

The understanding of the virus is evolving and several models have been proposed to find the basic reproducibility number (denoted by R_0), case fatality rate and case recovery rate. In this study, we use epidemic transmission dynamics based compartmental model called *SIRD model* to study the progression of the disease for different countries (in particular, Germany and India). We use the discrete approximations as in [1] to estimate the R_0 , case mortality and case recovery rates and analyse the changes as time progresses. Furthermore, we fit the model to the epidemic curves of these countries to make forecasts about the peak of the epidemic and the flattening of the curve.

```
confirmed <- read.csv("data/time_series_covid19_confirmed_global.csv")

deaths <- read.csv("data/time_series_covid19_deaths_global.csv")

recovered <- read.csv("data/time_series_covid19_recovered_global.csv")

head(confirmed, 5)

## Province.State Country.Region Lat Long X1.22.20 X1.23.20 X1.24.20
## 1 Afghanistan 33.0000 65.0000 0 0 0
## 2 Albania 41.1533 20.1683 0 0 0
## 3 Algeria 28.0339 1.6596 0 0 0
## 4 Andorra 42.5063 1.5218 0 0 0
## 5 Angola -11.2027 17.8739 0 0 0
## X1.25.20 X1.26.20 X1.27.20 X1.28.20 X1.29.20 X1.30.20 X1.31.20 X2.1.20
## 1 0 0 0 0 0 0 0 0
## 2 0 0 0 0 0 0 0 0
## 3 0 0 0 0 0 0 0 0
## 4 0 0 0 0 0 0 0 0
## 5 0 0 0 0 0 0 0 0
## X2.2.20 X2.3.20 X2.4.20 X2.5.20 X2.6.20 X2.7.20 X2.8.20 X2.9.20 X2.10.20
## 1 0 0 0 0 0 0 0 0
## 2 0 0 0 0 0 0 0 0
## 3 0 0 0 0 0 0 0 0
## 4 0 0 0 0 0 0 0 0
## 5 0 0 0 0 0 0 0 0
## X2.11.20 X2.12.20 X2.13.20 X2.14.20 X2.15.20 X2.16.20 X2.17.20 X2.18.20
## 1 0 0 0 0 0 0 0
## 2 0 0 0 0 0 0 0
## 3 0 0 0 0 0 0 0
## 4 0 0 0 0 0 0 0
## 5 0 0 0 0 0 0 0
## X2.19.20 X2.20.20 X2.21.20 X2.22.20 X2.23.20 X2.24.20 X2.25.20 X2.26.20
## 1 0 0 0 0 0 1 1 1
## 2 0 0 0 0 0 0 0 0
## 3 0 0 0 0 0 0 1 1
## 4 0 0 0 0 0 0 0 0
## 5 0 0 0 0 0 0 0 0
## X2.27.20 X2.28.20 X2.29.20 X3.1.20 X3.2.20 X3.3.20 X3.4.20 X3.5.20 X3.6.20
## 1 1 1 1 1 1 1 1 1
## 2 0 0 0 0 0 0 0 0 0
## 3 1 1 1 1 3 5 12 12 17
## 4 0 0 0 0 1 1 1 1 1
## 5 0 0 0 0 0 0 0 0 0
## X3.7.20 X3.8.20 X3.9.20 X3.10.20 X3.11.20 X3.12.20 X3.13.20 X3.14.20 X3.15.20
## 1 1 4 4 5 7 7 7 11 16
## 2 0 0 2 10 12 23 33 38 42
## 3 17 19 20 20 20 24 26 37 48
## 4 1 1 1 1 1 1 1 1 1
```

