

Inference of seasonal forcings on reproduction numbers during epidemics

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Abstract

Amidst the current COVID-19 epidemic, the prediction of its future spread and the length of the epidemic becomes increasingly significant. We investigate the effects of temperature and humidity on the reproductive rate R of COVID-19, as warmer climates in the summer have historically resulted in diminished spread of influenza. Early epidemiology research in COVID-19 have used simple linear models such as ordinary least squares regression to estimate the relationship between R and climate. We estimate the reproductive rate of the virus using a novel Bayesian inference model with MCMC sampling to daily case data of various climate regions. While results are largely inconclusive, they provide us with a weak posterior update that R decreases with temperature and relative humidity.

Introduction

Exponential growth of the number of symptomatic cases is usually observed in the initial stages of infectious disease epidemics. Typically, the rate of this growth is decomposed into two quantities:

- The serial interval: time between onset of symptoms of two adjacent individuals in a transmission chain.
- The reproductive number, R : the expected number of new infections a single symptomatic individual will cause.

Due to inadequate public data to compute the serial interval, we use the estimated serial interval of 4.7 days with standard deviation 2.9 days obtained from Nishiura et al. (2020) [1]. The estimation of the latter quantity is important for policy, because infectious disease outbreaks will switch from an exponential growth regime to an exponential decay regime when the reproductive number falls below 1. The goal of mitigation or suppression policies is therefore to effect change on the reproductive number R . Many infectious diseases are believed to be seasonal, including most respiratory viruses, like the SARS-CoV-2 virus implicated in the 2019–2020 coronavirus pandemic. Wang et al. (2020) [3] have analysed how the weather of locations within China correlates with local point estimates of the reproductive number R . However, because their analysis is frequentist in nature, their estimates of the seasonal forcing do not come with any estimate of uncertainty.

We first produce a baseline model, which predicts the reproductive rate using the EpiEstim package, based on previous work by Thompson et al. (2019) [2], and applied to COVID-19 data as in Abbot et al. (2020) [4]. We then apply ordinary least squares regression against climate variables as well as other underlying regional factors, such as population density, GDP per capita, and latitude, similar to the work produced by Wang et

al. (2020)[3]. We compare this against our novel Bayesian framework, and evaluate the significance of the explanatory variables in each model.

Since Italy was one of the first countries affected by the epidemic, it has a longer history of case data before intervention and lockdowns took place. Furthermore, due to the many provinces of Italy having diverse climates, we have access to a larger range of temperature and humidity. Therefore, we sourced the Italian government DPC dataset[5], which contains time series incidence for each province since mid February. Using the temperature and humidity data from weather stations sourced from NOAA, we are able to estimate the climate for over 80 distinct provinces in the Italian COVID-19 dataset.

Methodology

Thompson et al. (2019) outline a Bayesian framework for estimation of $R(t)$ using symptomatic case counts, decomposed into local and imported cases. The EpiEstim R package implements ideas from this paper. A simplified summary of their approach, ignoring the imported cases:

1. Estimation of (discrete) distribution of serial interval from line-list data with intervals reflecting date of infection.
2. Estimation of time-varying R from time-bucketed symptomatic case counts.

We developed a method to estimate the strength of various seasonal forcings on R through MCMC. This involves both substantial deviations from the second part of the Thompson et al. model and also less substantial adjustments for the nature of our data. Our method does not extend the first part of the Thompson et al. model.

To sanity-check the results of our method, we compare its conclusions against a point estimate for the seasonal forcings through linear regression with the EpiEstim estimates of R .

Incidence likelihood of the Thompson et al. model

In its estimation of time-varying R , the Thompson et al. model uses the following likelihood function for the number of new infections at time t :

$$I(t)|I(1), \dots, I(t-1), \mathbf{w}, R(t) \sim \text{Poisson} \left(R(t) \sum_{s=1}^t I(t-s)w_s \right)$$

where

- $I(t)$ is observed data containing the number of new infections at time t
- $R(t)$ is a parameter for the reproduction number at time t
- w_s denotes the proportion of serial intervals which take on discrete value s , i.e. $w_s = P(SI = s)$.

