

A comparison of node vaccination strategies to halt SIR epidemic spreading in real-world complex networks

Original

A comparison of node vaccination strategies to halt SIR epidemic spreading in real-world complex networks / Sartori, F., Turchetto, M., Bellingeri, M., Scotognella, F., Alfieri, R., Nguyen, N., Le, T., Nguyen, Q., Cassi, D.. - In: SCIENTIFIC REPORTS. - ISSN 2045-2322. - ELETTRONICO. - 12:(2022), pp. 1-13. [10.1038/s41598-022-24652-1]

Availability:

This version is available at: 11583/2981218 since: 2023-08-24T06:13:06Z

Publisher:

Springer Nature

Published

DOI:10.1038/s41598-022-24652-1

Terms of use:

This article is made available under terms and conditions as specified in the corresponding bibliographic description in the repository

Publisher copyright

(Article begins on next page)



OPEN

A comparison of node vaccination strategies to halt SIR epidemic spreading in real-world complex networks

F. Sartori¹✉, M. Turchetto^{1,4}, M. Bellingeri^{1,2,4}, F. Scotognella^{2,3}, R. Alfieri^{1,4}, N.-K.-K. Nguyen⁵, T.-T. Le⁶, Q. Nguyen^{7,8} & D. Cassi^{1,4}

We compared seven node vaccination strategies in twelve real-world complex networks. The node vaccination strategies are modeled as node removal on networks. We performed node vaccination strategies both removing nodes according to the initial network structure, i.e., non-adaptive approach, and performing partial node rank recalculation after node removal, i.e., semi-adaptive approach. To quantify the efficacy of each vaccination strategy, we used three epidemic spread indicators: the size of the largest connected component, the total number of infected at the end of the epidemic, and the maximum number of simultaneously infected individuals. We show that the best vaccination strategies in the non-adaptive and semi-adaptive approaches are different and that the best strategy also depends on the number of available vaccines. Furthermore, a partial recalculation of the node centrality increases the efficacy of the vaccination strategies by up to 80%.

Immunization through vaccination is an essential issue of research with obvious implications for public health^{1–3}. Network science plays a fundamental role in epidemiology, and many recent pieces of research modeled disease spreading using network models^{4–10}. A spreading disease can be described as a network where nodes represent the individuals and links (edges) represent the social contacts between them^{9,11–13}. The two classes of interventions that can be implemented to reduce the size of an outbreak can be divided into pharmacological interventions, PI, such as vaccinations, and non-pharmacological interventions, NPI, such as social distancing, washing hands, and lockdowns. NPIs are a set of changes in social behavior that aim to reduce the social interactions (links or connections) among individuals, like during lockdowns^{14,15}, or by reducing the risk that the disease is transmitted for each social interaction, like by wearing masks¹⁶. One of the fundamental problems in network epidemiology is finding the best vaccination strategy to halt epidemics^{6,7,13,17}. Node removal (attack) on complex social networks is an optimal framework to model vaccination strategies over a population where diseases can spread^{13,17–22}. As a first approach, the problem can be mapped to the classic percolation problem in graph theory, where nodes are immunized (removed) from the network to induce the fastest network dismantling^{6,7,17}.

In this context, it has been shown that removing highly connected nodes is an efficient strategy for determining a fast network dismantling^{6,7,17}. Nonetheless, focusing on how node removal affects the network's connectivity is a simple and static representation of vaccination effects on the disease spread. Fine-grained models should encompass further aspects of the epidemics, such as the temporal dynamics of the disease spreading. This can be done by solving the classical susceptible-infected-recovered (SIR) model²³ over a network^{10,12,24}. Coupling SIR epidemiological models and network structure provides further information about the disease spread, such as evaluating the effect of node vaccination/removal strategies (NVS) to reduce the pace of the epidemics, the peak of infected individuals, or the total number of infected at the end of the epidemic¹². For these reasons, understanding which node removal strategy is the most efficient for curbing SIR epidemics spreading

¹Dipartimento di Scienze Matematiche, Fisiche e Informatiche, Università di Parma, Via G.P. Usberti, 7/a, 43124 Parma, Italy. ²Dipartimento di Fisica, Politecnico di Milano, Piazza Leonardo da Vinci 32, 20133 Milano, Italy. ³Center for Nano Science and Technology@PoliMi, Istituto Italiano di Tecnologia, Via Giovanni Pascoli, 70/3, 20133 Milan, Italy. ⁴INFN, Gruppo Collegato di Parma, 43124 Parma, Italy. ⁵Faculty of Basic Science, Van Lang University, Ho Chi Minh City, Vietnam. ⁶John Von Neumann Institute, Vietnam National University Ho Chi Minh City, Ho Chi Minh City, Vietnam. ⁷Institute of Fundamental and Applied Sciences, Duy Tan University, Ho Chi Minh City 700000, Vietnam. ⁸Faculty of Natural Sciences, Duy Tan University, Da Nang City 550000, Vietnam. ✉email: fabio.sartori@unipr.it

in networks is a promising tool for developing effective vaccination strategies. In this paper, we implement seven NVSs over a set of twelve real-world networks from different fields of science. The NVSs adopted here are based on notions of node centrality from network science and graph theory. We studied two classes of NVSs, the one where the centrality of each node is calculated at the beginning of the simulations, and we refer to this class as non-adaptive, and the one where the node centrality is recalculated after a fraction r of nodes have been removed, as semi-adaptive.

To test the NVS efficacy, first, we analyze their ability to dismantle the network connectivity, i.e., to decrease the largest connected component (LCC)^{25,26}. Then, we study the NVS efficacy to curb two indicators of a SIR epidemic outbreak: the total number of infected individuals at the end of the outbreak (TI) and the maximum number of simultaneously infected individuals, or infection peak (IP). A non-stochastic approach that also includes the dynamical parameters of the SIR that relies on the Dynamic Message-Passing equations (DMP) allows calculating the set of individuals that minimize the TI. We find that in case of limited vaccine availability, the best strategy depends on the topology of the network where the outbreak is evolving and on the fraction of individuals that can be vaccinated. Furthermore, we showed that using a semi-adaptive algorithm reduces the three SI up to 75% for the same vaccination target. Finally, we showed that the vaccine requirement needed for the 10% threshold of acceptable TI and IP is lowered from 50 to 60% if a non-random NVS is implemented.

Results

SIR and vaccination strategies. To understand how node vaccination strategies (NVS) in case of limited vaccine doses affect the spread of an epidemic in real-world complex networks, we considered the SIR dynamic epidemic model. SIR is a compartmental model, where every individual can be in one of three states/compartments: Susceptible to be infected **S**, Infected individual **I**, and Removed (or Recovered) from the infection dynamic **R** individuals²⁷. We used the implementation of the SIR model on a network as in⁹. In this frame, each network node represents an individual, and the adjacency matrix **A** dictates which nodes are connected to whom and, therefore, how the outbreak will evolve in the network^{9,12,28,29}. Two parameters regulate the dynamic of the epidemic: β represents the probability that a Susceptible node connected to an Infected one will become Infected, and γ is the probability that an Infected node will overcome the infection becoming Recovered. Furthermore, we assume that a vaccinated individual falls in the same class as a Recovered one and cannot infect any of its neighbors.

The optimal vaccination strategy would be to vaccinate all the individuals in the network, but that is often not possible, both because of limited availability of vaccines, money, doctors, and time or due to fragile subjects that cannot be vaccinated. In this work, we aim to answer the question: what is the best vaccination strategy if only a fraction q of the population can be vaccinated?

To quantify the NVS efficacy on real-world complex networks, we analyzed three spread indicators (SIs) for 20 vaccination targets q . The first is the size of the network's largest connected component (LCC). The LCC accounts for the maximum number of connected nodes in the network, and it is a commonly used quantity to evaluate the efficacy of node vaccination (removal) strategies^{7,12,20}. The LCC is a purely topological indicator of the epidemic spreading, accounting solely for the maximum number of nodes that can be infected. The second metric we used is the total infected (TI) individuals, i.e., the number of infected individuals at the end of the outbreak event. If all the individuals in the network share the same risk factor, the TI is directly proportional to the number of hospitalizations and the total number of casualties at the end of the outbreak event. The third metric is the infected peak (IP), i.e., the maximum number of simultaneously infected individuals during the outbreak; IP is an important indicator to evaluate the possible overload of the health system, which would increase the mortality rate of both the epidemic itself and other diseases.

