

# Package ‘SimInf’

June 8, 2022

**Title** A Framework for Data-Driven Stochastic Disease Spread Simulations

**Version** 9.1.0

**Description** Provides an efficient and very flexible framework to conduct data-driven epidemiological modeling in realistic large scale disease spread simulations. The framework integrates infection dynamics in subpopulations as continuous-time Markov chains using the Gillespie stochastic simulation algorithm and incorporates available data such as births, deaths and movements as scheduled events at predefined time-points. Using C code for the numerical solvers and 'OpenMP' (if available) to divide work over multiple processors ensures high performance when simulating a sample outcome. One of our design goals was to make the package extendable and enable usage of the numerical solvers from other R extension packages in order to facilitate complex epidemiological research. The package contains template models and can be extended with user-defined models. For more details see the paper by Widgren, Bauer, Eriksson and Engblom (2019) [doi:10.18637/jss.v091.i12](https://doi.org/10.18637/jss.v091.i12)>. The package also provides functionality to fit models to time series data using the Approximate Bayesian Computation Sequential Monte Carlo ('ABC-SMC') algorithm of Toni and others (2009) [doi:10.1098/rsif.2008.0172](https://doi.org/10.1098/rsif.2008.0172)>.

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**License** GPL-3

**URL** <https://github.com/stewid/SimInf>

**BugReports** <https://github.com/stewid/SimInf/issues>

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**LazyData** true

**Biarch** true

**NeedsCompilation** yes

**SystemRequirements** GNU Scientific Library (GSL)

**Depends** R(>= 4.0)

**Imports** digest, graphics, grDevices, MASS, methods, stats, utils,  
Matrix

**Suggests** knitr, rmarkdown

**Collate** 'C-generator.R' 'check\_arguments.R' 'init.R' 'valid.R'  
'classes.R' 'SimInf\_model.R' 'SEIR.R' 'SIR.R' 'SIS.R' 'SISe.R'  
'SISe3.R' 'SISe3\_sp.R' 'SISe\_sp.R' 'SimInf.R' 'SimInf\_events.R'  
'run.R' 'density\_ratio.R' 'abc.R' 'abc\_support.R' 'degree.R'  
'distance.R' 'distributions.R' 'match\_compartments.R'  
'mparse.R' 'n.R' 'openmp.R' 'package\_skeleton.R' 'pfilter.R'  
'plot.R' 'prevalence.R' 'print.R' 'punchcard.R' 'trajectory.R'  
'update.R'

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---

abc *Approximate Bayesian computation*

---

**Description**

Approximate Bayesian computation

**Usage**

```

abc(
  model,
  priors = NULL,
  npart = NULL,
  ninit = NULL,
  distance = NULL,
  tolerance = NULL,
  ...,
  verbose = getOption("verbose", FALSE),
  post_gen = NULL
)

## S4 method for signature 'SimInf_model'
abc(
  model,
  priors = NULL,

```

```

npart = NULL,
ninit = NULL,
distance = NULL,
tolerance = NULL,
...,
verbose = getOption("verbose", FALSE),
post_gen = NULL
)

```

## Arguments

<code>model</code>	The <code>SimInf_model</code> object to generate data from.
<code>priors</code>	The priors for the parameters to fit. Each prior is specified with a formula notation, for example, <code>beta ~ uniform(0, 1)</code> specifies that <code>beta</code> is uniformly distributed between 0 and 1. Use <code>c()</code> to provide more than one prior, for example, <code>c(beta ~ uniform(0, 1), gamma ~ normal(10, 1))</code> . The following distributions are supported: <code>gamma</code> , <code>normal</code> and <code>uniform</code> . All parameters in <code>priors</code> must be only in either <code>gdata</code> or <code>ldata</code> .
<code>npart</code>	An integer ( $>1$ ) specifying the number of particles to approximate the posterior with.
<code>ninit</code>	Specify a positive integer ( $>npart$ ) to adaptively select a sequence of tolerances using the algorithm of Simola and others (2021). The initial tolerance is adaptively selected by sampling <code>ninit</code> draws from the prior and then retain the <code>npart</code> particles with the smallest distances. Note there must be enough initial particles to satisfactorily explore the parameter space, see Simola and others (2021). If the tolerance parameter is specified, then <code>ninit</code> must be <code>NULL</code> .
<code>distance</code>	A function for calculating the summary statistics for a simulated trajectory. For each particle, the function must determine the distance and return that information. The first argument, <code>result</code> , passed to the <code>distance</code> function is the result from a run of the model with one trajectory attached to it. The second argument, <code>generation</code> , to <code>distance</code> is an integer with the generation of the particle(s). Further arguments that can passed to the <code>distance</code> function comes from <code>...</code> in the <code>abc</code> function. Depending on the underlying model structure, data for one or more particles have been generated in each call to <code>distance</code> . If the <code>model</code> only contains one node and all the parameters to fit are in <code>ldata</code> , then that node will be replicated and each of the replicated nodes represent one particle in the trajectory (see ‘Examples’). On the other hand if the model contains multiple nodes or the parameters to fit are contained in <code>gdata</code> , then the trajectory in the <code>result</code> argument represents one particle. The function can return a numeric matrix (number of particles $\times$ number of summary statistics). Or, if the <code>distance</code> contains one summary statistic, a numeric vector with the length equal to the number of particles. Note that when using adaptive tolerance selection, only one summary statistic can be used, i.e., the function must return a matrix (number of particles $\times$ 1) or a numeric vector.
<code>tolerance</code>	A numeric matrix (number of summary statistics $\times$ number of generations) where each column contains the tolerances for a generation and each row contains a sequence of gradually decreasing tolerances. Can also be a numeric vector if there is only one summary statistic. The tolerance determines the number

	of generations of ABC-SMC to run. If the <code>ninit</code> parameter is specified, then tolerance must be <code>NULL</code> .
<code>...</code>	Further arguments to be passed to <code>fn</code> .
<code>verbose</code>	prints diagnostic messages when <code>TRUE</code> . The default is to retrieve the global option <code>verbose</code> and use <code>FALSE</code> if it is not set.
<code>post_gen</code>	An optional function that, if non- <code>NULL</code> , is applied after each completed generation. The function must accept one argument of type <code>codeSimInf_abc</code> with the current state of the fitting process. This function can be useful to, for example, save and inspect intermediate results.

### Value

A `SimInf_abc` object.

### References

T. Toni, D. Welch, N. Strelkowa, A. Ipsen, and M. P. H. Stumpf. Approximate Bayesian computation scheme for parameter inference and model selection in dynamical systems. *Journal of the Royal Society Interface* **6**, 187–202, 2009. doi: [10.1098/rsif.2008.0172](https://doi.org/10.1098/rsif.2008.0172)

U. Simola, J. Cisewski-Keche, M. U. Gutmann, J. Corander. Adaptive Approximate Bayesian Computation Tolerance Selection. *Bayesian Analysis*, **16**(2), 397–423, 2021. doi: [10.1214/20-BA1211](https://doi.org/10.1214/20-BA1211)

### Examples

```
## Not run:
## Let us consider an SIR model in a closed population with N = 100
## individuals of whom one is initially infectious and the rest are
## susceptible. First, generate one realisation (with a specified
## seed) from the model with known parameters \code{beta = 0.16} and
## \code{gamma = 0.077}. Then, use \code{abc} to infer the (known)
## parameters from the simulated data.
model <- SIR(u0 = data.frame(S = 99, I = 1, R = 0),
            tspan = 1:100,
            beta = 0.16,
            gamma = 0.077)

## Run the SIR model and plot the number of infectious.
set.seed(22)
infectious <- trajectory(run(model), "I")$I
plot(infectious, type = "s")

## The distance function to accept or reject a proposal. Each node
## in the simulated trajectory (contained in the 'result' object)
## represents one proposal.
distance <- function(result, ...) {
  ## Extract the time-series of infectious in each node as a
  ## data.frame.
  sim <- trajectory(result, "I")

  ## Split the 'sim' data.frame by node and calculate the sum of the
```

```

## squared distance at each time-point for each node.
dist <- tapply(sim$I, sim$node, function(sim_infectious) {
  sum((infectious - sim_infectious)^2)
})

## Return the distance for each node. Each proposal will be
## accepted or rejected depending on if the distance is less than
## the tolerance for the current generation.
dist
}

## Fit the model parameters using ABC-SMC and adaptive tolerance
## selection. The priors for the parameters are specified using a
## formula notation. Here we use a uniform distribution for each
## parameter with lower bound = 0 and upper bound = 1. Note that we
## use a low number particles here to keep the run-time of the example
## short. In practice you would want to use many more to ensure better
## approximations.
fit <- abc(model = model,
  priors = c(beta ~ uniform(0, 1), gamma ~ uniform(0, 1)),
  npart = 100,
  ninit = 1000,
  distance = distance,
  verbose = TRUE)

## Print a brief summary.
fit

## Display the ABC posterior distribution.
plot(fit)

## End(Not run)

```

---

as.data.frame.SimInf\_abc

*Coerce to data frame*


---

## Description

Coerce to data frame

## Usage

```
## S3 method for class 'SimInf_abc'
as.data.frame(x, ...)
```

## Arguments

x                    any R object.  
...                    additional arguments to be passed to or from methods.

---

```
as.data.frame.SimInf_events
```

*Coerce events to a data frame*

---

### Description

Coerce events to a data frame

### Usage

```
## S3 method for class 'SimInf_events'
as.data.frame(x, ...)
```

### Arguments

x                    any R object.  
 ...                 additional arguments to be passed to or from methods.

---

```
boxplot, SimInf_model-method
```

*Box plot of number of individuals in each compartment*

---

### Description

Produce box-and-whisker plot(s) of the number of individuals in each model compartment.