Because the Thompson et al. model uses a gamma-distributed prior for $R(t)$ (the conjugate prior for a Poisson likelihood) and because they chose a convenient family of serial interval distributions, the Bayesian posterior in the Thompson et al. model can be described by a closed form analytic expression.

Extension of Thompson et al. for R subject to forcings

Our Bayesian model extends the method of Thompson et al. to multiple locations, each with its own incidence time series $I_1(t), \dots, I_n(t)$ such that

$$R_i(t) = \exp(\theta(t) + \lambda_i + \beta^T x_i(t) + \varepsilon_i(t))$$

where:

- $x_i(t)$ is data of observed seasonal factors at time t and location i , with each feature standardized to mean 0 and variance 1
- β is a parameter describing the locally linear effect of each seasonal forcing
- $\theta(t)$ is a parameter for logarithm of country-wide R independent of modeled seasonal forcings
- λ_i is a parameter for local variation of log R independent of modeled seasonal forcings
- $\varepsilon_i(t)$ is a parameter for remaining fluctuation in log R not modeled above

Unlike the Thompson et al., our analysis fundamentally estimates the logarithm of R instead of R itself. This has some practical advantages: a linear model for R could predict negative values for sufficiently extreme climate conditions, but negative values for R do not make sense epidemiologically.

We maintain a very similar incidence likelihood function as the Thompson et al. model, but impose an additional correction for importations:

$$I_i(t) | P_i, x_i(t), I_i(1), \dots, I_i(t-1), \mathbf{w}, \theta(t), \lambda_i, \beta, \varepsilon(t), \gamma \sim \text{Poisson} \left(\gamma P_i + R_i(t) \sum_{s=1}^t I_i(t-s) w_s \right)$$

where in addition to the parameters described (with $R_i(t)$ given by the equation above), the following additional data and parameters are introduced:

- P_i is data giving the population of location i
- γ is a parameter estimating the nationwide rate of imports per capita per day, on which we impose prior $\gamma \sim \text{Gamma}(2, 10^{-5})$

The only substantive change to this likelihood function, beyond the decomposition of $R_i(t)$ into the various parameters influencing it, is the addition of importation risk. The reason for this adjustment is that the dataset we are using does not distinguish between importations and local cases. Thus, the Thompson likelihood would be 0 upon detection of the first case, had we not modeled the importations separately.

The most important feature of our model is the parameter β which are the coefficients of the linear model of log R . This feature is assumed to be global across all locations and times, and thus models inherent properties of the contagiousness of the outbreak, as opposed to local deviations in space or time.

To avoid the model ascribing deviations in R to the seasonal forcings without widespread evidence across locations and times, three other features of our model are the parameters $\theta(t)$, λ_i , and $\varepsilon_i(t)$. The first parameter, $\theta(t)$, describes the logarithm country-wide R before adjusting for location and climate and local idiosyncratic factors. That is, it can be thought of as a baseline R value affected simply by the national circumstances at the given time; such as national policies or human behavioral changes. This will encapsulate

various unmodeled behaviours such as weekends or interventions. At time zero, our prior belief is that this R will match the consensus of international estimates for R_0 , which vary widely but tend to cluster between 2 or 3 [TK: Citation needed?]. Because of the great uncertainty, there is significant variance in this assumption, so the prior is quite weak (note that the $\mu \pm 2\sigma$ values, when transformed exponentially, are below 1 and above 6 respectively).

$$\theta(0) \sim \text{Normal}(\log 2.5, 0.5^2)$$

The remaining days of θ are modeled using a simple autoregressive model:

$$\theta(t+1) \sim \text{Normal}(\theta(t), \theta^\sigma)$$

where θ^σ is a parameter describing the spread of daily drifts in θ . The prior on θ^σ is weakly informative:

$$\theta^\sigma \sim \text{Gamma}(2, 0.01)$$

The location-specific deviation from the country average in $\log R$ independent of modeled seasonal forcings is encoded in λ_i . The existence of this parameter helps the model explain the large variations of apparent R across different locations, some of which appear to be affected much more so than others. The remaining parameter $\varepsilon_i(t)$ encapsulates all other unexplained deviations across both time and location. All these parameters (and β) are given the following weakly informative prior:

$$\lambda_i, \varepsilon_i(t), \beta_k \sim \text{Normal}(0, 0.5)$$

Our technique is general and can be used for any epidemic and any combination of potential seasonal forcings.