## 4	1	1	1	1	1	1	1	1	1
## 5	0	0	0	0	0	0	0	0	0
##	X3.16.20	X3.17.20	X3.18.20	X3.19.20	X3.20.20	X3.21.20	X3.22.20	X3.23.20	
## 1	21	22	22	22	24	24	40	40	
## 2	51	55	59	64	70	76	89	104	
## 3	54	60	74	87	90	139	201	230	
## 4	2	39	39	53	75	88	113	133	
## 5	0	0	0	0	1	2	2	3	
##	X3.24.20	X3.25.20	X3.26.20	X3.27.20	X3.28.20	X3.29.20	X3.30.20	X3.31.20	
## 1	74	84	94	110	110	120	170	174	
## 2	123	146	174	186	197	212	223	243	
## 3	264	302	367	409	454	511	584	716	
## 4	164	188	224	267	308	334	370	376	
## 5	3	3	4	4	5	7	7	7	
##	X4.1.20	X4.2.20	X4.3.20	X4.4.20	X4.5.20	X4.6.20	X4.7.20	X4.8.20	X4.9.20
## 1	237	273	281	299	349	367	423	444	484
## 2	259	277	304	333	361	377	383	400	409
## 3	847	986	1171	1251	1320	1423	1468	1572	1666
## 4	390	428	439	466	501	525	545	564	583
## 5	8	8	8	10	14	16	17	19	19
##	X4.10.20	X4.11.20	X4.12.20	X4.13.20	X4.14.20	X4.15.20	X4.16.20	X4.17.20	
## 1	521	555	607	665	714	784	840	906	
## 2	416	433	446	467	475	494	518	539	
## 3	1761	1825	1914	1983	2070	2160	2268	2418	
## 4	601	601	638	646	659	673	673	696	
## 5	19	19	19	19	19	19	19	19	
##	X4.18.20	X4.19.20	X4.20.20	X4.21.20	X4.22.20	X4.23.20	X4.24.20	X4.25.20	
## 1	933	996	1026	1092	1176	1279	1351	1463	
## 2	548	562	584	609	634	663	678	712	
## 3	2534	2629	2718	2811	2910	3007	3127	3256	
## 4	704	713	717	717	723	723	731	738	
## 5	24	24	24	24	25	25	25	25	
##	X4.26.20	X4.27.20	X4.28.20	X4.29.20	X4.30.20	X5.1.20	X5.2.20	X5.3.20	X5.4.20
## 1	1531	1703	1828	1939	2171	2335	2469	2704	2894
## 2	726	736	750	766	773	782	789	795	803
## 3	3382	3517	3649	3848	4006	4154	4295	4474	4648
## 4	738	743	743	743	745	745	747	748	750
## 5	26	27	27	27	27	30	35	35	35
##	X5.5.20	X5.6.20	X5.7.20	X5.8.20	X5.9.20				
## 1	3224	3392	3563	3778	4033				
## 2	820	832	842	850	856				
## 3	4838	4997	5182	5369	5558				
## 4	751	751	752	752	754				
## 5	36	36	36	43	43				

We can see the data starts from January 22, 2020 and gives us the number of confirmed cases on each date since then till now.

Below we plot the progression of the disease (confirmed, recovered, deaths) in a few selected countries.

□□□

We can see from the plots that South Korea has successfully flattened the curve and countries like Germany and Italy have started flattening. On the other hand, India and Singapore are still far from flattening the curve.

3. Formulation of SIRD model

In this section, we discuss the basics of the SIRD model to simulate epidemic progression with time.

3.1 Discrete model

Within this model of the evolution of an epidemic outbreak, people can be divided into different classes. In the susceptible (S), infected (I), recovered (R), dead (D) scheme (SIRD), any individual in the fraction of the overall population that will eventually get sick belongs to one of the aforementioned classes. Let N be the size of the initial population of susceptible people. The discrete SIRD model can be written as follows:

$$\begin{aligned} S(t) - S(t-1) &= -\frac{\alpha}{N} S(t-1)I(t-1), \quad I(t) - I(t-1) = \frac{\alpha}{N} S(t-1)I(t-1) - \beta I(t-1) - \gamma I(t-1), \quad R(t) - R(t-1) = \beta I(t-1), \quad D(t) - D(t-1) = \gamma I(t-1), \end{aligned}$$

