as a valuable benchmark for the performance of moment closure approximations.

We have systematically evaluated the effect of the choice of moment closure on the quality of the resultant predictions. We compared five different forms of order-2 closure, several of them introduced from studies in continuous space and extended here to discrete space. Whereas other variations of the classical pair approximation have been developed, we could not consider all of them, since either they cannot be used to account for longer-range correlations (Sato et al., 1994; Filipe and Gibson, 2001; Bauch, 2005) or they are very similar to the approximations we have studied here (e.g., Van Baalen, 2000, or Keeling, 1999, have proposed forms close to the Bethe approximation, taking into account the clustering coefficient). For all the moment closures we have considered and for which this was feasible, we derived the corresponding extensions based on accounting for longer-range correlations. We found that, (1) for $\beta \approx \beta_c$ with $\beta < \beta_c$, transients towards extinction are often poorly predicted by classical moment closure approximation, (2) for $\beta \approx \beta_c$ with $\beta > \beta_c$, all considered approximations overestimate the disease prevalence both at equilibrium and during transients, and (3) for $\beta \gg \beta_c$, all considered approximations predict disease prevalence at equilibrium with good accuracy but perform less well in predicting the transients. Such a dependence in the performance of moment closures on different parameter regimes (subcritical, critical, and metastable) has already been pointed out (e.g., Krishnarajah et al. 2005).

Our systematic comparisons then demonstrated that the two moment closures leading to the best fit for disease prevalence, both at equilibrium and during transients, are the power-1 closure without extension and the normalized Bethe approximation with distance-3 correlations. Even though these two closures do not always predict the critical value of transmission rate with high accuracy, they provide reasonable estimations. If the main objective is to explore the influence of a graph's characteristics, rather than to obtain precise quantitative predictions, the normalized Bethe approximation with distance-3 correlations can be relied upon for good results.

Our study has also demonstrated that the length of loops in a contact graph may offer a valuable lead for selecting the most suitable moment closure. It is well known from statistical physics and from the study of Bayesian networks that dynamics on graphs without loops, commonly known as trees, are far simpler to analyze. Several approximation methods are rigorous on trees, but still work well on graphs with loops (e.g., Yedidia et al., 2005). On regular graphs with homogeneous degree, as those studied here, loops can be triangles, squares, or may be longer. While some approximations that do not account for long-range correlations provide good predictions when loops are short (resulting in a large clustering coefficient, such as for the triangular grid), the new method proposed here, by adding information about longer-range correlations, significantly improves the estimation of the critical transmission rate β_c when loops are longer than triangles. The improvements are particularly striking when minimal loops are squares (so that the clustering coefficient vanishes, such as for the square grid).

Based on these considerations, we offer the following suggestions for selecting suitable moment closures for approximating dynamics in discrete space: (1) mean field-approximation when studying random graphs far from the critical point or graphs with high mean degree; (2) pair approximation when predictions are needed closer to a critical point for graphs without loops (trees); (3) classical order-2 closures that distinguish between open and closed triplets when dynamics are studied close to a critical point and the graph's clustering coefficient is intermediate to high; (4) normalized Bethe approximation extended to account for longer-range correlations when studying dynamics close to a critical point for graphs that have small clustering coefficients and thus exhibit longer loops. In particular, it should be appreciated that our new method yields improvements under conditions that are known to cause the mean-field approximation's failure: our method offers satisfactory predictions close to critical points and for graphs with low homogeneous degree.