Finding the optimal set of nodes to vaccinate to reduce the size of the LCC below a threshold C is an NP-complete problem³⁰. Therefore, we are following a heuristic approach by assuming that the most central node is the one whose removal minimizes the three SI most. The NVSs we are comparing in this work are seven ways to define the centrality of each node, which we listed in Table 3 in the Methods section.

In Fig. 1, we give a graphical representation of the outcomes of three of the NVS to the BE3 network³¹, a road network in Beijing city. In the dashed boxes, we show the reduced BE3 network after removing the fraction q of nodes with the highest centrality, with $q = 0, 0.08, 0.16, \text{ and } 0.24$. In Fig. 1, we see that the more nodes we remove (vaccinate), the fewer links (representing the infective social interactions) remain in the network, and, therefore, the smaller the capability of an outbreak to spread within the network. This is clearer for the semi-adaptive, $r = 0.005$, that triggers a visible fragmentation of the LCC.

Non-adaptive strategies. The first two strategies we implemented are the *Random* strategy (RAN), where the centrality of each node is a random value between 0 and 1, and the *Degree* strategy (DEG), where the centrality of each node is the number of first neighbors that such node has. In Fig. 2A and B, we show how the three SIs decrease when the network is subjected to RAN and DEG, respectively. However, unsurprisingly DEG, which contains information about the network topology, is more effective at reducing all three SI (see Fig. 2C for a direct comparison).

The RAN strategy is not always worse than the DEG at minimizing the three SIs, as shown in Fig. 2D and E. In the BE3 network, for instance, the DEG strategy is more effective at minimizing the TI and the IP but, surprisingly, not at minimizing the LCC. In Fig. 2F, bottom line, we show that in the 12 analyzed real-world networks, only BE3 has this unintuitive behavior; in the same figure, we also show how often the other five NVS are more effective than DEG at minimizing the LCC, while Betweenness (BET) is the one that on a specific network performs better, it shows high variability. *Pagerank* (PAG), instead, performs consistently better than DEG.

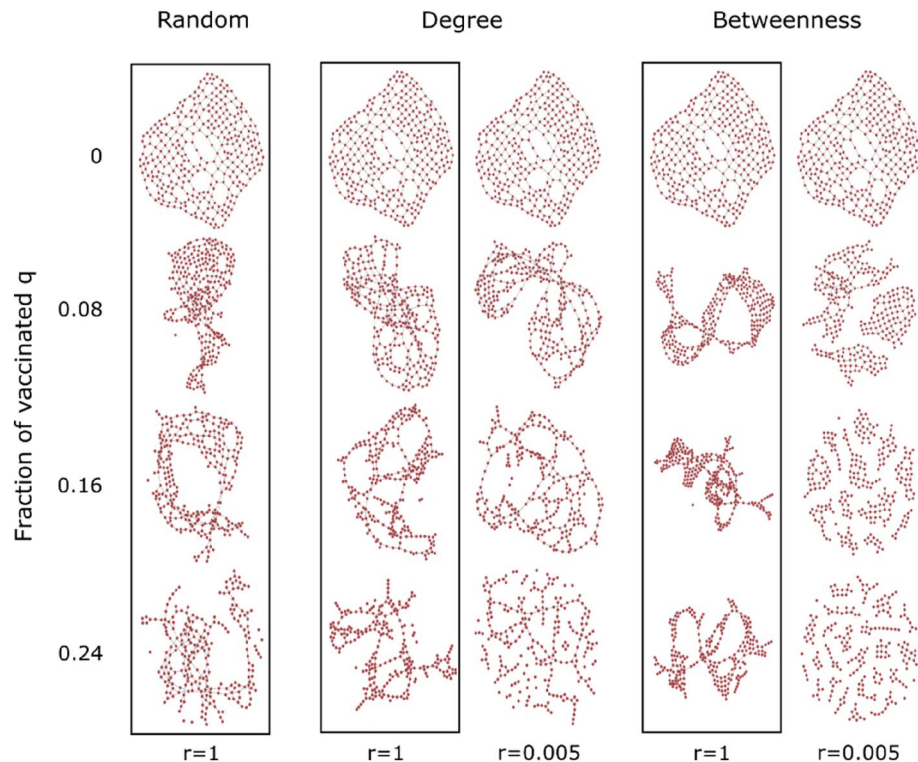


Figure 1. Effect of three nodes vaccination strategies for four fractions of vaccinated individuals q : 0, 0.08, 0.16, and 0.24 over the BE3 network. The boxed graphs correspond to the reduced network obtained following a non-adaptive algorithm, while non-boxed ones to the reduced network obtained from a semi-adaptive algorithm, with recalculation steps $r = 0.005$.

In Fig. 2G and H, we show the values of the three SIs averaged over the 12 analyzed networks, and in Fig. 2I, the average score of the seven strategies averaged both over the networks and the fraction of vaccinated individual q . When considering the average over the network, the best NVS to minimize the LCC is PAG; the best strategies for minimizing TI and IP are PAG and BET. Interestingly the DEG efficacy, which only contains information about the number of adjacent neighbors of a node, is comparable to BET and significantly outperforms *Closeness* (CLO) in minimizing the LCC despite BET and CLO containing more information about the topology of the network.

Semi-adaptive strategies. In the non-adaptive NVSs we used in Fig. 2, the centrality of each node is calculated on the initial network and is no longer updated. After each vaccination (node removal), the network structure changes, and the difference between the node centralities calculated on the initial network and the updated one increases. The difference between the degree calculated on the original network and the reduced one increases is smaller or equal to the number of removed nodes. For non-local NVS, such as the BET and the CLO, this difference varies in a non-linear way.

In Fig. 3, we show the outcomes from semi-adaptive NVS. In Fig. 3A, we show that for low values of the fraction of vaccinated q , when the difference between the initial centralities and their real value is small, the BET strategy outperforms, on average, PAG. Therefore, we introduced a new class of algorithms that we called semi-adaptive, where after a fraction of nodes r is removed from the network, recalculate the centrality of each node. Because the number of recalculations is fixed, the algorithm complexity of such semi-adaptive strategies is the same as the non-adaptive ones. For the BET strategy, moving from a non-adaptive to a semi-adaptive strategy can reduce the LCC, the TI, and the IP up to a factor of 3 to 4, as shown in Fig. 3B, C, and D, for a recalculation factor $r = 0.005$.

The consequences of implementing a semi-adaptive strategy on a single network can be seen in Fig. 1 for $r = 1$ (no recalculation) and $r = 0.005$. The reduced networks obtained by removing the fraction q of vaccinated individuals are visibly more fragmented for the same vaccination target for the semi-adaptive NVS.

In Fig. 3E, F, and G, we show the percental variation induced by going from a non-adaptive algorithm to a semi-adaptive one with $r = 0.005$, of the three SIs averaged over the 12 networks for three NVS. We show that CLO is the one that improves the most when implementing recalculation. When considering a recalculation factor $r = 0.005$, the NVS that, on average, triggers the best minimization of the three SIs is BET, as shown in Fig. 3H.

In Fig. 4A, C, and E, we show that in 5 out of the 12 networks, one of the three SIs is minimized by an additional strategy, while in 2 networks (ECO, BE3) the SIs are minimized by different NVS. In Fig. 4B, D, and F, we