### Usage

```
## S4 method for signature 'SimInf_model'
boxplot(x, compartments = NULL, index = NULL, ...)
```

### Arguments

x                    The model to plot  
 compartments       specify the names of the compartments to extract data from. The compartments can be specified as a character vector e.g. compartments = c('S', 'I', 'R'), or as a formula e.g. compartments = ~S+I+R (see 'Examples'). Default (compartments=NULL) includes all compartments.  
 index                indices specifying the nodes to include when plotting data. Default index = NULL include all nodes in the model.  
 ...                 Additional arguments affecting the plot produced.



**Examples**

```
## Create an 'SIR' model with 10 nodes and initialise
## it with 99 susceptible individuals and one infected
## individual. Let the model run over 100 days.
model <- SIR(u0 = data.frame(S = rep(99, 10),
                             I = rep(1, 10),
                             R = rep(0, 10)),
            tspan = 1:100,
            beta = 0.16,
            gamma = 0.077)

## Run the model and save the result.
result <- run(model)

## Create a boxplot that includes all compartments in all nodes.
boxplot(result)

## Create a boxplot that includes the S and I compartments in
## nodes 1 and 2.
boxplot(result, ~S+I, 1:2)
```

---

 continue

---

*Run more generations of ABC SMC*


---

**Description**

Run more generations of ABC SMC

**Usage**

```
continue(object, ...)

## S4 method for signature 'SimInf_abc'
continue(
  object,
  tolerance = NULL,
  ...,
  verbose = getOption("verbose", FALSE),
  post_gen = NULL
)
```

**Arguments**

object	The SimInf_abc to continue from.
...	Further arguments to be passed to the SimInf_abc@fn.

tolerance	A numeric matrix (number of summary statistics $\times$ number of generations) where each column contains the tolerances for a generation and each row contains a sequence of gradually decreasing tolerances. Can also be a numeric vector if there is only one summary statistic. The tolerance determines the number of generations of ABC-SMC to run.
verbose	prints diagnostic messages when TRUE. The default is to retrieve the global option verbose and use FALSE if it is not set.
post_gen	An optional function that, if non-NULL, is applied after each completed generation. The function must accept one argument of type codeSimInf_abc with the current state of the fitting process. This function can be useful to, for example, save and inspect intermediate results.

**Value**

A SimInf\_abc object.

---

C_code	<i>Extract the C code from a SimInf_model object</i>
--------	--

---

**Description**

Extract the C code from a SimInf\_model object

**Usage**

```
C_code(model)
```

**Arguments**

model            The SimInf\_model object to extract the C code from.

**Value**

Character vector with C code for the model.

**Examples**

```
## Use the model parser to create a 'SimInf_model' object that
## expresses an SIR model, where 'b' is the transmission rate and
## 'g' is the recovery rate.
model <- mparse(transitions = c("S -> b*S*I/(S+I+R) -> I", "I -> g*I -> R"),
               compartments = c("S", "I", "R"),
               gdata = c(b = 0.16, g = 0.077),
               u0 = data.frame(S = 99, I = 1, R = 0),
               tspan = 1:10)

## View the C code.
C_code(model)
```

---

distance_matrix	<i>Create a distance matrix between nodes for spatial models</i>
-----------------	--

---

**Description**

Calculate the euclidian distances between coordinates for all coordinates within the cutoff.

**Usage**

```
distance_matrix(x, y, cutoff, min_dist = NULL)
```

**Arguments**

x	Projected x coordinate
y	Projected y coordinate
cutoff	The distance cutoff
min_dist	The minimum distance to separate two nodes. If the coordinates for two nodes are identical, the min_dist must be assigned or an error is raised. Default is NULL i.e. to raise an error.

**Value**

dgCMatrix

**Examples**

```
## Generate a grid 10 x 10 and place one node in each cell
## separated by 100m.
nodes <- expand.grid(x = (0:9) * 100, y = (0:9) * 100)
plot(y ~ x, nodes)

## Define the cutoff to only include neighbors within 300m.
d <- distance_matrix(x = nodes$x, y = nodes$y, cutoff = 301)

## View the first 10 rows and columns in the distance matrix
d[1:10, 1:10]
```

---

events	<i>Extract the events from a SimInf_model object</i>
--------	--

---

**Description**

Extract the scheduled events from a SimInf\_model object.

**Usage**

```
events(model)

## S4 method for signature 'SimInf_model'
events(model)
```

**Arguments**

model            The model to extract the events from.

**Value**

[SimInf\\_events](#) object.

**Examples**

```
## Create an SIR model that includes scheduled events.
model <- SIR(u0 = u0_SIR(),
            tspan = 1:(4 * 365),
            events = events_SIR(),
            beta = 0.16,
            gamma = 0.077)

## Extract the scheduled events from the model and display summary
summary(events(model))

## Extract the scheduled events from the model and plot them
plot(events(model))
```

---

events\_SEIR

*Example data to initialize events for the 'SEIR' model*

---

**Description**

Example data to initialize scheduled events for a population of 1600 nodes and demonstrate the [SEIR](#) model.

**Usage**

```
events_SEIR()
```

**Details**

Example data to initialize scheduled events (see [SimInf\\_events](#)) for a population of 1600 nodes and demonstrate the [SEIR](#) model. The dataset contains 466692 events for 1600 nodes distributed over 4 \* 365 days. The events are divided into three types: 'Exit' events remove individuals from the population (n = 182535), 'Enter' events add individuals to the population (n = 182685), and 'External transfer' events move individuals between nodes in the population (n = 101472). The vignette contains a detailed description of how scheduled events operate on a model.

**Value**

A data.frame

**Examples**

```
## Create an 'SEIR' model with 1600 nodes and initialize
## it to run over 4*365 days. Add one infected individual
## to the first node.
u0 <- u0_SEIR()
u0$I[1] <- 1
tspan <- seq(from = 1, to = 4*365, by = 1)
model <- SEIR(u0      = u0,
              tspan   = tspan,
              events  = events_SEIR(),
              beta    = 0.16,
              epsilon = 0.25,
              gamma   = 0.01)

## Display the number of individuals affected by each event type
## per day.
plot(events(model))

## Run the model to generate a single stochastic trajectory.
result <- run(model)
plot(result)

## Summarize the trajectory. The summary includes the number of
## events by event type.
summary(result)
```

---

events\_SIR

*Example data to initialize events for the 'SIR' model*

---

**Description**

Example data to initialize scheduled events for a population of 1600 nodes and demonstrate the [SIR](#) model.

**Usage**

```
events_SIR()
```

**Details**

Example data to initialize scheduled events (see [SimInf\\_events](#)) for a population of 1600 nodes and demonstrate the [SIR](#) model. The dataset contains 466692 events for 1600 nodes distributed over 4 \* 365 days. The events are divided into three types: 'Exit' events remove individuals from the population (n = 182535), 'Enter' events add individuals to the population (n = 182685), and 'External transfer' events move individuals between nodes in the population (n = 101472). The vignette contains a detailed description of how scheduled events operate on a model.

**Value**

A data.frame

**Examples**

```
## Create an 'SIR' model with 1600 nodes and initialize
## it to run over 4*365 days. Add one infected individual
## to the first node.
u0 <- u0_SIR()
u0$I[1] <- 1
tspan <- seq(from = 1, to = 4*365, by = 1)
model <- SIR(u0      = u0,
             tspan   = tspan,
             events  = events_SIR(),
             beta    = 0.16,
             gamma   = 0.01)

## Display the number of individuals affected by each event type
## per day.
plot(events(model))

## Run the model to generate a single stochastic trajectory.
result <- run(model)
plot(result)

## Summarize the trajectory. The summary includes the number of
## events by event type.
summary(result)
```

---

events\_SIS

*Example data to initialize events for the 'SIS' model*

---

**Description**

Example data to initialize scheduled events for a population of 1600 nodes and demonstrate the [SIS](#) model.

**Usage**

```
events_SIS()
```

**Details**

Example data to initialize scheduled events (see [SimInf\\_events](#)) for a population of 1600 nodes and demonstrate the [SIS](#) model. The dataset contains 466692 events for 1600 nodes distributed over 4 \* 365 days. The events are divided into three types: 'Exit' events remove individuals from the population (n = 182535), 'Enter' events add individuals to the population (n = 182685), and 'External transfer' events move individuals between nodes in the population (n = 101472). The vignette contains a detailed description of how scheduled events operate on a model.

**Value**

A data.frame

**Examples**

```
## Create an 'SIS' model with 1600 nodes and initialize
## it to run over 4*365 days. Add one infected individual
## to the first node.
u0 <- u0_SIS()
u0$I[1] <- 1
tspan <- seq(from = 1, to = 4*365, by = 1)
model <- SIS(u0      = u0,
             tspan   = tspan,
             events  = events_SIS(),
             beta    = 0.16,
             gamma   = 0.01)

## Display the number of individuals affected by each event type
## per day.
plot(events(model))

## Run the model to generate a single stochastic trajectory.
result <- run(model)
plot(result)

## Summarize the trajectory. The summary includes the number of
## events by event type.
summary(result)
```

---

events\_SISe

*Example data to initialize events for the 'SISe' model*

---

**Description**

Example data to initialize scheduled events for a population of 1600 nodes and demonstrate the [SISe](#) model.

**Usage**

```
events_SISe()
```

**Details**

Example data to initialize scheduled events (see [SimInf\\_events](#)) for a population of 1600 nodes and demonstrate the [SISe](#) model. The dataset contains 466692 events for 1600 nodes distributed over 4 \* 365 days. The events are divided into three types: 'Exit' events remove individuals from the population (n = 182535), 'Enter' events add individuals to the population (n = 182685), and 'External transfer' events move individuals between nodes in the population (n = 101472). The vignette contains a detailed description of how scheduled events operate on a model.

