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Analytically Solvable Model of Spreading Dynamics with Non-Poissonian Processes

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Non-Poissonian bursty processes are ubiquitous in natural and social phenomena, yet little is known about their effects on the large-scale spreading dynamics. In order to characterize these effects, we devise an analytically solvable model of susceptible-infected spreading dynamics in infinite systems for arbitrary inter-event time distributions and for the whole time range. Our model is stationary from the beginning, and the role of the lower bound of inter-event times is explicitly considered. The exact solution shows that for early and intermediate times, the burstiness accelerates the spreading as compared to a Poisson-like process with the same mean and same lower bound of inter-event times. Such behavior is opposite for late-time dynamics in finite systems, where the power-law distribution of inter-event times results in a slower and algebraic convergence to a fully infected state in contrast to the exponential decay of the Poisson-like process. We also provide an intuitive argument for the exponent characterizing algebraic convergence.

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I. INTRODUCTION

Events of the dynamical processes of various complex systems are often not distributed homogeneously in time but have intermittent or bursty character. This is ubiquitously witnessed in processes of nature like earthquake statistics [1], solar flare [2], and firing of neurons [3], but also in social processes like financial interactions [4] and human communication activities [5]. In these examples, the distribution of inter-event times follows a power-law behavior [1–5], in contrast to the homogeneous Poissonian processes showing exponential distribution.

Dynamical processes of complex systems can be considered to take place on a network formed by pairwise interactions between the constituents of the system [6,7]. In the recently developed approach of temporal networks [8], a link between two nodes is considered to exist only at the moment of interaction. One of the most interesting dynamical processes on networks is the spreading [9–13] that takes place on temporal networks, and the statistics of events strongly influences its most important feature, namely, the speed of propagation. This feature is of pivotal interest and importance, e.g., for halting epidemic outbreaks or promoting diffusion of innovations.

Recently much effort has been devoted to clarifying how burstiness of events influences the spreading speed, partly by using empirical data analysis [14–19] and partly by model calculations [8,14,20–22]. In those studies, the bursty character of an event sequence was found to slow down the late-time dynamics of spreading, evidenced also by a heavy tail in the inter-event time distribution. However, for the early-time dynamics, conflicting results have been reported [23]. In studies by Vazquez et al. [14] and Karsai et al. [15], the burstiness is found to slow down spreading, while other works point towards the opposite direction [16,18,21]. This calls attention to the importance of small inter-event times or to the role of lower bounds of inter-event times. The effect of lower bounds has been largely ignored, although it is present in empirical phenomena and it is important for understanding the early-time behavior of reference systems.

Model studies usually aim to reproduce some empirical observations to uncover the main mechanisms of the real underlying processes. Here, we take the perspective of constructing a tractable and analytically solvable model, where the effects of different parts of the inter-event time distribution can be studied explicitly and understood in detail. The model we consider is without correlations found in realistic data sets, except for the correlation due to the inter-event time distribution. In this way, we hope to provide a reference system, which can serve as a starting point for later studies.

II. MODEL

In order to model bursty spreading phenomena, we study deterministic susceptible-infected (SI) dynamics taking place on a temporal network of infinite size. Each node remains inactive for an inter-event time, denoted by \( l \), before becoming instantaneously active, and then it turns inactive for another inter-event time period, and so on. The
The probability of choosing a susceptible node is $1$ in the case if the chosen node is susceptible (see Fig. 1). Here, a node becomes active, it chooses a random node and infects it if it is susceptible (S) or infected (I). Whenever any infected node is active, the node takes a different approach than ours by using inhomogeneous activities of nodes. If it is susceptible (S) or infected (I), it chooses a random node and infects it if the chosen node is susceptible (S) or infected (I). Whenever any infected node becomes active, it chooses a random node and infects it if the chosen node is susceptible (S) or infected (I). Whenever any infected node becomes active, it chooses a random node and infects it if the chosen node is susceptible (S) or infected (I). Whenever any infected node becomes active, it chooses a random node and infects it if the chosen node is susceptible (S) or infected (I). Whenever any infected node becomes active, it chooses a random node and infects it if the chosen node is susceptible (S) or infected (I). Whenever any infected node becomes active, it chooses a random node and infects it if the chosen node is susceptible (S) or infected (I). Whenever any infected node becomes active, it chooses a random node and infects it if the chosen node is susceptible (S) or infected (I). Whenever any infected node becomes active, it chooses a random node and infects it if the chosen node is susceptible (S) or infected (I).