The basic reproduction number R_0 is then defined as
$$R_0 := \frac{\alpha}{\beta + \gamma}.$$

Since the number of susceptible people is hard to determine and depends on the population, lockdown measures, social distancing etc, we take a different approach

to estimate R_0 as mentioned in Ref 1. Let us denote $\Delta^* X^*(t) := X^*(t) - X^*(t-1)$ for $X^* = I^*, R^*, D^*$. Now we define,
$$\sum_{t=1}^T \Delta X(t) \text{ \textit{, and} } \mathbf{C \Delta X}(T) := [C \Delta X(1), C \Delta X(2), \dots, C \Delta X(T)]^T. \end{aligned}$$

Here C stands for cumulative. Using the approximation $S(t-1) \approx N$ (true if susceptible population is much less than the population of the country), we can get
$$R_{-0} = \frac{\alpha}{\beta + \gamma} = \frac{I(t) - I(t-1) + R(t) - R(t-1) + D(t) - D(t-1)}{R(t) - R(t-1) + D(t) - D(t-1)}. \end{aligned}$$
 Summing this equation over time we get,
$$\frac{C \Delta I(t) + C \Delta R(t) + C \Delta D(t)}{C \Delta R(t) + C \Delta D(t)} = R_{-0}. \end{aligned}$$
 Based on this, we can get a coarse estimate for R_0 by finding a least squares solution to the following regression problem:

$$\mathbf{C \Delta I(t) + C \Delta R(t) + C \Delta D(t)} = \mathbf{C \Delta R(t) + C \Delta D(t)} R_{-0}, \end{aligned}$$
 with solution given by
$$\hat{R}_{-0} = \frac{(\mathbf{C \Delta R(t) + C \Delta D(t)})^T \mathbf{C \Delta I(t) + C \Delta R(t) + C \Delta D(t)}}{(\mathbf{C \Delta R(t) + C \Delta D(t)})^T \mathbf{C \Delta R(t) + C \Delta D(t)}}. \end{aligned}$$

Similarly, the case fatality rate ($\hat{\beta}$) and case recovery rate ($\hat{\gamma}$) can be estimated as:
$$\hat{\beta} = \frac{(\mathbf{C \Delta I(t)})^T \mathbf{C \Delta I(t)}}{(\mathbf{C \Delta I(t)})^T \mathbf{C \Delta R(t)}}, \quad \hat{\gamma} = \frac{(\mathbf{C \Delta I(t)})^T \mathbf{C \Delta D(t)}}{(\mathbf{C \Delta I(t)})^T \mathbf{C \Delta D(t)}}. \end{aligned}$$

3.2. Continuous model

In the continuous, the number of people in each class is a function of continuous time. So $S(t)$ denotes the susceptible people at a time t . The mean-field kinetics of the SIRD epidemic evolution is described by the following system of differential equations:

$$\frac{dS}{dt} = -\frac{\alpha}{N} S(t)I(t), \quad \frac{dI}{dt} = \frac{\alpha}{N} S(t)I(t) - \beta I(t) - \gamma I(t), \quad \frac{dR}{dt} = \beta I(t), \quad \frac{dD}{dt} = \gamma I(t), \end{aligned}$$

with initial condition $[S(t_0), I(t_0), R(t_0), D(t_0)]$ for some initial time t_0 . The parameter α is the infection rate, i.e. the probability per unit time that a susceptible individual contracts the disease when entering in contact with an infected person. The parameters β and γ denote, respectively, the recovery and death rates. This scheme has good chances to capture at least the gross features of the full time course of the outbreak.

SIRD model (Above figures shows SIRD model classes and change per unit time shown above arrows.)

4 Results

In this section we present our results for basic reproduction number, case fatality rate, case recovery ratios and time series forecasting for Germany and India.