**Value**

A data.frame

**Examples**

```
## Create an 'SISe' model with 1600 nodes and initialize
## it to run over 4*365 days. Add one infected individual
## to the first node.
u0 <- u0_SISe()
u0$I[1] <- 1
tspan <- seq(from = 1, to = 4*365, by = 1)
model <- SISe(u0 = u0, tspan = tspan, events = events_SISe(),
             phi = 0, upsilon = 1.8e-2, gamma = 0.1, alpha = 1,
             beta_t1 = 1.0e-1, beta_t2 = 1.0e-1, beta_t3 = 1.25e-1,
             beta_t4 = 1.25e-1, end_t1 = 91, end_t2 = 182,
             end_t3 = 273, end_t4 = 365, epsilon = 0)

## Display the number of individuals affected by each event type
## per day.
plot(events(model))

## Run the model to generate a single stochastic trajectory.
result <- run(model)

## Summarize the trajectory. The summary includes the number of
## events by event type.
summary(result)
```

---

events\_SISe3

*Example data to initialize events for the 'SISe3' model*

---

**Description**

Example data to initialize scheduled events for a population of 1600 nodes and demonstrate the [SISe3](#) model.

**Usage**

```
data(events_SISe3)
```

**Format**

A data.frame



## Details

Example data to initialize scheduled events (see [SimInf\\_events](#)) for a population of 1600 nodes and demonstrate the [SISe3](#) model. The dataset contains 783773 events for 1600 nodes distributed over  $4 * 365$  days. The events are divided into three types: ‘Exit’ events remove individuals from the population ( $n = 182535$ ), ‘Enter’ events add individuals to the population ( $n = 182685$ ), sQuoteInternal transfer events move individuals between compartments within one node e.g. ageing ( $n = 317081$ ), and ‘External transfer’ events move individuals between nodes in the population ( $n = 101472$ ). The vignette contains a detailed description of how scheduled events operate on a model.

## Examples

```
## Create an 'SISe3' model with 1600 nodes and initialize
## it to run over 4*365 days. Add one infected individual
## to the first node.
data("u0_SISe3", package = "SimInf")
data("events_SISe3", package = "SimInf")
u0_SISe3$I_1[1] <- 1
tspan <- seq(from = 1, to = 4*365, by = 1)
model <- SISe3(u0 = u0_SISe3, tspan = tspan, events = events_SISe3,
              phi = rep(0, nrow(u0_SISe3)), epsilon_1 = 1.8e-2,
              epsilon_2 = 1.8e-2, epsilon_3 = 1.8e-2,
              gamma_1 = 0.1, gamma_2 = 0.1, gamma_3 = 0.1,
              alpha = 1, beta_t1 = 1.0e-1, beta_t2 = 1.0e-1,
              beta_t3 = 1.25e-1, beta_t4 = 1.25e-1, end_t1 = 91,
              end_t2 = 182, end_t3 = 273, end_t4 = 365, epsilon = 0)

## Display the number of individuals affected by each event type
## per day.
plot(events(model))

## Run the model to generate a single stochastic trajectory.
result <- run(model)

## Summarize the trajectory. The summary includes the number of
## events by event type.
summary(result)
```

---

gdata

---

*Extract global data from a SimInf\_model object*


---

## Description

The global data is a numeric vector that is common to all nodes. The global data vector is passed as an argument to the transition rate functions and the post time step function.

## Usage

```
gdata(model)
```



**Examples**

```
## Create an SIR model
model <- SIR(u0 = data.frame(S = 99, I = 1, R = 0),
            tspan = 1:5, beta = 0.16, gamma = 0.077)

## Set 'beta' to a new value
gdata(model, "beta") <- 2

## Extract the global data vector that is common to all nodes
gdata(model)
```

---

indegree	<i>Determine in-degree for each node in a model</i>
----------	---

---

**Description**

The number of nodes with inward *external transfer* events to each node.

**Usage**

```
indegree(model)
```

**Arguments**

model                    determine in-degree for each node in the model.

**Value**

vector with in-degree for each node.

**Examples**

```
## Create an 'SIR' model with 1600 nodes and initialize
## it with example data.
model <- SIR(u0 = u0_SIR(), tspan = 1:1460, events = events_SIR(),
            beta = 0.16, gamma = 0.077)

## Display indegree for each node in the model.
plot(indegree(model))
```

---

ldata	<i>Extract local data from a node</i>
-------	---------------------------------------

---

**Description**

The local data is a numeric vector that is specific to a node. The local data vector is passed as an argument to the transition rate functions and the post time step function.

**Usage**

```
ldata(model, node)

## S4 method for signature 'SimInf_model'
ldata(model, node)
```

**Arguments**

model	The model to get local data from.
node	index to node to extract local data from.

**Value**

a numeric vector

**Examples**

```
## Create an 'SISe' model with 1600 nodes.
model <- SISe(u0 = u0_SISe(), tspan = 1:100, events = events_SISe(),
             phi = 0, upsilon = 1.8e-2, gamma = 0.1, alpha = 1,
             beta_t1 = 1.0e-1, beta_t2 = 1.0e-1, beta_t3 = 1.25e-1,
             beta_t4 = 1.25e-1, end_t1 = c(91, 101), end_t2 = c(182, 185),
             end_t3 = c(273, 275), end_t4 = c(365, 360), epsilon = 0)

## Display local data from the first two nodes.
ldata(model, node = 1)
ldata(model, node = 2)
```

---

logLik, SimInf_pfilter-method	<i>Log likelihood</i>
-------------------------------	-----------------------

---

**Description**

Extract the estimated log likelihood from a SimInf\_pfilter object.

**Usage**

```
## S4 method for signature 'SimInf_pfilter'
logLik(object)
```

**Arguments**

object            The SimInf\_pfilter object.

**Value**

the estimated log likelihood.

---

mparse

*Model parser to define new models to run in SimInf*


---

**Description**

Describe your model in a logical way in R. mparse creates a [SimInf\\_model](#) object with your model definition that is ready to [run](#).

**Usage**

```
mparse(
  transitions = NULL,
  compartments = NULL,
  ldata = NULL,
  gdata = NULL,
  u0 = NULL,
  v0 = NULL,
  tspan = NULL,
  events = NULL,
  E = NULL,
  N = NULL,
  pts_fun = NULL
)
```

**Arguments**

transitions      character vector containing transitions on the form "X -> ... -> Y". The left (right) side is the initial (final) state and the propensity is written in between the ->-signs. The special symbol @ is reserved for the empty set. For example, transitions = c("S -> k1\*S\*I -> I", "I -> k2\*I -> R") expresses a SIR model.

compartments    contains the names of the involved compartments, for example, compartments = c("S", "I", "R").



```

    compartments = c("S", "I", "R"),
    gdata = c(beta = 0.16, gamma = 0.077),
    u0 = data.frame(S = 100, I = 1, R = 0),
    tspan = 1:100)

## Run and plot the result
set.seed(22)
result <- run(model)
plot(result)

## End(Not run)

```

---

nodes

*Example data with spatial distribution of nodes*


---

### Description

Example data to initialize a population of 1600 nodes and demonstrate various models.

### Usage

```
data(nodes)
```

### Format

A data.frame

### Examples

```

## Create an 'SIR' model with 1600 nodes and initialize
## it to run over 4*365 days. Add one infected individual
## to the first node.
u0 <- u0_SIR()
u0$I[1] <- 1
tspan <- seq(from = 1, to = 4*365, by = 1)
model <- SIR(u0      = u0,
             tspan   = tspan,
             events  = events_SIR(),
             beta    = 0.16,
             gamma   = 0.077)

## Run the model to generate a single stochastic trajectory.
result <- run(model)

## Determine nodes with one or more infected individuals in the
## trajectory. Extract the 'I' compartment and check for any
## infected individuals in each node.
infected <- colSums(trajectory(result, ~ I, format = "matrix")) > 0

## Display infected nodes in 'blue' and non-infected nodes in 'yellow'.

```

```
data("nodes", package = "SimInf")
col <- ifelse(infected, "blue", "yellow")
plot(y ~ x, nodes, col = col, pch = 20, cex = 2)
```

---

n_generations	<i>Determine the number of generations</i>
---------------	--

---

### Description

Determine the number of generations

### Usage

```
n_generations(object)

## S4 method for signature 'SimInf_abc'
n_generations(object)
```

### Arguments

object            the SimInf\_abc object to determine the number of generations for.

### Value

an integer with the number of generations.

---

n_nodes	<i>Determine the number of nodes in a model</i>
---------	---

---

### Description

Determine the number of nodes in a model

### Usage

```
n_nodes(model)

## S4 method for signature 'SimInf_model'
n_nodes(model)
```

### Arguments

model            the model object to extract the number of nodes from.

### Value

the number of nodes in the model.



**Examples**

```
## Create an 'SIR' model with 100 nodes, with 99 susceptible,
## 1 infected and 0 recovered in each node.
u0 <- data.frame(S = rep(99, 100), I = rep(1, 100), R = rep(0, 100))
model <- SIR(u0 = u0, tspan = 1:10, beta = 0.16, gamma = 0.077)

## Display the number of nodes in the model.
n_nodes(model)
```

---

outdegree	<i>Determine out-degree for each node in a model</i>
-----------	--

---

**Description**

The number nodes that are connected with *external transfer* events from each node.

**Usage**

```
outdegree(model)
```

**Arguments**

model            determine out-degree for each node in the model.

**Value**

vector with out-degree for each node.

**Examples**

```
## Create an 'SIR' model with 1600 nodes and initialize
## it with example data.
model <- SIR(u0 = u0_SIR(), tspan = 1:1460, events = events_SIR(),
            beta = 0.16, gamma = 0.077)

## Display outdegree for each node in the model.
plot(outdegree(model))
```

---

package\_skeleton      *Create a package skeleton from a SimInf\_model*

---

### Description

Describe your model in a logical way in R, then `mparse` creates a `SimInf_model` object with your model definition that can be installed as an add-on R package.

