Estimating $\mathcal{R}(t)$ from Covid-19 case counts

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Abstract

Here we attempt to give an interpretation of the effective reproduction number, \mathcal{R}_t that arises from the Susceptible-Infected-Recovered (SIR) model for the spread of infectious disease. We also investigate how \mathcal{R}_t may be estimated from daily case count data.

1 Introduction

The Susceptible-Infected-Recovered (SIR) model [3] is often used to study the spread of infectious disease by tracking the number (S) of people susceptible to the disease, the number (I) of people infected with the disease, and the number (R) of people who have had the disease and are now no longer infectious.

Based on the model, the only way that a person can leave the susceptible group is to become infected, and the only way that a person can leave the infected group is to recover or die. It is further assumed that those who have recovered or died from the disease are no longer susceptible. It is also assumed that all those who have not had the disease are equally susceptible and that the probability of their contracting the disease at time t is proportional to the product of I(t) and S(t).

These assumptions lead us to a set of three ordinary differential equations for S(t), I(t), and R(t):

$$\frac{dS}{dt} = -\gamma \mathcal{R}_t I(t) \tag{1}$$

$$\frac{dI}{dt} = \gamma \mathcal{R}_t I(t) - \gamma I(t) \tag{2}$$

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$$\frac{dR}{dt} = \gamma I(t). \tag{3}$$

Here $\gamma \geq 0$ is called the **recovery rate**. It is the reciprocal of the length of the infectious period which in the case of Covid-19 is estimated to be about one week, so $\gamma = \frac{1}{7}$.

 $\gamma \mathcal{R}_t \geq 0$ measures the likelihood of transmitting the disease when an infected and a susceptible come in contact. Hence \mathcal{R}_t is the number of transmissions caused by an infected person during his infectious period.

sectionBehavior of Groups over Time From equations (1-3) one can analyze equilibria - where dS/dt = dI/dt = dR/dt = 0 - using techniques in [2, Sec. 9.2]. Equilibria occur when I=0. To see if they are stable we will linearize the equations about $(I_*, S_*) = (0, S_*)$. [Note that equation (3) is really equivalent to R(t) = N - I(t) - S(t). Since R is not involved in the other two equations (1-2), we can ignore it for this analysis.] Writing $S(t) = S_* + u(t)$ and $I(t) = I_* + v(t) = v(t)$, we see that u and v satisfy dS/dt = du/dt = f(u, v) and dI/dt = dv/dt = g(u, v), where $f(u,v) = -\beta(S_* + u)v$ and $g(u,v) = [\beta(S_* + u) - k]v$. Expanding f and g in Taylor series about (0,0) and dropping terms involving u^2 , uv, v^2 , etc., we can write

$$f(u,v) \approx f(0,0) + \frac{\partial f}{\partial u}(0,0) \cdot u + \frac{\partial f}{\partial v}(0,0) \cdot v = -\beta S_* v$$

$$g(u,v) \approx g(0,0) + \frac{\partial g}{\partial u}(0,0) \cdot u + \frac{\partial g}{\partial v}(0,0) \cdot v = (\beta S_* - k)v.$$

This leads to the linearized equations for u and v:

$$\frac{du}{dt} = -\beta S_* v \tag{4}$$

$$\frac{du}{dt} = -\beta S_* v$$

$$\frac{dv}{dt} = (\beta S_* - k) v$$
(5)

It follows from (5) that if v is a small positive number, then if $\beta S_* - k > 0$ then dv/dt is positive and hence v will increase, moving further away from its equilibrium value of 0. Then we have an unstable equilibrium. On the other hand, if $\beta S_* - k < 0$, then dv/dt is negative and so v will decrease back towards its equilibrium value of 0. Also, du/dt is negative and hence moves back towards its equilibrium value of 0 provided u > 0; if u < 0, however, then it becomes more negative. In the notation of [2, pp. 164-165], we have $p = \operatorname{trace}(A) = \beta S_* - k$ and $q = \det(A) = 0$, so this lies on the borderline between a "stable node" and a "saddle point".

The results of our analysis are inconclusive, but it seems that the quantity $\beta S(0) - k$ may play a critical role in the behavior of the system, depending on whether it is positive or negative. Further investigation of the literature [1] shows that the quantity $(\beta/k)S(0)$ is called the "basic reproductive ratio" and the behavior of the system is different according to whether it is greater than 1 or less than 1.

