

# Bayesian Inference Final Project

## Introduction

The reproduction number  $R_t$  is a common measure of transmissibility of an epidemic disease. By definition,  $R_t$  is the average number of secondary cases caused by an infected individual.  $R_t$  falling below 1 indicates the disease is unlikely to sustain. The change of  $R_t$  across time can be used as a proxy for epidemic trajectories. Following the method proposed by Cori et al.,  $R_t$  can be calculated by using Bayesian parametric estimation. In this project, I use the same method to compute the  $R_t$  of COVID-19 in New York before the presence of effective vaccination (Mar 2020 to Dec 2020).

```
library(dplyr)
library(lubridate)
library(rstan)
```

```
df <-
  read.csv("NY_cases.csv")
df<-df%>%select(state,submission_date,new_case, tot_cases)
```

## Method

Assuming the number of reported incident cases follows a poisson process, we have:

$$P(I_t|I_0, I_1, \dots, I_{t-1}, w_s, R_t) = \frac{(R_t \Lambda_t(w_s))^t \exp(-R_t \Lambda_t(w_s))}{I_t!},$$

where  $I_t$  is the number of incident cases arising at time  $t$  and  $\Lambda_t(w_s)$  is the overall infectivity and can be computed as:

$$\Lambda_t(w_s) = \sum_{s=1}^t I_{t-s} w_s.$$

$w_s$  is the serial interval distribution, which is the time between the onset of symptoms in a primary case and he onset of symptoms in secondary cases. It serves as the weight for reported new cases at each time step before time  $t$ . I choose the distribution of the serial period of COVID-19 follows a gamma distribution with mean of 5.9 and standard deviation of 3.9, according to the paper of Liu et al.. I also choose the prior of  $R_t$  to follow a gamma distribution with mean = 1.5 and standard deviation of 2.

With the above assumptions and procedure, I use Stan to yield the posterior distribution of  $R_t$  on the last day of 2020 (Dec 31st, 2020) with all previous reported cases.

```
#get the weight of previous days using a gamma distribution
lastday <- df %>%
  mutate(days_away=rev(row_number())-1,
         weight=dgamma(days_away,shape=(5.9/3.9)^2,rate=5.9/3.9^2))

data { /* these are known and passed as a named list from R */
  int<lower = 0> I;           // number of cases in day t
  real<lower = 0> infect;    // weighted sum of previous infectivity
  real<lower = 0> alpha;     //shape parameter of gamma prior
```

```

    real<lower = 0> beta;// rate parameter of gamma prior
    int<lower = 0, upper = 1> prior_only;
}

parameters {
    real<lower=0> Rt; // Reproduction number
}

model {
    if (!prior_only) {
        target +=poisson_lpmf(I | Rt*infect); // log-likelihood
    }
    target += gamma_lpdf(Rt | alpha, beta); //prior of Rt
}

#calculating overall infectivity
overallinfectivity=sum(lastday$new_case*lastday$weight)
#indicate new case on Dec 31,2020
It=lastday$new_case[nrow(lastday)]
#choose prior parameters (shape and rate)
a=9/16
b=3/8

post <- stan("reproduction_num.stan",
             data = list(infect=overallinfectivity,I=It, prior_only = 0, alpha = a, beta=b))

post

## Inference for Stan model: reproduction_num.
## 4 chains, each with iter=2000; warmup=1000; thin=1;
## post-warmup draws per chain=1000, total post-warmup draws=4000.
##
##      mean se_mean   sd  2.5%   25%   50%   75%  97.5% n_eff Rhat
## Rt    1.57     0.00 0.01   1.54   1.56   1.57   1.58   1.60  1332    1
## lp__ -7.39     0.01 0.68  -9.33  -7.56  -7.12  -6.95  -6.91  2265    1
##
## Samples were drawn using NUTS(diag_e) at Sun May 15 21:48:23 2022.
## For each parameter, n_eff is a crude measure of effective sample size,
## and Rhat is the potential scale reduction factor on split chains (at
## convergence, Rhat=1).