As the model is much more complex than the one in the Thompson et al. paper, we are unable to derive an analytic expression to compute the posterior. Instead, we use **rstan** to run 4 chains of MCMC for 2000 iterations each.

Using OLS and EpiEstim to check results

The baseline model we build estimates the reproductive rate using a parametric SI model, where we specify the mean and standard deviation of the serial interval. Because we do not have access to line-list SI data for COVID-19, we are unable to effectively estimate the serial interval from data, thus we take the findings of Nishiura et al. (2020) [1], which estimated the mean serial interval to be 4.7 and the standard deviation to be 2.4. We use the model detailed in Wallinga and Teunis (2004) [8] for estimating R from bayesian inference on a fixed serial interval distribution and time series data, which can be easily implemented using the EpiEstim package produced by Cori et al. (2013) [7].

For each province, we take a 7 day rolling sum on the number of new cases each day, and feed the data into the parametric SI model, which is given distribution $w(\tau)$ From there, the model uses a likelihood based approach to estimate R at each discrete timestep, computing the likelihood that a person at timestep i is

infected by a person infected at timestep j , and summing the likelihood for each j across all its infectees to compute the R value at timestep j . Thus, for each province, we produce a daily time series estimate for R on each day. This method is outlined in detail in Wallinga and Teunis (2004)[8].

For each province, day pair, we have $\bar{x} = (x_1, x_2, x_3, x_4), y$ where:

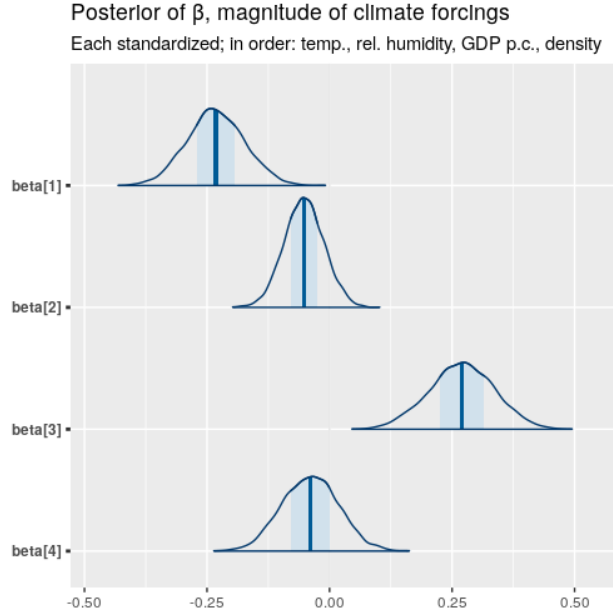
- x_1 : standardized GDP per capita
- x_2 : standardized population density
- x_3 : standardized air temperature
- x_4 : standardized relative humidity
- y : logarithm of estimated R

We then perform ordinary least squares regression to find $\bar{\beta}$:

$$y = \beta_0 + \beta_1 x_1 + \beta_2 x_2 + \beta_3 x_3 + \beta_4 x_4 + \varepsilon$$

Results

Applying the methods depicted from the methodology section, we obtain the following results.



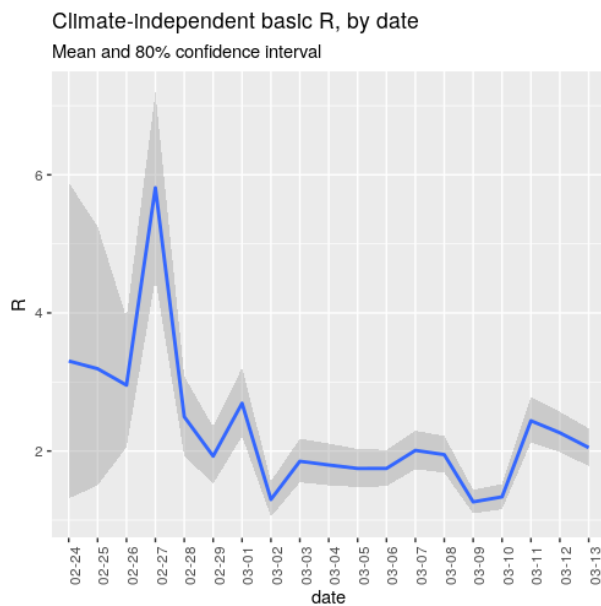
[TK: Beta Values] Fitting our Bayesian model, which extends the method of Thompson et. al, values of beta were estimated for temperature, relative humidity, Gross Domestic Product (GDP) and population density, respectively. Thus, our model estimates $\beta^T = [, , ,]$ in

$$R_i(t) = \exp(\theta(t) + \lambda_i + \beta^T x_i(t) + \varepsilon_i(t))$$

Note that each feature's data is standardized to mean 0 and variance 1.

