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Epidemic spreading in modular time-varying networks

Matthieu Nadini¹,⁶, Kaiyuan Sun², Enrico Ubaldi³, Michele Starnini⁴,⁵, Alessandro Rizzo⁶ & Nicola Perra⁷

We investigate the effects of modular and temporal connectivity patterns on epidemic spreading. To this end, we introduce and analytically characterise a model of time-varying networks with tunable modularity. Within this framework, we study the epidemic size of Susceptible-Infected-Recovered, SIR, models and the epidemic threshold of Susceptible-Infected-Susceptible, SIS, models. Interestingly, we find that while the presence of tightly connected clusters inhibits SIR processes, it speeds up SIS phenomena. In this case, we observe that modular structures induce a reduction of the threshold with respect to time-varying networks without communities. We confirm the theoretical results by means of extensive numerical simulations both on synthetic graphs as well as on a real modular and temporal network.

Network thinking has become a prominent and convenient paradigm to unveil the properties of complex systems. Such a paradigm has been rapidly enriched, giving rise to variants that account for inherent features of real complex systems inferred by the availability of large, often time-resolved datasets¹–³. Three are the main features that have captured the attention of researchers in the area. The first is heterogeneity in the statistical distributions of key topological properties such as the number of connections per node (degree) and the intensity of interactions (weight)⁴,⁵. This property is one of the hallmarks of complexity and is linked to a range of non-trivial dynamics⁶,⁷. For example, heterogeneity in the connectivity patterns makes networks extremely fragile to the spreading of infectious diseases and malicious attacks⁶,⁷. The second feature regards the presence of modules and communities⁸. Available datasets have highlighted that real networks are organized in modules, or communities, whereby the density of links within the community is much greater than the density of links between communities. On the one hand, communities can be treated as fairly independent entities within a large network, like the behaviour of different organs within the same body. On the other hand, from time to time, phenomena originating in a community may involve a huge portion, if not all, of the network. This is for example the case of pandemics originating from local outbreaks⁹–¹². Nevertheless, the role played communities is still ambivalent. For example, the presence of communities might slow down or speed up the propagation of a disease and facilitate the spreading of social norms¹³–²⁰. Detecting communities in a real system is not a trivial task, due to their fuzzy, vanishing, and overlapping nature. Also, in most of the available datasets communities are not explicitly labeled, so the validation of community detection algorithms cannot often be rigorously carried out²¹. Finally, networks are characterised by non trivial temporal dynamics²²,²³. The propensity of nodes to initiate and to attract interactions per unit time is typically heterogeneously distributed²⁴,²⁵. The same applies to the duration and the time interval between connections²⁶,²⁷. Furthermore, the creation or renewal of interactions might be correlated, and the dynamics driving the temporal evolution of networks are function of the time-scale considered²⁸,²⁹. However, the large majority of studies on dynamical processes unfolding on networks have been conducted under the hypothesis of time scale separation which effectively neglects all such features. In particular, the evolution of the process and the evolution of network are considered to take place at well-distinct time scales. Within this paradigm two opposite limits have been considered. In the first case, the dynamical process is assumed to be much faster than the evolution of the network. This is the limit of quenched/static networks²⁹. Here, networks are fully characterised from an adjacency

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matrix, \( A \), whose entries are non-zero for all connected pairs \( i \rightarrow j \). The second case instead, is the opposite limit where an annealed version of the network can be considered and averaging (mean-field) techniques can be applied\(^{30–32}\). Annealed networks are fully characterised by an average adjacency matrix \( \overline{A}_{k,k} \) describing the probability of connection for nodes of degree \( k \), and degree \( \overline{k} \). In our case instead, the time scale regulating the dynamical process and the evolution of the network are comparable, the time scale approximation is not valid. This is the regime of time-varying networks\(^{23}\). Interestingly, the temporal nature of interactions might inhibit or facilitate spreading processes evolving at comparable time-scales\(^{11,24,34–46}\). The effects introduced by communities and time-varying connectivity patterns on dynamical processes have been mostly scrutinised separately. However, as few recent works pointed out, the two attributes are connected and their interplay introduces non-trivial effects, such as segregated behaviours and formation of hierarchical structures\(^{47}\); or the intricate competition between topological and temporal correlations\(^{48}\). The presence of groups, think for example the interaction network of students in a school, introduces specific dynamics that deeply affect spreading processes. A thorough modelling and study of these phenomena may be useful to define prioritisation of interventions and containment strategies in epidemic spreading\(^{49}\).

Altogether, these observations call for a general modelling framework aimed at characterising both features and single out their effects on real networks. The model presented in this paper leverages on the paradigm of Activity Driven Networks (ADNs) to model realistic temporal networks where the node and link dynamics co-evolve at comparable time scales\(^{34,50}\) and includes the modularity phenomena, whereby connection patterns can be set to preferentially occur within a given community, rather than outside the community toward the rest of the population. While the coevolution of node and link dynamics is inherently considered in ADNs by construction, modularity is here modelled by a single parameter that regulates the interplay between the link formation within and outside of the community. In the context of epidemic processes on time-varying networks, our model is first characterised analytically. Then it is used to study the behaviour of different contagion processes on synthetic networks, and on a large, time-resolved dataset of scientific collaborations. Results and methods are discussed in detail in the following sections.

**Results**

Here, we study the effect of modularity (i.e., the presence of communities in the network) on time-varying networks. To this extent, we introduce a model of time-varying networks with tunable modularity, able to capture several features of real temporal graphs. We derive an analytical characterisation of the model, and we study the behaviour of the Susceptible-Infected-Recovered (SIR) and the Susceptible-Infected-Susceptible (SIS) epidemic processes unfolding on its fabric\(^{51}\). Remarkably, while the presence of tightly connected clusters inhibits SIR processes, it favours the spreading of SIS-like diseases lowering the epidemic threshold. Interestingly, similar results have been recently obtained in models of time-varying networks characterised by correlated topological features induced by reinforcement of specific ties\(^{42}\). We confirm the theoretical picture emerging from synthetic networks by means of extensive simulations on a real word dataset of scientific collaborations within the American Physical Society (APS). Our results contribute to characterise the mechanisms, and their interplay, behind the complex, and often contradictory, behaviour of dynamical processes unfolding on real networks.