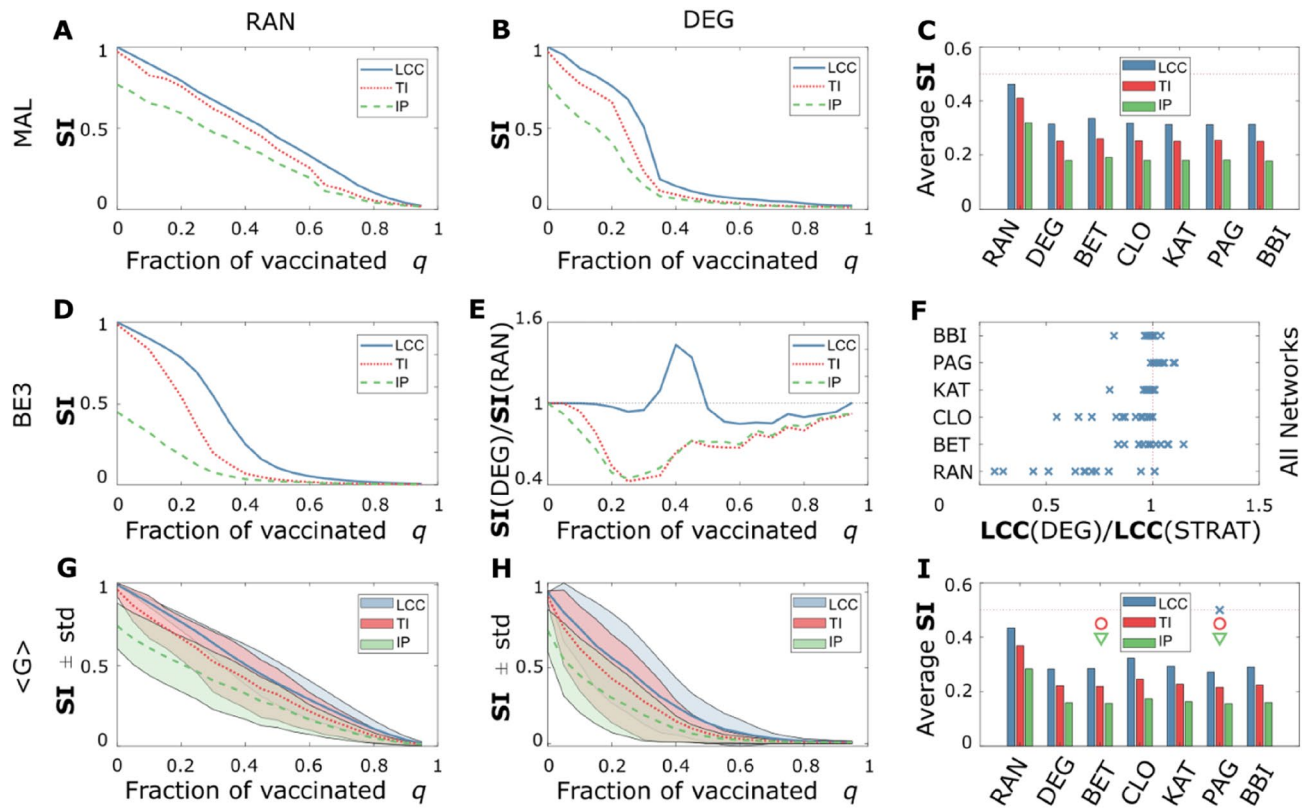


Figure 2. Comparison of non-adaptive strategies. (A, B) spread indicators **SI** for MAL network⁵⁰ as a function of the fractions of vaccinated q , for the RAN and the DEG strategies, respectively; (C) average **SI**s, defined as the average value that the three **SI**s assume in the 20 fractions of vaccinated q in the MAL network; (D) we show the dependency of the **SI**s for BE3; (E) **SI**s values obtained for the DEG strategy normalized by the ones obtained by the RAN strategy, a value smaller than one indicates that the DEG is more effective than RAN at minimizing that **SI**; (F), number of times in which the DEG strategy is more effective than the strategy on the y-axis at minimizing the LCC; (G, H) $\text{SI} \pm 1 \sigma$ confidence averaged over the 12 analysed networks; (I) average **SI**s, defined as the average value that the three **SI**s assume in the 12 networks and the 20 fractions of vaccinated q . (I) the overlaid symbols, blue cross, red circle, and green triangle, indicate the optimal strategies to minimize the LCC, the TI, and the IP, respectively; in (C), all the NVS but RAN and BET are equally effective in minimizing the three **SI**s, namely their relative difference is smaller than $1/50$: $\left(\frac{\text{STRAT}}{\min(\text{SI}(\text{STRAT}))} - 1 < \frac{1}{50}\right)$. The notation $\langle G \rangle$ indicates that the quantities in (G, H, I) are averaged over the 12 analysed networks.

show that the difference between the efficacy of the optimal strategies is more limited in the semi-adaptive case, with $r = 0.005$, in 2 out of the 12 networks (NET, WP1), two of the **SI**s are minimized by an additional strategy, and in one network (ECO) the **SI**s are minimized by two different NVS.

For $q > 0.5$, in most of the semi-adaptive cases, with $r = 0.005$, there are at least two NVS that perform equally. Therefore, Supplementary Table S1 shows the percentage of cases where each NVS is among the best to minimize the corresponding **SI**. For higher values of q , the strategy that more often minimizes the three **SI**s in the non-adaptive scenario is PAG, and in the semi-adaptive one is CLO. Interestingly, in the semi-adaptive approach, while BET is the worse of the non-random strategy in the $q > 0.5$ regime, it outperforms the others in the $q < 0.5$ region, and it becomes the best overall NVS when averaging across all the vaccination targets q .

We show a summary of the best strategies as a function of the recalculation step r , the **SI**, and the network in Table 1.

Vaccination requirements. One of the main concerns of governments regarding the COVID-19 pandemic has been to minimize the total number of infected individuals and to slow down the infection rate, not to overwhelm the health systems³². To achieve these aims, the governments immediately implemented non-pharmacological interventions (NPIs), like enforcing social distancing and lockdowns^{14,33}, and began immunizing individuals with a vaccination campaign³⁴. To study this problem, we investigate what percentage of the population needs to be vaccinated to keep the **TI** and the **IP** below a certain threshold θ .

In Fig. 5, we show that the NVS minimizing the vaccination requirement is the PAG, for a non-adaptive approach, and for a semi-adaptive approach with a low acceptable **SI**s threshold, namely higher vaccination requirements, is the CLO, and for a higher acceptable threshold, is the BET. Furthermore, going from a random vaccination strategy (RAN) to a non-random one reduces by 49–61% the number of vaccines to keep the spread of the disease below the threshold $\theta = 0.10$. Further, going from a non-adaptive algorithm to a semi-adaptive

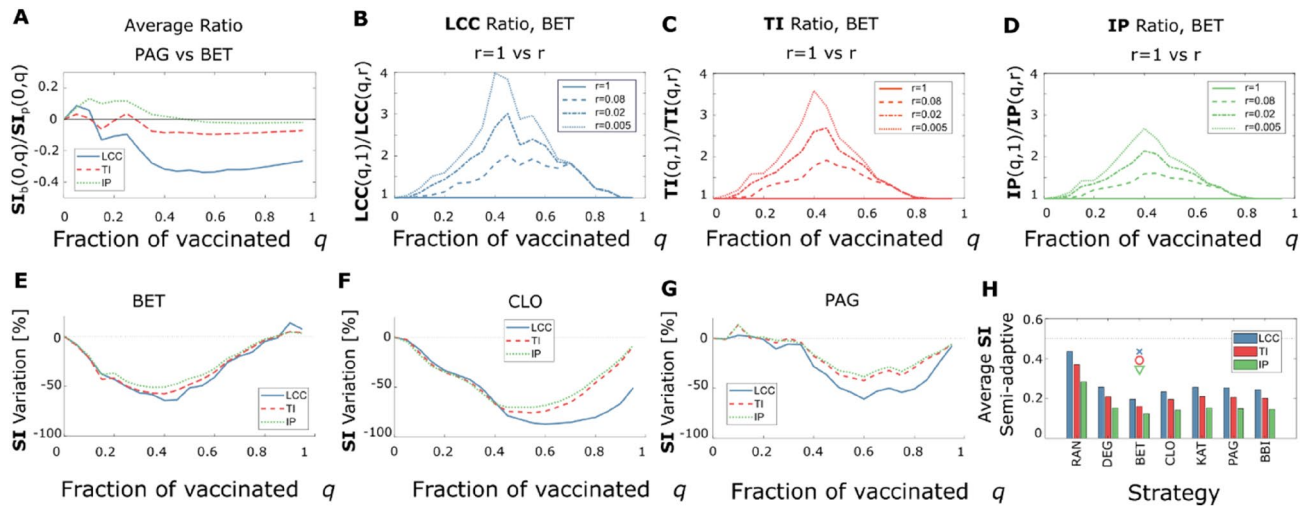


Figure 3. Comparison of Semi-Adaptive Strategies. In this figure, we compare the three SI obtained for non-adaptive strategies with the ones obtained for semi-adaptive strategies. (A) SI values averaged between 0 and q , obtained with BET and normalized by the ones obtained with PAG. A value bigger than one means that BET outperforms PAG, and a value smaller than one means that PAG outperforms BET, non-adaptive strategies; (B, C, D) relative decrease of LCC, TI, and IP respectively moving from a non-adaptive algorithm, solid line, to refined recalculation steps, dashed for $r = 0.08$, dot-dashed for $r = 0.02$, and dotted for $r = 0.005$; (E, F, G) percental reduction of the three SI going from a non-adaptive algorithm to a 0.005-adaptive algorithm, for three the BET, CLO, and PAG strategies respectively; (H) average SIs, defined as the average value that the three SIs assume in the 12 networks and the 20 fractions of vaccinated q , for a 0.005-adaptive algorithm.

one reduces the vaccination requirement of the best strategy to keep the spread of the disease below the same threshold by 20%.

