The effect of delayed awareness and fatigue on the efficacy of self-isolation in epidemic control

Giulia de Meijere, Vittoria Colizza, Eugenio Valdano, Claudio Castellano

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The isolation of infectious individuals is a key measure of public health for the control of communicable diseases. However, involving a strong perturbation of daily life, it often causes psychosocial distress, and severe financial and social costs. These may act as mechanisms limiting the adoption of the measure in the first place or the adherence throughout its full duration. In addition, difficulty of recognizing mild symptoms or lack of symptoms may impact awareness of the infection and further limit adoption. Here, we study an epidemic model on a network of contacts accounting for limited adherence and delayed awareness to self-isolation, along with fatigue causing overhasty termination. The model allows us to estimate the role of each ingredient and analyze the tradeoff between adherence and duration of self-isolation. We find that the epidemic threshold is very sensitive to an effective compliance that combines the effects of imperfect adherence, delayed awareness and fatigue. If adherence improves for shorter quarantine periods, there exists an optimal duration of isolation, shorter than the infectious period. However, heterogeneities in the connectivity pattern, coupled to a reduced compliance for highly active individuals, may almost completely offset the effectiveness of self-isolation measures on the control of the epidemic.

I. INTRODUCTION

A pillar of non-pharmaceutical interventions for the control of COVID-19 pandemic is the isolation of individuals testing positive for SARS-CoV-2 infection. The aim is to avoid onward propagation of the disease, while contacts are traced to further break the chains of transmission [1]. This measure, however, is met with a set of challenges, as it has no immediate benefit for the index case, but a number of downsides. It often causes psychosocial distress [2], and it may have severe financial and social costs impacting daily life, if a structured support program is not in place.

Ideally, the measure should cover the entire duration of the infectivity period. In practice, isolation may start when a person is already infectious, typically at the onset of symptoms, or when alerted by a contact tracing investigation. Also, the length of the infectious period may be strongly variable across individuals [3, 4]. Additional factors may undermine the effectiveness of isolation. Mild symptoms or lack of symptoms may ruin the motivation to respect it, as physical conditions are not an impediment to carry out the daily routine. The legal enforcement of the measure may create tradeoffs discouraging individuals to self-declare as cases [5]. Survey data report that adherence is low [6–8]. Among the reported reasons for non-adherence are lower socioeconomic grade, psychological distress, inadequate information and long quarantine duration [2].

During COVID-19 pandemic, the duration of isolation has been one flexible component that authorities adapted from initial estimates of 14 days [9] to shorter periods to make the measure more bearable, at the first signs in summer 2020 showing the difficulty of implementation of the measure [10, 11]. Variable durations mark a tradeoff between a long enough period of isolation to prevent onward transmission, and a short enough period that is acceptable by the population. Some countries went as low as 5 to 7 days to increase adherence [12, 13], especially in countries where self-isolation was not legally compulsory. Further changes (extension to 10 days [14]) occurred later because of the circulation of the Alpha variant (B.1.1.7 lineage), showing the complexity of the biological and social aspects of setting this public health measure [15].

As gaps in any of these aspects may undermine the effectiveness of isolation in aiding epidemic control, here we study through mathematical modeling the role of delayed awareness in entering into isolation and fatigue inducing early release of the measure. Our model is a variation of the standard susceptible-infected-susceptible (SIS) compartmental model for infectious disease dynamics [16, 17], allowing for three additional compartments: an isolated (Q) compartment, an undecided (U) compartment, and a fatigued (F) compartment. Here we borrow the classical notation Q commonly used in compartmental models to define the isolation of infectious individuals, notably with the susceptible-infected-quarantined-susceptible (SIQS) model [18–23]. We do not consider the quarantine as the preventive isolation of suspect cases or of contacts of confirmed cases [24–26], and in the following we will use the terms quarantine and self-isolation as synonyms. The existence of (temporary) immunity against SARS-CoV-2 would suggest the consideration of a SIR-like dynamics. We prefer to consider a SIS-based modeling framework as the absence of an immune state allows us to keep analytical derivations simpler while still providing (a worse case scenario) intuition on the behavioral mechanisms related to the self-isolation measure. We expect that similar
results would be obtained for a SIR-like model.