**Modular activity driven networks.** The system under investigation is composed by \( N \) nodes. Each node \( i \) is characterised by an activity rate \( a_i \), that describes its propensity to engage in social interactions with other nodes. To capture empirical observations performed in a wide set of systems ranging from R&D to online interactions networks\(^{25,30–32}\), we consider activity rates heterogeneously distributed, and extracted from a continuous functional form \( F(a) = B a^{-\gamma} \), where \( a \in [\varepsilon, 1] \) and \( \varepsilon = 10^{-3} \) to avoid divergence in the distribution. Furthermore, each node is assigned to only one group/community. To take into account empirical evidences, the size of each community is extracted from a heavy-tailed distribution, i.e. \( P(s) = Cs^{-\gamma} \) with \( s \in [s_{\text{min}}, \sqrt{N}] \)\(^{53}\). Therefore, we do not limit ourselves in studying a fixed number of modules\(^{39}\), whilst their number is driven from the model’s parameters. The assignment of nodes to communities is done as follow. A community of size \( s \) is extracted. The ID of \( s \) nodes is progressively assigned to the community. The processes is repeated until all \( N \) nodes are assigned to one community. Very rarely, all nodes can be perfectly assigned to the extracted community. However, in the last extraction we might have available only a fraction of the nodes necessary to fill the community. This is the regime of time-varying networks\(^{23}\). Interestingly, the temporal nature of interactions might inhibit or facilitate spreading processes evolving at comparable time-scales\(^{11,24,34–46}\). The effects introduced by communities and time-varying connectivity patterns on dynamical processes have been mostly scrutinised separately. However, as few recent works pointed out, the two attributes are connected and their interplay introduces non-trivial effects, such as segregated behaviours and formation of hierarchical structures\(^{47}\); or the intricate competition between topological and temporal correlations\(^{48}\). The presence of groups, think for example the interaction network of students in a school, introduces specific dynamics that deeply affect spreading processes. A thorough modelling and study of these phenomena may be useful to define prioritisation of interventions and containment strategies in epidemic spreading\(^{49}\).

All the interactions have a constant duration \( \Delta t \). In the model, neither self-loops nor multiple edges are allowed. In the following, without loss of generality, we fix \( \Delta t = 1 \). Furthermore, we consider the case \( m = 1 \).

Given an heterogeneous distribution of activity, at each time step, the model generates a random, structureless network in which few nodes are active. The modular features of the network emerge integrating connections in time. Such time-integrated properties, at different time regimes, can be computed analytically. In the following,
we will report the results for the evolution of the average number of connections of each node $\langle k_i(t) \rangle$ (average degree) and the overall degree distribution $\rho(k)$. The complete set of results is shown in the Supplementary Information (SI).

To solve the average degree’s dynamics, let us introduce the effective activity $\bar{a}_i = a_i + \langle a \rangle$ (where $\langle a \rangle$ is the average value of the activity distribution) and the mixing parameter $\mu' = 1 - \mu$. We refer to the degree of node $i$ at time $t$ as $k_i(a, s, t)$, where $s$ is the node’s community size. By defining an activity class as the group of nodes featuring similar activity values $a$, we set the average in-community degree $\langle k_c(a, s, t) \rangle$ to be the average number of connections that nodes belonging to the activity class $a$ and falling in communities of size $s$ have toward nodes of their same community. The latter grows as

$$\tau(a, s) \sim -\frac{t}{\tau(a, s)}$$

where $\tau(a, s)$ is the characteristic time that it takes for the degree $k_c(a, s, t)$ of nodes of activity $a$ belonging to a community of size $s$ to be $k_c(a, s, t) \sim (s - 1)$, being $s - 1$ the maximum value of the in-community degree (see the Supplementary Information for the evaluation of $\tau(a, s)$).

Similarly, we can define the average out-community degree $\langle k_o(a, t) \rangle$ as the number of connections that nodes of activity class $a$ have outside of their communities at time $t$. We expect this quantity to be independent of the nodes’ community size $s$ so that, for large networks, we can write:

$$\langle k_o(a, t) \rangle = \mu' \bar{a} t$$

The average total degree $\langle k(a, s, t) \rangle$ can be computed as the simple sum between the two previous equations, obtaining

$$\langle k(a, s, t) \rangle = \langle k_c(a, s, t) \rangle + \langle k_o(a, t) \rangle \simeq \begin{cases} \bar{a} t & t \ll \tau(a, s) \\ \mu' \bar{a} t + (s - 1) t & t \sim \tau(a, s) \\ \mu' \bar{a} t & t \gg \tau(a, s) \end{cases}$$

Three regimes are readily identified: an initial growth in which both the in-community and the out-community degrees are growing linearly in time, followed by the slowing down of the in-community degree, which saturates to $s - 1$, and then a further linear regime driven only by the out-community degree growth. Figure 2 shows that the numerical simulations perfectly match with the theoretical formulas (see the SI for details).

Noticeably, the long time evolution of the node degree is linear in time and proportional to its activity class $a$, so that we find the asymptotic degree distribution of the system to feature the same functional form of $F(a) \propto a^{-\nu}$, as found in non-modular activity driven networks:

$$F(a) \propto a^{-\nu}$$

Figure 1. Schematic representation of the model. In red, we show active nodes. Straight lines and arcs describe links connecting nodes in the same or in different communities respectively. In the bottom right panel we show the integrated network obtained as the union of $G_1$, $G_2$, $G_3$. In Fig. 3, we integrate the network for $T = 10^5$ and we plot the three degree distributions. As expected, the out-community $\rho(k_o)$ and the total $\rho(k)$ degree distributions fall as power laws with exponent $-\nu$. On the other
hand, the in-community degree $\rho(k_c)$ saturates to the community size distribution $P(s)$, as all the nodes reach
their maximum in-community degree value ($s-1$), being that the modules’ size is far smaller than the network
size $N/s\ll N$. On the contrary, the out-community degree takes longer times to saturate to its maxi-
mum value $N/s$.

It is worth stressing that the results presented in this section apply to the networks obtained integrating links
over time. A process unfolding on such networks, in general, will be affected by the time-aggregated features of
the graph. The extent to which this is true, is function of the interplay between the time-scale describing its evo-
lution, $\tau_P$ and the various $\tau(a, s)$. In the limit $\tau_P \ll \tau(a, s)$ the process would effectively evolve on the instantaneous,
annealed networks that are characterised by a small average degree and modularity. In the opposite limit
instead, the process would effectively unfold on static networks obtained integrating links over longer time char-
acterised by high average degree and low modularity. Indeed, the average degree in this regime will be dominated
by out-community links that make the connections between different communities increasingly stronger, thus
increasingly destroying the identity of communities. In the limit $\tau_P \sim \tau(a, s)$ the process would effectively evolve
on maximally modular networks (for a given set of parameters). Arguably, this is the most interesting regime that
we will consider in the following.

**Epidemic processes on modular activity driven networks.** Let us turn our attention on the dynamical
properties of SIR and SIS processes (see the Methods section for a detailed definition of the two) unfolding on the
proposed model. Although similar, the two processes are intrinsically different. Indeed, SIR processes are
always characterised by the so called disease-free equilibrium, provided $d_i N = 0$. The illness eventually disappear,
i.e., $I = 0$ for $t \to \infty$. SIS models instead allow the existence of an endemic state where a finite and constant frac-
tion of infected individuals permanently colonise the population, i.e., $I > 0$ for $t \to \infty$. Here, we focus on a central
concept of contagion phenomena: the epidemic threshold. This quantity defines the conditions necessary for the
spreading of the illness. In static graphs the expression is given by the principle eigenvalue of the adjacency matrix. In
time-varying networks instead, the threshold is determined by the interplay between the time-scales of the conta-
gion and network evolution processes. In the case of SIR models, we also consider another important
quantity: the epidemic size $R_\infty$ which is defined as the final ratio of recovered nodes. This describes the fraction of nodes affected by the disease.

