

Package ‘eSIR’

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Type Package

Title Extended state-space SIR models

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Description An implementation of extended state-space SIR models developed by Song Lab at UM school of Public Health

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Depends rjags, scales, ggplot2, chron, gtools

RoxygenNote 6.1.1

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qh.eSIR	<i>Extended state-space SIR with quarantine</i>
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Description

Fit an extended state-space SIR model being reduced by in-home hospitalization.

Usage

```
qh.eSIR(Y, R, phi0 = NULL, change_time = NULL,
  begin_str = "01/13/2020", T_fin = 200, nchain = 4,
  nadapt = 10000, M = 500, thn = 10, nburnin = 200, dic = FALSE,
  death_in_R = 0.02, casename = "qh.eSIR", beta0 = 0.2586,
  gamma0 = 0.0821, R0 = beta0/gamma0, gamma0_sd = 0.1, R0_sd = 1,
  file_add = character(0), save_files = FALSE, save_mcmc = FALSE)
```

Arguments

<code>Y</code>	the time series of daily observed infected compartment proportions.
<code>R</code>	the time series of daily observed removed compartment proportions, including death and recovered.
<code>phi0</code>	a vector of values of the dirac delta function ϕ_t . Each entry denotes the proportion that will be quarantined at each change time point. Note that all the entries lie between 0 and 1, its default is <code>NULL</code> .
<code>change_time</code>	the change points over time corresponding to <code>phi0</code> , to formulate the dirac delta function ϕ_t ; its default value is <code>NULL</code> .
<code>begin_str</code>	the character of starting time, the default is "01/13/2020".
<code>T_fin</code>	the end of follow-up time after the beginning date <code>begin_str</code> , the default is 200.
<code>nchain</code>	the number of MCMC chains generated by <code>rjags</code> , the default is 4.
<code>nadapt</code>	the iteration number of adaptation in the MCMC. We recommend using at least the default value 1e4 to obtained fully adapted chains.
<code>M</code>	the number of draws in each chain, with no thinning. The default is $M=5e2$ but suggest using $5e5$.
<code>thn</code>	the thinning interval between mixing. The total number of draws thus would become $\text{round}(M/\text{thn}) * \text{nchain}$. The default is 10.
<code>nburnin</code>	the burn-in period. The default is $2e2$ but suggest $2e5$.
<code>dic</code>	logical, whether compute the DIC (deviance information criterion) for model selection.
<code>death_in_R</code>	the numeric value of average of cumulative deaths in the removed compartments. The default is 0.4 within Hubei and 0.02 outside Hubei.
<code>casename</code>	the string of the job's name. The default is "qh.eSIR".
<code>beta0</code>	the hyperparameter of average transmission rate, the default is the one estimated from the SARS first-month outbreak (0.2586).
<code>gamma0</code>	the hyperparameter of average removed rate, the default is the one estimated from the SARS first-month outbreak (0.0821).
<code>R0</code>	the hyperparameter of the mean reproduction number R_0 . The default is thus the ratio of <code>beta0/gamma0</code> , which can be specified directly.
<code>gamma0_sd</code>	the standard deviation for the prior distribution of the removed rate γ , the default is 0.1.
<code>R0_sd</code>	the standard deviation for the prior distribution of R_0 , the default is 1.
<code>file_add</code>	the string to denote the location of saving output files and tables.
<code>save_mcmc</code>	logical, whether save (<code>TRUE</code>) all the MCMC outputs or not (<code>FALSE</code>). The output file will be an <code>.RData</code> file named by the <i>casename</i> . We include arrays of prevalence values of the three compartments with their matrices of posterior draws up to the last date of the collected data as <code>theta_p[, , 1]</code> and afterwards as <code>theta_pp[, , 1]</code> for θ_t^S , <code>theta_p[, , 2]</code> and <code>theta_pp[, , 2]</code> for θ_t^I , and <code>theta_p[, , 3]</code> and <code>theta_pp[, , 3]</code> for θ_t^R . The posterior draws of the prevalence process of the quarantine compartment can be obtained via <code>thetaQ_p</code> and <code>thetaQ_pp</code> . Moreover, the input and predicted proportions <code>Y</code> , <code>Y_pp</code> , <code>R</code> and <code>R_pp</code> can also be retrieved. The prevalence and predicted proportion matrices have rows for MCMC replicates, and columns for days. The MCMC posterior draws of other parameters including <code>beta</code> , <code>gamma</code> , <code>R0</code> , and variance controllers <code>k_p</code> , <code>lambdaY_p</code> , <code>lambdaR_p</code> are also available.