The undecided compartment U corresponds to an inter-
mediate state, following infection, during which awareness
arises around the knowledge of being infected, involving
a delay before the decision to comply with isolation. This
state may also be interpreted as the time between infec-
tion and testing (thus corresponding to logistical delays
in accessing and performing a test, and obtaining the
test results), or to the time between infection and symp-
toms onset (thus corresponding to a pre-symptomatic
phase) [1, 27]. Another addition to the standard SIQS
model is the possibility that the individual exits self-
isoation before its full duration and while still infectious.
The compartmental model and transitions are fully ex-
plained in the next section.

We investigate the model on a networked population
with a mean-field approach, highlighting how the key pa-
rameters describing the epidemic dynamics – i.e. the epi-
demic threshold and the prevalence of infected individuals
in the endemic state – depend on the different durations
associated to these states. We then consider increasingly
more accurate mean-field types of approach, allowing to
analyze in detail how individual heterogeneities influence
collective properties of the system. We therefore rely on
proven effective analytical and numerical tools in order
to quantitatively uncover the role of various kinds of in-
perfections of self-isolation in the spread of a pathogen,
which can be of public health relevance to the control of
the currently ongoing pandemic.

II. EPIDEMIC MODEL WITH QUARANTINE,
DELAY AND FATIGUE

The model we consider is a modification of the usual SIS
dynamics [16], based on the existence of three additional
compartments, beyond the standard S (susceptible) and
I (infected) states.

The contact of a susceptible individual with an infec-
tious one leads to a transmission of the pathogen at rate
\(\beta\). The newly infected individual enters the state U (un-
decked) preceding the decision on whether to self-isolate
or not. The decision process is assumed to be Poissonian
with rate \(\mu_U\). After the decision, the individual enters
the quarantined state Q with probability \(p_Q\) (quarantine
probability). With complementary probability he/she
instead enters the infected state I. In the latter case the
individual behaves as in the standard infected state of
the SIS model. In the Q state instead the individual has
no contacts, and does not transmit the infection. Compli-
ance to isolation may however end before full recovery, as
fatigue sets in. We model this assuming that each quar-
tained individual transitions to compartment F (fatigued)
at rate \(\mu_F\). Individuals in state F are infectious with the
same transmissibility \(\beta\) of those in state I.

As the progression of the disease does not depend on
the isolation status, spontaneous recovery transitions occur
from states U, I, Q and F to the susceptible state S,
at same rate \(\mu\). As a consequence of this choice, the
average time spent before reaching the S compartment
from any of the infected states U, I, Q and F is \(1/\mu\),
also when there exist multiple paths to recovery. The
computation of the average time should indeed receive the
contributions of all the possible paths, each weighted by
the probability of being chosen: when multiple transitions
out of a compartment are possible, the average time of
each transition must be conditioned on the fact that the
other possible transitions were not undergone.

Our model overall depends on five independent param-
eters \(p_Q, \beta, \mu_U, \mu_Q\) and \(\mu\). The compartments and the
transitions allowed between them are depicted in Fig. 1.
All transitions are spontaneous except the one taking indi-
muals in state S to the undecided state U, which occurs
because of a contact between a susceptible individual and
an infectious one. Note that in this model individuals
in compartments I, F, U are all infectious with the same
transmissibility.

III. MEAN-FIELD APPROACH

Let us define as \(S(t), U(t), I(t), Q(t), F(t)\), the proba-
bilities for an individual to be in the respective compart-
ments. The sum of these probabilities equals 1, leaving
only 4 independent quantities. We assume a homoge-
neous pattern of interactions, with average number of
contacts \(k\). Then, the differential equations describing
the evolution of the aforementioned probabilities read as
follows:

\[
\begin{align*}
\dot{U} &= \beta \langle k \rangle (1 - I - U - F - Q)(I + U + F) - (\mu_U + \mu)U \\
\dot{I} &= \mu_U (1 - p_Q)U - \mu I \\
\dot{Q} &= \mu_UPQ - (\mu_Q + \mu)Q \\
\dot{F} &= \mu_QQ - \mu F
\end{align*}
\] (1)

The disease-free state \((S, U, I, Q, F) = (1, 0, 0, 0, 0)\) is
always an equilibrium solution of the system. Linearization
around it shows that it is not stable if \(\lambda = \beta/\mu\) is
above the critical value \( \lambda_c \), marking the existence of an endemic state:

\[
\lambda_c = \frac{1}{\langle k \rangle} \frac{1}{1 - p_Q \frac{p_Q}{1 + \frac{p_Q T_Q}{1 + \frac{p_Q}{1 + \frac{T_Q}{1 + \frac{T}{1 + \frac{T}{T}}}}}}.
\]