Our derivation of extensions of classical closures enables us to draw attention to several new geometric characteristics of graphs. We have shown that these are important for simplifying the spatiotemporal complexity of dynamics on graphs in general and for studying the structural determinants of disease spread in particular. In addition to the above-mentioned clustering coefficient, we have identified several new coefficients: the square clustering coefficients affecting distance-2 correlations, as well as further clustering coefficients affecting distance-3 correlations. In Caldarelli et al. (2004), the authors pointed out that the clustering coefficient alone may not suffice for understanding the key features of contact networks and therefore considered extensions based on more complex patterns of nodes. Here we have formally linked such descriptions with the spatiotemporal dynamics of the contact process. Exploring the dynamical implications of clustering and square clustering coefficients is easy based on the extended normalized Bethe approximation, and yields results that are valid for whole classes of graphs sharing given combinations of these coefficients. In this way we have found that the influence of these coefficients, and thus of the corresponding features of graph geometry, is strong for low to intermediate values of transmission rate, as these coefficients serve as indicators of path redundancy in a graph. For graphs with homogeneous degree, analytical computation of these coefficients is straightforward. When considering more complex contact networks, appropriate algorithms are readily available for evaluating such coefficients (Schreiber and Schwöbbermeyer, 2005 and references therein).

While this study has focused on graphs with homogeneous degree, real interaction networks (except in some particular cases such as orchards) tend to be complex (which includes social contact networks, networks of airline routes, and ecological networks; Newman, 2003; Proulx et al., 2005) and are characterized by more general degree distributions. The methodology presented here can be extended without conceptual problems to these situations and could thus help account for geometric features specific to non-homogeneous graphs. Pair approximation and Bethe approximations have already been derived for a graph with a general degree distribution (Peyrard and Franc, 2005). Combining the influence of degree distributions and of long-range correlations in a unified framework can be expected to provide increasingly accurate moment closure approximations of real spatiotemporal dynamics even close to critical points.

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Appendix

1 Dynamics of $P^1(01)$

Pairs of nodes at distance 1 in state 01 can originate from a node in a 00 -pair becoming infected or from a node in a 11-pair becoming susceptible. 01-pairs can disappear either by infection of the susceptible node or by recovery of the infected one. Denoting by $T^1(z_1, z_2 \rightarrow z_3, z_4)$ the transition rate of a pair of nodes at distance 1 from state (z_1, z_2) to state (z_3, z_4) , these considerations yield the dynamics of $P^1(01)$,

$$\frac{dP^{1}(01)}{dt} = P^{1}(00)T^{1}(00 \to 01) + P^{1}(11)T^{1}(11 \to 01) -P^{1}(01)T^{1}(01 \to 11) - P^{1}(01)T^{1}(01 \to 00).$$

The transition rates $T^1(11 \rightarrow 01)$ and $T^1(01 \rightarrow 00)$ are equal to 1, while the transition rates $T^1(00 \rightarrow 01)$ and $T^1(01 \rightarrow 11)$ depend on the state of the neighborhood of the node moving from state 0 to state 1.

To see this, let us consider $T^1(01 \rightarrow 11)$ and introduce some suitable notations. We will refer to as *i* the node changing from 0 to 1 and as *j* the node staying in state 1. With N(i) representing the set of the *h* nodes linked with *i* by an edge, and with $z_{N(i)}^t$ denoting the state of the nodes in the neighborhood of *i* at time *t*, then the number of neighbors of node *i* that are infected at time *t* is $a_i(z_{N(i)}^t) = \sum_{l \in N(i)} \delta_{1,z_l^t}$, with $\delta_{1,0} = 0$ and $\delta_{1,1} = 1$. Since the state of node *j* is known, we can rewrite this expression as $1 + \sum_{l \in N(i) \setminus \{j\}} \delta_{1,z_l^t}$. Using properties of the mathematical expectation *E*, we thus obtain

$$T^{1}(01 \to 11) = E[\beta a_{i}(z_{N(i)}^{t}) | z_{i}^{t} = 0, z_{j}^{t} = 1]$$

$$= \beta \left(1 + \sum_{l \in N(i) \setminus \{j\}} E[\delta_{1, z_{l}^{t}} | z_{i}^{t} = 0, z_{j}^{t} = 1] \right)$$

$$= \beta \left(1 + \sum_{l \in N(i) \setminus \{j\}} P(z_{l}^{t} = 1 | z_{i}^{t} = 0, z_{j}^{t} = 1) \right)$$