Discussion and conclusion

This research has focused on finding an effective way to curb epidemic spreading in real-world complex networks via vaccinations. Finding the optimal set of nodes to minimize the size of the LCC is an NP-complete problem approached in the literature both in random networks^{24,25,35} and in real-world networks^{25,36}. We followed the approach in²⁵ by using established centrality measures to determine the node vaccination priority. Here we analyzed two classes of NVSs, the non-adaptive, where the centrality is calculated on the original adjacency matrix, and the semi-adaptive ones, where the centrality of each node is updated after a fraction r of nodes is removed. The computational complexity of a semi-adaptive NVS is the same as the non-adaptive equivalent because the centrality calculation is repeated a fixed number of times. In contrast, in the adaptive approach, where nodes rank is computed after each node removal³⁷, the computational complexity increases with the number of nodes in the network.

Random NVSs are generally less effective at curbing the epidemic than a strategy containing information about the network nodes' social structure^{22,30}. We showed that the vaccination requirement to keep the SIs below 10% of the total population dramatically benefits from a non-random NVS, saving an average of 49 and 60% of the vaccines, compared to the random NVS for the non-adaptive and the semi-adaptive approach, respectively. These outcomes give important insights into performing effective vaccination policies. On the one hand, these outcomes show that when implementing vaccination strategies, it is fundamental to consider the network's structure. On the other hand, they would suggest that gathering information about the local structure of the network nodes, such as sampling the number of contacts of individuals, might significantly improve the efficacy of the vaccination process. This would be particularly important since many vaccination campaigns used to curb the COVID19 spreading neglected the network structure^{38–40}. We explored six non-random NVSs and found that in the non-adaptive approach, both the betweenness centrality, BET, and the PageRank centrality, PAG, are the two most effective strategies for minimizing the size of the LCC, but also the other two spread indicators. In the semi-adaptive approach, the two best NVSs are based on the betweenness (BET) and closeness (CLO) centrality of the nodes. The former is very effective at minimizing all three SIs when the fraction of available vaccines is below 45%, while the second performs better in the 45–70% range, above which their performances are comparable. These outcomes indicate that, when possible, considering the global and whole structure of the network, as required to implement the PAGERANK (PAG), betweenness (BET), and closeness (CLO) strategies, it would be an optimal way to vaccinate the population.

We found that implementing a semi-adaptive (recalculated) NVS significantly reduces the vaccination requirement compared to a non-adaptive approach. This finding would indicate that when implementing vaccination policies in real-world networks, considering the changes in the network structure along the vaccination process may significantly improve the efficacy of the vaccination process to curb epidemics. Noteworthy, we improved the non-adaptive NVS by partially recalculating the node centralities. In the semi-adaptive NVS, we recalculated the

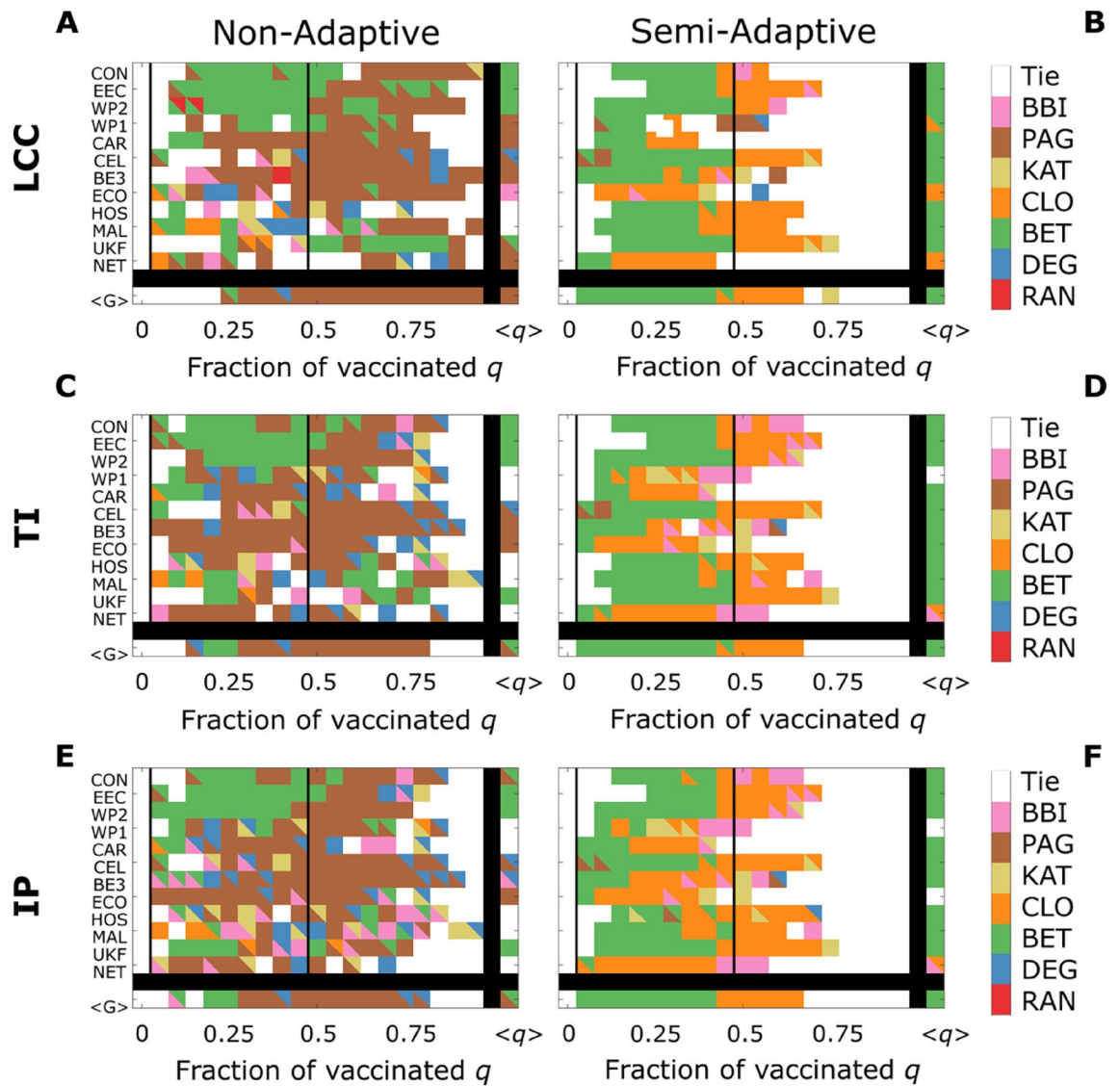


Figure 4. Best Strategy comparison. In this figure, we show, for both non-adaptive and semi-adaptive strategies, the best NVSs to minimize each SIs for the 12 networks. (A, C, E), optimal strategies for the non-adaptive approach; (B, D, F) optimal strategies for the semi-adaptive approach with a recalculation step $r = 0.005$. (A, B) best strategies to minimize LCC; (C, D) best strategies to minimize TI; (E, F) best strategies to minimize IP. The bottom line in each subplot shows the strategy that, on average, minimizes the SI for that value of q , while the leftmost column the strategy that, on average, minimizes the corresponding SI of that network. The colour code is the same in the six subpanels, a description of the acronym of the strategies can be found in Table 3, and a summary of the acronyms of the networks in Table 2; in case of a tie between two strategies, both colours are indicated, and in case of a tie between three or more strategies the colour white is used. A tie happens when the relative difference between the best NVS and the second-best one is smaller than $1/50$. The notation $\langle G \rangle$ indicates that the quantities are averaged over the 12 analysed networks and the notation $\langle q \rangle$ indicates that the quantities are averaged over the 20 analysed fraction of vaccinated q .

