

Compartmental epidemic models with nonlinear recovery

DE, DC, JD

July 14, 2016 @ 12:57

Contents

1	The model	1
2	Equilibria	2
2.1	Disease free equilibria (DFE)	2
2.2	Endemic equilibria (EE)	2
3	Infectious period distribution at EE	3
4	Infectious period distribution near DFE (invasion)	3
5	Infectious period distribution at any time	3

1 The model

Assume the recovery rate, $\mathcal{K}(I)$, is a smooth, non-negative function on $[0, 1]$.

$$\frac{dS}{d\tau} = \varepsilon(1 - S) - \rho SI \quad (1a) \quad \{\mathbf{E}; \mathbf{SIR}; \mathbf{S}\}$$

$$\frac{dI}{d\tau} = \rho SI - \mathcal{K}(I)I \quad (1b) \quad \{\mathbf{E}; \mathbf{SIR}; \mathbf{I}\}$$

$$\frac{dR}{d\tau} = I - \varepsilon(1 - S) \quad (1c) \quad \{\mathbf{E}; \mathbf{SIR}; \mathbf{R}\}$$

where $S + I + R = 1$. In terms of the usual SIR parameters, the dimensionless parameters

16 above are

$$17 \quad \varepsilon = \frac{\mu}{\gamma + \mu}, \quad (2a) \quad \{\mathbf{E:eps}\}$$

$$18 \quad \rho = \frac{\beta N}{\gamma + \mu}. \quad (2b) \quad \{\mathbf{E:rho}\}$$

20 If $\mathcal{K}(I) \equiv 1$ then ε is the mean time in the infectious class as a fraction of the of the mean
 21 lifetime and ρ is \mathcal{R}_0 . Note that $0 \leq \varepsilon < 1$ necessarily. I will write initial conditions as
 22 (S_i, I_i, R_i) .

23 **2 Equilibria**

24 There are typically two equilibria:

25 **2.1 Disease free equilibria (DFE)**

26 As for the standard KM SIR model:

27 Unique if $\varepsilon > 0$: $(S, I) = (1, 0)$.

28 Continuum of DFEs if $\varepsilon = 0$: $(S_0, 0)$ is a DFE for any $S_0 \in [0, 1]$.

29 **2.2 Endemic equilibria (EE)**

30 $(S, I) = (\hat{S}, \hat{I})$.

$$31 \quad 0 = \varepsilon(1 - \hat{S}) - \rho\hat{S}\hat{I} \quad (3) \quad \{\mathbf{E:}\}$$

$$32 \quad 0 = \rho\hat{S} - \mathcal{K}(\hat{I}) \quad (4)$$

34 Hence

$$35 \quad \hat{S} = \frac{\mathcal{K}(\hat{I})}{\rho} \quad (5) \quad \{\mathbf{E:Shat}\}$$

$$36 \quad \mathcal{K}(\hat{I}) = \frac{\varepsilon}{\hat{I} + (\varepsilon/\rho)} \quad (6)$$

38 To solve for \hat{I} we need a specific form for $\mathcal{K}(I)$. In principle there can be multiple EEs.

39 For the standard KM SIR model,

$$40 \quad \hat{I} = \varepsilon \left(1 - \frac{1}{\rho}\right), \quad \mathcal{K}(I) \equiv 1. \quad (7) \quad \{\mathbf{E:IhatKM}\}$$

41 If the recovery rate is density-dependent, it is most natural to assume $\mathcal{K}(I)$ is decreasing,
 42 since this means that higher prevalence reduces recovery rates and hence lengthens infectious

43 periods. Here are some examples:

$$44 \quad \hat{I} = \begin{cases} \frac{\varepsilon}{\rho} \cdot \frac{a\rho-1}{1-b\varepsilon} & \mathcal{K}(I) = \frac{1}{a+bI}, \quad \rho > \frac{1}{a}, \quad \varepsilon < \frac{1}{b} \\ \frac{a}{2} - \frac{\varepsilon}{2\rho} - \sqrt{\left(\frac{a}{2} - \frac{\varepsilon}{2\rho}\right)^2 - \varepsilon\left(1 - \frac{a}{\rho}\right)} & \mathcal{K}(I) = a - I, \quad a \geq 1 \\ -\frac{1}{b}W(-b\varepsilon e^{-b\varepsilon/\rho}) - \frac{\varepsilon}{\rho} & \mathcal{K}(I) = e^{-bI}, \quad b > 0 \end{cases} \quad (8) \quad \{\mathbf{E:Ihat}\}$$

45 Here, W is Lambert's W function. For the above cases, **Figure 1** shows the dependence of
46 the EE on ρ for $\varepsilon = 0.1$.