To develop a deeper understanding, let us derive the mean-field level dynamical equations describing the contagion process in modular activity driven networks. We define the activity block variables $S_{a,s}$, $I_{a,s}$, and $R_{a,s}$ as the number of susceptible, infected and recovered individuals, respectively, in the class of activity $a$ and community of size $s$ at time $t$ (to enhance readability, we omit to note the dependence on time). This allows us to write the mean-field evolution of the number of infected individuals, for a SIR process, in each group of nodes with activity $a$ as:

$$
\frac{dI_{a,s}}{dt} = -\gamma I_{a,s} + \lambda a \mu a_s I_{a,s} \left[ \frac{I_{a,s}}{s} + (1 - \mu) \frac{I}{N} \right] + \lambda \sum_{a'} a' a \mu I_{a',s} I_{a,s} \left[ \frac{I_{a,s}}{s} + (1 - \mu) \frac{I_{a',s}}{N} \right],
$$

where $I$ and $I'$ are the number of infected in communities of size $s$ and in the whole network, respectively. The first term in the r.h.s. accounts for the recovery of infected individuals. The other four terms account for the probability that a Susceptible node in a community of size $s$ connects to an Infected node inside (first) or outside (second) its community acquiring the infection, and for the probability that an Infected node of class $a'$ connects to a Susceptible node inside (third) or outside (forth) a community of size $s$, contracting the disease. For simplicity, we consider that $N - s - N$ and, at least initially, $I - I_s - I_a$. Summing over all the activities and community sizes, and considering only the first order terms in $I$, $I_s$, and their products, we obtain

$$
\frac{dI}{dt} = -\gamma I + \lambda \langle a^2 \rangle I + \lambda \Theta + \lambda \mu \sum_s \langle (a) I_s - (a) I \rangle I_s,
$$

$$
\frac{d\Theta}{dt} = -\gamma \Theta + \lambda \langle a^2 \rangle I + \lambda \langle a \Theta \rangle + \lambda \mu \sum_s \left[ \langle (a) I_s \rangle - \langle a \rangle \langle I_s \rangle \right],
$$

where we defined $\Theta = \sum_s a \mu I_s$ and $\Theta_s = \sum_s a \mu I_s$. The term $\langle a^2 \rangle = \sum_s N_s a^2 / s$ describes the moments of the activity distribution in any community of size $s$. The second, auxiliary, equation is obtained from the first by multiplying both sides by $a$ and summing over all $s$ and $a$. The epidemic threshold, in principle, can be derived evaluating the principle eigenvalue of the Jacobian matrix of the system of differential equations in $I$ and $\Theta$. In general, a closed expression for the threshold does not exist. However, we can point out some interesting observations.

First of all, the terms associated to $R_{a,s}$ vanish, implying that, at the first order, the thresholds of both SIR and SIS are equal. Furthermore, the terms in $\mu$ weight a comparison between the moments of the activity distribution in the network with the corresponding quantities evaluated inside each community. If fluctuations of these terms are negligible, due for example to very large community sizes or to narrow distribution of activity, the equations become equivalent to the case $\mu = 0$. In the limit $\mu \to 0$ the network has no modular structure, and the threshold, for both SIR and SIS, becomes $\beta_1/\gamma \geq 2((1 + \sqrt{\gamma}) \mu a_s a^2 / s$ as derived with different approaches in ref.\cite{24,41,45,69,70,26}. We defined $\gamma = \langle a^2 \rangle / (a)^2$, where the moments are evaluated over the whole network. As expected, the spreading condition is determined by the interplay between the time-scale of the contagion process and the time-scale of the network. Furthermore, the threshold is significantly larger with respect to the case in which the disease would spread in static or annealed networks generated integrating connections over time\cite{24,46,73}. Indeed, the concurrency of contacts in the two time scale separation regimes drastically facilitates the spreading of diseases\cite{24,74}. It is important to notice how the threshold is not function of $m$ even in the case of $m > 1$. This is due to the fact that we absorbed the contact rates in the definition of $\beta$. As detailed in the methods section, this is defined as the per capita rate of infection. By adopting such definition, we are able to estimate the spreading power of a disease independently of possible differences in contacts rates. The expression could be easily changed to explicitly account this aspect obtaining $\lambda / \gamma \geq m(a) \frac{1}{m(a) (1 + \sqrt{\gamma})}$\cite{24,46,73}. In the opposite limit $\mu \to 1$ networks are extremely modular, fluctuations become important and the symmetry between SIR and SIS breaks. In order to understand this limit, let us consider first a SIR process started from a single infected node in a community of size $s$. The large majority of connections are towards a small number of vertices in the same group. As soon as the disease starts to spread within the community the number of infected and then recovered nodes grows, thus the probability of links $I - I$ and $I - R$ increases. Such connections cannot help the spreading of the disease. In fact they hamper the contagion process. In case instead of a network characterised by smaller values of modularity the growth of infected and recovered nodes has a much smaller effect. Indeed, the connectivity between communities would guarantee access to larger pool of susceptible nodes to sustain the spreading. From these simple observations we can expect that SIR processes are inhibited by highly modular connectivity patterns. Except for few exceptions in particular topologies\cite{26}, this is the case on static and annealed graphs\cite{26,46}. As we will show below, the same arguments hold also in time-varying connectivity patterns. On the other hand, in case of SIS processes, the repetition of contacts does not lead to such "pair annihilation": contacts between infected nodes do not help the spreading of the disease, but they are only temporary (eventually, all infected nodes become susceptible again). Thus, we expect that modularity plays a different role in SIS dynamics. This is what has been found also in the case of one annealed network model\cite{19}. Below we will show how this applies also in the case of time-varying networks. In order to numerically characterise SIR models, we study the epidemic size, $R_\infty$, as a function of $\beta / \gamma$. This quantity
acts as the order parameter of a second-order phase transition. For SIS processes instead, the order parameter is the final fraction of infected individuals, . The numerical estimation of this quantity is challenging, since it requires the precise determination of endemic states. For these reasons, we follow refs. , measuring the lifetime of the disease, , that acts as the susceptibility in phase transitions. This quantity is defined as the average time it takes for the disease to either die out or reach a macroscopic fraction, , of the populations. Without loss of generality, we start our simulations by setting 1% of randomly selected nodes as initial infected seed. Other parameters are set as: , , , , and (see SI for similar plots obtained fixing ).

Figure 4. Panel (A) as a function of , for selected values of and . Vertical black line represents the theoretical value of the epidemic threshold for as derived in refs. . Panel (B) , i.e. the max value of , as a function of . In red curves we set , in blue curves . In solid curves, we draw community sizes directly from the community size distribution . In dashed curves, we fix the community sizes as equal to the average value of for all communities. The 95% confidence interval is in grey. Each point is an average of independent simulations.

Figure 5. Panel (A) Lifetime of the disease as a function of , for selected values of and when . Vertical lines are the epidemic threshold. Panel (B) Ratio in correspondence of , as a function of . In red curves we set , blue curves . Each point is an average of independent simulations. Note that we avoid to simulate because the criterion we follow for the estimation of the threshold does not hold for a network with many connected components.