### Usage

```
package_skeleton(  
  model,  
  name = NULL,  
  path = ".",  
  author = NULL,  
  email = NULL,  
  maintainer = NULL,  
  license = "GPL-3"  
)
```

### Arguments

<code>model</code>	The model <code>SimInf_model</code> object with your model to create the package skeleton from.
<code>name</code>	Character string with the package name. It should contain only (ASCII) letters, numbers and dot, have at least two characters and start with a letter and not end in a dot. The package name is also used for the class name of the model and the directory name of the package.
<code>path</code>	Path to put the package directory in. Default is '.' i.e. the current directory.
<code>author</code>	Author of the package.
<code>email</code>	Email of the package maintainer.
<code>maintainer</code>	Maintainer of the package.
<code>license</code>	License of the package. Default is 'GPL-3'.

### Value

invisible NULL.

### References

Read the *Writing R Extensions* manual for more details.

Once you have created a *source* package you need to install it: see the *R Installation and Administration* manual, [INSTALL](#) and [install.packages](#).

---

 pairs, SimInf\_model-method

*Scatterplot of number of individuals in each compartment*


---

### Description

A matrix of scatterplots with the number of individuals in each compartment is produced. The  $i$ th scatterplot contains  $x[, i]$  plotted against  $x[, j]$ .

### Usage

```
## S4 method for signature 'SimInf_model'
pairs(x, compartments = NULL, index = NULL, ...)
```

### Arguments

x	The model to plot
compartments	specify the names of the compartments to extract data from. The compartments can be specified as a character vector e.g. <code>compartments = c('S', 'I', 'R')</code> , or as a formula e.g. <code>compartments = ~S+I+R</code> (see 'Examples'). Default ( <code>compartments=NULL</code> ) includes all compartments.
index	indices specifying the nodes to include when plotting data. Default <code>index = NULL</code> include all nodes in the model.
...	Additional arguments affecting the plot produced.

### Examples

```
## Create an 'SIR' model with 10 nodes and initialise
## it with 99 susceptible individuals and one infected
## individual. Let the model run over 100 days.
model <- SIR(u0 = data.frame(S = rep(99, 10),
                             I = rep(1, 10),
                             R = rep(0, 10)),
            tspan = 1:100,
            beta = 0.16,
            gamma = 0.077)

## Run the model and save the result.
result <- run(model)

## Create a scatter plot that includes all compartments in all
## nodes.
pairs(result)

## Create a scatter plot that includes the S and I compartments in
## nodes 1 and 2.
pairs(result, ~S+I, 1:2)
```

---

pfilter

*Bootstrap particle filter*


---

### Description

The bootstrap filtering algorithm. Systematic resampling is performed at each observation.

### Usage

```
pfilter(model, obs_process, data, npart)

## S4 method for signature 'SimInf_model'
pfilter(model, obs_process, data, npart)
```

### Arguments

model	The SimInf_model object to simulate data from.
obs_process	Specification of the stochastic observation process. The obs_process can be specified as a formula if the model contains only one node and there is only one data point for each time in data. The left hand side of the formula must match a column name in the data data.frame and the right hand side of the formula is a character specifying the distribution of the observation process, for example, Iobs ~ poisson(I). The following distributions are supported: $x \sim \text{binomial}(\text{size}, \text{prob})$ , $x \sim \text{poisson}(\text{rate})$ and $x \sim \text{uniform}(\text{min}, \text{max})$ . The observation process can also be a function to evaluate the probability density of the observations given the simulated states. The first argument passed to the obs_process function is the result from a run of the model and it contains one trajectory with simulated data for a time-point. The second argument to the obs_process function is a data.frame containing the rows for the specific time-point that the function is called for. Note that the function must return the log of the density.
data	A data.frame holding the time series data.
npart	An integer with the number of particles (> 1) to use at each timestep.

### Value

A SimInf\_pfilter object.

### References

N. J. Gordon, D. J. Salmond, and A. F. M. Smith. Novel Approach to Nonlinear/Non-Gaussian Bayesian State Estimation. *Radar and Signal Processing, IEE Proceedings F*, **140**(2) 107–113, 1993. doi: [10.1049/ipf2.1993.0015](https://doi.org/10.1049/ipf2.1993.0015)

**Examples**

```

## Not run:
## Let us consider an SIR model in a closed population with N = 100
## individuals of whom one is initially infectious and the rest are
## susceptible. First, generate one realisation (with a specified
## seed) from the model with known parameters 'beta = 0.16' and
## 'gamma = 0.077'. Then, use 'pfilter' to apply the bootstrap
## particle algorithm on the simulated data.
model <- SIR(u0 = data.frame(S = 99, I = 1, R = 0),
            tspan = seq(1, 100, by = 3),
            beta = 0.16,
            gamma = 0.077)

## Run the SIR model to generate simulated observed data for the
## number of infected individuals.
set.seed(22)
infected <- trajectory(run(model), "I")[, c("time", "I")]
colnames(infected) <- c("time", "Iobs")

## Use a Poisson observation process for the infected individuals, such
## that 'Iobs ~ poisson(I + 1e-6)'. A small constant '1e-6' is added to
## prevent numerical errors, since the simulated counts 'I' could be
## zero, which would result in the Poisson rate parameter being zero,
## which violates the conditions of the Poisson distribution. Use 1000
## particles.
pf <- pfilter(model,
             obs_process = Iobs ~ poisson(I + 1e-6),
             data = infected,
             npart = 1000)

## Print a brief summary.
pf

## Compare the number infected 'I' in the filtered trajectory with the
## infected 'Iobs' in the observed data.
plot(pf, ~I)
lines(Iobs ~ time, infected, col = "blue", lwd = 2, type = "s")

## End(Not run)

```

---

plot, SimInf\_abc-method

*Display the ABC posterior distribution*


---

**Description**

Display the ABC posterior distribution

**Usage**

```
## S4 method for signature 'SimInf_abc'
plot(x, y, ...)
```

**Arguments**

x	The SimInf_abc object to plot.
y	The generation to plot. The default is to display the last generation.
...	Additional arguments affecting the plot.

---

plot,SimInf\_events-method

*Display the distribution of scheduled events over time*

---

**Description**

Display the distribution of scheduled events over time

**Usage**

```
## S4 method for signature 'SimInf_events'
plot(x, frame.plot = FALSE, ...)
```

**Arguments**

x	The events data to plot.
frame.plot	Draw a frame around each plot. Default is FALSE.
...	Additional arguments affecting the plot

---

plot,SimInf\_model-method

*Display the outcome from a simulated trajectory*

---

**Description**

Plot either the median and the quantile range of the counts in all nodes, or plot the counts in specified nodes.

**Usage**

```
## S4 method for signature 'SimInf_model'
plot(
  x,
  y,
  level = 1,
  index = NULL,
  range = 0.5,
  type = "s",
  lwd = 2,
  frame.plot = FALSE,
  legend = TRUE,
  ...
)
```

**Arguments**

x	The model to plot.
y	Character vector or formula with the compartments in the model to include in the plot. Default includes all compartments in the model. Can also be a formula that specifies the compartments that define the cases with a disease or that have a specific characteristic (numerator), and the compartments that define the entire population of interest (denominator). The left-hand-side of the formula defines the cases, and the right-hand-side defines the population, for example, $I \sim S + I + R$ in a 'SIR' model (see 'Examples'). The . (dot) is expanded to all compartments, for example, $I \sim .$ is expanded to $I \sim S + I + R$ in a 'SIR' model (see 'Examples').
level	The level at which the prevalence is calculated at each time point in <code>tspan</code> . 1 (population prevalence): calculates the proportion of the individuals (cases) in the population. 2 (node prevalence): calculates the proportion of nodes with at least one case. 3 (within-node prevalence): calculates the proportion of cases within each node. Default is 1.
index	Indices specifying the nodes to include when plotting data. Plot one line for each node. Default ( <code>index = NULL</code> ) is to extract data from all nodes and plot the median count for the specified compartments.
range	Show the quantile range of the count in each compartment. Default is to show the interquartile range i.e. the middle 50% of the count in transparent color. The median value is shown in the same color. Use <code>range = 0.95</code> to show the middle 95% of the count. To display individual lines for each node, specify <code>range = FALSE</code> .
type	The type of plot to draw. The default <code>type = "s"</code> draws stair steps. See base plot for other values.
lwd	The line width. Default is 2.
frame.plot	a logical indicating whether a box should be drawn around the plot.
legend	a logical indicating whether a legend for the compartments should be added to the plot. A legend is not drawn for a prevalence plot.
...	Other graphical parameters that are passed on to the plot function.

**Examples**

```

## Create an 'SIR' model with 100 nodes and initialise
## it with 990 susceptible individuals and 10 infected
## individuals in each node. Run the model over 100 days.
model <- SIR(u0 = data.frame(S = rep(990, 100),
                             I = rep(10, 100),
                             R = rep(0, 100)),
            tspan = 1:100,
            beta = 0.16,
            gamma = 0.077)

## Run the model and save the result.
result <- run(model)

## Plot the median and interquartile range of the number
## of susceptible, infected and recovered individuals.
plot(result)

## Plot the median and the middle 95% quantile range of the
## number of susceptible, infected and recovered individuals.
plot(result, range = 0.95)

## Plot the median and interquartile range of the number
## of infected individuals.
plot(result, "I")

## Use the formula notation instead to plot the median and
## interquartile range of the number of infected individuals.
plot(result, ~I)

## Plot the number of susceptible, infected
## and recovered individuals in the first
## three nodes.
plot(result, index = 1:3, range = FALSE)

## Use plot type line instead.
plot(result, index = 1:3, range = FALSE, type = "l")

## Plot the number of infected individuals in the first node.
plot(result, "I", index = 1, range = FALSE)

## Plot the proportion of infected individuals (cases)
## in the population.
plot(result, I ~ S + I + R)

## Plot the proportion of nodes with infected individuals.
plot(result, I ~ S + I + R, level = 2)

## Plot the median and interquartile range of the proportion
## of infected individuals in each node
plot(result, I ~ S + I + R, level = 3)

```



```
## Plot the proportion of infected individuals in the first
## three nodes.
plot(result, I ~ S + I + R, level = 3, index = 1:3, range = FALSE)
```

---

```
plot,SimInf_pfilter-method
```

*Diagnostic plot of a particle filter object*

---

## Description

Diagnostic plot of a particle filter object

## Usage

```
## S4 method for signature 'SimInf_pfilter'
plot(x, y, ...)
```

## Arguments

x	The SimInf_pfilter object to plot.
y	If y is NULL or missing (default), the filtered trajectory (top) and the effective sample size (bottom) are displayed. If y is a character vector or a formula, the plot function for a SimInf_model object is called with the filtered trajectory, see <a href="#">plot,SimInf_model-method</a> for more details about the specification a plot.
...	Other graphical parameters that are passed on to the plot function.