(2)

where the temporal scales \( T = 1/\mu \), \( T_Q = 1/\mu Q \) and \( T_U = 1/\mu U \) are the average times spent in the corresponding states. Eq. (2) contains several known results for limit values of its parameters. \( p_Q = 0 \) – individuals never in isolation – yields the well-known SIS Mean-field result \( \lambda_c = 1/\langle k \rangle \). The same limit is recovered if the quarantine has vanishing duration \( (T/T_Q \to \infty) \) or when the time to take a decision diverges \( (T_U/T \to \infty) \).

For generic values of the parameters Eq. (2) can be written as

\[
\lambda_c = \frac{1}{\langle k \rangle} \frac{1}{1 - p_Q^{eff}}
\]

(3)

where

\[
p_Q^{eff} = \frac{p_Q}{\left(1 + \frac{T_U}{T_Q}\right) \left(1 + \frac{T_Q}{1 + \frac{T}{T}}\right)}.
\]

(4)

The quantity \( p_Q^{eff} \) is a scaling law turning the effect of compartments \( U, F \) into an effective probability to self-isolate in a SIQS model. It is smaller than \( p_Q \) and reflects the reduction in the efficacy of the quarantine due to undecidedness and fatigue. Note that, even for full compliance with the quarantine prescription \( (p_Q = 1) \), any value \( T_Q < \infty \) or \( T_U > 0 \) is sufficient to make the threshold finite. For relatively large decision time (compared with recovery time) or small quarantine duration, the factor multiplying \( p_Q \) in Eq. (4) is small and therefore the increase of the epidemic threshold for a full quarantine probability \( (p_Q = 1) \) compared to none \( (p_Q = 0) \), might be very limited.

In the endemic state, the densities of individuals in the various compartments are given by

\[
\begin{align*}
U^* &= \frac{1}{1 + T/T_U} \frac{\lambda - \lambda_c}{\lambda_c} \\
I^* &= \frac{1}{1 - p_Q} \frac{1}{1 + T_U/T} \frac{\lambda - \lambda_c}{\lambda_c} \\
Q^* &= \frac{1}{1 + T_U/T} \frac{1}{1 + T_Q/T} \frac{\lambda - \lambda_c}{\lambda_c} \\
F^* &= \frac{1}{1 + T_U/T} \frac{1}{1 + T_Q/T} \frac{\lambda - \lambda_c}{\lambda_c}\end{align*}
\]

(5)

where the quantity \( \frac{\lambda - \lambda_c}{\lambda_c} \) is the total density of infected individuals \( 1 - S^* = U^* + I^* + Q^* + F^* \). The total density of infectious individuals instead is

\[
I_{tot}^* = I^* + U^* + F^* = \frac{1}{\langle k \rangle} \left( \frac{1}{\lambda_c} - \frac{1}{\lambda} \right).
\]

(6)

For any \( \lambda, I_{tot}^* \) depends on \( p_Q, T_Q \) and \( T_U \) only via the value of the epidemic threshold. Equation (6) then indicates that, for a given \( \lambda \), changing parameters in order to increase the epidemic threshold simultaneously reduces the overall prevalence of infectious individuals. Therefore maximizing the epidemic threshold, minimizing \( I_{tot}^* \) and maximizing \( S^*/\lambda \) are equivalent procedures.

In general, the two parameters \( p_Q \) and \( T_Q \) are likely to be dependent, as the perspective of long isolation may discourage people from isolating. We assume that the quarantine probability is a function \( p_Q(T_Q) \) of its duration. In particular we expect that \( p_Q \) decreases as \( T_Q \) increases. This leads to the existence of an optimal quarantine duration \( T_Q^* \). For small values of \( T_Q \) adherence to quarantine is high, but its duration is too short, so that people exit from it when they are still infectious. For large values of \( T_Q \) isolated individuals recover when in isolation, but compliance is low. An optimal tradeoff exists between these two limits. We want to find the optimal duration of self-isolation \( T_Q^* \) that maximizes pathogen circulation, i.e. it either minimizes \( I_{tot}^* \) or maximizes the epidemic threshold. This occurs when the denominator of the epidemic threshold (Eq. (2)) attains its minimum, i.e. for \( T_Q = T_Q^* \) such that

\[
\frac{dp_Q}{dT_Q} \bigg|_{T_Q = T_Q^*} = -\frac{p_Q(T_Q)}{T_Q^*} \frac{1}{1 + \frac{T_Q^*}{T}}.
\]