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Results obtained from SIR models are represented in Fig. 4A,B, whilst results from SIS models are shown in Fig. 5A,B. In Figs 4B and 5B we study different community structure, either by considering a constant community size (dashed curves) or by drawing community sizes directly from the community size distribution (solid curves). In general, red curves represents a network with bigger communities than the one represented with blue curves.
For SIR models, Fig. 4A tells us that, as expected, the higher $\beta/\gamma$ the higher the epidemic size. The figure also confirms the intuitions about the threshold. Indeed, we observe a dependence on $\mu$: the higher $\mu$ the higher the threshold. However, it is important to notice how such dependence is weak especially when compared with the SIS case (see below). Moreover, the higher the fraction of links created between pair of nodes sharing the same community (i.e. the higher $\mu_c$), the lower the epidemic size. This second observation is confirmed studying different community structures, as done in Fig. 4B, in which we plot the maximum epidemic size (corresponding to the largest value of $\beta/\mu$ in our settings), $R_{max}$, as a function of $\mu$. In the limit $\mu \rightarrow 0$, we observe that the disease impact is the same: the networks behave as if no community structure was present. Instead, when $\mu \rightarrow 1$, the modular structure influences the spread of the disease. As mentioned before, repeating contacts within communities significantly narrows the chances of having new infected individuals. Indeed, in SIR models, once a node recovers, it cannot be infected again. Repeating contacts with nodes already recovered does not favour the spread of the disease. Overall, the main observations are four. (i) Increasing the modularity reduces the epidemic size. (ii) A network with, on average, larger modules is likely to yield a higher epidemic size. (iii) The larger the modules the weaker the dependence on $\mu$ of the epidemic size. (iv) In case of small modules, the distribution of community sizes seems to influence the spreading of the disease. In particular, a network organised in small groups of constant sizes leads to smaller epidemic size respect to a network in which the average community size is the same, but individual sizes are extracted from a power-law distribution.

For SIS models, the lower $\mu$, the lower the life time $\tau$ (see Fig. 5). Inter-community links speed up the disease spreading and an endemic state, i.e. $Y = 0.5$, is reached faster. Moreover, the higher $\mu_c$, the lower the epidemic threshold. This last observation, which implies that increasing values of modularity favour the survival of the disease, is confirmed in Fig. 5B where we also test the effects of different community structures. In the limit $\mu_c \rightarrow 0$, there is no community structure and the curves converge to the same epidemic threshold. On the contrary, when $\mu_c \rightarrow 1$, the community structure becomes increasingly important and influences the spreading. Qualitatively, higher levels of modularity diminish the epidemic threshold. This is due to the repetition of the same contacts within a community which becomes increasingly more likely. Indeed, in SIS models, reinfection is allowed and nodes can become infected many times: communities act as a reservoir for the disease and favour the contagion process pushing the epidemic threshold to smaller values. Besides this last point, there are two main observations. (i) A network with larger modules is likely to have a higher epidemic threshold. (ii) In case of communities with smaller average sizes and high values of modularity, having the community size extracted from a power-law seems to slightly increase the threshold. Thus, the disease is able to spread more easily in modular networks with communities of similar or equal sizes. With the exception of one data point, we observed for $\mu > 0.5$ (see the dashed blue line in Fig. 5B).

**Real networks.** Although the modelling framework presented captures realistic activity and community size distributions of real networks, it neglects other important features such as burstiness and more complex temporal/structural correlations. It is then crucial grounding the picture emerging from synthetic models with a real world system. To this extent, we consider a temporal and modular network about scientific collaborations in the American Physical Society (APS). We study 96940 scholars connected by 692667 links (see the Supplementary Information for more details). We focus on ten years of data (January 1997–December 2006) coarse-grained at a time resolution of one month. To single out the effects introduced by communities on contagion processes, we consider also a randomised version of the dataset. Here, the interactions at each time are shuffled, destroying the community structure, but the sequence of activation times for each node and the degree distribution at each time step are preserved. In order to make sure that the randomisation process removes topological structures, we integrate the two networks over all time steps and we use OSLOM to find the communities. The modularity of the real APS network is $Q = 0.6685$, and of its randomised counterpart $Q = 0.0937$. As expected, the degree preserving randomisation reduces the modularity significantly. Using these two networks, we study the dynamical properties of SIR and SIS processes unfolding on their structure. In Fig. 6A,B we present the results. The modular properties of the real network do not influence the threshold of SIR models. Considering the weak dependence on the modularity observed in synthetic networks this results is not surprising. Even more, in this case the maximum value of modularity is defined by the data. We cannot increase it manually as done in our model. Nevertheless, the presence of communities reduces the impact of the disease, i.e. lowers the epidemic size. In the case of SIS processes instead, communities have a larger effect shifting the threshold to smaller values. These results qualitatively confirm what observed in synthetic systems.

**Discussion**

Real networks are characterised by heterogeneous statistical distributions of crucial topological features; they are organised in modules/communities; they are subject to non trivial temporal dynamics. It has long been acknowledged that such attributes have critical effects on contagion processes evolving on systems’ fabric. In particular, the heterogeneity in the connectivity patterns makes static/annealed networks extremely fragile to the spreading of infectious diseases. Moreover, the presence of communities in static/annealed graphs might either slow down or facilitate the propagation of a disease. In particular, community structures are most likely to inhibit SIR-like processes. However, in particular networks, such as Autonomous System Graphs, peculiar topological properties might have the opposite effect. The study of SIS processes in the context of modular networks has received much less attention. However, at least in the case of one artificial network model, modularity has been found to help the spreading. A note of caution is however important. Indeed, in this paper, modular networks are compared with random graphs of different degree distribution. Thus, it is hard to disentangle the real effect of modularity on SIS processes in this case. The temporal features of networks have been found to either facilitate or hamper the spreading of contagion processes. We refer the reader to ref. for a recent compendium of epidemic spreading on time-varying networks. The study of the effects of modularity on epidemic spreading...
unfolding on time-varying networks has been very limited. At the best of our knowledge only one paper so far tackled directly this issue. In this work, an activity-driven network organised in two communities of equal size has been considered. In these settings, the threshold is not function of the modularity as each community is a good representation of the full network, thus the critical behaviour is the same independently of the number of connections between the two modules. Furthermore, the authors found a weak correlation between modularity and epidemic size. Starting from all these results, here we aimed to characterise the interplay between modularity and temporal dynamics of networks considering realistic community structures. To this end, we proposed a model of temporal networks with tunable modularity and heterogeneous activity distributions. We provided an analytical description of time-aggregated properties of such networks, and studied the impact of modular and temporal features on epidemic spreading processes. In synthetic networks, we found that modularity reduces the epidemic size and threshold in SIR models, slowing down the spreading process. The effect of modularity on the threshold is weak and appreciable only for large values of it. In SIS models, modularity reduces the epidemic threshold making the system more prone to disease spreading. The dependence on modularity is in this case much stronger. The repetition of the same contacts between nodes belonging to the same community acts as a reservoir for SIS-like diseases and allows the pathogen to reach an endemic state more easily. Our work is not exempt by caveats. The most important limitation, perhaps, is represented by the adoption of Poissonian activation dynamics, which has been shown to be unrealistic in empirical temporal networks which are characterised by bursty behaviours (heterogenous inter-event time distributions). Furthermore, not only links are dynamical entities being created and terminated, but also nodes may appear and disappear during the dynamics. Our model of modular activity-driven networks does not capture all these aspects of real temporal networks. Thus, we confirmed the picture emerging in synthetic graphs by numerical simulations of SIR and SIS models on real network, characterised by both modular and temporal features. We selected a co-authorship network in which nodes describe authors and links between them capture scientific collaborations. Clearly, such network is not of direct epidemiological relevance. Unfortunately, datasets more suitable for the study of the behaviour of spreading processes on temporal networks, such as face-to-face interactions, are scarce and quite small in size. Thus, they are not the optimal choice to study how temporal and topological properties affect epidemic thresholds which are analytically defined in the limit of $N \to \infty$. However, it is important to notice that social networks, in many different contexts, are characterised by the wide range of features missing in our simple model. Thus, in the spirit of a qualitative comparison, studying the behaviour of contagion processes on co-authorship networks is a meaningful exercise. In conclusion, our findings show that on time-varying networks modularity can have opposite effects on different classes of spreading processes. Dynamical processes unfolding on real networks have been show to exhibit a rich and complex phenomenology, depending on the topological and temporal properties of the underlying substrate as well as on the characteristics of the process under investigation. Such complexity may be addressed by both the definition of proper generative models, that allows to control the desired features, and the study of the processes behaviour in real-world network. Our work is set within this framework, and it contributes to shed light on the impact of modular and temporal properties of real networks on epidemic spreading dynamics.