---

```
prevalence
```

*Generic function to calculate prevalence from trajectory data*

---

## Description

Calculate the proportion of individuals with disease in the population, or the proportion of nodes with at least one diseased individual, or the proportion of individuals with disease in each node.

## Usage

```
prevalence(model, formula, level = 1, index = NULL, ...)
```

**Arguments**

model	The model with trajectory data to calculate the prevalence from.
formula	A formula that specifies the compartments that define the cases with a disease or that have a specific characteristic (numerator), and the compartments that define the entire population of interest (denominator). The left-hand-side of the formula defines the cases, and the right-hand-side defines the population, for example, $I \sim S+I+R$ in a ‘SIR’ model (see ‘Examples’). The <code>.</code> (dot) is expanded to all compartments, for example, $I \sim .$ is expanded to $I \sim S+I+R$ in a ‘SIR’ model (see ‘Examples’). The formula can also contain a condition (indicated by <code> </code> ) for each node and time step to further control the population to include in the calculation, for example, $I \sim . \mid R == 0$ to calculate the prevalence when the recovered is zero in a ‘SIR’ model. The condition must evaluate to TRUE or FALSE in each node and time step. Note that if the denominator is zero, the prevalence is NaN.
level	The level at which the prevalence is calculated at each time point in <code>tspan</code> . 1 (population prevalence): calculates the proportion of the individuals (cases) in the population. 2 (node prevalence): calculates the proportion of nodes with at least one case. 3 (within-node prevalence): calculates the proportion of cases within each node. Default is 1.
index	indices specifying the subset of nodes to include when extracting data. Default (index = NULL) is to extract data from all nodes.
...	Additional arguments, see <a href="#">prevalence, SimInf_model-method</a>

---

```
prevalence, SimInf_model-method
```

*Calculate prevalence from a model object with trajectory data*

---

**Description**

Calculate the proportion of individuals with disease in the population, or the proportion of nodes with at least one diseased individual, or the proportion of individuals with disease in each node.

**Usage**

```
## S4 method for signature 'SimInf_model'
prevalence(model, formula, level, index, format = c("data.frame", "matrix"))
```

**Arguments**

model	The model with trajectory data to calculate the prevalence from.
formula	A formula that specifies the compartments that define the cases with a disease or that have a specific characteristic (numerator), and the compartments that define the entire population of interest (denominator). The left-hand-side of the formula defines the cases, and the right-hand-side defines the population, for example, $I \sim S+I+R$ in a ‘SIR’ model (see ‘Examples’). The <code>.</code> (dot) is expanded

to all compartments, for example,  $I \sim .$  is expanded to  $I \sim S + I + R$  in a ‘SIR’ model (see ‘Examples’). The formula can also contain a condition (indicated by  $|$ ) for each node and time step to further control the population to include in the calculation, for example,  $I \sim . | R == 0$  to calculate the prevalence when the recovered is zero in a ‘SIR’ model. The condition must evaluate to TRUE or FALSE in each node and time step. Note that if the denominator is zero, the prevalence is NaN.

level	The level at which the prevalence is calculated at each time point in <code>tspan</code> . 1 (population prevalence): calculates the proportion of the individuals (cases) in the population. 2 (node prevalence): calculates the proportion of nodes with at least one case. 3 (within-node prevalence): calculates the proportion of cases within each node. Default is 1.
index	indices specifying the subset of nodes to include when extracting data. Default ( <code>index = NULL</code> ) is to extract data from all nodes.
format	The default ( <code>format = "data.frame"</code> ) is to generate a <code>data.frame</code> with one row per time-step with the prevalence. Using <code>format = "matrix"</code> returns the result as a matrix.

### Value

A `data.frame` if `format = "data.frame"`, else a matrix.

### Examples

```
## Create an 'SIR' model with 6 nodes and initialize
## it to run over 10 days.
u0 <- data.frame(S = 100:105, I = c(0, 1, 0, 2, 0, 3), R = rep(0, 6))
model <- SIR(u0 = u0, tspan = 1:10, beta = 0.16, gamma = 0.077)

## Run the model to generate a single stochastic trajectory.
result <- run(model)

## Determine the proportion of infected individuals (cases)
## in the population at the time-points in 'tspan'.
prevalence(result, I ~ S + I + R)

## Identical result is obtained with the shorthand 'I~.'
prevalence(result, I ~ .)

## Determine the proportion of nodes with infected individuals at
## the time-points in 'tspan'.
prevalence(result, I ~ S + I + R, level = 2)

## Determine the proportion of infected individuals in each node
## at the time-points in 'tspan'.
prevalence(result, I ~ S + I + R, level = 3)

## Determine the proportion of infected individuals in each node
## at the time-points in 'tspan' when the number of recovered is
## zero.
prevalence(result, I ~ S + I + R | R == 0, level = 3)
```

---

punchcard<-                      *Set a template for where to record result during a simulation*

---

## Description

Using a sparse result matrix can save a lot of memory if the model contains many nodes and time-points, but where only a few of the data points are of interest for post-processing.

## Usage

```
punchcard(model) <- value

## S4 replacement method for signature 'SimInf_model'
punchcard(model) <- value
```

## Arguments

model	The model to set a template for where to record result.
value	A data.frame that specify the nodes, time-points and compartments to record the number of individuals at tspan. Use NULL to reset the model to record the number of individuals in each compartment in every node at each time-point in tspan.

## Details

Using a sparse result matrix can save a lot of memory if the model contains many nodes and time-points, but where only a few of the data points are of interest for post-processing. To use this feature, a template has to be defined for which data points to record. This is done using a data.frame that specifies the time-points (column 'time') and nodes (column 'node') to record the state of the compartments, see 'Examples'. The specified time-points, nodes and compartments must exist in the model, or an error is raised. Note that specifying a template only affects which data-points are recorded for post-processing, it does not affect how the solver simulates the trajectory.

## Examples

```
## Create an 'SIR' model with 6 nodes and initialize it to run over 10 days.
u0 <- data.frame(S = 100:105, I = 1:6, R = rep(0, 6))
model <- SIR(u0 = u0, tspan = 1:10, beta = 0.16, gamma = 0.077)

## Run the model.
result <- run(model)

## Display the trajectory with data for every node at each
## time-point in tspan.
trajectory(result)

## Assume we are only interested in nodes '2' and '4' at the
## time-points '3' and '5'
```

```

df <- data.frame(time = c(3, 5, 3, 5),
                 node = c(2, 2, 4, 4),
                 S = c(TRUE, TRUE, TRUE, TRUE),
                 I = c(TRUE, TRUE, TRUE, TRUE),
                 R = c(TRUE, TRUE, TRUE, TRUE))
punchcard(model) <- df
result <- run(model)
trajectory(result)

## We can also specify to record only some of the compartments in
## each time-step.
df <- data.frame(time = c(3, 5, 3, 5),
                 node = c(2, 2, 4, 4),
                 S = c(FALSE, TRUE, TRUE, TRUE),
                 I = c(TRUE, FALSE, TRUE, FALSE),
                 R = c(TRUE, FALSE, TRUE, TRUE))
punchcard(model) <- df
result <- run(model)
trajectory(result)

## A shortcut to specify to record all of the compartments in
## each time-step is to only include node and time.
df <- data.frame(time = c(3, 5, 3, 5),
                 node = c(2, 2, 4, 4))
punchcard(model) <- df
result <- run(model)
trajectory(result)

## It is possible to use an empty 'data.frame' to specify
## that no data-points should be recorded for the trajectory.
punchcard(model) <- data.frame()
result <- run(model)
trajectory(result)

## Use 'NULL' to reset the model to record data for every node at
## each time-point in tspan.
punchcard(model) <- NULL
result <- run(model)
trajectory(result)

```

---

run

*Run the SimInf stochastic simulation algorithm*


---

## Description

Run the SimInf stochastic simulation algorithm

## Usage

```
run(model, ...)
```

```
## S4 method for signature 'SimInf_model'  
run(model, solver = c("ssm", "aem"), ...)  
  
## S4 method for signature 'SEIR'  
run(model, solver = c("ssm", "aem"), ...)  
  
## S4 method for signature 'SIR'  
run(model, solver = c("ssm", "aem"), ...)  
  
## S4 method for signature 'SIS'  
run(model, solver = c("ssm", "aem"), ...)  
  
## S4 method for signature 'SISe'  
run(model, solver = c("ssm", "aem"), ...)  
  
## S4 method for signature 'SISe3'  
run(model, solver = c("ssm", "aem"), ...)  
  
## S4 method for signature 'SISe3_sp'  
run(model, solver = c("ssm", "aem"), ...)  
  
## S4 method for signature 'SISe_sp'  
run(model, solver = c("ssm", "aem"), ...)  
  
## S4 method for signature 'SimInf_abc'  
run(model, ...)
```

### Arguments

model	The SimInf model to run.
...	Additional arguments.
solver	Which numerical solver to utilize. Default is 'ssm'.

### Value

[SimInf\\_model](#) object with result from simulation.