(7)

We make the relationship between \( p_Q \) and \( T_Q \) explicit, making the following minimal assumption:

\[
p_Q(T_Q) = \frac{1}{1 + c T_Q/T},
\]

(8)

where the parameter \( c \) determines how quickly the probability to enter quarantine decays with its duration. This way compliance tends to be perfect \( (p_Q \to 1) \) for extremely short quarantine, while virtually nobody decides to isolate \( (p_Q \to 0) \) if the duration of self-isolation is much longer than the average recovery time.

The optimal quarantine duration, solution of Eq. (7), is then

\[
\frac{T_Q^*}{T} = \frac{1}{\sqrt{c}},
\]

(9)

corresponding to an optimal probability to quarantine \( p_Q^* = 1/(1 + \sqrt{c}) \) and to the maximum threshold

\[
\lambda^*_c = \frac{1}{\langle k \rangle} \frac{1}{1 - \frac{1}{\left(1 + \sqrt{c}\right)^2 \left(1 + \frac{T}{T_Q}\right)}}.
\]

(10)

We find that for the optimal duration of quarantine \( T_Q^* \) the threshold is a decreasing function of the parameter \( c \) (see Fig. 2).

IV. HETEROGENEOUS MEAN FIELD APPROACH

We now allow for the more realistic assumptions of individuals to have heterogeneous contact rates. We start
A nontrivial solution $\Theta > 0$ only exists if the derivative with respect to $\Theta$ of the r.h.s. of Eq. (15) [where we substitute $U^*_k$ by its explicit dependence on $\Theta$ using Eq. (14)] evaluated for $\Theta = 0$, is larger than 1. This condition allows us to determine the epidemic threshold

$$\lambda_c = \frac{1}{1 - \frac{q^2 P_{\text{inf}}}{(1 + \frac{q}{\mu})(1 + \frac{q}{\mu})}}. \tag{16}$$

We observe that this threshold is simply the HMF threshold for SIS [28] modulated by a factor that takes into account quarantine probability, duration as well as delay. The effect of topology factorizes. For a homogeneous network Eq. (2) is recovered, since $\langle k^2 \rangle = \langle k \rangle^2$.

We perform numerical checks of these predictions, by simulating SIS dynamics (using a Gillespie optimized algorithm [32]) on networks built according to the uncorrelated configuration model [33]. In this model, we consider an upper cutoff on the degrees $k \in [k_{\text{min}} = 3, k_{\text{max}} = \sqrt{N}]$ – in order to have an uncorrelated network without multiple and self connections. The epidemic threshold is estimated by finding the value of $\lambda = \beta/\mu$ at which the susceptibility of the system reaches a maximum value [30]. Such susceptibility is computed for the network Eq. (2) is recovered, since $\langle k^2 \rangle = \langle k \rangle^2$.

At stationarity we have

$$\begin{cases}
0 = \beta(1 - I_k - U_k - Q_k - F_k)k\Theta - (\mu_U + \mu)U_k \\
0 = \mu_U(1 - pQ)U_k - \mu I_k \\
0 = \mu_U pQ U_k - (\mu Q + \mu) Q_k \\
0 = \mu Q Q_k - \mu F_k
\end{cases} \tag{13}$$

whose solution reads

$$\begin{align*}
U_k^* & = \frac{\lambda_c \Theta}{(1 + \mu_U/\mu)(1 + \lambda_c \Theta)} \\
I_k^* & = \frac{\mu_U}{(1 - pQ)}U_k^* \\
Q_k^* & = \frac{\mu_U pQ U_k^*}{\mu Q + \mu} \\
F_k^* & = \frac{\mu Q Q_k^*}{\mu Q + \mu}
\end{align*} \tag{14}$$

Inserting the stationary values into Eq. (12) we find

$$\Theta = \sum_{k'} \frac{k' P(k')}{\langle k \rangle^2} \left[ 1 - \frac{pQ}{(1 + \mu_U/\mu)(1 + \mu_U/\mu)} \right] (1 + \mu_U/\mu) U_k^* \tag{15}$$