**Methods**

**SIS and SIR models.** In both processes nodes are divided in different classes according to their disease status. In SIR models nodes are either Susceptible (S), Infected (I) or Recovered (R). Susceptible nodes describe healthy individuals. Infected nodes contract the disease and are infectious. Recovered nodes are no longer

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**Figure 6.** Panel (A) $R_\infty$ as a function of $\beta/\gamma$ for SIR processes diffusing on APS (cyan circles) and on the randomized APS dataset (green circles). Panel (B) $L$ as a function of $\beta/\gamma$ for a SIS models evolving on the same two networks. Each point is the average of $10^5$ independent simulations started from 1% of random seeds. We fix $\gamma = 0.05$. 

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infected and acquire complete immunity to the illness. The model is fully characterized by two transitions: \( S + I \xrightarrow{\beta} 2I \) and \( I \xrightarrow{\gamma} R \). The first describes the infection propagation and \( \beta \) is the capita infection rate. This quantity is defined by the average contacts per node \( \langle k \rangle \) and by the per contact probability of transmission \( \lambda \), i.e. \( \beta = \lambda \langle k \rangle \). The second transition describes the recovery process. Infected individuals recover spontaneously and permanently with rate \( \gamma \). In SIS models instead we have just Susceptible and Infected nodes. While the contagion process is equivalent to the SIR case, the recovery is different and described by the following transition: \( I \xrightarrow{\gamma} S \). Infected nodes spontaneously return in the susceptible compartment with rate \( \gamma \).

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Author Contributions
N.P. conceived the research, M.N. and K.S. conducted the numerical simulations, E.U. developed the analytical calculations, all authors analysed the results, wrote and reviewed the manuscript.
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Supplementary Information: epidemic spreading in modular time-varying networks

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ABSTRACT

In this Supplementary Information, we present in a more detailed and comprehensive way the definition of the model, its key properties and the results (both analytical and numerical) found. Furthermore, we show additional simulations studied in synthetic networks for SIR and SIS models. Finally, an explanation of the APS dataset is given.

1 The Model

The network is defined by means of the following parameters:

- the total number $N$ of nodes in the network;
- the activity distribution parameters, i.e. the lower cut-off $\epsilon$ and the leading exponent $\nu$, so that $F(a) \propto a^{-\nu}$ for $a \in [\epsilon, 1]$;
- the lower cut-off $s_{\text{min}}$, the upper limit $s_{\text{max}}$ and the exponent $\omega$ describing the community size distribution, i.e. $P(s) \propto s^{-\omega}$ for $s \in [s_{\text{min}}, s_{\text{max}}]$;
- $\mu$ is the probability that, once active, a node will connect to a node inside the community, so that $\mu' = 1 - \mu$ is the probability to fire outside the community;

We initialize the network extracting $N$ activity values from the activity distribution $F(a)$ and we then group the nodes in communities of size $s$ drawn from a size-distribution $P(s)$. Once we initialized the network we let it evolve following the time-varying activity driven framework. At each time step we start with $N$ disconnected nodes. Each node gets active with probability $a_i dt$ at each time step $dt$ and fire to a randomly chosen node inside (outside) its own community with probability $\mu$ ($\mu'$). At time $t + dt$ we delete all the edges and repeat the above procedure. Each node $i$ will then have a set of neighbors that have been contacted or have contacted the node during the network growth. The size of such a set is the integrated degree $k$ of the node $i$. Of these $k$ neighbors, $k_c$ will be inside the $i$’s community (in-community degree) and $k_o = k - k_c$ will be external to the community (out-community degree).

2 The Network growth

In the integrated network, each node $i$ has a set of neighbors that have been contacted or have contacted the node during the network growth. The size of such a set is the integrated degree $k$. Of these $k$ neighbors, $k_c$ are inside the $i$’s community (i.e. the in-community degree) and $k_o = k - k_c$ are external to the community (i.e. the out-community degree). Since the model is memoryless, the in-community degree $k_c$ and the out-community degree $k_o$ are decoupled and can, in fact, be treated separately. Even the activity potential $a_i$ of each node can be “split” in two components: the in-community activity $\mu a_i = (1 - \mu') a_i$...
and the complementary out-community $\mu a_i$. Indeed, each node points, on average, a fraction $\mu$ of its own events toward the community, while the remaining $\mu'$ are directed outside the community itself. Each node experiences a mean field of activity, $\mu(a)$, coming from the community (provided that the community is large enough) and a supplementary external field $\mu'(a)$ coming from the rest of the network.

2.1 The network time scales

As a first insight, let us note that the in-degree time dependence can be easily approximated with a probabilistic consideration. Each node $i$ of activity $a_i$, within a community of size $s$, has $s - 1$ available edges. Now, for each time step of the dynamics, the edge $e_{ij}$ is created with probability $\mu(a_i + a_j)/(s - 1)$. Then, on average, each edge emanating from $i$ is activated with probability $c(a_i, \mu)/(s - 1) = \mu(a_i + (a' ))/(s - 1)$, where $c(a_i, \mu)$ is the number of edges intra-community. The probability $P(a_i, \mu, s, t)$ for an edge pointing to $i$ not to be activated after $t$ time steps then reads:

$$P(a_i, \mu, s, t) = \left(1 - \frac{c(a_i, \mu)}{s - 1}\right)^t.$$  

(1)

Since the in-degree $k_c \in [0, s - 1]$, we can write:

$$k_c(a_i, \mu, s, t) = (s - 1)\left(1 - P(a_i, \mu, s, t)\right).$$  

(2)

In the following, we always have the dependency on $a_i, \mu$ and $t$, so to simplify the notation we drop most of those parameters: $P(a_i, \mu, s, t) = P(s)$, to avoid confusions with the community size distribution $P(s)$. Also, $k_c(a_i, \mu, s) = k_c(s)$ and $c(a_i, \mu) = c$.