The inter-event time distribution $P_0(l)$ is assumed to be the same for all nodes, implying a homogeneous population. The activation pattern of a node is independent of whether it is susceptible (S) or infected (I). Whenever any infected node becomes active, it chooses a random node and infects it if the chosen node is susceptible (S) or infected (I). Whenever any infected node becomes active, it chooses a random node and infects it if the chosen node is susceptible (S) or infected (I). Whenever any infected node becomes active, it chooses a random node and infects it if the chosen node is susceptible (S) or infected (I). Whenever any infected node becomes active, it chooses a random node and infects it if the chosen node is susceptible (S) or infected (I). Whenever any infected node becomes active, it chooses a random node and infects it if the chosen node is susceptible (S) or infected (I). Whenever any infected node becomes active, it chooses a random node and infects it if the chosen node is susceptible (S) or infected (I). Whenever any infected node becomes active, it chooses a random node and infects it if the chosen node is susceptible (S) or infected (I). Whenever any infected node becomes active, it chooses a random node and infects it if the chosen node is susceptible (S) or infected (I). Whenever any infected node becomes active, it chooses a random node and infects it if the chosen node is susceptible (S) or infected (I). Whenever any infected node becomes active, it chooses a random node and infects it if the chosen node is susceptible (S) or infected (I).

The generating function for $I_0(t)$ is defined as $F_0(z,t) = \sum_{k=0}^{\infty} \Pr[I_0(t) = k] z^k$, and we get

$$F_0(z,t) = \left\{ \begin{array}{ll} z & \text{if } t < l, \\ F_0(z,t-l)F_1(z,t-l) & \text{if } t \geq l. \end{array} \right.$$  

Here, $F_1(z,t)$ is the generating function defined for $I_1(t)$. By taking the expectation over $l$ with $P_0(l)$, one obtains

$$F_0(z,t) = z \int_0^\infty P_0(l)dl + \int_0^t F_0(z,t-l)F_1(z,t-l)P_0(l)dl.$$  

Then, the average number of $I_0(t)$ is calculated as

$$n_0(t) = \langle I_0(t) \rangle = \frac{\partial F_0(z,t)}{\partial z} \bigg|_{z=1}$$  

where $n_0(t) \equiv \langle I_0(t) \rangle$. Taking the Laplace transform, we get

$$\tilde{n}_0(s) = \frac{1 - \tilde{P}_0(s)}{s} + [\tilde{n}_0(s) + \tilde{n}_1(s)]\tilde{P}_0(s),$$  

$$\tilde{n}_1(s) = \frac{1 - \tilde{P}_1(s)}{s} + [\tilde{n}_0(s) + \tilde{n}_1(s)]\tilde{P}_1(s),$$  

which straightforwardly leads to

$$\tilde{n}_0(s) = \frac{1 + \tilde{P}_0(s)}{s} \left(1 - \frac{\tilde{P}_0(s)}{s - \mu^{-1}}[1 - \tilde{P}_0(s)] \right),$$  

where we have used the relation $\tilde{P}_1(s) = \frac{1}{\mu} [1 - \tilde{P}_0(s)]$. Then, $n_0(t)$ can be calculated by taking the inverse Laplace
transform of \( \hat{n}_0(s) \) analytically or numerically if necessary. Note that this solution has been obtained for arbitrary inter-event time distributions and for the whole time range, enabling us to evaluate the effect of burstiness on spreading at any stage of dynamics. In contrast to this case of infinite system size, the late-time behavior of finite systems cannot be investigated analytically.