### References

- S. Widgren, P. Bauer, R. Eriksson and S. Engblom. **SimInf**: An R Package for Data-Driven Stochastic Disease Spread Simulations. *Journal of Statistical Software*, **91**(12), 1–42. doi: [10.18637/jss.v091.i12](https://doi.org/10.18637/jss.v091.i12). An updated version of this paper is available as a vignette in the package.
- P. Bauer, S. Engblom and S. Widgren. Fast Event-Based Epidemiological Simulations on National Scales. *International Journal of High Performance Computing Applications*, **30**(4), 438–453, 2016. doi: [10.1177/1094342016635723](https://doi.org/10.1177/1094342016635723)
- P. Bauer and S. Engblom. Sensitivity Estimation and Inverse Problems in Spatial Stochastic Models of Chemical Kinetics. In: A. Abdulle, S. Deparis, D. Kressner, F. Nobile and M. Picasso (eds.),

*Numerical Mathematics and Advanced Applications - ENUMATH 2013*, pp. 519–527, Lecture Notes in Computational Science and Engineering, vol 103. Springer, Cham, 2015. doi: [10.1007/9783319107059\\_51](https://doi.org/10.1007/9783319107059_51)

## Examples

```
## Create an 'SIR' model with 10 nodes and initialise
## it to run over 100 days.
model <- SIR(u0 = data.frame(S = rep(99, 10),
                             I = rep(1, 10),
                             R = rep(0, 10)),
            tspan = 1:100,
            beta = 0.16,
            gamma = 0.077)

## Run the model and save the result.
result <- run(model)

## Plot the proportion of susceptible, infected and recovered
## individuals.
plot(result)
```

---

SEIR

*Create an SEIR model*

---

## Description

Create an SEIR model to be used by the simulation framework.

## Usage

```
SEIR(u0, tspan, events = NULL, beta = NULL, epsilon = NULL, gamma = NULL)
```

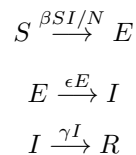
## Arguments

<code>u0</code>	A <code>data.frame</code> with the initial state in each node (see ‘Details’).
<code>tspan</code>	A vector (length $\geq 1$ ) of increasing time points where the state of each node is to be returned. Can be either an integer or a Date vector. A Date vector is coerced to a numeric vector as days, where <code>tspan[1]</code> becomes the day of the year of the first year of <code>tspan</code> . The dates are added as names to the numeric vector.
<code>events</code>	a <code>data.frame</code> with the scheduled events, see <a href="#">SimInf_model</a> .
<code>beta</code>	A numeric vector with the transmission rate from susceptible to infected where each node can have a different beta value. The vector must have length 1 or <code>nrow(u0)</code> . If the vector has length 1, but the model contains more nodes, the beta value is repeated in all nodes.

epsilon	A numeric vector with the incubation rate from exposed to infected where each node can have a different epsilon value. The vector must have length 1 or <code>nrow(u0)</code> . If the vector has length 1, but the model contains more nodes, the epsilon value is repeated in all nodes.
gamma	A numeric vector with the recovery rate from infected to recovered where each node can have a different gamma value. The vector must have length 1 or <code>nrow(u0)</code> . If the vector has length 1, but the model contains more nodes, the beta value is repeated in all nodes.

### Details

The SEIR model contains four compartments; number of susceptible (S), number of exposed (E) (those who have been infected but are not yet infectious), number of infectious (I), and number of recovered (R). Moreover, it has three state transitions,



where  $\beta$  is the transmission rate,  $\epsilon$  is the incubation rate,  $\gamma$  is the recovery rate, and  $N = S + E + I + R$ .

The argument `u0` must be a `data.frame` with one row for each node with the following columns:

- S** The number of susceptible in each node
- E** The number of exposed in each node
- I** The number of infected in each node
- R** The number of recovered in each node

### Value

A `SimInf_model` of class SEIR

### Examples

```
## Create a SEIR model object.
model <- SEIR(u0 = data.frame(S = 99, E = 0, I = 1, R = 0),
             tspan = 1:100,
             beta = 0.16,
             epsilon = 0.25,
             gamma = 0.077)

## Run the SEIR model and plot the result.
set.seed(3)
result <- run(model)
plot(result)
```



---

SEIR-class	<i>Definition of the 'SEIR' model</i>
------------	---------------------------------------

---

### Description

Class to handle the SEIR [SimInf\\_model](#).

---

select_matrix	<i>Extract the select matrix from a SimInf_model object</i>
---------------	---

---

### Description

Utility function to extract events@E from a SimInf\_model object, see [SimInf\\_events](#)

### Usage

```
select_matrix(model)

## S4 method for signature 'SimInf_model'
select_matrix(model)
```

### Arguments

model            The model to extract the select matrix E from.

### Value

[dgMatrix](#) object.

### Examples

```
## Create an SIR model
model <- SIR(u0 = data.frame(S = 99, I = 1, R = 0),
            tspan = 1:5, beta = 0.16, gamma = 0.077)

## Extract the select matrix from the model
select_matrix(model)
```

---

```
select_matrix<-          Set the select matrix for a SimInf_model object
```

---

### Description

Utility function to set events@E in a SimInf\_model object, see [SimInf\\_events](#)

### Usage

```
select_matrix(model) <- value

## S4 replacement method for signature 'SimInf_model'
select_matrix(model) <- value
```

### Arguments

model            The model to set the select matrix for.  
value            A matrix.

### Examples

```
## Create an SIR model
model <- SIR(u0 = data.frame(S = 99, I = 1, R = 0),
            tspan = 1:5, beta = 0.16, gamma = 0.077)

## Set the select matrix
select_matrix(model) <- matrix(c(1, 0, 0, 1, 1, 1, 0, 0, 1), nrow = 3)

## Extract the select matrix from the model
select_matrix(model)
```

---

```
set_num_threads          Specify the number of threads that SimInf should use
```

---

### Description

Set the number of threads to be used in SimInf code that is parallelized with OpenMP (if available). The number of threads is initialized when SimInf is first loaded in the R session using optional environment variables (see ‘Details’). It is also possible to specify the number of threads by calling set\_num\_threads. If the environment variables that affect the number of threads change, then set\_num\_threads must be called again for it to take effect.

### Usage

```
set_num_threads(threads = NULL)
```

**Arguments**

threads            integer with maximum number of threads to use in functions that are parallelized with OpenMP (if available). Default is NULL, i.e. to use all available processors and then check for limits in the environment variables (see 'Details').

**Details**

The `omp_get_num_procs()` function is used to determine the number of processors that are available to the device at the time the routine is called. The number of threads is then limited by `omp_get_thread_limit()` and the current values of the environmental variables (if set)

- `Sys.getenv("OMP_THREAD_LIMIT")`
- `Sys.getenv("OMP_NUM_THREADS")`
- `Sys.getenv("SIMINF_NUM_THREADS")`

Additionally, the maximum number of threads can be controlled by the `threads` argument, given that its value is not above any of the limits described above.

**Value**

The previous value is returned (invisible).

---

<code>shift_matrix</code>	<i>Extract the shift matrix from a <code>SimInf_model</code> object</i>
---------------------------	---

---

**Description**

Utility function to extract the shift matrix `events@N` from a `SimInf_model` object, see [SimInf\\_events](#)

**Usage**

```
shift_matrix(model)

## S4 method for signature 'SimInf_model'
shift_matrix(model)
```

**Arguments**

model            The model to extract the shift matrix `events@N` from.

**Value**

A matrix.

**Examples**

```
## Create an SIR model
model <- SIR(u0 = data.frame(S = 99, I = 1, R = 0),
            tspan = 1:5, beta = 0.16, gamma = 0.077)

## Extract the shift matrix from the model
shift_matrix(model)
```

---

shift\_matrix<-                    *Set the shift matrix for a SimInf\_model object*

---

**Description**

Utility function to set events@N in a SimInf\_model object, see [SimInf\\_events](#)

**Usage**

```
shift_matrix(model) <- value

## S4 replacement method for signature 'SimInf_model'
shift_matrix(model) <- value
```

**Arguments**

model	The model to set the shift matrix events@N.
value	A matrix.

**Value**

SimInf\_model object

**Examples**

```
## Create an SIR model
model <- SIR(u0 = data.frame(S = 99, I = 1, R = 0),
            tspan = 1:5, beta = 0.16, gamma = 0.077)

## Set the shift matrix
shift_matrix(model) <- matrix(c(2, 1, 0), nrow = 3)

## Extract the shift matrix from the model
shift_matrix(model)
```

---

show, SimInf\_abc-method

*Print summary of a SimInf\_abc object*

---

### **Description**

Print summary of a SimInf\_abc object

### **Usage**

```
## S4 method for signature 'SimInf_abc'  
show(object)
```

### **Arguments**

object            The SimInf\_abc object.

### **Value**

invisible(object).

---

show, SimInf\_events-method

*Brief summary of SimInf\_events*

---

### **Description**

Shows the number of scheduled events.

### **Usage**

```
## S4 method for signature 'SimInf_events'  
show(object)
```

### **Arguments**

object            The SimInf\_events object

### **Value**

None (invisible 'NULL').

---

show,SimInf\_model-method

*Brief summary of SimInf\_model*

---

## Description

Brief summary of SimInf\_model

## Usage

```
## S4 method for signature 'SimInf_model'  
show(object)
```

## Arguments

object            The SimInf\_model object

## Value

None (invisible 'NULL').

## Examples

```
## Create an 'SIR' model with 10 nodes and initialise  
## it to run over 100 days.  
model <- SIR(u0 = data.frame(S = rep(99, 10),  
                              I = rep(1, 10),  
                              R = rep(0, 10)),  
            tspan = 1:100,  
            beta = 0.16,  
            gamma = 0.077)  
  
## Brief summary of the model  
model  
  
## Run the model and save the result  
result <- run(model)  
  
## Brief summary of the result. Note that 'U' and 'V' are  
## non-empty after running the model.  
result
```

---

 show, SimInf\_pfilter-method

*Brief summary of a SimInf\_pfilter object*


---

### Description

Brief summary of a SimInf\_pfilter object

### Usage

```
## S4 method for signature 'SimInf_pfilter'
show(object)
```

### Arguments

object            The SimInf\_pfilter object.