We note that Eq. 1 gives us an estimation of the characteristic time $\tau(s)$ that takes for a node of activity $a_i$ to saturate the in-degree $k_c \rightarrow (s - 1)$. Indeed, we can rewrite Eq. 1 as:

$$P'(s) = \exp\left[t \ln \left(1 - \frac{c}{s - 1}\right)\right] \Rightarrow \tau(s) = -\ln \left(1 - \frac{c}{s - 1}\right) \Rightarrow P_c(s) = \frac{2 s \mu \langle a \rangle}{s (s - 1)}.$$  

(3)

The probability for one edge not to be created after $t$ time steps is then:

$$\hat{P}_c(s) = \left(1 - \frac{2 \mu \langle a \rangle}{s - 1}\right)^t \Rightarrow \hat{P}_c(s) = \exp\left[\frac{t}{\tau_c(s)}\right] \Rightarrow \tau_c(s) = -\ln \left(1 - \frac{2 \mu \langle a \rangle}{s - 1}\right) \Rightarrow \tau_c(s) = -\ln \left(1 - \frac{2 \mu \langle a \rangle}{s - 1}\right).$$  

(5)

where $\tau_c(s)$ represents the typical time by which the majority of the nodes of a community has a degree $k_c \approx s$.

Note that, in the evaluation of both $\tau(s)$ and $\tau_c(s)$, we did not take into account the difference between edges pointing to a more active node and the ones pointing to a less active one. Nevertheless, this is a simple estimation that, as we will show later, correctly catches the general behaviour of the in-degree $k_c$ for any value of $\mu, a_i$ and $s$. Besides, when computing the key features of the evolving network, we are now able to distinguish the short time range $t \ll \tau_c(s)$ (in which $k_c \ll s$ for any activity value $a_i$) and the long time limit $t \gg \tau_c(s)$ (in which $k_c \sim s$ for any activity value $a_i$).

2.2 The Master Equation and the $P(a, k, t)$

We can now write down the Master Equation (ME) for the quantities $P_c(s, k_c)$ and $P_c(s, k_c)$, that is, the probability for a node of activity $a_i$ belonging to a community of size $s$ to have degree in (out) degree $k_c \sim k_o$ at time $t$. In general, $a_i\Delta t$ represents the probability the node $i$ is active, where $a_i$ is the activity rate of node $i$. Without loss of generality we will assume $\Delta t = 1$. To get the ME for the in-degree $k_c$ distribution, we exploit also the time-dependence for a couple of passages:

$$P_c(s, k_c, t + 1) = P_c(s, k_c, t) \left[1 - \mu \sum_j a_j\right] + P_c(k_c, t) \left[\frac{k_c}{s} \mu a_i + \frac{s - 1}{s} \mu \sum_j a_j + \mu \sum_j a_j\right] +$$

$$+ P_c(s, k_c - 1, t) \left[\frac{s - k_c}{s} \mu a_i + \frac{s - 1}{s} \mu \sum_j a_j\right].$$  

(6)
where $\sum_{j \neq i}$ and $\sum_{j=1}$ are respectively two contracted notations for the sum of all the first neighbors of node $i$ and the sum of all nodes but the neighbors of node $i$. The first parenthesis indicates the probability that none of the nodes in the network fire. Third (sixth) term is the probability a node $i$, in the instantaneous network, is active and fires to a node where, in the integrated counterpart, there is already (isn’t) a link. Four (seventh) term is the probability a node $j$ not linked to $i$ fires to any of the nodes but $i$ (fires to $i$). Fifth factor is the probability a node $j$ already linked to $i$ fires.

After some algebra, ME can be written as:

$$P_c(s, k_c, t + 1) - P_c(s, k_c, t) = -[P_c(s, k_c, t) - P_c(s, k_c - 1, t)] \left( \frac{s - k_c}{s} \mu a_i + \frac{\mu}{s} \sum_{j \neq i} a_j \right).$$

(7)

Now we pass to the continuum limit by considering $t \gg 1$ and $k \gg 1$. So the l.h.s becomes simply the time derivative with respect to $P_c(s, k_c)$ and, to obtain a proper convergence of the results, we can expand the probability with respect to the in-community degree up to second order.

In the regime $t \ll \tau(s)$, we can neglect $k_c \ll s$ and $1/s \sum_{j \neq i} a_j \approx \langle a \rangle$.

$$\frac{\partial P_c(s, k_c)}{\partial t} = (\mu + \mu \langle a \rangle) \left[ \frac{\partial P_c(s, k_c)}{\partial k_c} - \frac{1}{2} \frac{\partial^2 P_c(s, k_c)}{\partial k_c^2} \right],$$

(8)

where we dropped the $a_i$ index since we expect all the nodes of a given activity to behave in the same way. Now $a$ is an activation rate and in the treatment we assume it takes small values to avoid that two nodes become active together.

The solution of Eq. 8 reads:

$$P_c(s, k_c) = C \exp \left[ -\frac{(k_c - \mu \langle a \rangle \tau)^2}{2 \mu \langle a \rangle \tau} \right],$$

(9)

where $C$ is a normalization constant.

By following the same procedure, we recover the same results of Eq. 9 for the out-community degree $k_o(a, t)$, by substituting $\mu \to \mu'$ and $k_c \to k_o$:

$$P_o(s, k_o) = C \exp \left[ -\frac{(k_o - \mu' \langle a \rangle \tau)^2}{2 \mu' \langle a \rangle \tau} \right]$$

(10)

Since $N \gg s_{max}$, the out-degree $k_o \ll N$ for any time $t$ of the process, thus we assume that Eq. 10 is valid for all the time scales analyzed. Also note that, as expected, the net effect of the mixing parameter $\mu'$ is just a time rescaling of the out-community and in-community activity, respectively.

Then, in the $t \sim \tau(s)$ time range is not possible to find an analytic formula, however simulations will be run to provide, at least, a qualitative behavior. In the $t \gg \tau(s)$ time limit the $P_c(s, k_c)$ converges to the $\delta(k_c - (s - 1))$ distribution. In fact, all the nodes will have all their edges activated and the $P_c(s, k_c)$ time derivative goes to zero.

Let us now resume the results found in this section:

$$P_c(s, k_c) \propto \begin{cases} \exp \left[ -\frac{(k_c - \mu \langle a \rangle \tau)^2}{2 \mu \langle a \rangle \tau} \right] & \text{for } t \ll \tau(s) \\ \delta(k_c - (s - 1)) & \text{for } t \gg \tau(s) \end{cases}$$

(11a)

$$P_o(s, k_o) \propto \exp \left[ -\frac{(k_o - \mu' \langle a \rangle \tau)^2}{2 \mu' \langle a \rangle \tau} \right] \text{ for } \forall t$$

(11b)

where we now distinguish between the in-community degree distribution $P_c(s, k_c)$ and the out-community degree distribution $P_o(s, k_o)$. The latter however, is independent on the community size and we can then define $P_o(s, k_o) = P_o(k_o)$.

So far we treated the two probability functions separately when, in fact, $k_c$ and $k_o$ are bound by the relation $k = k_c + k_o$. The total degree distribution $P(s, k)$ will then be determined by the convolution of both the $P_c(s, k_c)$ and $P_o(s, k - k_c)$:

$$P(s, k) = \int_0^k dk_c P_c(s, k_c) P_o(k - k_c)$$

(13)
where we integrate over all the possible arrangements of the \( k_c \) edges. 