47 The (singular) recovery function $\mathcal{K}(I) = I^{-1/2}$ has been used a fair bit in the literature
48 even though it blows up at $I = 0$, so I'm mentioning it. This has two EEs if $\rho > 4/\varepsilon$ and
49 none if $\rho < 4/\varepsilon$:

$$50 \quad \hat{I} = \frac{\varepsilon}{2\rho} \left(\rho\varepsilon - 2 \pm \sqrt{\rho\varepsilon(\rho\varepsilon - 4)} \right), \quad \mathcal{K}(I) = I^{-1/2}, \quad \rho > 4/\varepsilon \quad (9) \quad \{\mathbf{E:Ihat-1/2}\}$$

51 This is quite peculiar and surely of mathematical interest only.

52 In case it is somehow helpful, here's an example with $\mathcal{K}(I)$ increasing:

$$53 \quad \hat{I} = \varepsilon \left(\sqrt{\frac{1}{\varepsilon} + \frac{1}{4\rho^2}} - 1 \right), \quad \mathcal{K}(I) = I \quad (10) \quad \{\mathbf{E:Ihatinc}\}$$

54 **3 Infectious period distribution at EE**

55 At the EE, prevalence is constant, so there is no change in the recovery rate over time.
56 Consequently, individuals recover according to

$$57 \quad \frac{dI}{d\tau} = -\mathcal{K}(\hat{I}) I, \quad (11) \quad \{\mathbf{E:}\}$$

58 *i.e.*, the infectious period distribution is exponential with mean $1/\mathcal{K}(\hat{I})$ rather than mean 1.

59 **4 Infectious period distribution near DFE (invasion)**

60 In this limit, we can use the linearized system to obtain an approximation $I_{\text{lin}}(\tau)$ initially.
61 Then, for as long as this approximation is valid, the recovery rate is $\mathcal{K}(I_{\text{lin}}(\tau))$, so we can
62 approximate the initial distribution of infectious by solving

$$63 \quad \frac{dI}{d\tau} = -\mathcal{K}(I_{\text{lin}}(\tau)) I, \quad (12) \quad \{\mathbf{E:invlim}\}$$

64 **5 Infectious period distribution at any time**

65 More generally, between the limits of invasion and endemic equilibrium, we could plug in
66 the exact numerically computed $I(\tau)$ rather than $I_{\text{lin}}(\tau)$ into Equation (12) and compute the
67 distribution of infectious periods at any time. However, this is a pain, not just because it is
68 numerical but because we really need to integrate to $\tau = \infty$. Still, this is an alternative to
69 agent-based simulations.

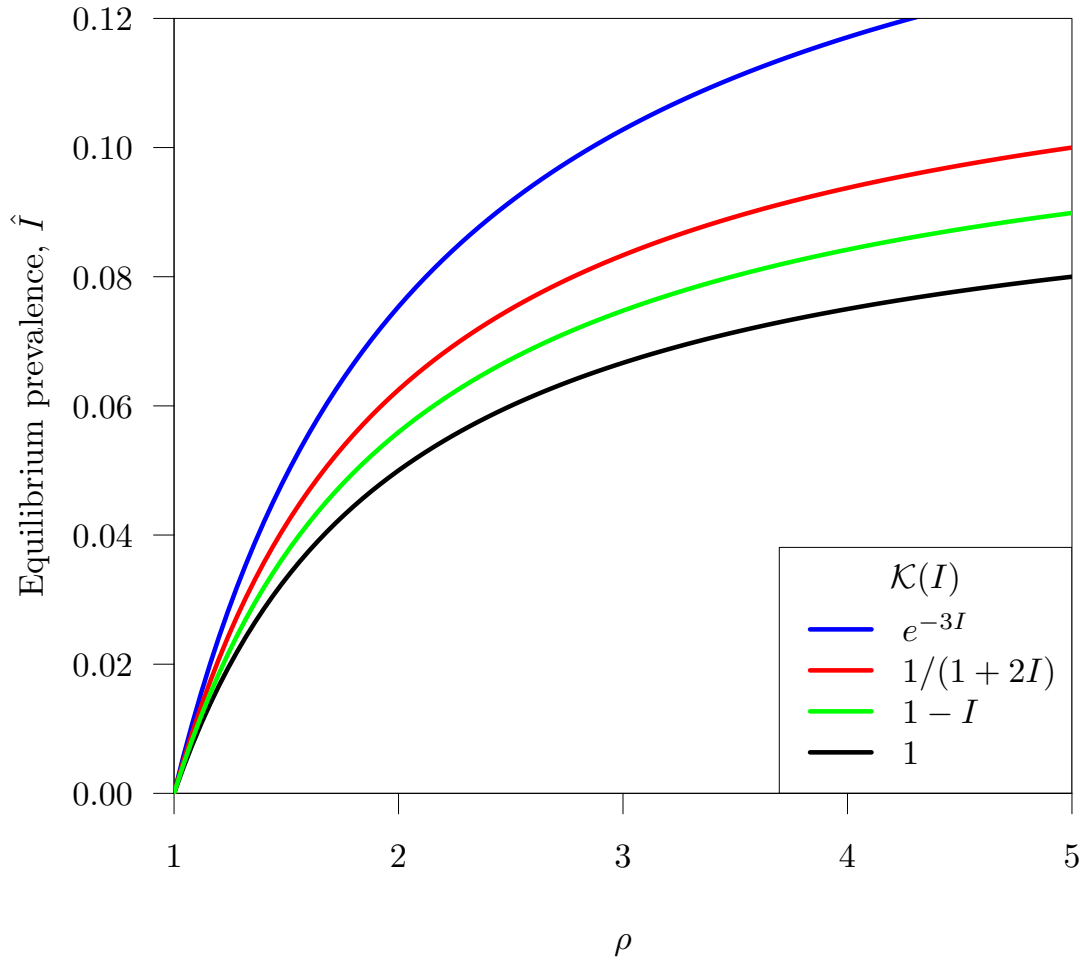


Figure 1: Equilibrium prevalence \hat{I} as a function of ρ for several recovery functions $\mathcal{K}(I)$, with $\varepsilon = 0.1$.