centrality of each node after 0.5%, 2%, and 8% of the node were removed, and this semi-adaptive approach does not increase the complexity of the algorithms. Some strategy, like CLO, profits from moving from a non-adaptive to an adaptive approach more than others, like PAG. We attributed this different behavior to the locality of the NVS considered. We argue that highly non-local measures may experience a higher increase in performance by passing from a non-adaptive to a semi-adaptive approach. On the contrary, the more local a centrality measure is, the less it will improve by implementing a semi-adaptive NVS. This is because if the centrality measure is local, the centrality of a node will be less affected by the removal of a random node in the network. For example, CLO is defined as the inverse of the harmonic average of the distance. The distance between points is a highly non-local measure that can vary considerably by removing a single node. On the other hand, PAG quantifies a node's authority by looking at the authority of the nodes pointing at it. Therefore, the information about distant nodes

SI	r	Network name											
		CON	EEC	WP2	WP1	CAR	CEL	BE3	ECO	HOS	MAL	UKF	NET
LCC	1	BET 0.34	BET 0.32	BET 0.40	PAG 0.20	PAG 0.19	DEG PAG 0.36	PAG 0.31	BBI 0.13	TIE 0.33	TIE 0.33	BET PAG 0.40	PAG 0.08
	0.08	TIE 0.34	BET 0.30	BET 0.32	KAT BBI 0.19	TIE 0.19	BET 0.29	DEG 0.26	BET 0.12	BET PAG 0.30	BET 0.28	BET CLO 0.35	CLO BBI 0.08
	0.02	BET BBI 0.32	BET 0.29	BET 0.27	BET CLO 0.18	BET 0.16	BET 0.28	BET 0.19	BET 0.12	BET 0.28	BET 0.24	BET 0.28	BET 0.07
	0.005	BET 0.30	BET 0.28	BET 0.23	BET CLO 0.18	BET 0.14	BET 0.24	BET 0.11	CLO 0.11	BET 0.28	BET 0.24	BET 0.28	CLO 0.06
TI	1	BET 0.29	BET 0.27	BET 0.30	PAG 0.14	TIE 0.15	DEG PAG 0.31	PAG 0.19	PAG 0.09	TIE 0.26	TIE 0.26	BET 0.35	TIE 0.05
	0.08	BET PAG 0.29	BET 0.26	BET 0.24	KAT 0.13	TIE 0.14	BET 0.24	DEG 0.18	TIE 0.09	TIE 0.25	BET 0.22	BET 0.31	TIE 0.05
	0.02	BET 0.28	BET 0.25	BET 0.21	BET CLO 0.13	BET 0.12	BET 0.23	BET 0.15	BBI 0.09	BET 0.23	BET 0.19	BET 0.26	CLO BBI 0.05
	0.005	BET 0.26	BET 0.24	BET 0.18	TIE 0.13	BET 0.11	BET 0.20	BET 0.09	BET 0.08	BET 0.23	BET 0.19	BET 0.26	CLO BBI 0.05
IP	1	BET PAG 0.23	BET 0.22	BET 0.21	DEG PAG 0.10	TIE 0.10	DEG PAG 0.23	DEG BBI 0.08	PAG 0.06	TIE 0.21	TIE 0.19	BET 0.27	TIE 0.03
	0.08	TIE 0.22	BET 0.21	BET 0.18	KAT BBI 0.09	TIE 0.10	BET 0.19	BBI 0.07	TIE 0.06	TIE 0.20	BET CLO 0.17	BET 0.25	TIE 0.03
	0.02	BET 0.21	BET 0.20	BET 0.16	TIE 0.09	BET 0.09	BET 0.18	BBI 0.06	BBI 0.06	BET 0.19	BET 0.15	BET 0.21	CLO BBI 0.03
	0.005	BET 0.20	BET 0.20	BET 0.14	TIE 0.09	BET 0.08	BET 0.16	BET 0.05	TIE 0.06	BET 0.19	BET 0.15	BET 0.21	CLO BBI 0.03

Table 1. Best vaccination strategy per network. For each recalculation step r and network, we indicate the best NVS to minimize the average SI over q , as described in the methods. For each box we indicate the acronym of the best NVS and the value of the average of that strategy. If the relative difference between two NVS is smaller than $1/50$, we indicate both the best strategies in bold, while when the three (or more) best strategies differ by less than $1/50$, we label that case with TIE.

is embedded in the adjacent nodes. Therefore, using a semi-adaptive algorithm always improves the efficacy of the NVS, and it is especially beneficial in the case of non-local NVS.

The outcomes of our research show that when planning to contain an epidemic event is important to consider the social structure of interaction between individuals. Knowing the network structure underlying social interactions is necessary to calculate the centrality measures discussed in this work, but it is a promising way to perform effective vaccination strategies. Unfortunately, considering the real-world social network structure is a challenging task. Nonetheless, in the last years, several methods have been developed to reconstruct the social interaction network in metropolitan areas like Berlin¹⁵ and to approximate the value of some of the centrality measures having only access to local and partial information²².

Another problem that became evident during the current COVID pandemic was the reluctance of a large fraction to get vaccinated or to wear masks; this suggests the importance of coupling two dynamics, one for the virus and one for the vaccination-predisposition⁴¹. As developed in Ref.⁴², the Dynamical Message-Passing method can be used to approach these coupled dynamics.

Finally, we showed that in 5 of the 12 networks, the best NVS to minimize each SIs is the same, and in 2 out of 12, the best strategies to minimize the three SIs are different. Since the LCC is a static and the most straightforward indicator of the maximal possible disease spreading, it is necessary to analyze the two dynamic spread indicators TI and IP, to contain the outbreak. This is particularly important considering that many studies in network science adopted the LCC to evaluate the efficacy of the NVS^{5,7,17,25}.

Methods

Simulations algorithm. To study the impact of different node vaccination strategies (NVS) on the dynamic of an epidemic outbreak, we used the following protocol: (1) We import one of the 12 real-world networks; (2) We infect one random node in the network; (3) We set a vaccination target q , namely the fraction of the network nodes that will be vaccinated; (4) We remove Nq nodes from the network; (5) We simulate on the reduced net-

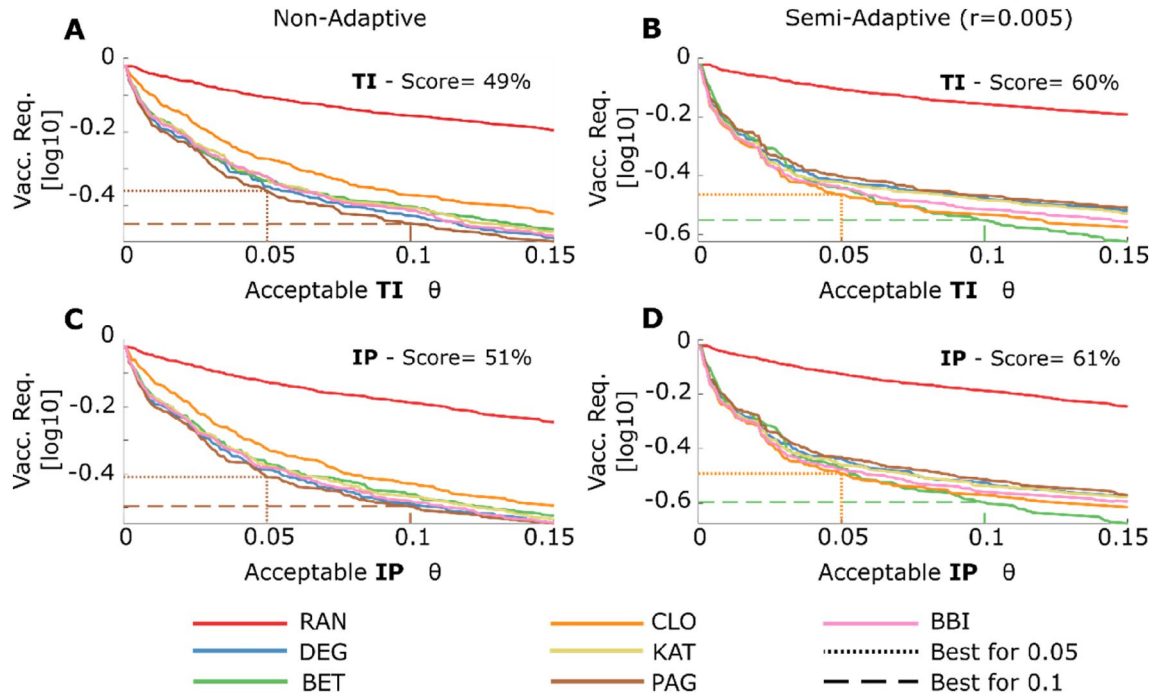


Figure 5. Vaccination Requirements. In this figure, we show the vaccination requirement to keep the TI and the IP below the acceptable threshold ϑ . (A, C) fraction of the population that needs to be vaccinated to keep the acceptable SIs value below a threshold ϑ , in case of non-adaptive algorithm ($r = 1$); (B, D) fraction of the population that needs to be vaccinated to keep the acceptable SIs value below a threshold ϑ in case of semi-adaptive algorithm with a recalculation step $r = 0.005$. The dashed and the dotted lines indicate the best strategy for $\vartheta = 0.1$, and $\vartheta = 0.05$ respectively. The score is the percental reduction in the vaccination requirement going from the random strategy (RAN) to the best one for $\vartheta = 0.1$. We show the y-axis in log scale to increase the readability of the figure.