4.1 Results for Germany

We first use the discrete model to find these parameters and predict time series for Germany. We use the time series data available and for the required vectors $\mathbf{C \Delta X}(T)$ for $X = I, R, D$.

```

country = "Germany" #Country chosen

#Extract country data from countries data
italy_cnf_melt = confirmed_countries_melt[which(confirmed_countries_melt$Country.Region == country), ]

#Extract confirmed number as a vector
italy_cnf = italy_cnf_melt$value

#Extract delta infected vector  $\text{delta\_cnf}(t) = I(t) - I(t-1)$ 
italy_delta_cnf = diff(italy_cnf)

#Cumulative sum of delta_cnf
italy_cum_delta_cnf = cumsum(italy_delta_cnf)

italy_deaths_melt = deaths_countries_melt[which(deaths_countries_melt$Country.Region == country), ]

#Extract deaths number as a vector
italy_deaths = italy_deaths_melt$value

#Extract delta deathsvector  $\text{delta\_deaths}(t) = D(t) - D(t-1)$ 
italy_delta_deaths = diff(italy_deaths)

#Cumulative sum of delta_deaths
italy_cum_delta_deaths = cumsum(italy_delta_deaths)

#Extract Italy data from countries recovered data
italy_recovered_melt = recovered_countries_melt[which(recovered_countries_melt$Country.Region == country), ]

#Extract recovered number as a vector
italy_recovered = italy_recovered_melt$value

#Extract delta recovered vector  $\text{delta\_recovered}(t) = R(t) - R(t-1)$ 
italy_delta_recovered = diff(italy_recovered)

#Cumulative sum of delta_recovered
italy_cum_delta_recovered = cumsum(italy_delta_recovered)

### Caluclating infected numbers from confirmed cases
#Extract infected number cases as a vector
italy_inf = italy_cnf - italy_recovered - italy_deaths

#Extract delta infected vector  $\text{delta\_inf}(t) = I(t) - I(t-1)$ 
italy_delta_inf = diff(italy_inf)

#Cumulative sum of delta_inf
italy_cum_delta_inf = cumsum(italy_delta_inf)

```

We then use different time windows to estimate the time evolution of these parameters.

```

###ESTIMATING CASE FATALITY RATIO

#Making data frame of cumulative data
italy_cum_data_full = data.frame(delta_inf= italy_delta_inf, cum_delta_inf=italy_cum_delta_inf, delta_recovered = italy_delta_recover

```

```

##VARY THE NUMBER OF DAYS CHOSEN FOR ANALYSIS
ndays = 65:97
gamma_data <- data.frame(matrix(ncol = 3, nrow = 0))
x <- c("est", "lwr", "upr")
colnames(gamma_data) <- x

beta_data <- data.frame(matrix(ncol = 3, nrow = 0))
x <- c("est", "lwr", "upr")
colnames(beta_data) <- x

R0_data <- data.frame(matrix(ncol = 3, nrow = 0))
x <- c("est", "lwr", "upr")
colnames(R0_data) <- x

#loop over days window chosen
for (days in ndays) {

  italy_cum_data = italy_cum_data_full[1:days, ]
  #View(italy_cum_data)

  #fitting a linear model for case fatality ratio
  italy_gamma <- lm(cum_delta_deaths ~ cum_delta_inf -1 , data=italy_cum_data) # build linear regression model on full data

  ###ESTIMATING CASE RECOVERY RATIO
  cor(italy_cum_delta_inf, italy_cum_delta_recovered)
  #high correlation
  #fitting a linear model for case recovery ratio
  italy_beta <- lm(cum_delta_recovered ~ cum_delta_inf -1 , data=italy_cum_data) # build linear regression model on full data with no

  ###ESTIMATING R0
  #fitting a linear model for case basic reproducibility number R0
  italy_R0 <- lm(cum_delta_deaths + cum_delta_recovered + cum_delta_inf ~ I(cum_delta_recovered + cum_delta_deaths) - 1 , data=italy

  ##Storing estimations and conf intervals

  conf = confint(italy_gamma)
  gamma_row <- list(est = summary(italy_gamma)$coefficients[1], lwr = conf[1], upr = conf[2])

  gamma_data = rbind(gamma_data, gamma_row)

  conf = confint(italy_beta)
  beta_row <- list(est = summary(italy_beta)$coefficients[1], lwr = conf[1], upr = conf[2])

  beta_data = rbind(beta_data, beta_row)

  conf = confint(italy_R0)
  R0_row <- list(est = summary(italy_R0)$coefficients[1], lwr = conf[1], upr = conf[2])

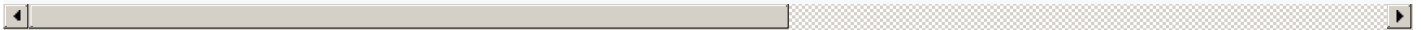
  R0_data = rbind(R0_data, R0_row)

}