As for the non-Poissonian bursty processes, they are often characterized by broad inter-event time distributions, such as Gamma and log-normal distributions [16] and power-law distributions with an exponential cutoff [14,21]. Since these distributions have a zero lower bound for inter-event times, the effect of the lower bound on the early stage of spreading dynamics has been ignored, despite the importance of the finite lower bound in empirical phenomena. In order to systematically investigate the effect of the lower bound as well as the heavy tails of inter-event times, we consider the shifted power-law distribution with an exponential cutoff:

\[
P_0(l) = \frac{\mu_{c}^{-1}}{\Gamma(1 - \alpha, \frac{l_0}{\mu_{c}})} l^{-\alpha} e^{-l/\mu_{c}} \theta(l - l_0),
\]

where \( \Gamma \) is the upper incomplete Gamma function and \( \theta \) is the Heaviside step function. \( l_0 \) and \( l_c \) denote the lower bound and the exponential cutoff of \( l \), respectively. In the case with \( l_c \to \infty \), the value of the power-law exponent \( \alpha \) should be larger than 2 to guarantee finite \( \mu \), i.e., the mean of \( l \). The mean is related to other parameters as follows:

\[
x = y \frac{\Gamma(1 - \alpha, y)}{\Gamma(2 - \alpha, y)},
\]

where \( x \equiv \frac{l_0}{\mu} \) and \( y \equiv \frac{l_0}{\mu_{c}} \). Here, \( 0 \leq x \leq 1 \) because the mean cannot be smaller than the lower bound. When \( y = 0 \), the relation reduces to \( x = \frac{\alpha}{\alpha - 1} \). Note that setting the power-law exponent \( \alpha = 0 \) reduces the distribution to the shifted Poissonian case.

### A. Poissonian processes

As the simplest case, the Poissonian process with \( P_0(l) = \mu^{-1} e^{-l/\mu} \) results in the solution

\[
n_0(t) = e^{t/\mu}.
\]

Generally, we consider the shifted Poissonian process by setting \( \alpha = 0 \) in Eq. (10), leading to

\[
P_0(l) = \frac{1}{\mu - l_0} \exp \left( \frac{-l - l_0}{\mu - l_0} \right) \theta(l - l_0).
\]

Here, we have used \( l_c = \mu - l_0 \) from Eq. (11). Then, we get

\[
\hat{n}_0(s) = \frac{1}{s} + \frac{1}{s - \mu^{-1}} \left( \frac{1}{s + 1} e^{s l_0} - 1 \right).
\]

For the early-time dynamics, by assuming that \( s \gg 1 \), we obtain

\[
\hat{n}_0(s) \approx \frac{1}{s} + \frac{1}{s - \mu^{-1}} \left( e^{-s l_0} \right)
\]

which results in the following solution:

\[
n_0(t) \approx 1 + \frac{1}{2} \left( e^{-\frac{t - l_0}{\mu}} - e^{-\frac{t}{\mu}} \right) \theta(t - l_0)
\]

\[
\approx 1 + \frac{t - l_0}{\mu} \theta(t - l_0).
\]

As for Eq. (17), the exponential functions for \( t \ll \mu \) have been expanded. The lower bound \( l_0 \) delays the first branching, while at times later than \( t = l_0 \), the spreading speeds up. Let us define a dimensionless spreading rate at the moment of the first branching as follows:

\[
C_0 = \mu \frac{d n_0}{dt} \bigg|_{t=l_0},
\]

\[
C^p_0(x) = \frac{1}{1 - x}.
\]