Details

In this function we allow it to characterize time-varying proportions of susceptible due to government-enforced stringent in-home isolation. We expanded the SIR model by adding a quarantine compartment with a time-varying rate of quarantine ϕ_t , the chance of a susceptible person being willing to take in-home isolation at time t .

Value

casename	the predefined casename.
incidence_mean	mean incidence.
incidence_ci	2.5%, 50%, and 97.5% quantiles of the incidences.
out_table	summary tables including the posterior mean of the prevalence processes of the 3 states compartments ($\theta_t^S, \theta_t^I, \theta_t^R, \theta_t^H$) at last date of data collected (t') decided by the lengths of your input data \mathbb{Y} and \mathbb{R}), and their respective credible intervals (ci); the respective means and ci's of the reproduction number (R_0), removed rate (γ), transmission rate (β).
plot_infection	plot of summarizing and forecasting for the infection compartment, in which the vertical blue line denotes the last date of data collected (t'), the vertical dark-gray line denotes the deacceleration point (first turning point) that the posterior mean first-derivative of infection prevalence $\dot{\theta}_t^I$ achieves the maximum, the vertical purple line denotes the second turning point that the posterior mean first-derivative infection proportion $\dot{\theta}_t^I$ equals zero, the darkgray line denotes the posterior mean of the infection prevalence θ_t^I and the red line denotes its posterior median.
plot_removed	plot of summarizing and forecasting for the removed compartment with lines similar to those in the <code>plot_infection</code> . The vertical lines are identical, but the horizontal mean and median correspond to the posterior mean and median of the removed process θ_t^R . An additional line indicates the estimated death prevalence from the input <code>death_in_R</code> .
first_tp_mean	the date t at which $\ddot{\theta}_t^I = 0$, calculated as the average of the time points with maximum posterior first-order derivatives $\dot{\theta}_t^I$; this value may be slightly different from the one labeled by the "darkgreen" lines in the two plots <code>plot_infection</code> and <code>plot_removed</code> , which indicate the stationary point such that the first-order derivative of the averaged posterior of θ_t^I reaches its maximum.
first_tp_mean	the date t at which $\ddot{\theta}_t^I = 0$, calculated as the average of the time points with maximum posterior first-order derivatives $\dot{\theta}_t^I$; this value may be slightly different from the one labeled by the "darkgreen" lines in the two plots <code>plot_infection</code> and <code>plot_removed</code> , which indicate the stationary point such that the first-order derivative of the averaged posterior of θ_t^I reaches its maximum.
first_tp_ci	fwth <code>first_tp_mean</code> , it reports the corresponding credible interval and median.
second_tp_mean	the date t at which $\theta_t^I = 0$, calculated as the average of the stationary points of all of posterior first-order derivatives $\dot{\theta}_t^I$; this value may be slightly different from the one labeled by the "purple" lines in the plots of <code>plot_infection</code> and <code>plot_removed</code> . The latter indicate stationary t at which the first-order derivative of the averaged posterior of θ_t^I equals zero.

`second_tp_ci` with `second_tp_mean`, it reports the corresponding credible interval and median.

`dic_val` the output of `dic.sample()` in `dic.sample`, computing deviance information criterion for model comparison.

Examples

```
NI_complete <- c( 41,41,41,45,62,131,200,270,375,444,549, 729,
                 1052,1423,2714,3554,4903,5806,7153,9074,11177,
                 13522,16678,19665,22112,24953,27100,29631,31728,33366)
RI_complete <- c(1,1,7,10,14,20,25,31,34,45,55,71,94,121,152,213,
                 252,345,417,561,650,811,1017,1261,1485,1917,2260,
                 2725,3284,3754)

N=58.5e6
R <- RI_complete/N
Y <- NI_complete/N- R #Jan13->Feb 11

change_time <- c("01/23/2020","02/04/2020","02/08/2020")
phi0 <- c(0.1,0.4,0.4)
res.q <- qh.eSIR (Y,R,begin_str="01/13/2020",death_in_R = 0.4,
                 phi0=phi0,change_time=change_time,
                 casename="Hubei_q",save_files = T,save_mcmc = F,
                 M=5e2,nburnin = 2e2)

res.q$plot_infection
#res.q$plot_removed

res.noq <- qh.eSIR (Y,R,begin_str="01/13/2020",death_in_R = 0.4,
                 T_fin=200,casename="Hubei_noq",
                 M=5e2,nburnin = 2e2)

res.noq$plot_infection
```

tvt.eSIR

Fit extended state-space SIR model with time-varying transmission rates

Description

Fit extended state-space SIR model with prespecified changes in the transmission rate, either step-wise or continuous, accomodating time-varying quarantine protocols.