A nontrivial solution $\Theta > 0$ only exists if the derivative with respect to $\Theta$ of the r.h.s. of Eq. (15) evaluated for $\Theta = 0$, is larger than 1. This condition allows us to determine the epidemic threshold

$$\lambda_c = \frac{1}{1 - \frac{q^2 P_{\text{inf}}}{(1 + \frac{q}{\mu})(1 + \frac{q}{\mu})}}. \tag{16}$$

We observe that this threshold is simply the HMF threshold for SIS [28] modulated by a factor that takes into account quarantine probability, duration as well as delay. The effect of topology factorizes. For a homogeneous network Eq. (2) is recovered, since $\langle k^2 \rangle = \langle k \rangle^2$.

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In the following, we consider networks with an exponent $\gamma = 2.25$ of the degree distribution and, unless otherwise specified, with a network size $N = 10^5$.

In Fig. 3 we plot the epidemic threshold as a function of the system size $N$ for several values of the probability $pQ$ to enter quarantine. We first note an excellent agreement between the theory (dashed lines) and the simulations

$$\Theta = \sum_{k'} \frac{k' P(k')}{\langle k \rangle} (I_{k'} + U_{k'} + F_{k'}). \tag{12}$$
Figure 3. Epidemic threshold $\lambda_c$ as a function of the system size $N$ for $\gamma = 2.25$ and several values of $p_Q$. Dashed lines represent the theoretical prediction, Eq (16), while symbols are the numerical estimates, obtained from the peak of the susceptibility [32]. $T = 1$, $T_U = 0.25$, $T_Q = 1$.

Figure 4. Epidemic threshold $\lambda_c$ as a function of $T_Q$ for $T = 1$, $T_U = 0.2$ and various $p_Q$. Symbols are the results of numerical simulations, dashed lines are the predictions of HMF theory, dotted lines are the predictions of QMF theory.

Figure 5. Epidemic threshold $\lambda_c$ as a function of $T_Q$ for $T = 1$, $p_Q = 0.7$ and various $T_U$. Dashed lines are the predictions of HMF theory and dotted lines are the predictions of QMF theory while symbols are the results of numerical simulations.

Figure 6. Density of individuals in the various compartments in the stationary state as a function of $\lambda$ for $T = 1$, $T_U = 0.2$, $T_Q = 0.33$ and $p_Q = 0.7$.

The effect is sizeable only provided $p_Q$ is quite large. In Fig. 5 we show the dependence of the threshold on the duration of quarantine for various delays $T_U$. Here too the threshold increases smoothly with $T_Q$. If the decision time is much smaller than the time to heal the effect becomes relevant also for reasonable values of $T_Q$.

Finally in Fig. 6 we report the dependence of the total density of individuals in the various compartments as a function of $\lambda$, showing a fair agreement between theory and numerics. As long as the parameters of the model are finite and strictly positive, each of these densities carries information on the global state of the epidemic, each of them simply being a fraction of the order parameter.
B. Degree-dependent epidemic parameters

Empirical evidence [34] suggests that people having more contacts or being more active tend to be more reluctant in reducing their interactions to prevent contagion. This may be due to the fact that each interrupted contact carries with it a substantial economical, social and/or psychological cost. Real data also suggest that individuals with high activity are more attractive, which may make it more difficult for them to self-isolate given the numerous solicitations they receive from others [23]. It is then quite natural to believe that also adherence to the prescription to self-isolate may be different (and in particular be suppressed) for people with a large number of contacts. In our framework it is possible to model such a realistic element by assuming that the probability to enter quarantine and/or its duration depend on the degree of the node, a proxy of individual activity. In particular it is reasonable to expect both \( p_Q \) and \( T_Q \) to decrease with \( k \).