ID	Network name	Acronym	Type	N	E	$\langle k \rangle$	\varnothing	Reference
1	Conference	CON	Social	257	3248	25.3	6	43
2	Email EU Core	EEC	Email	986	25,552	51.8	7	44
3	Workplace 2	WP2	Social	159	2180	27.4	7	43
4	Workplace 1	WP1	Social	47	138	5.9	7	45
5	Cargoship BB	CAR	Ship	821	4342	10.5	9	46
6	C-Elegans	CEL	Neural	297	2345	15.8	5	47
7	Beijing 3	BE3	Street	322	1075	6.7	27	31
8	E-Coli	ECO	Protein	1100	3637	6.6	11	48
9	Hospital	HOS	Social	43	358	16.7	5	49
10	Malawi	MAL	Social	84	346	8.2	5	50
11	UK Faculty	UKF	Social	81	817	20.2	4	51
12	Netscience	NET	Citation	397	914	4.8	17	52

Table 2. Real-world networks summary. Real-world networks used in this research with acronym, network type, reference, and basic statistical features: number of nodes (N), number of edges (E), average node degree ($\langle k \rangle$), and network diameter (\varnothing).

work a SIR dynamic, and for each step, we track the number of nodes in the three populations: S, I, and R. (6) We stop the simulation when no more nodes are in the state I.

Real-world networks summary. In this work, we analyzed twelve real-world networks from different fields. In Table 2 below, for each network, we summarized the type, the number of nodes, the number of links, the average node degree, and the network’s diameter. We implement the SIR over each network’s largest connected component (LCC). Four real-world networks (CON, WP1, WP2, and HOS) are face-to-face networks, where the interaction between individuals and the interaction time are accounted for⁴³. We consider only the interactions that happened in the first 12 h for these networks.

The epidemic SIR-dynamic we considered in this work, strictly speaking, can be properly applied to face-to-face networks. Nonetheless, coupling node removal and SIR-like dynamic could be useful to describe a variety of real problems in other types of networks, such as halting computer viruses or stopping epidemic spreading among airports or cities. For example, vaccinating vertices in CAR, and BE3, could be interpreted as a localized lockdown to prevent the spread of an epidemic in a geographical area and a district, respectively. For this reason, we also considered a variety of real-world networks that are not face-to-face.

Non-adaptive approach to vaccination. In the non-adaptive approach of the NVS, we rank the nodes according to the node centrality metrics at the beginning of the simulation (before any node removal), and we do not update the rank. The non-adaptive node removal (attack) strategies are also named ‘initial node attack’ or ‘not-recalculated node attack’^{25,53,54}. In the case of ties, i.e., nodes with equal centrality rank value, one of the nodes with the highest centrality nodes are removed at random.

Semi-adaptive approach to vaccination. In the semi-adaptive approach of the NVS, the node’s rank is updated during the removal process. First, we calculate the rank of each node, and after a fraction $r < q$ of nodes is removed, the rank of each node is calculated in the reduced network. We repeat this procedure until we vaccinate a fraction q of the nodes. In the last iteration, we vaccinate a fraction of nodes smaller than r to vaccinate exactly a fraction q of the population. In this work, we considered four values of r : $r = 0.005$, $r = 0.02$, $r = 0.08$, and $r = 1$.

Node vaccination strategies (NVS). In this work, we implemented seven NVSs; except for the random vaccination (RAN), the others require knowing the entire adjacency matrix of the network.

Random. The first strategy we approach is to assign to each node a random weight. This vaccination strategy has the significant advantage of not requiring prior knowledge of the network’s topology.

Degree. The degree centrality of a node is the number of edges (connections) to it^{26,53}:

$$C_D(v) = \sum_j A_{j,v}$$

where A is the adjacency matrix.

Betweenness. The betweenness centrality of a node v was introduced in⁵⁵, and it measures the fraction of the shortest paths connecting any two nodes i , and j passing through the node v . In formulas:

$$C_B(v) = \sum_{i \neq v \neq j} \frac{\sigma_{ij}(v)}{\sigma_{ij}}$$

where $\sigma_{i,j}$ is the total number of shortest paths connecting i and j , and $\sigma_{i,j}(v)$ the total number of shortest paths connecting i and j passing through v .

Closeness. The closeness centrality was introduced in⁵⁶ as the inverse of farness. The closeness of a node v is the inverse of the sum of the distance (or shortest path length) between node v and all the other nodes in the graph:

$$\tilde{C}_C(v) = \frac{1}{\sum_{j \neq v} d(j,v)}$$
 where $d(j, v)$ is the distance between node j and v .

The definition of closeness centrality we used in this paper is an extension of the one we show above. We implemented the harmonic closeness centrality introduced in⁵⁷, which assumes finite values also in disconnected. This measure is often called harmonic centrality, and it is defined as:

$$C_C(v) = \sum_{j \neq v} \frac{1}{d(j,v)}$$

Katz. The Katz centrality, introduced in⁵⁸, is defined as:

$$C_K = \alpha A C_K + \beta$$

where α is an attenuation factor, A the adjacency matrix, and β a personalization vector. Here, we assume $\alpha = 0.01$, and $\beta = 1$. An alternative definition of Katz centrality is:

$$C_K(v) = \sum_{k=1}^{\infty} \sum_{\{j=1\}}^n \alpha^k (A^k)_{vj} + \beta,$$

this second definition makes its interpretation more straightforward: it is counting the number of paths of length k connecting the node v to each other node j , and this value is weighted with the k -power of the attenuation factor α .

ID	Name	Acronym	Formula	Reference
1	Random	RAN	$C_R(v) = \xi \in [0, 1]$	64
2	Degree	DEG	$C_D(v) = \sum_j A_{j,v}$	55
3	Betweenness	BET	$C_B(v) = \sum_{i \neq v \neq j} \frac{\sigma_{ij}(v)}{\sigma_{ij}}$	55
4	Closeness	CLO	$C_C(v) = \sum_{j \neq v} \frac{1}{d(i,v)}$	57
5	Katz	KAT	$C_K = \alpha AW_K + \beta$	58
6	Pagerank	PAG	$C_P = \frac{1-\delta}{N} + \delta \sum_{u \in \Gamma[v]} \frac{C_P(u)}{C_D(u)}$	59
7	BB-Index	BBI	$C_{BB}(v) = \sum_{i \neq v} \frac{C_D(v)}{d(i,j)}$	60

Table 3. NVS summary. For each of the 7 considered NVS we used in this work, we report name, acronym, formula used to calculate, and the reference.

PageRank. PageRank Centrality, introduced in⁵⁹, was the first algorithm used by Google to rank the importance of a webpage. It is defined iteratively by:

$$C_P = \frac{1-\delta}{N} + \delta \sum_{u \in \Gamma[v]} \frac{C_P(u)}{C_D(u)}$$

where δ is a dumping factor, which we set to 0.85, and $\Gamma[v]$ is the set of first neighbors of v .

BB Index. The BB-index was introduced in⁶⁰, and it is defined as a closeness centrality rescaled by the degree of each node:

$$C_{BB}(v) = \sum_{i \neq v} \frac{C_D(v)}{d(i, v)}$$

where $C_d(v)$ is the degree centrality of the node v and $d(i, v)$ is the distance between nodes i and v .

The BB-Index was introduced to create a network descriptor that combines information about the connectivity of the network and its small-world properties.