Usage

```
tvt.eSIR(Y, R, pi0 = NULL, change_time = NULL, exponential = FALSE,
         lambda0 = NULL, begin_str = "01/13/2020", T_fin = 200,
         nchain = 4, nadapt = 10000, M = 500, thn = 10, nburnin = 200,
         dic = FALSE, death_in_R = 0.02, beta0 = 0.2586, gamma0 = 0.0821,
         R0 = beta0/gamma0, gamma0_sd = 0.1, R0_sd = 1,
         casename = "tvt.eSIR", file_add = character(0), save_files = FALSE,
         save_mcmc = FALSE)
```

Arguments

Y	the time series of daily observed infected compartment proportions.
R	the time series of daily observed removed compartment proportions, including death and recovered.
pi0	the time-dependent transission rate modifier $\pi(t)$ between 0 and 1.
change_time	the change points over time for step function pi, default value is NULL.
exponential	logical, whether $\pi(t)$ is exponential $\exp(-\lambda_0 t)$ or not; the default is FALSE.
lambda0	the rate of decline in the exponential survival function in $\exp(-\lambda_0 t)$.
begin_str	the character of starting time, the default is "01/13/2020".
T_fin	the end of follow-up time after the beginning date begin_str, the default is 200.
nchain	the number of MCMC chains generated by <code>rjags</code> , the default is 4.
nadapt	the iteration number of adaptation in the MCMC. We recommend using at least the default value 1e4 to obtained fully adapted chains.
M	the number of draws in each chain, with no thinning. The default is M=5e2 but suggest using 5e5.
thn	the thinning interval between mixing. The total number of draws thus would become $\text{round}(M/\text{thn}) * \text{nchain}$. The default is 10.
nburnin	the burn-in period. The default is 2e2 but suggest 2e5.
dic	logical, whether compute the DIC (deviance information criterion) for model selection.
death_in_R	the numeric value of average of cumulative deaths in the removed compartments. The default is 0.4 within Hubei and 0.02 outside Hubei.
beta0	the hyperparameter of average transmission rate, the default is the one estimated from the SARS first-month outbreak (0.2586).
gamma0	the hyperparameter of average removed rate, the default is the one estimated from the SARS first-month outbreak (0.0821).
R0	the hyperparameter of the mean reproduction number R0. The default is thus the ratio of beta0/gamma0, which can be specified directly.
gamma0_sd	the standard deviation for the prior distribution of the removed rate γ , the default is 0.1.
R0_sd	the standard deviation for the prior disbution of R0, the default is 1.
casename	the string of the job's name. The default is "tvt.eSIR".
file_add	the string to denote the location of saving output files and tables.
save_files	logical, whether save (TRUE) results or not (FALSE). This enables to save summary tables, trace plots, and plots of the posterior means of the first-order derivatives of the infection prevalence process θ_t^I .
save_mcmc	logical, whether save (TRUE) all the MCMC outputs or not (FALSE). The output file will be an .RData file named by the <i>casename</i> . We include arrays of prevalence values of the three compartments with their matrices of posterior draws up to the last date of the collected data as <code>theta_p[, , 1]</code> and afterwards as <code>theta_pp[, , 1]</code> for θ_t^S , <code>theta_p[, , 2]</code> and <code>theta_pp[, , 2]</code> for θ_t^I , and <code>theta_p[, , 3]</code> and <code>theta_pp[, , 3]</code> for θ_t^R . Moreover, the input and predicted proportions Y, Y_pp, R and R_pp can also be retrieved. The prevalence and predicated proportion matrices have rows for MCMC replicates, and columns for days. The MCMC posterior draws of other parameters including beta_p, gamma_p, R0_p, and variance controllers k_p, lambdaY_p, lambdaR_p are also available.

Details

We fit a state-space model with extended SIR, in which a time-varying transmission rate modifier $\pi(t)$ (between 0 and 1) is introduced to model. This allows us to accommodate quarantine protocol changes and ignored resources of hospitalization. The form of reducing rate may be a step-function with jumps at times of big policy changes or a smooth exponential survival function $\exp(-\lambda_0 t)$. The parameters of the function and change points, if any, should be predefined.