$$= \beta \left(1 + (h-1) \frac{P^{1, t}(101)}{P^{1}(01)} \right).$$

The same logic leads to

$$T^{1}(00 \rightarrow 01) = \beta(h-1) \frac{P^{1,1}(100)}{P^{1}(00)}$$

With these results, the dynamics of $P^{1}(01)$ is given by

$$\frac{dP^{1}(01)}{dt} = P^{1}(11) + \beta(h-1)P^{1,1}(100)$$
$$-P^{1}(01)\left[1 + \beta + \beta(h-1)\frac{P^{1,1}(101)}{P^{1}(01)}\right].$$

2 Dynamics of $P^2(01)$

To determine the dynamics of $P^2(01)$, we first consider the transition rate $T^2(01 \rightarrow 11)$ and specify that the node in the 01-pair that changes from 0 to 1 is called j and the other node is called k. To determine this rate, we have to evaluate the probability for each neighbor of j to be infected. One of these neighbors is the node in between j and k (since we know that the shortest path between j and k is of length 2). The ratio $\hat{P}^{1,1,2}(011)/P^2(01)$ approximates the probability of this intermediate node to be infected, while $\hat{P}^{1,2}(101)/P^2(01)$ approximates the probability of the h-1 other neighbors of j to be infected, so that

$$\hat{T}^{2}(01 \rightarrow 11) = \beta(h-1)\frac{\hat{P}^{1,2}(101)}{P^{2}(01)} + \beta\frac{\hat{P}^{1,1,2}(011)}{P^{2}(01)}$$

The triplet probability $\hat{P}^{1,1,2}(011)$ is obtained using Eq. (10). To determine $\hat{P}^{1,2}(101)$, we decompose $\hat{P}^{1,2}(z_i, z_j, z_k)$ with respect to the distance between node *i* and node *k*, in the same way we decomposed $\hat{P}^{1,1}(z_i, z_j, z_k)$ using the clustering coefficient θ . For this we recall that $P^{1,2}(z_i, z_j, z_k)$ is the probability to find three arbitrary nodes *i*, *j*, and *k* in configuration (z_i, z_j, z_k) , given that the distance between *i* and *j* is 1, that the shortest path between *j* and *k* is of length 2, and that *i* is not the intermediate node between *j* and *k*. The nodes *i* and *k* may be at distance 1 (Fig. 1a), distance 2 (Fig. 1b), or distance 3. A distance of 4 is not possible because of the triangular ine-

quality $d_{ik} \le d_{ij} + d_{jk} = 1 + 2 = 3$. As in Section 4.2, we denote by the square clustering coefficients α_1 , α_2 , and α_3 , respectively, the proportions of these three triplet configurations, so that we can decompose as follows,

$$\widehat{P}^{1,2}(z_i, z_j, z_k) = \alpha_1 \widehat{P}^{1,2,1}(z_i, z_j, z_k) + \alpha_2 \widehat{P}^{1,2,2}(z_i, z_j, z_k) + \alpha_3 \widehat{P}^{1,2,3}(z_i, z_j, z_k) .$$

Finally, we apply the distance-2 Bethe closure to determine the three approximate triplet probabilities above,

$$\hat{P}^{1,2}(z_i, z_j, z_k) = \alpha_1 \frac{P^1(z_i, z_j)P^2(z_j, z_k)P^1(z_i, z_k)}{P(z_i)P(z_j)P(z_k)} + \alpha_2 \frac{P^1(z_i, z_j)P^2(z_j, z_k)P^2(z_i, z_k)}{P(z_i)P(z_j)P(z_k)} + \alpha_3 \frac{P^1(z_i, z_j)P^2(z_j, z_k)}{P(z_j)},$$

where in the third term we have assumed that the states of the pair of nodes at distance 3 are independent.

References

Albert, R., Barabási, A.-L., 2002. Statistical mechanics of complex networks. Rev. Mod. Phys. 74, 47–97.