```

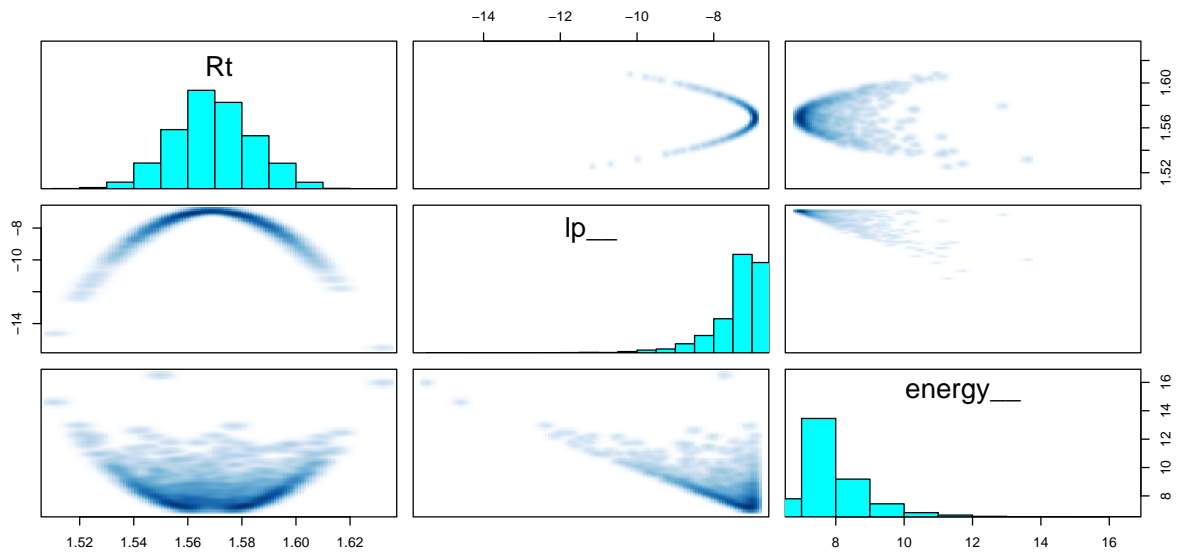
The mean of the posterior distribution of  $R_t$  on Dec.31st, 2020 in New York is 1.57 with a 0.95 credible interval of (1.54,1.60).

## Posterior Planes

```

pairs(post, pars = "p", include = FALSE)

```



```
draws <- as.data.frame(post) %>% select(-starts_with("p"))
mean(draws$Rt)
```

```
## [1] 1.569408
```

## Different Assumptions

I then modify the approach by using a negative binomial likelihood instead of a poisson likelihood since negative binomial distribution allows for overdispersion. The negative binomial distribution has an extra dispersion parameter  $\phi$  and it is assumed to follow a half cauchy distribution with location parameter =5 and scale = 5.

```
data { /* these are known and passed as a named list from R */
  int<lower = 0> I;           // number of cases in day t
  real<lower = 0> infect;     // weighted sum of previous infectivity
  real<lower = 0> alpha;      //shape parameter of gamma prior
  real<lower = 0> beta;       // rate parameter of gamma prior
  real<lower = 0> mu;         //location parameter of cauchy prior
  real<lower = 0> sigma;      //scale parameter of cauchy prior
  int<lower = 0, upper = 1> prior_only;
}

parameters {
  real<lower=0> Rt; // Reproduction number
  real<lower=0> phi; // dispersion parameter
}

model {
  if (!prior_only) {
    target += neg_binomial_2_lpmf(I | Rt*infect, phi); // log-likelihood
  }
}
```

```

}
target += gamma_lpdf(Rt | alpha, beta); //prior of Rt
target += cauchy_lpdf(phi | mu, sigma); //prior of phi
}

```

```

post_neg <- stan("reproduction_num2.stan",
  data = list(infect=overallinfectivity,I=It, prior_only = 0, alpha = a, beta=b, mu=5,sigma=