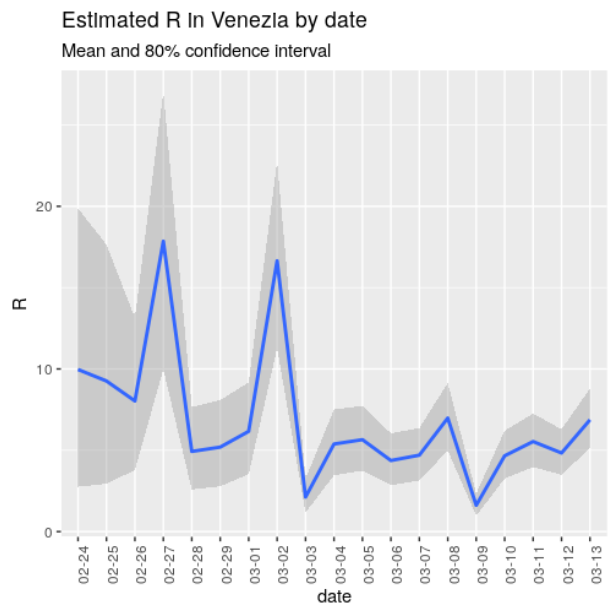
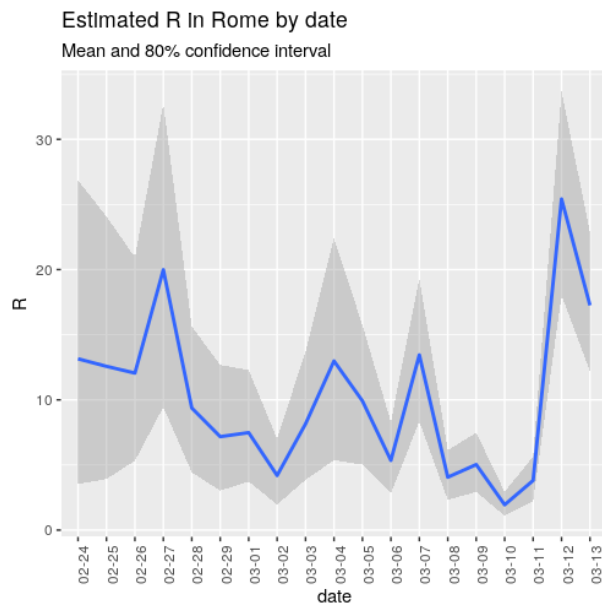
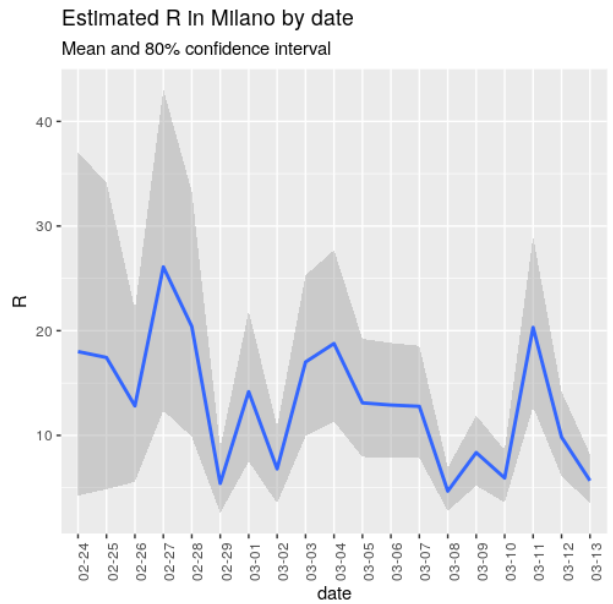
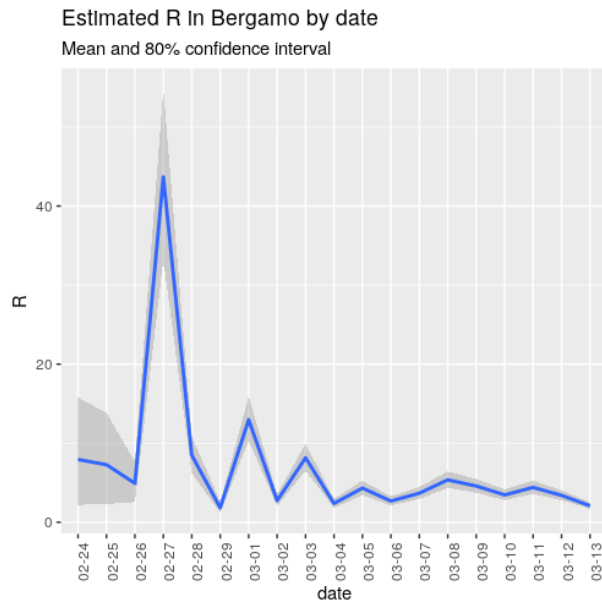
The model indicates that higher temperatures are observed with lower R_0 values, and higher GDP is observed with higher R_0 values, while relative humidity and population density have R_0 close to zero.