In the \( t \ll \tau(s) \) limit, by substituting Eq. 11a and 12 in Eq. 13, we sum the two exponents getting:

\[
P(s,k) = C \int_0^k dk_c \exp \left[ -\frac{(k_c - \mu(a + \langle a \rangle)t)^2}{2\mu(a + \langle a \rangle)t} - \frac{(k - k_c - \mu'(a + \langle a \rangle)t)^2}{2\mu'(a + \langle a \rangle)t} \right] \tag{14}
\]

where \( C \) is, again, a normalization constant. 

By combining the two terms and after some algebra we get:

\[
P(s,k) = C \int_0^k dk_c \exp \left[ -\frac{(k_c - \mu k)^2}{2\mu \mu'(a + \langle a \rangle)t} - \frac{k^2 - 2k(a + \langle a \rangle)t - (a + \langle a \rangle)^2 t^2}{2(a + \langle a \rangle)t} \right]. \tag{15}
\]

The integration over \( k_c \) gives:

\[
P(s,k) = C \left[ \text{Erf} \left( \frac{\mu' k}{\sqrt{2}\mu \mu'(a + \langle a \rangle)t} \right) - \text{Erf} \left( \frac{\mu k}{\sqrt{2}\mu \mu'(a + \langle a \rangle)t} \right) \right] \exp \left[ -\frac{k^2 - 2k(a + \langle a \rangle)t - (a + \langle a \rangle)^2 t^2}{2(a + \langle a \rangle)t} \right] \tag{16}
\]

where \( \text{Erf}(x) \) is the error function evaluated at \( x \).

In the small time limit, if we want to evaluate the \( P(k) = \int_{s \min}^{s \max} ds P(s) P(s,k) \), we have to consider \( t \ll \min_i (\tau_i(s)) \). In this way, for each community we can use Eq. 16 as the true value of the \( P(s,k) \). The integration over the different community size \( s \) is then straightforward since the terms are independent on it, giving \( P(k) = P(s,k) \). Note that this result holds for any value of \( \mu \), \( N \) and \( a \).

The computation of \( P(k) \) in the large time limit (i.e. \( t \gg \langle \tau_i(s) \rangle \)) is more complicated and we have to assume that \( k_c = s - 1 \) for each node in a community of size \( s \), otherwise Eq. 11b put everything equal to zero. The \( P_c(s,k_c) \) will be approximated by Eq. 12. The integral now reads:

\[
P(k) = \int_{s \min}^{s \max} ds P(s) P(s,k) = C \int_{s \min}^{s \max} ds P(s) \exp \left[ -\frac{(k - (s - 1) - \mu'(a + \langle a \rangle)t)^2}{2\mu'(a + \langle a \rangle)t} \right]. \tag{17}
\]

The exponential can be written as:

\[
\exp \left[ -\frac{(k - \mu'(a + \langle a \rangle)t)^2 + (s - 1)\langle (s - 1) - 2(k - \mu'(a + \langle a \rangle)t) \rangle}{2\mu'(a + \langle a \rangle)t} \right], \tag{18}
\]

where the rise of new terms proportional to \( s^2 \) and \( sk \) makes it difficult to perform the integral.

We can, however, give a solution for the simple case \( P(s) = \delta(s - \bar{s}) \), when all the communities have equal size. First of all, we have, for large times, \( P_c(s,k_c) = \delta(k_c - (s - 1)) \). Then:

\[
P(s,k) = \begin{cases} 
P_c(k - (s - 1)) & \text{for } k \geq s - 1 \\ 0 & \text{for } k < s - 1. \end{cases} \tag{19a}
\]

When \( k < s - 1 \), for sure \( k_c < s - 1 \) and the delta put everything equal to zero. In the other case \( k_c = s - 1 \) with a certain probability \( P_c(k - (s - 1)) \). Finally, equation 17 can be written as:

\[
P(k) = \begin{cases} 
P_c(k - (\bar{s} - 1)) & \text{for } k \geq \bar{s} - 1 \\ 0 & \text{for } k < \bar{s} - 1. \end{cases} \tag{20a}
\]

\[ \text{2.3 The average degree } \langle k(a,s,t) \rangle \]

We can provide a simple expression for the nodes average degree belonging in different classes. As we already showed in Eq. 2, \( \langle k_c(a,s,t) \rangle \) grows as:

\[
\langle k_c(a,s,t) \rangle = (s - 1) \left( 1 - \exp \left( -\frac{t}{\tau(s)} \right) \right) = (s - 1) \left( 1 - P(s) \right) = (s - 1) \left( 1 - \left( 1 - \frac{c}{s - 1} \right) \right). \tag{21}
\]
and the $\langle k_o \rangle$ grows as the mean value of the distribution given in equation 12, and turns out to be independent on $s$.

$$\langle k_o(a,t) \rangle = \mu'(a + \langle a \rangle)t \tag{22}$$

The average total degree $k(a,s,t)$ for nodes of activity $a$ belonging to communities of size $s$ depends on the time scale we analyse the problem. For small times (i.e. $t \ll \langle \tau(s) \rangle_s$):

$$\langle k(a,s,t) \rangle = \langle k_c(a,s,t) \rangle + \langle k_o(a,t) \rangle \approx (s-1) \left( 1 - \frac{\mu(a + \langle a \rangle)t}{s-1} \right) + (1 - \mu)(a + \langle a \rangle)t = (a + \langle a \rangle)t \tag{23}$$

As time grows toward the regime of times comparable to the average (i.e. $t \sim \langle \tau(a,s,t) \rangle$), we cannot approximate the in-community degree anymore but we use directly equation 2:

$$\langle k(a,s,t) \rangle = (s-1)(1 - P'(s)) + \langle k_o(a,t) \rangle. \tag{24}$$

Then, the regime of large times (i.e. $t \gg \langle \tau(s) \rangle$) is:

$$\langle k(a,s,t) \rangle \approx s - 1 + \mu'(a + \langle a \rangle)t \approx \mu'(a + \langle a \rangle)t \tag{25}$$

The above equations then predict that the average degree has a linear growth proportional to $a + \langle a \rangle$ for short time limit (equation 23), then a transition for $t \sim \langle \tau(s) \rangle_s$ is followed by a second linear growth, valid for large times, proportional to $\mu'(a + \langle a \rangle)$. These regimes correspond to the initial growth, in which both the in-community and the out-community degrees are growing linearly in time, followed by the slowing down of the in-community degree which is saturating to $s - 1$. Finally, the third regime is again linear and it is driven by the $\mu'(a + \langle a \rangle)$ coefficient: it means meaning that only the out-community degree is growing.

### 2.4 The degree distribution $\rho(k_{th})$

Now that we have the expression of the average degree, it is straightforward to write the degree distribution. At all the time scales we found $\langle k_{th} \rangle \propto Ct$, where $C$ is a time-independent coefficient. Then, $k_{th} \propto at$, and it results in an equal increment in the activity and in the degree values ($da = dk_{th}$). If we use the change of variable rule, we obtain:

$$F(a)da = \rho(k_{th})dk_{th} \Rightarrow a^{-\nu}da = k_{th}^{-2}dk_{th} \Rightarrow x = \nu, \tag{26}$$

i.e., the degree distribution has the same exponent $\nu$ as the activity distribution function.