Bauch, C.T., 2005. The spread of infectious diseases in spatially structured populations: an invasory pair approximation. Math. Biosci. 198, 217–237.

Bolker, B.M., Pacala, S.W., 1997. Using moment equations to understand stochastically driven spatial pattern formation in ecological systems. Theor. Popul. Biol. 52, 179–197.

Caldarelli, G., Pastor-Satorras, R., Vespignani, A., 2004. Structure of cycles and local ordering in complex networks. Eur. Phys. J. B 38, 183–186.

Colizza,V., Barrat, A., Barthélemy, M., Vespignani, A., 2006. The modeling of global epidemics: stochastic dynamics and predictability. Bull. Math. Biol. 68, 1893–1921.

Dickman, R., Martins de Oliveira, M., 2005. Quasi-stationary simulation of the contact process. Physica A, 357, 134–141.

Dieckmann, U., Law, R., 2000. Relaxation projections and the method of moments, Chapter 21 in The Geometry of Ecological Interactions: Simplifying Spatial Complexity, Dieckmann, U., Law, R. and Metz, J.A.J. (Eds.), pp. 412–455. Cambridge University Press.

Dieckmann, U., Law, R., Metz, J.A.J. editors, 2000. The Geometry of Ecological Interactions: Simplifying Spatial Complexity. Cambridge Studies in Adaptive Dynamics. Cambridge University Press, UK.

Drossel, B., Schwabl, F., 1992. Self-organized critical forest-fire model. Phys. Rev. Lett. 69, 1629–1632.

Durrett, R., Levin, S., 1994. The importance of being discrete (and spatial). Theor. Pop. Biol. 46, 363–394.

Eguiluz, V., Klemm, K., 2002. Epidemic threshold in structured scale-free networks. Phys. Rev. Lett. 89, 108701.

Filipe, J.A.N., Gibson, G.J., 1998. Studying and approximating spatio-temporal models for epidemic spread and control. Philos. Trans. R. Soc. Lond. 353, 2153–2162.

Filipe, J.A.N., Gibson, G.J., 2001. Comparing approximations to spatio-temporal mod-

els for epidemics with local spread. Bull. Math. Biol. 63, 603-624.

Franc, A., 2004. Metapopulation dynamics as a contact process on a graph. Ecol. Comp. 1, 49–63.

Harris, T.E., 1974. Contact interactions on a lattice. Ann. Prob. 2, 969–988.

Iwasa, Y., 2000. Lattice models and pair approximation in ecology, Chapter 13 in The Geometry of Ecological Interactions: Simplifying Spatial Complexity, Dieckmann, U., Law, R. and Metz, J.A.J. (Eds.), pp. 227–251. Cambridge University Press.

Keeling, M., 1999. The effects of local spatial structure on epidemiological invasions. Proc. Royal Soc. Lond. B 266, 859–867.

Keeling, M., Woolhouse, M., May, R., Davies, G., Grenfell, B., 2003. Modelling vaccination strategies against foot-and-mouth disease. Nature 421, 136–142.

Keeling, M., 2005. The implications of network structure for epidemic dynamics. Theor. Pop. Biol. 67, 1–8.

Kikuchi, R., 1951. A theory of cooperative phenomena. Phys. Rev. 81, 988–1003.

Krishnarajah, I., Cook, A., Marion, G., Gibson, G., 2005. Novel moment closure approximations in stochastic epidemics. Bull. Math. Biol. 67, 855–873.

Lavis, D.A., Bell, G.M., 1999. Statistical Mechanics of Lattice Systems: 1 Closed-Form and Exact Solutions. Springer Verlag.

Law, R., Dieckmann, U., 2000. A dynamical system for neighborhoods in plant communities. Ecology 81, 2137–2148.

Law, R., Murrell, D.J, Dieckmann, U., 2003. Population growth in space and time: Spatial logistic equations. Ecology 84, 252–262.

Liggett, T.M., 1985. Interacting Particle Systems. Springer, New York.