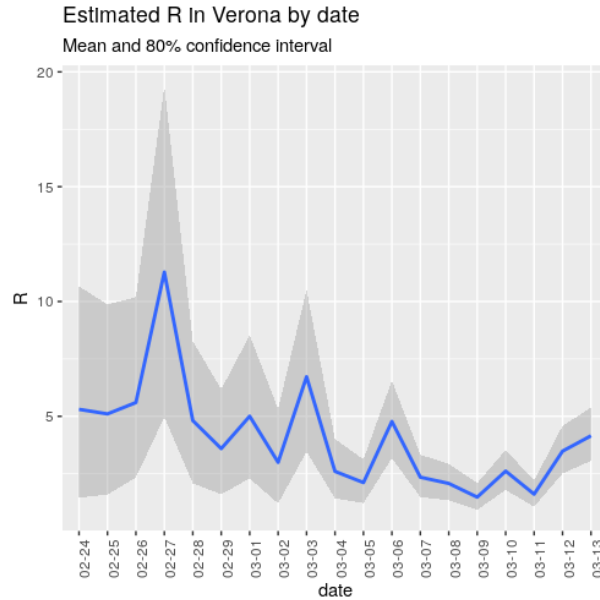
Climate-Independent Basic R , by Date



First, we observe that our computed value of R_0 in Italy matches what We observe that R_0 spikes in , with an early spike on February 27, potentially due to increased in testing, which captures cases before.

Estimated R in Various Provinces





Observe that in many cities, our estimated R_0 values are extremely high compared to the climate-independent plot shown earlier.

Discussion

Directions for Future Research

With our `cfc19i` library, it is

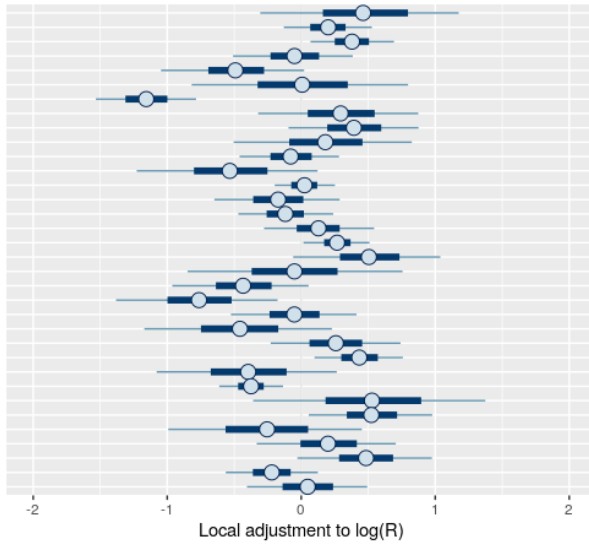
Looking at the climate and infection data in American and Canadian cities as the situation develops further. In particular, compare this with the results in Italy, and observe if the linear assumption between seasonal factors and R holds true across different climate regions. If not, try fitting while assuming a nonlinear relationship between R and seasonal factors.