By repeating the calculations already performed in the case with degree-independent parameters we easily find that the epidemic threshold reads

\[
\lambda_c = \frac{\langle k \rangle}{\left(1 - \frac{p_Q(k)}{\langle k \rangle P(k)} \left(1 + \frac{\lambda}{\langle k \rangle X(k)} \right)\right)^{1/2}},
\]

where \( \langle X(k) \rangle = \sum_k P(k) X(k) \). Hence, depending on how \( p_Q \) and \( T_Q \) behave for large \( k \), quarantine may reduce or not the vulnerability of scale-free networks to epidemics. We test this prediction again by performing simulations on networks built according to the uncorrelated configuration model. For reference, we compare with results obtained with degree-independent parameters tuned to have exactly the same average value of the degree-dependent case. We first check what happens assuming \( p_Q(k) = k_{\text{min}}/k \), so that compliance is perfect for nodes having minimal connectivity while it becomes very small for large \( k \). In Fig. 7 we report the behavior of the epidemic threshold as a function of \( T_Q \) for the degree-dependent case and for a degree independent case such that \( p_Q = \langle p_Q(k) \rangle \). It turns out that the threshold is smaller in the degree-dependent case and in particular that it grows much more slowly with \( T_Q \). The effect of a long self-isolation of less connected individuals is almost completely offset by the little compliance of nodes of large degree.

![Figure 7](image)

Figure 7. Epidemic threshold as a function of \( T_Q \) for \( T = 1, T_U = 0.25 \). In the heterogeneous case (circles and bottom lines) \( p_Q(k) = k_{\text{min}}/k \). In the homogeneous case (squares and top lines) \( p_Q \) is the same for all nodes: \( p_Q = \langle p_Q(k) \rangle = 0.66 \). Dashed lines are for the HMF predictions whereas dotted lines are for the QMF ones.

![Figure 8](image)

Figure 8. Epidemic threshold as a function of \( p_Q \) for \( T = 1, T_U = 0.25 \). In the heterogeneous case (circles and bottom lines) \( T_Q(k) = (T - T_U)(k_{\text{max}} - k)/(k_{\text{max}} - k_{\text{min}}) \). In the homogeneous case (squares and top lines) \( T_Q \) is the same for all nodes \( T_Q = \langle T_Q(k) \rangle = 0.71 \). Dashed lines are for the HMF predictions whereas dotted lines are for the QMF ones.

We then check what happens instead when \( T_Q(k) = (T - T_U)(k_{\text{max}} - k)/(k_{\text{max}} - k_{\text{min}}) \). Complying individuals with few contacts self-isolate until full recovery, whereas individuals with large connectivities spend a vanishing time in quarantine. This degree-dependent scenario is compared in Fig. 8 to the degree-independent one in such a way that in the latter \( T_Q = \langle T_Q(k) \rangle \). We find that the possibility for a few hubs to undergo shorter isolation periods than the average individual of the population lowers the epidemic threshold the more the larger the quarantine probability. The linear interpolation between the extreme behaviours of hubs and poorly-connected individuals seems to be slow enough not to completely offset the advantages put forward by the self-isolation prescription.
V. QUENCHED MEAN FIELD APPROACH

A. Epidemic parameters independent of $k$

A more refined approach to the model dynamics on networks is provided by the Quenched Mean Field (QMF) approximation [35-37] (also known as Individual Based mean-field approach), which takes into account the detailed structure of the network encoded in the adjacency matrix $A_{ij}$. Defining as $I_i(t), U_i(t), Q_i(t), F_i(t)$ the probabilities that node $i$ is in state I, U, Q and F respectively, the evolution of the model is described by the set of 4N equations

$$\begin{align*}
\dot{U}_i &= \lambda (1 - I_i - U_i - Q_i - F_i) \\
\dot{I}_i &= \mu_1 (1 - p_Q) U_i - \mu_1 I_i \\
\dot{Q}_i &= \mu_2 p_Q U_i - (\mu_1 + \mu_2) Q_i \\
\dot{F}_i &= \mu_2 Q_i - \mu_2 F_i
\end{align*}$$

(18)

By linearizing around the healthy state and imposing the largest eigenvalue of the Jacobian matrix to be equal to 0 one obtains the epidemic threshold

$$\lambda_c = 1 - \frac{1}{\rho(A)} \frac{1}{\rho(A)}$$

(19)

where $\rho(A)$ is the spectral radius of the adjacency matrix (i.e. its largest eigenvalue). This expression is perfectly analogous to the QMF result for standard SIS dynamics (corresponding to Eq. (19) for $p_Q = 0$), for which it is well known [16, 38] that the QMF threshold is a lower bound of the true threshold. We expect this to be true also for the present modification of the SIS model. Indeed Figs. 4 and 5 show that Eq. (19) is a tight lower bound, as it is the case for $\gamma = 2.25 < 5/2$. For larger values of $\gamma$ instead additional nontrivial effects [39] make the QMF estimate inaccurate for large networks. We note also that the HMF predictions are slightly closer than QMF to the numerical results. This better performance of HMF with respect to QMF (which occurs also for SIS [40]) is accidental: the additional approximation introduced by HMF partly cancels the error due to the QMF approximation. It is only for larger values of $\gamma > 3$ and larger system sizes that QMF reveals its more accurate qualitative behavior.