SIR on network. To model the temporal dynamics of the epidemic outbreak, we used a Monte Carlo agent-based Susceptible-Infected-Recovered compartmental model (SIR) to simulate the spread of a disease on real-world networks^{9,12,29,61}. In an agent-based SIR, each node of the network represents an individual, which at any given time can be in one of three states, susceptible (S), Infected (I), or Recovered (R). At any time step, a susceptible node is infected by an adjacent infected node with a probability of β , and every infected node has a probability of γ to recover from the infection. In our simulations, we used the values of β and γ that were recovered from the first months of the COVID pandemic in Italy: $\beta = 0.18$, and $\gamma = 0.037$ ⁶². Once an individual is immunized, it cannot be reinfected nor be infective to other individuals/nodes; this can happen if the individual undergoes vaccination or recovers from the infection. Therefore, we initialize the system by setting all the individuals to be susceptible, then we infect one random individual, and finally, we vaccinate a fraction q of the susceptible ones accordingly to one of the seven NVS. To simulate the SIR spreading, we parallelized the implementation of the above-mentioned process, realized using the graph-tool Python library⁶³. For each set of parameters and real-world network, we performed 1000 independent simulations where the initially infected node is randomly selected (Table 3).

Spreading indicators (SI) calculation. We considered three SIs in this work: the LCC, the TI, and the IP. For every set of simulation parameters (G, q, r, γ, β) where G represents the network, we run the simulation protocol 1000 times, and for each, we calculate the three SIs. Finally, we defined LCC, TI, and IP as the average of the SI values over the 1000 iterations.

The largest connected component (LCC) is the biggest connected subgraph of the network; here, we consider only its size, namely the number of nodes in it. The total infected (TI) is defined as the number of infected individuals at the end of the pandemic event. We obtained it as the difference between the number of Susceptible individuals at time zero and at the end of the simulation: $TI_i = S_i(0) - S_i(END)$. The Infected Peak (IP) is defined as the maximum number of simultaneously infected nodes. Operatively we defined it as the maximum of $I(t)$: $I_i = \max_t I_i(t)$. To simplify the comparison of the SIs between networks of different sizes, we normalized them by the size of the LCC of the initial network (i.e., before any removal).

Averaged SI calculations. When the average value of a SI is used in the text, it is defined in one of the three following ways: $E[SI] = \frac{1}{20} \sum_i SI(r, q = i, STR, G)$ if it is over the vaccination target q ; $E[SI] = \frac{1}{12} \sum_i SI(r, q, STR, G = i)$ if it is averaged over the networks; and $E[SI] = \frac{1}{20} \frac{1}{12} \sum_i \sum_j SI(r, q = i, STR, G = j)$ if it is averaged both over networks and over vaccination target.

Data availability

The datasets analysed during the current study are available in the ‘Netzschleuder’ repository [<https://networks.skewed.de/>], in the ‘Stanford Large Network Dataset Collection’ repository [<https://snap.stanford.edu/data/index.html>], in ‘The Colorado Index of Complex Networks (ICON)’ repository [<https://icon.colorado.edu/#/>], and in the ‘SocioPatterns’ repository [<http://www.sociopatterns.org/datasets/>].

Received: 18 July 2022; Accepted: 18 November 2022

Published online: 09 December 2022

References

- Britton, T. Epidemics in heterogeneous communities: estimation of R_0 and secure vaccination coverage. *J. R. Statist. Soc. Ser. B (Statist. Methodol.)* **63**(4), 705–715 (2001).
- Sultana, J. *et al.* Potential effects of vaccinations on the prevention of COVID-19: rationale, clinical evidence, risks, and public health considerations. *Expert Rev. Vacc.* **19**, 919–936 (2020).
- Mathieu, E. *et al.* A global database of COVID-19 vaccinations. *Nat. Hum. Behav.* **5**, 947–953 (2021).
- Moreno, Y., Pastor-Satorras, R. & Vespignani, A. Epidemic outbreaks in complex heterogeneous networks. *Eur. Phys. J. B* **26**, 521–529 (2002).
- Newman, M. E. Spread of epidemic disease on networks. *Phys. Rev. E* **66**(1), 016128 (2002).
- Cohen, R., Havlin, S. & Ben-Avraham, D. Efficient immunization strategies for computer networks and populations. *Phys. Rev. Lett.* **91**(24), 247901 (2003).
- Gallos, L. K., Liljeros, F., Argyrakis, P., Bunde, A. & Havlin, S. Improving immunization strategies. *Phys. Rev. E - Stat. Nonlinear, Soft Matter Phys.* **75**, (2007).
- Sun, Y., Liu, C., Zhang, C. X. & Zhang, Z. K. Epidemic spreading on weighted complex networks. *Phys. Lett. Sect A Gen. At. Solid State Phys.* **378**, 635–640 (2014).
- Pastor-Satorras, R., Castellano, C., Van Mieghem, P. & Vespignani, A. Epidemic processes in complex networks. *Rev. Modern Phys.* **87**(3), 925 (2015).
- Amaral, M. A., de Oliveira, M. M. & Javarone, M. A. An epidemiological model with voluntary quarantine strategies governed by evolutionary game dynamics. *Chaos Solitons Fractals* **143**, 110616 (2021).
- Bellingeri, M., Bevacqua, D., Scotognella, F., Alfieri, R. & Cassi, D. A comparative analysis of link removal strategies in real complex weighted networks. *Sci. Rep.* **10**(1), 1–15 (2020).
- Bellingeri, M. *et al.* Modeling the consequences of social distancing over epidemics spreading in complex social networks: from link removal analysis to SARS-CoV-2 prevention. *Front. Phys.* **9**, (2021).
- Saunders, H. A. & Schwartz, J. M. COVID-19 vaccination strategies depend on the underlying network of social interactions. *Sci. Rep.* **11**(1), 1–10 (2021).
- Müller, S. A., Balmer, M., Neumann, A. & Nagel, K. Mobility traces and spreading of COVID-19. (2020) <https://doi.org/10.14279/depositon-9835>.
- Nagel, K., Rakow, C. & Müller, S. A. Realistic agent-based simulation of infection dynamics and percolation. *Phys. A Statist. Mech. Its Appl.* **584**, 126322 (2021).
- Dönges, P. *et al.* Interplay between risk perception, behavior, and COVID-19 spread. *Front. Phys.* **10**, (2022).
- Holme, P. Efficient local strategies for vaccination and network attack. *EPL (Europhys. Lett.)* **68**(6), 908 (2004).
- Chen, Y., Paul, G., Havlin, S., Liljeros, F. & Stanley, H. E. Finding a better immunization strategy. *Phys. Rev. Lett.* **101**(5), 058701 (2008).
- Hadidjojo, J. & Cheong, S. A. Equal graph partitioning on estimated infection network as an effective epidemic mitigation measure. *PLoS One* **6**(7), e22124 (2011).
- Schneider, C. M., Mihaljev, T., Havlin, S. & Herrmann, H. J. Suppressing epidemics with a limited amount of immunization units. *Phys. Rev. E* **84**(6), 061911. <https://doi.org/10.1103/PhysRevE.84.061911> (2011).
- Bellingeri, M., Agliari, E. & Cassi, D. Optimization strategies with resource scarcity: from immunization of networks to the traveling salesman problem. *Modern Phys. Lett. B* **29**(29), 1550180 (2015).
- Wang, Z., Zhao, D. W., Wang, L., Sun, G. Q. & Jin, Z. Immunity of multiplex networks via acquaintance vaccination. *EPL (Europhys. Lett.)* **112**(4), 48002 (2015).
- Kudryashov, N. A., Chmykhov, M. A. & Vigdorowitsch, M. Analytical features of the SIR model and their applications to COVID-19. *Appl. Math. Model.* **90**, 466–473 (2021).
- Hartvigsen, G., Dresch, J. M., Zielinski, A. L., Macula, A. J. & Leary, C. C. Network structure, and vaccination strategy and effort interact to affect the dynamics of influenza epidemics. *J. Theor. Biol.* **246**, 205–213 (2007).
- Wandelt, S., Sun, X., Feng, D., Zanin, M. & Havlin, S. A comparative analysis of approaches to network-dismantling. *Sci. Rep.* **8**(1), 1–15 (2018).
- Nguyen, Q., Nguyen, N. K. K., Cassi, D. & Bellingeri, M. New Betweenness Centrality Node Attack Strategies for Real-World Complex Weighted Networks. *Complexity* **2021**, (2021).
- Kendall, D. G. *Deterministic and stochastic epidemics in closed populations.* (1956).
- Moreno, Y., Gómez, J. B. & Pacheco, A. F. Epidemic incidence in correlated complex networks. *Phys. Rev. E* **68**(3), 035103 (2003).
- Kitsak, M. *et al.* Identification of influential spreaders in complex networks. *Nat. Phys.* **6**, 888–893 (2010).
- Braunstein, A., Dall’Asta, L., Semerjian, G. & Zdeborová, L. Network dismantling. *Proc. Natl. Acad. Sci. U. S. A.* **113**, 12368–12373 (2016).
- Guo, X. L. & Lu, Z. M. Urban road network and taxi network modeling based on complex network theory. *J. Inform. Hiding Multimed. Signal Process.* **7**(3), 558–568 (2016).
- Remuzzi, A. & Remuzzi, G. COVID-19 and Italy: What next?. *The Lancet* **395**, 1225–1228. [https://doi.org/10.1016/S0140-6736\(20\)30627-9](https://doi.org/10.1016/S0140-6736(20)30627-9) (2020).
- Thunström, L., Newbold, S. C., Finnoff, D., Ashworth, M. & Shogren, J. F. The benefits and costs of using social distancing to flatten the curve for COVID-19. *J. Benefit-Cost Anal.* **11**, 179–195 (2020).
- Matrajt, L. *et al.* Optimizing vaccine allocation for COVID-19 vaccines shows the potential role of single-dose vaccination. *Nat. Commun.* **12**(1), 1–18 (2021).
- Kobayashi, T. & Masuda, N. Fragmenting networks by targeting collective influencers at a mesoscopic level. *Sci. Rep.* **6**(1), 1–12 (2016).
- Jahanpour, E. & Chen, X. Analysis of complex network performance and heuristic node removal strategies. *Commun. Nonlinear Sci. Numer. Simul.* **18**, 3458–3468 (2013).
- Lordan, O. & Albareda-Sambola, M. Exact calculation of network robustness. *Reliab. Eng. Syst. Saf.* **183**, 276–280 (2019).
- Scabini, L. F. *et al.* Social interaction layers in complex networks for the dynamical epidemic modeling of COVID-19 in Brazil. *Phys. A Statist. Mech. Appl.* **564**, 125498 (2021).