Look into statistical simulation models which consider intercity travel and geographic proximity.

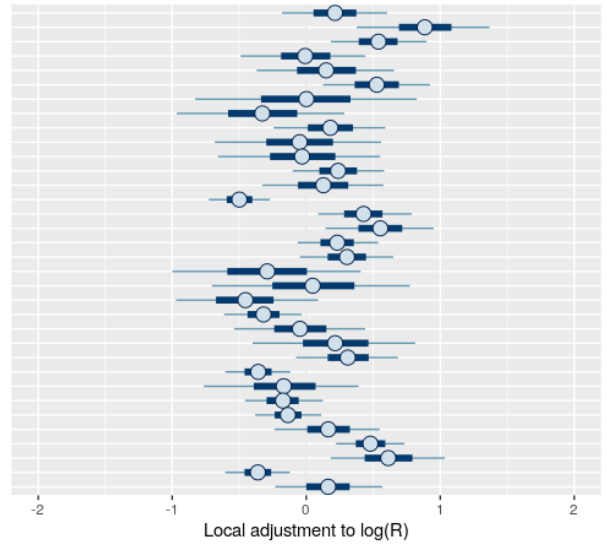
Appendix

Posterior of Lambda, location-dependent adjustment to $\log(R)$

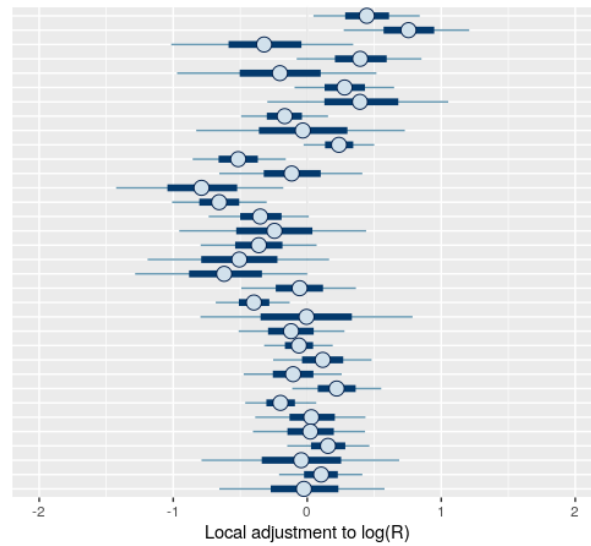
Posterior of λ , location-dependent adjustment to $\log(R)$
Mean, 50% and 90% confidence intervals for locations 1 through 34



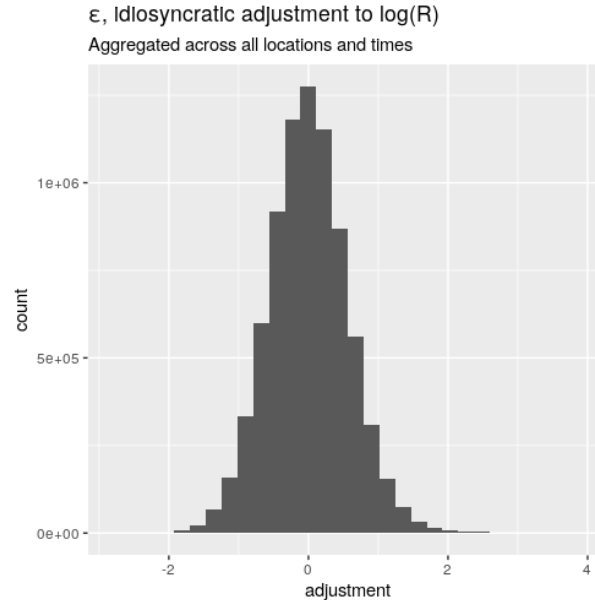
Posterior of λ , location-dependent adjustment to $\log(R)$
Mean, 50% and 90% confidence intervals for locations 35 through 68



Posterior of λ , location-dependent adjustment to $\log(R)$
Mean, 50% and 90% confidence intervals for locations 69 through 102



Epsilon, Idiosyncratic adjustment to $\log(R)$



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