### Value

invisible(object).

---

SimInf

*A Framework for Data-Driven Stochastic Disease Spread Simulations*


---

### Description

The SimInf package provides a flexible framework for data-driven spatio-temporal disease spread modeling, designed to efficiently handle population demographics and network data. The framework integrates infection dynamics in each subpopulation as continuous-time Markov chains (CTMC) using the Gillespie stochastic simulation algorithm (SSA) and incorporates available data such as births, deaths or movements as scheduled events. A scheduled event is used to modify the state of a subpopulation at a predefined time-point.

### Details

The [SimInf\\_model](#) is central and provides the basis for the framework. A [SimInf\\_model](#) object supplies the state-change matrix, the dependency graph, the scheduled events, and the initial state of the system.

All predefined models in SimInf have a generating function, with the same name as the model, for example [SIR](#).

A model can also be created from a model specification using the [mparse](#) method.

After a model is created, a simulation is started with a call to the [run](#) method and if execution is successful, it returns a modified [SimInf\\_model](#) object with a single stochastic solution trajectory attached to it.

SimInf provides several utility functions to inspect simulated data, for example, `show`, `summary` and `plot`. To facilitate custom analysis, it provides the `trajectory`, `SimInf_model-method` and `prevalence` methods.

One of our design goal was to make SimInf extendable and enable usage of the numerical solvers from other R extension packages in order to facilitate complex epidemiological research. To support this, SimInf has functionality to generate the required C and R code from a model specification, see `package_skeleton`

## References

S. Widgren, P. Bauer, R. Eriksson and S. Engblom. **SimInf**: An R Package for Data-Driven Stochastic Disease Spread Simulations. *Journal of Statistical Software*, **91**(12), 1–42. doi: [10.18637/jss.v091.i12](https://doi.org/10.18637/jss.v091.i12). An updated version of this paper is available as a vignette in the package.

---

SimInf_abc-class	Class "SimInf_abc"
------------------	--------------------

---

## Description

Class "SimInf\_abc"

## Slots

`model` The `SimInf_model` object to estimate parameters in.

`priors` A data.frame containing the four columns `parameter`, `distribution`, `p1` and `p2`. The column `parameter` gives the name of the parameter referred to in the model. The column `distribution` contains the name of the prior distribution. Valid distributions are `'gamma'`, `'normal'` or `'uniform'`. The column `p1` is a numeric vector with the first hyperparameter for each prior: `'gamma'`) `shape`, `'normal'`) `mean`, and `'uniform'`) `lower bound`. The column `p2` is a numeric vector with the second hyperparameter for each prior: `'gamma'`) `rate`, `'normal'`) `standard deviation`, and `'uniform'`) `upper bound`.

`target` Character vector (`gdata` or `ldata`) that determines if the ABC-SMC method estimates parameters in `model@gdata` or in `model@ldata`.

`pars` Index to the parameters in `target`.

`nprop` An integer vector with the number of simulated proposals in each generation.

`fn` A function for calculating the summary statistics for the simulated trajectory and determine the distance for each particle, see `abc` for more details.

`tolerance` A numeric matrix (number of summary statistics  $\times$  number of generations) where each column contains the tolerances for a generation and each row contains a sequence of gradually decreasing tolerances.

`x` A numeric array (number of particles  $\times$  number of parameters  $\times$  number of generations) with the parameter values for the accepted particles in each generation. Each row is one particle.

`weight` A numeric matrix (number of particles  $\times$  number of generations) with the weights for the particles `x` in the corresponding generation.



**distance** A numeric array (number of particles  $\times$  number of summary statistics  $\times$  number of generations) with the distance for the particles  $x$  in each generation. Each row contains the distance for a particle and each column contains the distance for a summary statistic.

**ess** A numeric vector with the effective sample size (ESS) in each generation. The effective sample size is computed as

$$\left( \sum_{i=1}^N (w_g^{(i)})^2 \right)^{-1},$$

where  $w_g^{(i)}$  is the normalized weight of particle  $i$  in generation  $g$ .

### See Also

[abc](#) and [continue](#).

---

SimInf\_events

Create a [SimInf\\_events](#) object

---

### Description

The argument `events` must be a data.frame with the following columns:

**event** Four event types are supported by the current solvers: *exit*, *enter*, *internal transfer*, and *external transfer*. When assigning the events, they can either be coded as a numerical value or a character string: *exit*; 0 or 'exit', *enter*; 1 or 'enter', *internal transfer*; 2 or 'intTrans', and *external transfer*; 3 or 'extTrans'. Internally in **SimInf**, the event type is coded as a numerical value.

**time** When the event occurs i.e., the event is processed when time is reached in the simulation. Can be either an integer or a Date vector. A Date vector is coerced to a numeric vector as days, where `t0` determines the offset to match the time of the events to the model `tspan` vector.

**node** The node that the event operates on. Also the source node for an *external transfer* event.  $1 \leq \text{node}[i] \leq \text{Number of nodes}$ .

**dest** The destination node for an *external transfer* event i.e., individuals are moved from node to dest, where  $1 \leq \text{dest}[i] \leq \text{Number of nodes}$ . Set `event = 0` for the other event types. `dest` is an integer vector.

**n** The number of individuals affected by the event.  $n[i] \geq 0$ .

**proportion** If `n[i]` equals zero, the number of individuals affected by `event[i]` is calculated by sampling the number of individuals from a binomial distribution using the `proportion[i]` and the number of individuals in the compartments. Numeric vector.  $0 \leq \text{proportion}[i] \leq 1$ .

**select** To process an `event[i]`, the compartments affected by the event are specified with `select[i]` together with the matrix `E`, where `select[i]` determines which column in `E` to use. The specific individuals affected by the event are sampled from the compartments corresponding to the non-zero entries in the specified column in `E[, select[i]]`, where `select` is an integer vector.

**shift** Determines how individuals in *internal transfer* and *external transfer* events are shifted to enter another compartment. The sampled individuals are shifted according to column `shift[i]` in matrix `N` i.e., `N[, shift[i]]`, where `shift` is an integer vector. See above for a description of `N`. Unused for the other event types.

**Usage**

```
SimInf_events(E = NULL, N = NULL, events = NULL, t0 = NULL)
```

**Arguments**

E	Each row corresponds to one compartment in the model. The non-zero entries in a column indicates the compartments to include in an event. For the <i>exit</i> , <i>internal transfer</i> and <i>external transfer</i> events, a non-zero entry indicate the compartments to sample individuals from. For the <i>enter</i> event, all individuals enter first non-zero compartment. E is sparse matrix of class <code>dgCMatrix</code> .
N	Determines how individuals in <i>internal transfer</i> and <i>external transfer</i> events are shifted to enter another compartment. Each row corresponds to one compartment in the model. The values in a column are added to the current compartment of sampled individuals to specify the destination compartment, for example, a value of 1 in an entry means that sampled individuals in this compartment are moved to the next compartment. Which column to use for each event is specified by the <code>shift</code> vector (see below). N is an integer matrix.
events	A data.frame with events.
t0	If <code>events\$time</code> is a Date vector, then <code>t0</code> determines the offset to match the time of the events to the model <code>tspan</code> vector, see details. If <code>events\$time</code> is a numeric vector, then <code>t0</code> must be NULL.

**Value**

S4 class `SimInf_events`

**Examples**

```
## Let us illustrate how movement events can be used to transfer
## individuals from one node to another. Use the built-in SIR
## model and start with 2 nodes where all individuals are in the
## first node (100 per compartment).
u0 <- data.frame(S = c(100, 0), I = c(100, 0), R = c(100, 0))

## Then create 300 movement events to transfer all individuals,
## one per day, from the first node to the second node. Use the
## fourth column in the select matrix where all compartments
## can be sampled with equal weight.
events <- data.frame(event      = rep("extTrans", 300),
                     time      = 1:300,
                     node      = 1,
                     dest      = 2,
                     n          = 1,
                     proportion = 0,
                     select     = 4,
                     shift     = 0)

## Create an SIR model without disease transmission to
## demonstrate the events.
```

```

model <- SIR(u0      = u0,
            tspan   = 1:300,
            events   = events,
            beta    = 0,
            gamma   = 0)

## Run the model and plot the number of individuals in
## the second node. As can be seen in the figure, all
## individuals have been moved to the second node when
## t = 300.
plot(run(model), index = 1:2, range = FALSE)

## Let us now double the weight to sample from the 'I'
## compartment and rerun the model.
model@events@E[2, 4] <- 2
plot(run(model), index = 1:2, range = FALSE)

## And much larger weight to sample from the I compartment.
model@events@E[2, 4] <- 10
plot(run(model), index = 1:2, range = FALSE)

## Increase the weight for the R compartment.
model@events@E[3, 4] <- 4
plot(run(model), index = 1:2, range = FALSE)

```

---

SimInf\_events-class    Class "SimInf\_events"

---

## Description

Class to hold data for scheduled events to modify the discrete state of individuals in a node at a pre-defined time  $t$ .

## Slots

**E** Each row corresponds to one compartment in the model. The non-zero entries in a column indicates the compartments to include in an event. For the *exit*, *internal transfer* and *external transfer* events, a non-zero entry indicate the compartments to sample individuals from. For the *enter* event, all individuals enter first non-zero compartment. E is sparse matrix of class `dgCMatrix`.

**N** Determines how individuals in *internal transfer* and *external transfer* events are shifted to enter another compartment. Each row corresponds to one compartment in the model. The values in a column are added to the current compartment of sampled individuals to specify the destination compartment, for example, a value of 1 in an entry means that sampled individuals in this compartment are moved to the next compartment. Which column to use for each event is specified by the shift vector (see below). N is an integer matrix.

