The role of quarantine for containing the epidemic

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My work is concerned to study a degenerated susceptible, infected, quarantine, recovered (SIQR) epidemic model with spatial heterogeneity, and generalized nonlinear incidence functional. The problem associated to this work is:

$$\frac{\partial S}{\partial t} = d_1 \Delta S + \Lambda(x) - f(x, S, I) - \mu(x)S, \qquad t > 0, \ x \in \Omega,$$

$$\frac{\partial I}{\partial t} = d_2 \Delta I + f(x, S, I) - (\mu(x) + \kappa_1(x) + \theta(x))I + \gamma(x)Q, \quad t > 0, \ x \in \Omega,$$

$$\frac{\partial Q}{\partial t} = \theta(x)I - (\mu(x) + \kappa_2(x) + \gamma(x))Q, \qquad t > 0, \ x \in \Omega,$$

$$\frac{\partial R}{\partial t} = d_3 \Delta R + \kappa_1(x)I + \kappa_2(x)Q - \mu(x)Q, \qquad t > 0, \ x \in \Omega,$$

$$\frac{\partial S}{\partial \eta} = 0, \frac{\partial I}{\partial \eta} = 0, \qquad x \in \partial\Omega.$$
(1)

By the generalized Krein-Ruthman theorem we identify the basic reproduction number R_0 , with it threshold role. For $R_0 \leq 1$, the disease will die out, which is guaranteed by the global asymptotic stability of the disease-free steady state, and for $R_0 > 1$, the disease will persist.

Lemma 1. (i) $\mathcal{R}_0 - 1$ has the same sign as s(A), where s(A) is the spectral bound of A.(to obtain this we use Theorem 3.5, [1])

(ii) According to [2] we can prove that \mathcal{R}_0 is defined as

$$\mathcal{R}_{0} = \sup_{\phi \in H^{1}(\Omega), \phi \neq 0} \frac{\int_{\Omega} \frac{\partial f}{\partial I}(., \hat{S}, 0)\phi^{2} dx}{\int_{\Omega} \left(d_{2} |\nabla \phi|^{2} + (\mu + \kappa_{1} + \theta \frac{\mu + \kappa_{2}}{\mu + \kappa_{2} + \gamma})\phi^{2} \right) dx}$$
(2)

(iii) $\mathcal{R}_0 - 1$ and s(A) have the same sign as η^0 , where η^0 is the principal eigenvalue of

$$\begin{cases}
d_2 \Delta \phi - (\mu + \kappa_1 + \theta \frac{\mu + \kappa_2}{\mu + \kappa_2 + \gamma})\phi + \frac{\partial f}{\partial I}(., \hat{S}, 0)\phi = \eta^0 \phi \quad x \in \Omega, \\
\frac{\partial \phi}{\partial \eta} = 0 \qquad \qquad x \in \partial\Omega,
\end{cases}$$
(3)

Theorem 1. Suppose that $R_0 \leq 1$. The disease free steady state is globally asymptotically stable

Theorem 2. Assume that $\mathcal{R}_0 > 1$, then for any initial data $u_0 = (S_0, I_0, Q_0) \in \mathbb{X}^+ := C(\bar{\Omega}, \mathbb{R}^3_+)$ with $I_0 \neq 0$, or $Q_0 \neq 0$, then there exists $\delta > 0$ such that the solution u = (S, I, Q) of (1) satisfies

$$\min\left\{\liminf_{t\to\infty} S(.,t), \ \liminf_{t\to\infty} I(.,t), \ \liminf_{t\to\infty} Q(.,t)\right\} \ge \delta, \ uniformly \ for \ all \ x \in \bar{\Omega}.$$
(4)

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The main question that can be put in our study (due to the fact that the individuals in quarantine can repulse into the infected stage, and start infecting again): Is the quarantine effective in this case? To understand the role of quarantine better we compared our problem with SIR model. Numerical simulations are used to validate the theoretical finding.

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