## 3 Comparison with numerical simulations

To check the analytical predictions of Section 2 we performed numerical simulation. In particular we realized 100 representations of a network featuring:

- $N = 10^5$ nodes with modularity $\mu = 0.9$ evolving for $10^5$ evolution steps;
- activity potential distributed following the $F(a) \propto a^{-\nu}$ with $\nu = 2.1$ and $a \in [10^{-3}, 1]$ interval;
- power-law distributed community sizes $P(s) \propto s^{-\omega}$ with $\omega = 2.1$ and $s \in [10, \sqrt{N}]$.

In order to analyze the collective behavior of the nodes we group them by their activity and community size, thus defining $b$ classes of nodes. We average over the representations of the network and for each class of nodes $b$ we evaluate:

- $P_i(s,k), P_o(s,k), P(s,k)$ for $t \ll \tau_i(s)$ and $t \gg \tau_o(s)$;
- the average degree $\langle k_c(s) \rangle, \langle k_o \rangle$ and $\langle k(s) \rangle$;
- the degree distribution $\rho(k_{th})$.

In the main discussion, we already showed that some of the above measures have a great agreement with analytical predictions. To complete the discussion, we add below Fig. 1 which proves that also the in-community probability and the in-community degree perfectly matches our expectation. In panel B) we display $P_i(s,k)$ for $t \sim \tau$, even if it was impossible to obtain an exact result, the comparison demonstrates that the in-community probability starts to deviate from a Gaussian distribution.
Figure 1. Panels A-C) rescaled \( P_c(k_c) \) probability distribution as found for a selected node class at different times (legends). Functions in panels A-B) are rescaled accordingly to the theoretical distribution given in Eq. 11a, i.e., by sending \( k_c \rightarrow \tilde{k}_c = (k_c - \langle k_c \rangle) / \langle k_c \rangle^{1/2} \) and plotting \( P_c(\tilde{k}_c \langle k_c \rangle^{1/2}) \). Panel C) is rescaled accordingly to the theoretical distribution given in Eq. 11b, i.e., by sending \( k_c \rightarrow k'_c = k_c / \langle k_c \rangle \) and plotting \( P_c(k'_c \langle k_c \rangle) \). In panel D) we plot \( \langle k_c \rangle \) for nodes featuring different activity potential and belonging to communities of different size. The data are rescaled sending the time \( t \rightarrow t/\tau \) and then plotting \( \langle k_c(t/\tau) \rangle / s \). The analytical prediction of Eq. 2 computed for a node of activity \( a = \langle a \rangle \) and belonging to a community of size \( s = \langle s \rangle \). Each point is an average over \( 10^2 \) independent simulations, parameters used are: \( N = 10^5 \), \( \omega = 2.1 \), \( \nu = 2.1 \), \( m = 1 \), \( s_{\text{min}} = 10 \) and \( \mu = 0.9 \).
4 SIR and SIS processes on modular activity driven networks

We present together all the results about SIR and SIS models obtained in synthetic networks. We start our simulations by setting 1% of randomly selected nodes as initial infected seeds, the other parameters are fixed as $\gamma = 0.01$, $m = 3$, $\nu = 2.1$, $N = 10^5$ and $Y = 0.5$. Panels A) and C) set the exponent of the distribution of community sizes $\omega = 1.5$, while panels B) and D) have $\omega = 2.1$. The qualitative picture is unchanged due to selecting a different value of $\omega$. The modular structure becomes irrelevant for $\mu \to 0$, whilst it significantly modifies the spread of the disease when $\mu \gg 0$. Modularity slows down contagion processes in SIR models, it favors the disease outbreak in SIS models. Moreover, quantitatively, in SIR models the presence of larger communities lead to higher epidemic size, in SIS models it lower the epidemic threshold. Moreover, in SIS models and when $\mu \to 1$, the epidemic threshold is likely to increase due to major limitations in reaching an endemic size. This phenomenon is particularly visible in Panel C), solid blue curve.

Figure 2. Panels A-B) take $R_{\text{max}}$ of each $R_\infty$ curve and plot it as a function of $\mu$. Panels C-D) $\xi_{\text{SIS}}$, that is $\beta/\gamma$ in correspondence of $L_{\text{max}}$, and plot it as a function of the modularity. In red curves we set $s_{\text{min}} = 100$, in blue curves $s_{\text{min}} = 10$. Solid curves are obtained by drawing community sizes from a power law distribution, 95% confidence interval is in gray. Dashed curves have a constant community size equal to the average value of the power law. Each point is an average of $10^2$ independent simulations.
5 Real network: APS dataset

In the data each author of an article is described as a node. An undirected link between two different authors is drawn if they collaborated in the same article. We used the dataset from Ref\(^1\) which spans a period between 1893 and 2006. To have the average degree in each instantaneous network as comparable as possible (see Fig. 3), we select a period of ten years, from January 1997 to December 2006. In this time window we register 96940 scholars who create 692667 connections. When we simulate SIR and SIS models on top of APS temporal network, we use periodic boundary conditions to let the disease dynamics evolve without late-time constrains.

![Figure 3](image)

**Figure 3.** Average degree for each month in the selected subset of APS dataset (red circles), compared with the global average of all the ten years considered (solid black line). We label each month with increasing integer numbers from 1 to 120, where 1 represents the beginning of our sample, January 1997, and 120 the end, December 2006. We observe an increasing number of collaborations through years.

Using OSLOM, in Fig.4A we show that the integrated APS network is modular. Then, we apply the following degree-preserving randomization technique to destroy the network’s community structure. We choose randomly a source node \(S_1\) and, among its neighbors, we select randomly a target node \(T_1\). We do the same for other two nodes \(S_2\) and \(T_2\). If the two pairs are equivalent or if a multi-edge will be created, we start back by selecting \(S_1\) again and repeating the instructions. Otherwise, we swap \(T_1\) and \(T_2\) to have the new undirected links: \(S_1 - T_2\) and \(S_2 - T_1\). The edges are chosen within the same temporal network and the number of swaps is equal to the number of edges in that instantaneous network. So, the above procedure is applied for each instantaneous network. At the end, we integrate the randomized temporal network and use OSLOM to detect the community structure. In Fig.4B, we qualitatively prove that our degree preserving randomization destroys the network’s community structure.

The options used in OSLOM are: -uw (to study undirected networks); -cp0.99 (to have communities as large as possible); -hr (to avoid to consider hierarchies); -r100 (to repeat 100 times the community detection). Since OSLOM finds communities in a non-deterministic way, last option is useful to get rid of stochastic fluctuations and have a more reliable community structure. Finally, we evaluate quantitatively the modularity of the two networks. For the original APS network, \(Q = 0.6685\), and for its randomized counterpart \(Q = 0.0937\).
Figure 4. Panel A) Community sizes probability density function in the original network. Number of communities found is 10,825, minimum community size is 1, maximum 91. Panel B) Community sizes probability density function after having applied the degree-preserving randomization. Number of communities found is 1,489, minimum community size is 1, maximum 222. Note also that the shape of the distribution is completely different from panel A).

References
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