Value

casename	the predefined casename.
incidence_mean	mean incidence.
incidence_ci	2.5%, 50%, and 97.5% quantiles of the incidences.
out_table	summary tables including the posterior mean of the prevalence processes of the 3 states compartments ($\theta_t^S, \theta_t^I, \theta_t^R$) at last date of data collected (t') decided by the lengths of your input data Y and R), and their respective credible intervals (ci); the respective means and ci's of the reproduction number (R_0), removed rate (γ), transmission rate (β).
plot_infection	plot of summarizing and forecasting for the infection compartment, in which the vertical blue line denotes the last date of data collected (t'), the vertical dark-gray line denotes the deacceleration point (first turning point) that the posterior mean first-derivative of infection prevalence θ_t^I achieves the maximum, the vertical purple line denotes the second turning point that the posterior mean first-derivative infection proportion θ_t^I equals zero, the darkgray line denotes the posterior mean of the infection prevalence θ_t^I and the red line denotes its posterior median.
plot_removed	plot of summarizing and forecasting for the removed compartment with lines similar to those in the <code>plot_infection</code> . The vertical lines are identical, but the horizontal mean and median correspond to the posterior mean and median of the removed process θ_t^R . An additional line indicates the estimated death prevalence from the input <code>death_in_R</code> .
first_tp_mean	the date t at which $\ddot{\theta}_t^I = 0$, calculated as the average of the time points with maximum posterior first-order derivatives $\dot{\theta}_t^I$; this value may be slightly different from the one labeled by the "darkgreen" lines in the two plots <code>plot_infection</code> and <code>plot_removed</code> , which indicate the stationary point such that the first-order derivative of the averaged posterior of θ_t^I reaches its maximum.
first_tp_ci	with <code>first_tp_mean</code> , it reports the corresponding credible interval and median.
second_tp_mean	the date t at which $\theta_t^I = 0$, calculated as the average of the stationary points of all of posterior first-order derivatives $\dot{\theta}_t^I$; this value may be slightly different from the one labeled by the "purple" lines in the plots of <code>plot_infection</code> and <code>plot_removed</code> . The latter indicate stationary t at which the first-order derivative of the averaged posterior of θ_t^I equals zero.
second_tp_ci	with <code>second_tp_mean</code> , it reports the corresponding credible interval and median.
dic_val	the output of <code>dic.sample()</code> in <code>dic.sample</code> , computing deviance information criterion for model comparison.

Examples

```

NI_complete <- c( 41,41,41,45,62,131,200,270,375,444,549, 729,
                  1052,1423,2714,3554,4903,5806,7153,9074,11177,
                  13522,16678,19665,22112,24953,27100,29631,31728,33366)
RI_complete <- c(1,1,7,10,14,20,25,31,34,45,55,71,94,121,152,213,
                  252,345,417,561,650,811,1017,1261,1485,1917,2260,
                  2725,3284,3754)

N=58.5e6
R <- RI_complete/N
Y <- NI_complete/N- R #Jan13->Feb 11
### Step function of pi(t)
change_time <- c("01/23/2020","02/04/2020","02/08/2020")
pi0<- c(1.0,0.9,0.5,0.1)
res.step <- tvt.eSIR(Y,R,begin_str="01/13/2020",death_in_R = 0.4,
                    T_fin=200,pi0=pi0,change_time=change_time,dic=T,
                    casename="Hubei_step",save_files = T,
                    save_mcmc=F,M=5e2,nburnin = 2e2)
res.step$plot_infection
res.step$plot_removed
res.step$dic_val

### continuous exponential function of pi(t)
res.exp <- tvt.eSIR(Y,R,begin_str="01/13/2020",death_in_R = 0.4,
                   T_fin=200,exponential=TRUE,dic=F,lambda0=0.05,
                   casename="Hubei_exp",save_files = F,save_mcmc=F,
                   M=5e2,nburnin = 2e2)
res.exp$plot_infection
#res.exp$plot_removed

### without pi(t), the standard state-space SIR model without intervention
res.nopi <- tvt.eSIR(Y,R,begin_str="01/13/2020",death_in_R = 0.4,
                   T_fin=200,casename="Hubei_nopi",save_files = F,
                   M=5e2,nburnin = 2e2)
res.nopi$plot_infection
#res.nopi$plot_removed

```

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