Next, we study the late-time dynamics, where the late time for infinite-size systems corresponds to the intermediate time for finite-size systems. Since it is evident that \( n_0(t) \sim e^{t/\mu} \) for large \( t \), we characterize the asymptotic behavior by defining the coefficient of the leading exponential term as

\[
C_\infty = \lim_{t \to \infty} n_0(t) e^{-t/\mu}
\]

\[
= \lim_{s \to 0} \hat{n}_0(s + \mu^{-1}).
\]

Here, we have used the final value theorem [25]

\[
\lim_{t \to \infty} f(t) = \lim_{s \to 0} \hat{f}(s),
\]

with \( f(t) = n_0(t) e^{-t/\mu} \). By plugging Eq. (14) into Eq. (21), we obtain

\[
C_\infty(x) = \frac{1}{(2 - x) e^{x} + 1}.
\]

This result implies that the finite lower bound suppresses the late-time spreading dynamics.
B. Non-Poissonian processes

Now we consider the general form of $P_0(l)$ in Eq. (10). The Laplace transform of $P_0(l)$ is obtained as follows:

$$\tilde{P}_0(s) = (sl_c + 1)^{\alpha-1} \frac{\Gamma(1-\alpha,y(sl_c + 1))}{\Gamma(1-\alpha,y)}.$$  \hfill (24)

To investigate the early-time dynamics of $n_0(t)$, we consider the case with $s \gg 1$. By expanding the incomplete gamma function, we obtain

$$\tilde{P}_0(s) \approx \frac{y^{-\alpha}e^{-y}}{s^{\alpha-1}} e^{-ys},$$  \hfill (25)

$$\tilde{n}_0(s) \approx \frac{1}{s} + A \frac{1}{s} \frac{e^{-ys}}{s^{\alpha-1}} e^{-ys},$$  \hfill (26)

with $A = \frac{1}{s^{\alpha-1} \Gamma(1-\alpha,y)}$, leading to

$$n_0(t) \approx 1 + A \left( e^{-\alpha} - e^{-\alpha t} \right) \theta(t - t_0).$$  \hfill (27)

The spreading rate at $t = t_0^+$, i.e., $C_0$, is obtained as

$$C_0(x,y,\alpha) = \frac{1}{x} \frac{y^{-\alpha}e^{-y}}{\Gamma(1-\alpha,y)}. $$  \hfill (28)

Note that $x$, $y$, and $\alpha$ are not independent by means of Eq. (11) and that $C_0(x,y,0) = \frac{1}{x^\alpha} = C_0^p(x)$. For the comparison to shifted Poissonian processes, we define the ratio of spreading rates as

$$r_0(x,y,\alpha) = \frac{C_0(x,y,\alpha)}{C_0^p(x)},$$  \hfill (29)

which turns out to be exactly the same as the ratio of

$$\frac{P_0(l = t_0^+)}{P_0^p(l = t_0^+)}. $$  \hfill (30)

This indicates that the probability of having $l = t_0^+$ determines the early-time spreading dynamics. In addition to $r_0(x,y,0) = 1$, by definition, it is found that $r_0 > 1$ for $\alpha > 0$ and $r_0 < 1$ for $\alpha < 0$ [see Fig. 2(a)]. Provided that $\alpha > 0$, one can conclude that the non-Poissonian bursty activity always accelerates the early-time spreading dynamics as compared to the shifted Poissonian case with the same mean $\mu$ and the same lower bound $l_0$.