Marro, J., Dickman, R., 1999. Nonequilibrium Phase Transitions in Lattice Models. Monographs and Texts in Statistical Physics. Cambridge University Press, UK.

Matsuda, H., Ogita, A., Sasaki, N., Sato, K., 1992. Statistical mechanics of population: the lattice Lotka-Volterra model. Progr. Theor. Phys. 88, 1035–1049.

Morita, T., 1994. Formal structure of the cluster variation method. Prog. Theor. Phys. (Supplement) 115, 27–39.

Murrel, D., Dieckmann, U., Law, R., 2004. On moment closures for population dynamics in continuous space. J. Theor. Biol. 229, 421–432.

Newman, M.E.J., 2003. The structure and function of complex networks. SIAM Rev. 45, 167–256.

Pastor-Satorras, R., Vespignani, A., 2001. Epidemic spreading in scale-free networks. Phys. Rev. Lett. 86, 3200–3203.

Petermann, T., De Los Rios, P., 2004. Cluster approximations for epidemic processes: a systematic description of correlations beyond the pair level. J. Theor. Biol. 229, 1–11.

Petermann, T., De Los Rios, P., 2004. The role of clustering and gridlike ordering in epidemic spreading. Phys. Rev. E 69, 066116.

Peyrard, N., Franc, A., 2005. Cluster variational approximations for a contact process living on a graph. Physica A 358, 575–592.

Proulx, S., Promislow, D., Phillips, P., 2005. Network thinking in ecology and evolution. Trends Ecol. Evol. 20, 345–353.

Sata, K., Iwasa, Y., 2000. Pair approximations for lattice-based ecological models, Chapter 18 in The Geometry of Ecological Interactions: Simplifying Spatial Complexity, Dieckmann, U., Law, R. and Metz, J.A.J. (Eds.), pp. 341–358. Cambridge University Press.

Sato, K., Matsuda, H., Sasaki, A., 1994. Pathogen invasion and host extinction in lattice structured populations. J. Math. Biol 32, 251–268.

Schreiber, S., Schwöbbermeyer, H., 2005. Frequency concepts and pattern detection for the analysis of motifs in networks. Trans. Comput. Sys. Biol. 3 (LNBI 3737), 89–104.

Singer, A., 2004. Maximum entropy formulation of the Kirkwood superposition approximation. J. Ch. Phys. 121, 3657–3666.

Snyder, R., Nisbet, R., 2000. Spatial structure and fluctuations in the contact process and related models. Bull. Math. Biol. 62, 959–975.

Van Baalen, M., 2000. Pair approximations for different spatial geometries, Chapter 19 in The Geometry of Ecological Interactions: Simplifying Spatial Complexity, Dieckmann, U., Law, R. and Metz, J.A.J. (Eds.), pp. 359–387. Cambridge University Press.

Van Baalen, M., Rand, D., 1998. The unit of selection in viscous populations and the

evolution of altruism. J. Theor. Biol. 193, 631–648.

Yedidia, J., Freeman, W., Weiss, Y., 2005. Constructing free energy approximations and generalized belief propagation algorithms, IEEE Trans. Inf. Theor. 51, 2282–2312.

Captions of figures and tables

Table 1. Statistics characterizing geometric features of the three studied graphs: *h* is the degree; θ is the clustering coefficient; α_1 , α_2 , and α_3 are the square clustering coefficients affecting distance-2 closures, and γ_2 , γ_3 , and γ_4 are the coefficients affecting distance-3 closures.

Table 2. Estimations of the critical value β_c of the contact process transmission rate from simulations and predictions obtained by solving the ODE systems resulting from the six different moment closure approximation without (classical) and with extension to distance-2 and distance-3 correlations. (a) 4-regular graph, (b) square grid, (c) triangular grid. No value is reported when a closure leads to a negative value for the singlet probability at equilibrium.

Fig. 1. Illustration on the triangular grid of configurations of triplets of nodes *i*, *j*, *k* that fall into the categories measured by α_1 and α_2 (top row), and by γ_2 and γ_3 (bottom row). Enlarged nodes are part of the triplet, while thickened lines depict (not necessarily unique) paths between each two of the three nodes that have shortest length and thus define their distance.