```

We plot the estimates and the corresponding 99% confidence intervals for R_0, β, γ as below.

```
ggplot(R0_data, aes(ndays, est)) + geom_point() + geom_line(aes(ndays, est)) + geom_ribbon(aes(ymin=lwr,ymax=upr), alpha=0.5) + xlab("Days since first case")
```



□

```
ggplot(beta_data, aes(ndays, est)) + geom_point() + geom_line(aes(ndays, est)) + geom_ribbon(aes(ymin=lwr,ymax=upr), alpha=0.5) + xlab("Days since first case")
```



□

```
ggplot(gamma_data, aes(ndays, est)) + geom_point() + geom_line(aes(ndays, est)) + geom_ribbon(aes(ymin=lwr,ymax=upr), alpha=0.5) + xlab("Days since first case")
```



□

From these, we can see that the R_0 for Germany has come down from 8.2 to 1.6 (99% CI: [1.5 1.7]). This is good news and indicates flattening of the curve. Note that if $R_0 < 1$, the disease stops spreading. The case recovery ratio is going up and has reached approximately 0.9 while the case fatality ratio is about 0.049. We note here that since the estimate for case recovery and fatality only includes infected cases, hence the estimates for β and γ are only reliable for the early stages of the epidemic.

Since R_0 is estimated by a linear model, for curiosity we carry out some diagnostics to see how if the data actually fits to the linear plot. We plot out fitted slope (which is equal to R_0) and observe that a rather poor fit which is expected because R_0 changes with time.

```
##Predicted R0 using model
# add 'fit', 'lwr', and 'upr' columns to dataframe (generated by predict)
R0_predict <- cbind(italy_cum_data, predict(italy_R0, interval = 'confidence'))
R0_prediction = predict(italy_R0)
# plot the points (actual observations), regression line, and confidence interval
p <- ggplot(R0_predict, aes(cum_delta_recovered + cum_delta_deaths, cum_delta_recovered + cum_delta_deaths + cum_delta_inf))
p <- p + geom_point()
p <- p + geom_line(aes(cum_delta_deaths + cum_delta_recovered, R0_prediction))
p <- p + geom_ribbon(aes(ymin=lwr,ymax=upr), alpha=0.5) + ggtitle("Germany: Fitted R0 line and actual data")
p
```

□

Now, we solve the actual differential equation to fit the evolution of the disease and forecast the time series. We initialize the values below. We ignore the initial data because of noisiness and low scale testing.

```
library(deSolve)
library(RColorBrewer)

Infected <- italy_inf[50:97]
Recovered <- italy_recovered[50:97]
Deaths <- italy_deaths[50:97]
Confirmed <- italy_cnf[50:97]
day <- 0:(length(Infected)-1)
N <- 830000

###edit 1: use different boundary condition
###init <- c(S = N-1, I = 1, R = 0)
init <- c(S = N-Infected[1] - Recovered[1] - Deaths[1], I = Infected[1], R = Recovered[1], D = Deaths[1])
```

Then, we define the differential changes in the quantities with respect to time.

```
SIR <- function(time, state, parameters) {
  par <- as.list(c(state, parameters))
  #####edit 2; use equally scaled variables
  with(par, { dS <- -alpha * (S/N) * I
    dI <- alpha * (S/N) * I - beta * I - gamma * I
    dR <- beta * I
    dD <- gamma * I
    list(c(dS, dI, dR, dD))
  })
}
```

