



Mathematical modelling of COVID-19 and analysis of Rio de Janeiro city's outbreak in 2020

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Highlights

- Model the spread of the coronavirus SARS-CoV-2 (COVID-19) in 2020 with non-pharmaceutical interventions.
- Analyze the data from the first outbreak in the Rio de Janeiro city.
 Estimate relevant parameters with emphasis to the unreported rate and the effective reproduction curve, which are relevant to measure the real extent and impact of the disease.

COVID-19 in Rio de Janeiro

In March 2020, the state of Rio de Janeiro declared public health emergency and social distancing measures, but the outbreak could not be avoided: New COVID-19 cases in Rio de Janeiro

For the fitting presents a good matching: Curve fitting for the cumulated cases data -2.250

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Every reported case by the health system corresponds 9–10 not detected cases.

SEIAQR model

Compartmental model simplified from [1] without the lockdown for non-essential workers. The diagram flow is





Methods

Fit the **SEIAQR model** with $\beta(t), \mu(t)$ and α being unknown.



- We had evidence that the residuals are normally distributed but correlated, which requires more attention in future works.
- > We estimated $\hat{\alpha} = 0.9$ with confidence interval (0.85, 0.93). We verified that the values of the non-estimated parameters do not impact the estimation of α .
- The evolution of the non-tested positive cases over the total number of positive cases is

Ratio between unreported infections and total infections





- > The parameter $\beta(t)$ measures the **effective contact rate**, which depends on the average contact rate and the probability of infection given a contact.
- ρ is the testing rate among infected people with mild or no symptoms. We estimated it using IBGE data.
- The mortality rate among detected cases is μ(t) and varies in periods of distress for the health system and lack of testing kits.
- The other rate parameters are related to the pathogen and taken from the literature.
- > The **basic reproduction number** is

> The functions β and μ are written in a basis of polynomials of degree k - 1, the **B**-splines:

 $\beta(t) \approx \sum_{i=1}^{3} \beta_j B_{j,k}(t), \qquad \mu(t) \approx \sum_{i=1}^{r} \mu_j B_{j,k}(t),$

equally spaced knots (where the polynomials meet). The order and the number of coefficients are chosen using the Akaike Information Criterion: four coefficients for both with k = 2 for β and k = 1 for μ.
> The parameter estimation assumes that the observations have an uncorrelated normally distributed error:

 $\begin{cases} NewCases(i) = T(i) - T(i-1) + er_T(i) \\ NewDeaths(i) = D(i) - D(i-1) + er_D(i)' \\ \text{and minimizes a weighted quadratic error, balancing } \\ new detected cases and deaths. \end{cases}$

A parametric Bootstrap approach is used to calculate confidence intervals. The parameters are estimated in new datasets generated by sampling the errors from the normal distribution and summing them to the observed new cases/deaths.

Identifiability

Discussion and conclusions

 \succ The estimated range of values for α and the curve

 $R_0 = \frac{1}{2} \left(\varphi + \sqrt{\varphi^2 + \frac{4\sigma\alpha}{\rho + \gamma_1} \varphi} \right), \varphi = \frac{\beta\tau}{(\rho\delta + \tau)(\sigma + \rho)}.$

➢ As the epidemic evolves, the effective reproduction curve R_t has the same expression of R_0 with the change that

 $\varphi(t) = \frac{\beta(t)\tau S(t)}{(\rho\delta + \tau)(\sigma + \rho)}.$

Can the parameters **be estimated in a unique way** from the available measurements?

- **1 Structural:** theoretical analysis of the structure of the model. We could not guarantee structural identifiability using the available methods in the literature.
- **2 Practical:** considers the noisy data. We analyzed the correlation between the estimated parameters using the **Fisher Information Matrix (FIM)** and expected low values for the entries of its inverse.
- R_t over time are in line with other works.
- The identifiability of the model is a matter to study in the future.
- Data analysis techniques have been deployed to deal with the low quality of data.
- ➤ The rate of unreported positive cases is about 90% (85%-93%), that is, every case reported by the health system corresponds to about 9-10 cases that were not detected.
- This estimation provides a useful tool to determine periods of growth or decrease of the epidemic force.

References

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