Fig. 2. The three studied graphs with homogeneous degree: (a) triangular grid with h = 6, (b) 4-regular graph with h = 4, (c) square grid with h = 4.

Fig. 3. Predictions of the singlet probability ρ at equilibrium close to β_c by *NBA* with extension to longer-range correlations and by *P1* without extension. Continuous curves with error bars: mean and 90% confidence intervals from simulations; continuous curves with symbols 1 to 3: *NBA* based on distance-1 (classical), distance-2, and distance-3 correlations, respectively; dash-dotted curve: *P1* without extension to longer-range correlations. (a) 4-regular graph, (b) square grid.

Fig. 4. Predictions of the singlet probability ρ during transients for the 4-regular graph by *NBA* with extension to longer-range correlations and by *P1* without extension. Continuous curves: three independent realizations of the contact process; dashed curve: *NBA* with extension to distance-3 correlations; dash-dotted curve: *P1* without extension to longer-range correlations. (a) $\beta = 0.39$, (b) $\beta = 0.49$, (c) $\beta = 1.3$.

Fig. 5. Predictions of the influence of the clustering coefficient θ on the probability ρ at equilibrium for any graph with homogeneous degree h = 4 and different transmission rates β .

Fig. 6. Predictions of the influence of the square clustering coefficients α_1 and α_2 on the probability ρ at equilibrium for any graph with homogeneous degree h = 4 and a transmission rate $\beta = 0.47$.

| T 11 | 1 |
|-------------|---|
| Table | |
| Table | T |

| Graph | h | θ | α_1 | α_2 | α_2 | γ_2 | γ_3 | γ_4 |
|-----------------|---|-----|------------|------------|------------|------------|------------|------------|
| Triangular grid | 6 | 2/5 | 2/15 | 6/15 | 7/15 | 2/15 | 6/15 | 7/15 |
| 4-regular graph | 4 | 2/5 | 4/45 | 4/15 | 29/45 | 2/45 | 12/45 | 31/45 |
| Square grid | 4 | 0 | 2/9 | 0 | 7/9 | 2/5 | 0 | 2/5 |

| Ta | b | le | 2 |
|----|---|----|---|
| | | | |

| Simulation | 0.458 | | |
|---------------------|-----------|------------|------------|
| Closure | Classical | Distance-2 | Distance-3 |
| Mean field | 1/4 | | |
| Pair approximation | 1/3 | | |
| Bethe approximation | 0.37 | 0.40 | 0.36 |
| Normalized BA | 0.37 | 0.41 | 0.42 |
| Power-1 | 0.44 | - | - |
| Power-2 | 0.30 | 0.32 | 0.33 |

(a)

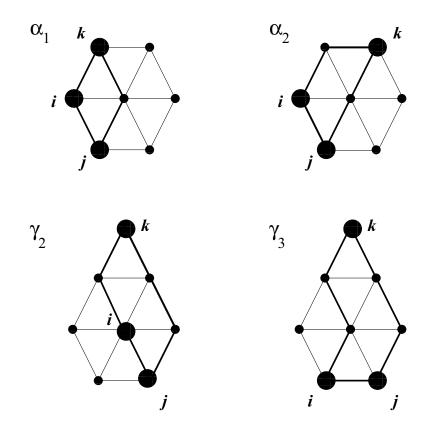
| Simulation | 0.40 | | |
|---------------------|-----------|------------|------------|
| Closure | Classical | Distance-2 | Distance-3 |
| Mean field | 1/4 | | |
| Pair approximation | 1/3 | | |
| Bethe approximation | 1/3 | 0.37 | 0.355 |
| Normalized BA | 1/3 | 0.375 | 0.395 |
| Power-1 | 0.33 | - | - |
| Power-2 | 0.26 | 0.26 | 0.27 |

(b)