For the late-time dynamics, we focus on the asymptotic behavior characterized by $C_\infty$ in Eq. (20). Similarly to the Poissonian case, one gets the following general result:

$$C_\infty(x,y,\alpha) = \frac{(1 + \frac{x}{y})^{\alpha-1} \Gamma(1-\alpha,x+y)}{\Gamma(1-\alpha,x+y) - (1 + \frac{x}{y})^{\alpha-1} \Gamma(1-\alpha,x+y)}. $$  \hfill (31)

FIG. 2. Exact solutions of (a) the ratio of initial spreading rates $r_0(x,y,\alpha)$ in Eq. (29) and (b) the ratio of asymptotic coefficients $r_\infty(x,y,\alpha)$ in Eq. (32). We note that $x = \frac{y}{2}$ and $y = \frac{y}{2}$, with the lower bound $l_0$, the mean $\mu$, and the cutoff $l_c$ of the inter-event time distribution in Eq. (10). We plot $r_0(x,y)$ and $r_\infty(x,y)$ for various values of $y$ and, in the insets, $r_0(y,\alpha)$ and $r_\infty(y,\alpha)$ for various values of $\alpha$. By definition, the ratios have the value of 1 for $\alpha = 0$. In the limits of $x \rightarrow 1$ and/or $y \rightarrow \infty$, one gets $P_0(l) \rightarrow \delta(l - \mu)$; thus, $r_0$ and $r_\infty$ have the value of 1.

Note that $C_\infty(x,y,0) = \frac{1}{(1-x)e^{-\alpha} - 1} = C_\infty^p(x)$. Similarly to $r_0$, we define the ratio

$$r_\infty(x,y,\alpha) \equiv \frac{C_\infty(x,y,\alpha)}{C_\infty^p(x)}.$$  \hfill (32)

In addition to $r_\infty(x,y,0) = 1$, by definition, it is found that $r_\infty > 1$ for $\alpha > 0$ and $r_\infty < 1$ for $\alpha < 0$ [see Fig. 2(b)]. This implies that the non-Poissonian bursty activity accelerates the late-time dynamics as compared to the corresponding Poissonian processes. Our analysis is confirmed by the numerical simulations as depicted in Fig. 3(a).

The correction term to the exponential growth for late-time dynamics is obtained for the case of $y = 0$ such that

$$n_0(t) \approx C_\infty e^{t/\mu (1 - Be^{-\lambda t})}.$$  \hfill (33)

Both $B$ and $\lambda$ can be analytically obtained by taking the Laplace transform of the above equation for $s > \mu^{-1}$. By defining $\epsilon \equiv s - \mu^{-1} > 0$, Eq. (33) is transformed to

$$e^{\epsilon \mu^{-1} - \epsilon} = C_\infty \left( 1 - \frac{Be}{\lambda + \epsilon} \right). $$  \hfill (34)

By expanding both sides up to the order of $\epsilon^2$ for small $\epsilon$ and comparing the coefficients, one can get $\lambda$ and $B$ as functions of $x$ and $\alpha$. In the limit of $x \rightarrow 0$ ($\alpha \rightarrow 2$), we obtain $\lambda \rightarrow \mu^{-1}$ and $B \rightarrow 1$, resulting in
FIG. 3. Numerical simulation results of Poisson-like and non-Poissonian cases for systems with infinite size (a) and finite size $N = 4 \times 10^3$ (b–e). In (a,b), we set $\mu = 1$ and $l_0 \approx 0.106$ for all cases, from which we get $\alpha \approx 2.118$ for $l_c \to \infty$ (power law) and $\alpha = 1.5$ for $l_c = 100 l_0$ (power law with cutoff). Each curve is averaged over up to $1.5 \times 10^6$ runs. In (c), we plot the fractions of susceptible nodes, $1 - n_0(t)$, using the same results as in (b), as well as for various values of $\alpha$ in the power-law case with $\mu = 1$ (thin reddish curves). The Poisson-like case is fitted with $e^{-\lambda t}$, while the power-law cases are fitted with $\Gamma^{-\alpha}$. The estimated $\beta$ as a function of $\alpha$ is presented in (d), comparable with $\beta = \alpha - 1$. For understanding this scaling relation, in (e) we plot the individual runs with different initial inter-event times (from blue to red curves) with their average (black solid curve) in the case with $\alpha \approx 2.118$.

$$n_0(t) = C_\infty e^{\alpha t/\mu} - C_\infty.$$

The constant term implies the existence of residual times that are effectively infinite because of the non-normalizability of $P_\lambda(w)$, whose tail is characterized by the exponent $\alpha - 1$.