To see what happens for different values of $(\beta/k)S(0)$, one can solve equations (1-3) numerically using Matlab routine ode45, for example. Starting with 10 infected people in a population of 1000, we set k=0.1 and solved these equations for four different values of β : $\beta=0.005$ so that $(\beta/k)S(0)=49.5$; $\beta=0.0005$ so that $(\beta/k)S(0)=4.95$; $\beta=0.0002$ so that $(\beta/k)S(0)=1.98$; and $\beta=0.00005$ so that $(\beta/k)S(0)=0.495$. The results are shown in Figure 1.

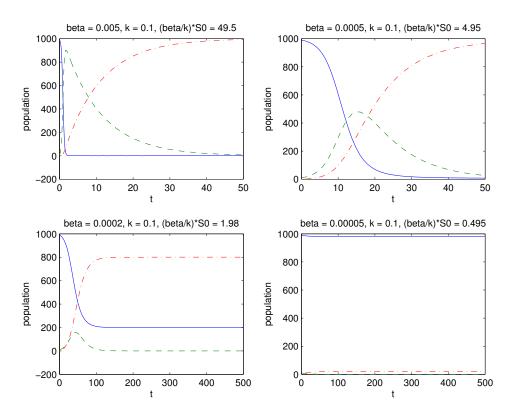


Figure 1: Number of Susceptibles (solid), Infected (dashed), and Recovered/Dead (dash-dot) for different basic reproductive ratios $(\beta/k)S(0)$.

From the figure, one can see that if $(\beta/k)S(0)$ is much greater than 1, then the entire population quickly becomes infected and then recovered/dead. For $(\beta/k)S(0)$ less than 1, the infection seems to die out after

only a small percentage of the population has been infected. For values somewhat greater than 1, the number infected and hence later recovered/dead seems to be a larger fraction but not necessarily all of the population. The Matlab code used to produce these plots can be found in Appendix A.

2 Conclusions

The results of this simple model look fairly reasonable and so they might be used to decide on a strategy for curtailing an epidemic through vaccinations, quarantines, etc. Many enhancements to the model can be made. For a discussion of some of these, see, for example, [3].

A Matlab Code for Producing Plots

```
global betaglob % Global variables to be supplied to function sirdot.
global kglob
                 % Give them funny names so they won't be used elsewhere.
kglob = 0.1;
                           % Set k.
betas = [.005; .0005; .0002; .0001];
for kase=1:4,
  betaglob = betas(kase); % Set beta.
  tfinal = 50; if kase > 2, tfinal = 500; end;
  [tout,sout] = ode45('sirdot',[0 tfinal], [990; 10; 0]); % Solve with ode45.
  subplot(2,2,kase)
  plot(tout,sout(:,1),'-', tout,sout(:,2),'--', tout,sout(:,3),'-.');
  xlabel('t'), ylabel('population')
  if kase==1,
    title('beta = 0.005, k = 0.1, (beta/k)*S0 = 49.5');
  elseif kase==2,
    title('beta = 0.0005, k = 0.1, (beta/k)*S0 = 4.95');
  elseif kase==3,
    title('beta = 0.0002, k = 0.1, (beta/k)*S0 = 1.98');
    title('beta = 0.00005, k = 0.1, (beta/k)*S0 = 0.495');
  end;
end;
function sirprime = sirdot(t,sir)
global betaglob
global kglob
beta = betaglob; k = kglob;
sirprime(1,1) = -beta*sir(1)*sir(2);
sirprime(2,1) = beta*sir(1)*sir(2) - k*sir(2);
sirprime(3,1) = k*sir(2);
```

References

- [1] Johnson, Teri, MathematicalModelingofDiseases: $Susceptible ext{-}Infected ext{-}Recovered$ (SIR) Model,Math 4901 of Minnesota, 2009. Seminar, University Spring http://www.morris.umn.edu/academic/math/Ma4901/Sp09/Final/Teri-Johnson-Final.pdf
- [2] Tung, K. K., *Topics in Mathematical Modeling*, Princeton University Press, 2007.
- [3] Wikipedia, Compartmental models in epidemiology, http://en.wikipedia.org/wiki/Compartmental_models_in_epidemiology

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