B. Degree-dependent epidemic parameters

If the parameter $p_Q$ depends on $k$ we obtain that the threshold is

$$\lambda_c = \frac{1}{\rho(A)}$$

(20)

The quantity $\tilde{A}$ is a modification of the adjacency matrix of the system

$$\tilde{A} = A \left(1 - \mathbb{I}_Q \frac{1}{\rho(A)} \frac{1}{\rho(A)} \right)$$

(21)

where $\mathbb{I}_Q = diag(p_Q(k(i)))$.

If instead the parameter $T_Q$ depends on $k$ we obtain that the threshold is

$$\lambda_c = \frac{1}{\rho(A)}$$

(22)

$\tilde{A}$ is now

$$\tilde{A} = A \left(1 - \mathbb{I}_Q \frac{1}{\rho(A)} \frac{1}{\rho(A)} \right)$$

(23)

where $\mathbb{I}_Q = diag(T_Q(k(i))))$.

VI. CONCLUSIONS

In this paper we have defined and used an epidemic model to study the role of delayed case detection/infection awareness, compliance to self-isolation, and fatigued-induced early drop-out on the effectiveness of self-isolation as a non-pharmaceutical intervention. We found that adherence to the prescription to self-isolate once infected scaled up the epidemic threshold compared to the simple SIS result, and delay into entering isolation or early release from it resulted in a reduction of effective adherence to self-isolation. If the propensity to enter self-isolation and the time spent isolated decrease with the individual number of contacts (degree), the low adherence of few well socially connected individuals may undermine the effectiveness of the entire non-pharmaceutical measure against the epidemic. This is a key result as it suggests that these phenomena, empirically found [6, 41], may strongly limit the impact of isolation programs on the pandemic, unless specific measures are implemented to overcome these barriers [6].

The applicability of this model to real case scenarios would take advantage from being informed by real data for both the parameters describing the COVID-19 dynamics and behavioral parameters. Whereas the former can be obtained by fitting case incidence time-series, the latter rely on a complex interaction between top-down regulations and behavioral adaptations and are hence harder to be inferred from data. The behavioral parameters are therefore explored rather than fit to data. We plan to include data from surveys in a future development of this study. Also additional ingredients must be considered in the model to increase the applicability of the model to more realistic scenarios. First, a more detailed compartmental structure accounting for the different phases of COVID-19 disease progression, to better
account for the interplay of different time periods and include asymptomatic and paucisymptomatic states. These may also result in different behaviors, reducing adherence and increasing early drop-outs, compared to symptomatic cases. Our simplified approach has, however, the advantage of being analytically tractable, therefore providing an immediate solution under certain approximation and offering an intuition into the behavior of the system. In this perspective, the SIS model was preferred to a model offering an intuition into the behavior of the system. In this context, we expect that this approximation may impact the recovery process from the state F differently than from the state I.

Further directions can be considered to expand this approach in future work. Here, we assumed that isolation prevents all contacts. In reality, isolation is never 100% effective, due both to behavioral aspects, and hard living constraints (e.g., household crowding [44]). Different degrees of imperfect isolation can be considered in terms of approximations altering the contact pattern. We did not consider in this study the role of quarantine as preventive isolation of suspect cases or contacts of confirmed cases. There is now a large body of literature on the role of contact tracing in combination to isolation and testing in COVID-19 control [15], and the importance of speeding up this process through digital tools [24]. Beside contact tracing, the introduction of a compartment describing individuals uncertain about their infection status, but still with recommended self-isolation, constitutes an additional component in limiting adherence, as motivation to self-isolate is reduced in absence of symptoms or of a test result confirmation. These processes are likely to be governed by different parameters of quarantine probability and duration. Finally, adherence to self-isolation may be the result of an individual component, explored here, along with a population component defined by a level of awareness and of risk perception that may evolve over time [45], depending on the evolving epidemic context. This may be an important component contributing to the observed relaxation effects after COVID-19 pandemic wave, possibly resulting in case resurgences [46–48].