C. Finite-size effects

Finally, we consider the effect of the finite system size $N$ on the spreading dynamics. Whenever an infected node becomes active at time $t$, it chooses a random node and infects it if the chosen node is susceptible. In other words, the infection occurs with probability $\frac{N - n_0(t)}{N - 1} \approx 1 - \frac{n_0(t)}{N}$, but otherwise it does not occur. Since the exact solution could not be obtained, we perform numerical simulations to obtain the spreading dynamics shown in Figs. 3(b), 3(c), 3(d), and 3(e). The early- and intermediate-time dynamics are consistent with the early- and late-time dynamics of the infinite system, respectively. For the late-time dynamics, the non-Poissonian bursty activity results in a slower algebraic convergence to the fully infected state, i.e., $\sim t^{-\beta}$ with $\beta = \alpha - 1$ for the power-law case, in contrast to the exponential decay of the Poisson-like process, i.e., $\sim e^{-t}$.

We provide an intuitive argument for the relation $\beta = \alpha - 1$ in the case of power-law inter-event time distributions. While the average fraction of susceptible nodes decays algebraically, the fraction of susceptible nodes for each run remains close to 1 and then suddenly decays exponentially, as shown in Fig. 3(e). The period staying close to 1 must be governed mostly by the inter-event time initially given to the first infected node. Therefore, the average fraction of susceptible nodes can be obtained as the fraction of runs that did not reach the fully infected state at time $t$. Such a fraction of runs is equal to the probability of having $l > t$, which is proportional to $t^{-\beta - 1}$, leading to $\beta = \alpha - 1$. Considering the dominant role of inter-event time given to the first infected node, this argument should be valid for the SI dynamics with a power-law inter-event time distribution on a broad class of networks.

III. CONCLUSIONS

We have introduced an analytically solvable model for studying the effect of non-Poissonian bursty inter-event time distributions on the SI spreading dynamics. Our model is devised to be stationary from the beginning. For this, we make a realistic assumption that for each infection event the infecting node should wait another inter-event time and the newly infected node should wait a residual time before they become active again. With this assumption, we can obtain the analytic solution of spreading dynamics in infinite systems for arbitrary inter-event time distributions, but more importantly for the whole time range. With our analytic solution, the role of the lower bound of inter-event times has been exactly compared for Poisson-like and non-Poissonian processes. We also note that, as done in the case of null models, randomizing or shuffling the event timings to destroy temporal correlations can eliminate the lower bound of inter-event times. Hence, for systematic comparison between the original situation and the null model, one needs to employ a shuffling method that conserves the lower bound.

Let us next discuss apparently conflicting results for the early-stage spreading dynamics presented in Refs. [15,18]. The early-stage dynamics is mainly driven by small inter-event times, which generally leads to a faster spreading for non-Poissonian cases than for Poisson-like cases. This is the case when any infected node finds a susceptible node without a topological limit. However, it is well known that the mobile-phone-call network (MCN) in Ref. [15] has the community structure accompanying the bottleneck effect due to weak links between communities [26]. The large inter-event times associated with such weak links are the...
main reason for slowing down the spreading on the MCN, while the role of small inter-event times coupled with local topological structure is still important for spreading within communities. This leads us to study more realistic models as a future work. On the other hand, the spreading is enhanced by the burstiness on the sexual network. This might be because the sexual network has different community and/or temporal structures from the MCN, so the spreading on the sexual network can be better understood by our model to some extent.

As a followup, our model can be extended to incorporate a number of other complex situations, such as susceptible-infected-recovered spreading dynamics and cascading phenomena. Our results should be of interest beyond the community of network scientists because non-Poissonian processes are ubiquitous and yet little is known about their impact on the large-scale dynamics.

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