Then we define an optimizer to find the optimum parameters to fit the confirmed cases curve with an initial guess. For this we also define a misfit function which is a simple L2 error function. The code is given below.

```
RSS.SIR <- function(parameters) {
  names(parameters) <- c("alpha", "beta", "gamma")
  out <- ode(y = init, times = day, func = SIR, parms = parameters)
  fit <- out[, 3] + out[, 4] + out[, 5]
  RSS <- sum((Confirmed- fit)^2)
  return(RSS)
}

lower = c(0, 0, 0)
upper = c(10, 1, 1) ###Limit box for parameters for L-BFGS-B

optimstart <- c(0.7, 0.4, 0.2) #initial guess for parameters

set.seed(12)
Opt <- optim(optimstart, RSS.SIR, method = "L-BFGS-B", lower = lower, upper = upper,
            hessian = TRUE)
#Opt$par
```

Once we have optimized, we can predict the case evolution as follows.

```
Opt_par <- Opt$par
names(Opt_par) = c("alpha", "beta", "gamma")
t <- 0:120
fit <- data.frame(ode(y = init, times = t, func = SIR, parms = Opt_par))
predict <- fit$I + fit$D + fit$R
plot(t, predict, col="green", xlab="Days since March 11", ylab="Confirmed cases")
lines(day, Confirmed, col="red")
title("Germany: Green is confirmed cases predicted by our model, red is actual data.")
```

□

The results indicate that by the end of June, the confirmed cases will peak and the epidemic will end in Germany.

4.2 Results for India

Now we repeat the same analysis for India. Note India is still in the early phase of the disease.

We plot the estimates and the corresponding 99% confidence intervals for R_0, β, γ as below.

□□

From these, we can see that the R_0 for India has come down from 10.6 to 4.09 (99% CI : [3.94 4.25]). Thus India is quite far away from the end of the epidemic. Note that if $R_0 < 1$, the disease stops spreading. The case recovery ratio is going up and has reached approximately 0.26 while the case fatality ratio is about 0.041.

Since R_0 is estimated by a linear model, for curiosity we carry out some diagnostics to see how if the data actually fits to the linear plot. We plot the fitted slope (which is equal to R_0) and observe that a rather poor fit which is expected because R_0 changes with time.

□

We solve the differential equations system and fit the parameters to do time series forecasting for the India data.

Once we have optimized, we can predict the case evolution as follows.

□

The results indicate that the curve will peak in India by the end of July.

5. Conclusions and Future work

In this study, we used SIRD model commonly used in epidemiology to estimate the evolution of basic reproducibility number, case fatality ratio and case recovery ratio as the disease progresses. We also use it to fit the parameters to the data coming from Germany and India and use it to predict the evolution of disease. Our results indicate that the epidemic should end in Germany by the end of June. India might have to wait till the end of July before the peak is reached. The recovery ratios obtained for India (about 26%) are consistent with the government estimates recently reported in [business-standard](#). The prediction of beginning of curve flattening by July end are in line with the recent WHO estimates given [WHO envoy interview](#).

6. Limitations

However, we must state here that there are several limitations to our predictions. First, the model itself is a simplistic model to study the disease as it assumes constant transmission rates among different classes. Also, the model does not take into account the asymptomatic cases which may be contributing to spreading the disease. Secondly, the data itself might be unreliable as there might be severe underreporting of infected people because of lack of testing. Thirdly, the uplifting of

lockdown may accelerate the spread of disease. A more complex model with these factors taken into account would be desirable for a better forecasting.

7. References

[1] Cleo Anastassopoulou ,Lucia Russo,Athanasios Tsakris, Constantinos Siettos," Data-based analysis, modelling and forecasting of the COVID-19 outbreak", PLOS ONE (2020) (<https://doi.org/10.1371/journal.pone.0230405>)