39. Silvestris, N., Brunetti, O., Bernardini, R. & Cinieri, S. COVID vaccination in cancer patients: What vaccination priority strategies should there be?. *Front. Oncol.* **11**, 641388 (2021).
40. Wagner, C. E., Saad-Roy, C. M. & Grenfell, B. T. Modelling vaccination strategies for COVID-19. *Nat. Rev. Immunol.* **22**, 139–141. <https://doi.org/10.1038/s41577-022-00687-3> (2022).
41. Andrey, L. Y. & David, S. Optimal deployment of resources for maximizing impact in spreading processes. *Proceedings of the National Academy of Sciences* **114**(39), E8138–E8146 (2017).
42. Sun, H., David, S., & Andrey, L. Y. Competition, collaboration, and optimization in multiple interacting spreading processes. *Physical Review X* **11**(1), 011048 (2021).
43. Génois, M. & Barrat, A. Can co-location be used as a proxy for face-to-face contacts?. *EPJ Data Sci.* **7**(1), 1–18 (2018).
44. Leskovec, J., Kleinberg, J. & Faloutsos, C. Graph evolution: densification and shrinking diameters. *ACM Trans. Knowl. Discov. Data (TKDD)* **1**(1), 2 (2007).
45. Génois, M. *et al.* Data on face-to-face contacts in an office building suggest a low-cost vaccination strategy based on community linkers. *Netw. Sci.* **3**, 326–347 (2015).
46. Kaluza, P., Kölzsch, A., Gastner, M. T. & Blasius, B. The complex network of global cargo ship movements. *J. R. Soc. Interface* **7**(48), 1093–1103. <https://doi.org/10.1098/rsif.2009.0495> (2010).
47. White, J. G., Southgate, E., Thomson, J. N. & Brenner, S. The structure of the nervous system of the nematode *Caenorhabditis elegans*: the mind of a worm. *Philos. Trans. R. Soc. London* **314**(1), 340 (1986).
48. Serrano, M. Á., Boguñá, M. & Sagués, F. Uncovering the hidden geometry behind metabolic networks. *Mol. Biosyst.* **8**, 843–850 (2012).
49. Vanhems, P. *et al.* Estimating potential infection transmission routes in hospital wards using wearable proximity sensors. *PLoS One* **8**(9), e73970 (2013).
50. Ozella, L. *et al.* Using wearable proximity sensors to characterize social contact patterns in a village of rural Malawi. *EPJ Data Sci.* **10**(1), 46 (2021).
51. Nepusz, T., Petróczy, A., Négyessy, L. & Bazsó, F. Fuzzy communities and the concept of bridgeness in complex networks. *Phys. Rev. E* **77**(1), 016107 (2008).
52. Newman, M. E. Analysis of weighted networks. *Phys. Rev. E* **70**(5), 056131 (2004).
53. Iyer, S., Killingback, T., Sundaram, B. & Wang, Z. Attack robustness and centrality of complex networks. *PLoS One* **8**(4), e59613 (2013).
54. Bellingeri, M., Cassi, D. & Vincenzi, S. Efficiency of attack strategies on complex model and real-world networks. *Phys. A Stat. Mech. its Appl.* **414**, 174–180 (2014).
55. Linton, F. C. A set of measures of centrality based on betweenness. *Sociometry* **40**, 35–41 (1977).
56. Bavelas, A. Communication patterns in task-oriented groups. *J. Acoust. Soc. Am.* **22**, 725–730 (1950).
57. Marchiori, M. & Latora, V. Harmony in the small-world. *Phys. A Statist. Mech. Appl.* **285**(3–4), 539–546 (2000).
58. Katz, L. A new status index derived from sociometric analysis. *Psychometrika* **18**(1), 39–43 (1953).
59. Brin, S. & Page, L. The anatomy of a large-scale hypertextual web search engine. *Comput. Netw. ISDN Syst.* **30**(1–7), 107–117 (1998).
60. Bonchev, D. & Buck, G. A. From molecular to biological structure and back. *J. Chem. Inf. Model.* **47**, 909–917 (2007).
61. Thurner, S., Klimek, P. & Hanel, R. A network-based explanation of why most COVID-19 infection curves are linear. *Proc. Natl. Acad. Sci.* **117**(37), 22684–22689. <https://doi.org/10.1073/pnas.2010398117/-/DCSupplemental> (2020).
62. Cooper, I., Mondal, A. & Antonopoulos, C. G. A SIR model assumption for the spread of COVID-19 in different communities. *Chaos Solitons Fractals* **139**, 110057 (2020).
63. Peixoto, T. P. The graph-tool python library. at (2014).
64. Albert, R. & Barabási, A. L. Statistical mechanics of complex networks. *Rev. Modern Phys.* **74**(1), 47 (2002).

Acknowledgements

F.S., M.B., M.T., D.C., and R.A. acknowledge the Italian Ministry of Foreign Affairs and International Cooperation. This research is funded by a grant from the Italian Ministry of Foreign Affairs and International Cooperation. This project has received funding from the European Research Council (ERC) under the European Union's Horizon 2020 research and innovation programme (Grant agreement No. [816313]). This work is supported by Vietnam's Ministry of Science and Technology (MOST) under the Vietnam-Italy scientific and technological cooperation program for the period of 2021–2023. This work is supported by the Vietnam National University Ho Chi Minh City (VNU-HCM), Ho Chi Minh City, Vietnam under grant number B2018-42-01. We are greatly thankful to Van Lang University, Vietnam for providing the budget for this study.

Author contributions

F.S., M.B., D.C., and Q.N. designed the research; M.T., R.A., and F.Sar. wrote the simulation code; M.T., parallelized the algorithm; F.Sar., analyzed the data; F.Sar., and M.B. wrote the manuscript with the contribution of F.S., K.N.N.K., and T.L.T.; all authors reviewed the manuscript.

Competing interests

The authors declare no competing interests.

Additional information

Supplementary Information The online version contains supplementary material available at <https://doi.org/10.1038/s41598-022-24652-1>.

Correspondence and requests for materials should be addressed to F.S.

Reprints and permissions information is available at www.nature.com/reprints.

Publisher's note Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.



Open Access This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons licence, and indicate if changes were made. The images or other third party material in this article are included in the article's Creative Commons licence, unless indicated otherwise in a credit line to the material. If material is not included in the article's Creative Commons licence and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this licence, visit <http://creativecommons.org/licenses/by/4.0/>.

© The Author(s) 2022