```

```
post_neg
```

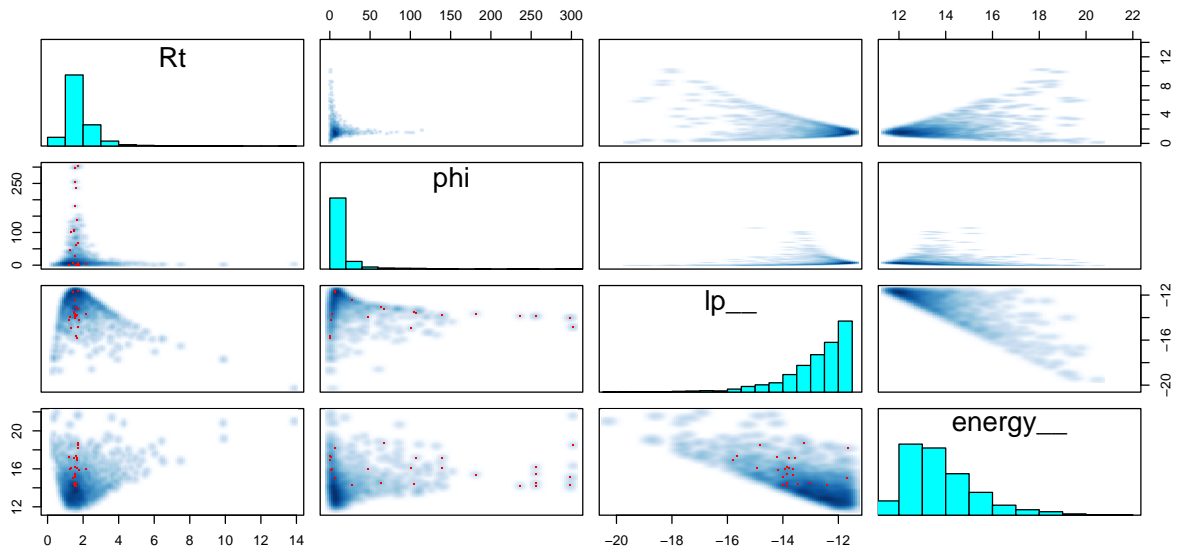
```

## Inference for Stan model: reproduction_num2.
## 4 chains, each with iter=2000; warmup=1000; thin=1;
## post-warmup draws per chain=1000, total post-warmup draws=4000.
##
##      mean se_mean   sd  2.5%  25%   50%   75%  97.5% n_eff Rhat
## Rt      1.80    0.03  0.88  0.74  1.30  1.61  2.03   3.96 1177 1.00
## phi     14.12   1.11 24.54  0.96  4.35  7.52 13.44 83.07  488 1.01
## lp__    -12.78   0.04  1.20 -15.89 -13.27 -12.44 -11.90 -11.54  797 1.00
##
## Samples were drawn using NUTS(diag_e) at Sun May 15 21:48:45 2022.
## For each parameter, n_eff is a crude measure of effective sample size,
## and Rhat is the potential scale reduction factor on split chains (at
## convergence, Rhat=1).

```

After modification, the mean of the posterior distribution of  $R_t$  on Dec.31st, 2020 in New York is 1.70 with a 0.95 credible interval of (0.77,3.41). The negative binomial likelihood gives a wider credible interval which implies higher uncertainty in estimation.

```
pairs(post_neg, pars = "p", include = FALSE)
```



Essentially, we can compute the posterior distribution of  $R_t$  for all time points in the data set, assuming the same serial interval distribution. Using the mean of the posterior distributions, a graph of  $R_t$  can be plotted.

```

for (i in 2:nrow(df)){
  dt<-df[1:i,] %>%
  mutate(days_away=rev(row_number())-1,
         weight=dgamma(days_away,shape=5.9^2/3.9,rate=5.9/3.9))
  overall=sum(dt$new_case*dt$weight)
  I_t=dt$new_case[nrow(dt)]
  a=9/16
  b=3/8
  posterior <- stan("reproduction_num.stan",
                  data = list(infect=overall,I=I_t, prior_only = 0, alpha = a, beta=b))
  post.df <- as.data.frame(posterior) %>% select(-starts_with("p"))
  df$Rt[i]<-mean(post.df$Rt)
}

```

## Reference

- Anne Cori, Neil M. Ferguson, Christophe Fraser, Simon Cauchemez, A New Framework and Software to Estimate Time-Varying Reproduction Numbers During Epidemics, American Journal of Epidemiology, Volume 178, Issue 9, 1 November 2013, Pages 1505–1512, <https://doi.org/10.1093/aje/kwt133>
- Liu, X., Xu, X., Li, G., Xu, X., Sun, Y., Wang, F., Shi, X., Li, X., Xie, G., & Zhang, L. (2021). Differential impact of non-pharmaceutical public health interventions on COVID-19 epidemics in the United States. BMC Public Health, 21(1), 965. <https://doi.org/10.1186/s12889-021-10950-2>