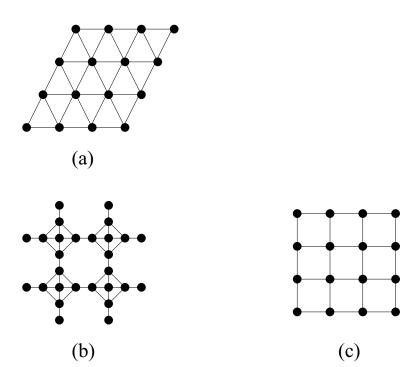
| Simulation | | | |
|---------------------|-----------|------------|------------|
| Closure | Classical | Distance-2 | Distance-3 |
| Mean Field | 1/6 | | |
| Pair approximation | 1/5 | | |
| Bethe approximation | | | |
| Normalized BA | | | |
| Power-1 | | | |
| Power-2 | | | |

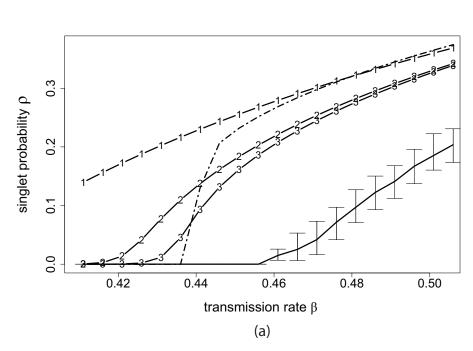
(c)











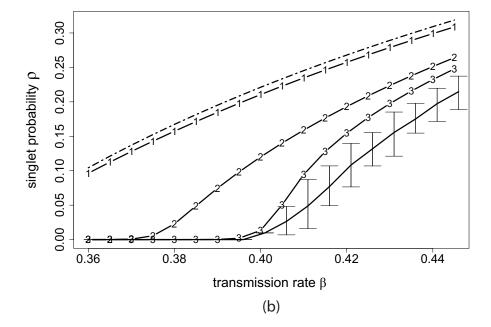
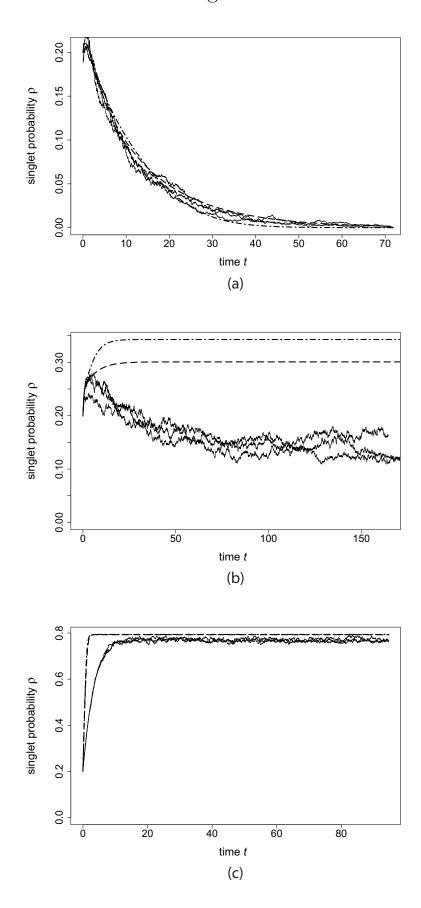


Figure 3

Figure 4





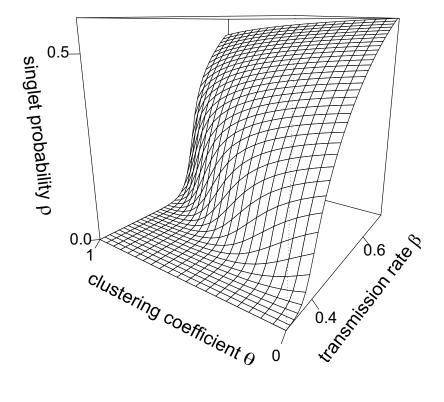


Figure 6