**event** Type of event: 0) *exit*, 1) *enter*, 2) *internal transfer*, and 3) *external transfer*. Other values are reserved for future event types and not supported by the current solvers. Integer vector.

- time** Time of when the event occurs i.e., the event is processed when time is reached in the simulation. `time` is an integer vector.
- node** The node that the event operates on. Also the source node for an *external transfer* event. Integer vector.  $1 \leq \text{node}[i] \leq \text{Number of nodes}$ .
- dest** The destination node for an *external transfer* event i.e., individuals are moved from `node` to `dest`, where  $1 \leq \text{dest}[i] \leq \text{Number of nodes}$ . Set `event = 0` for the other event types. `dest` is an integer vector.
- n** The number of individuals affected by the event. Integer vector.  $n[i] \geq 0$ .
- proportion** If  $n[i]$  equals zero, the number of individuals affected by `event[i]` is calculated by sampling the number of individuals from a binomial distribution using the `proportion[i]` and the number of individuals in the compartments. Numeric vector.  $0 \leq \text{proportion}[i] \leq 1$ .
- select** To process `event[i]`, the compartments affected by the event are specified with `select[i]` together with the matrix `E`, where `select[i]` determines which column in `E` to use. The specific individuals affected by the event are proportionally sampled from the compartments corresponding to the non-zero entries in the specified column in `E[, select[i]]`, where `select` is an integer vector.
- shift** Determines how individuals in *internal transfer* and *external transfer* events are shifted to enter another compartment. The sampled individuals are shifted according to column `shift[i]` in matrix `N` i.e., `N[, shift[i]]`, where `shift` is an integer vector. See above for a description of `N`. Unsued for the other event types.

---

 SimInf\_model

 Create a SimInf\_model
 

---

## Description

Create a SimInf\_model

## Usage

```

SimInf_model(
  G,
  S,
  tspan,
  events = NULL,
  ldata = NULL,
  gdata = NULL,
  U = NULL,
  u0 = NULL,
  v0 = NULL,
  V = NULL,
  E = NULL,
  N = NULL,
  C_code = NULL
)

```

**Arguments**

G	Dependency graph that indicates the transition rates that need to be updated after a given state transition has occurred. A non-zero entry in element $G[i, i]$ indicates that transition rate $i$ needs to be recalculated if the state transition $j$ occurs. Sparse matrix ( $Nt \times Nt$ ) of object class <code>dgMatrix</code> .
S	Each column corresponds to a transition, and execution of state transition $j$ amounts to adding the $S[, j]$ to the state vector of the node where the state transition occurred. Sparse matrix ( $Nc \times Nt$ ) of object class <code>dgMatrix</code> .
tspan	A vector (length $\geq 1$ ) of increasing time points where the state of each node is to be returned. Can be either an integer or a Date vector. A Date vector is coerced to a numeric vector as days, where $tspan[1]$ becomes the day of the year of the first year of $tspan$ . The dates are added as names to the numeric vector.
events	A data.frame with the scheduled events.
ldata	local data for the nodes. Can either be specified as a data.frame with one row per node. Or as a matrix where each column $ldata[, j]$ contains the local data vector for the node $j$ . The local data vector is passed as an argument to the transition rate functions and the post time step function.
gdata	A numeric vector with global data that is common to all nodes. The global data vector is passed as an argument to the transition rate functions and the post time step function.
U	The result matrix with the number of individuals in each disease state in every node ( $N_n N_c \times \text{length}(tspan)$ ). $U[, j]$ contains the number of individuals in each disease state at $tspan[j]$ . $U[1:Nc, j]$ contains the state of node 1 at $tspan[j]$ . $U[(Nc + 1):(2 * Nc), j]$ contains the state of node 2 at $tspan[j]$ etc.
u0	The initial state vector. Either a matrix ( $N_c \times N_n$ ) or a data.frame with the number of individuals in each compartment in every node.
v0	The initial continuous state vector in every node. ( $\text{dim}(ldata)[1] \times N_n$ ). The continuous state vector is updated by the specific model during the simulation in the post time step function.
V	The result matrix for the real-valued continuous compartment state ( $N_n \text{dim}(ldata)[1] \times \text{length}(tspan)$ ). $V[, j]$ contains the real-valued state of the system at $tspan[j]$ .
E	Sparse matrix to handle scheduled events, see <a href="#">SimInf_events</a> .
N	Sparse matrix to handle scheduled events, see <a href="#">SimInf_events</a> .
C_code	Character vector with optional model C code. If non-empty, the C code is written to a temporary C-file when the run method is called. The temporary C-file is compiled and the resulting DLL is dynamically loaded. The DLL is unloaded and the temporary files are removed after running the model.

**Value**

[SimInf\\_model](#)

---

SimInf\_model-class      Class "SimInf\_model"

---

### Description

Class to handle data for the SimInf\_model.

### Slots

- G Dependency graph that indicates the transition rates that need to be updated after a given state transition has occurred. A non-zero entry in element  $G[i, i]$  indicates that transition rate  $i$  needs to be recalculated if the state transition  $j$  occurs. Sparse matrix ( $Nt \times Nt$ ) of object class `dgCMatrix`.
- S Each column corresponds to a state transition, and execution of state transition  $j$  amounts to adding the  $S[, j]$  column to the state vector  $u[, i]$  of node  $i$  where the transition occurred. Sparse matrix ( $Nc \times Nt$ ) of object class `dgCMatrix`.
- U The result matrix with the number of individuals in each compartment in every node.  $U[, j]$  contains the number of individuals in each compartment at  $tspan[j]$ .  $U[1:Nc, j]$  contains the number of individuals in node 1 at  $tspan[j]$ .  $U[(Nc + 1):(2 * Nc), j]$  contains the number of individuals in node 2 at  $tspan[j]$  etc. Integer matrix ( $N_n N_c \times \text{length}(tspan)$ ).
- U\_sparse If the model was configured to write the solution to a sparse matrix (`dgCMatrix`) the `U_sparse` contains the data and `U` is empty. The layout of the data in `U_sparse` is identical to `U`. Please note that `U_sparse` is numeric and `U` is integer.
- V The result matrix for the real-valued continuous state.  $V[, j]$  contains the real-valued state of the system at  $tspan[j]$ . Numeric matrix ( $N_n \text{dim}(ldata)[1] \times \text{length}(tspan)$ ).
- V\_sparse If the model was configured to write the solution to a sparse matrix (`dgCMatrix`) the `V_sparse` contains the data and `V` is empty. The layout of the data in `V_sparse` is identical to `V`.
- ldata A matrix with local data for the nodes. The column  $ldata[, j]$  contains the local data vector for the node  $j$ . The local data vector is passed as an argument to the transition rate functions and the post time step function.
- gdata A numeric vector with global data that is common to all nodes. The global data vector is passed as an argument to the transition rate functions and the post time step function.
- tspan A vector of increasing time points where the state of each node is to be returned.
- u0 The initial state vector ( $N_c \times N_n$ ) with the number of individuals in each compartment in every node.
- v0 The initial value for the real-valued continuous state. Numeric matrix ( $\text{dim}(ldata)[1] \times N_n$ ).
- events Scheduled events `SimInf_events`
- C\_code Character vector with optional model C code. If non-empty, the C code is written to a temporary C-file when the run method is called. The temporary C-file is compiled and the resulting DLL is dynamically loaded. The DLL is unloaded and the temporary files are removed after running the model.

---

SimInf\_pfilter-class    *Class "SimInf\_pfilter"*

---

### Description

Class "SimInf\_pfilter"

### Slots

`model` A `SimInf_model` object with one filtered trajectory attached.

`npart` An integer with the number of particles that was used at each timestep.

`loglik` The estimated log likelihood.

`ess` A numeric vector with the effective sample size (ESS). The effective sample size is computed as

$$\left( \sum_{i=1}^N (w_t^i)^2 \right)^{-1},$$

where  $w_t^i$  is the normalized weight of particle  $i$  at time  $t$ .

---

SIR                                      *Create an SIR model*

---

### Description

Create an SIR model to be used by the simulation framework.

### Usage

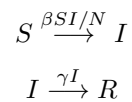
`SIR(u0, tspan, events = NULL, beta = NULL, gamma = NULL)`

### Arguments

<code>u0</code>	A data.frame with the initial state in each node (see ‘Details’).
<code>tspan</code>	A vector (length $\geq 1$ ) of increasing time points where the state of each node is to be returned. Can be either an integer or a Date vector. A Date vector is coerced to a numeric vector as days, where <code>tspan[1]</code> becomes the day of the year of the first year of <code>tspan</code> . The dates are added as names to the numeric vector.
<code>events</code>	a data.frame with the scheduled events, see <a href="#">SimInf_model</a> .
<code>beta</code>	A numeric vector with the transmission rate from susceptible to infected where each node can have a different beta value. The vector must have length 1 or <code>nrow(u0)</code> . If the vector has length 1, but the model contains more nodes, the beta value is repeated in all nodes.
<code>gamma</code>	A numeric vector with the recovery rate from infected to recovered where each node can have a different gamma value. The vector must have length 1 or <code>nrow(u0)</code> . If the vector has length 1, but the model contains more nodes, the beta value is repeated in all nodes.

**Details**

The SIR model contains three compartments; number of susceptible (S), number of infectious (I), and number of recovered (R). Moreover, it has two state transitions,



where  $\beta$  is the transmission rate,  $\gamma$  is the recovery rate, and  $N = S + I + R$ .

The argument `u0` must be a `data.frame` with one row for each node with the following columns:

**S** The number of susceptible in each node

**I** The number of infected in each node

**R** The number of recovered in each node

**Value**

A `SimInf_model` of class `SIR`

**Examples**

```
## Create an SIR model object.
model <- SIR(u0 = data.frame(S = 99, I = 1, R = 0),
            tspan = 1:100,
            beta = 0.16,
            gamma = 0.077)

## Run the SIR model and plot the result.
set.seed(22)
result <- run(model)
plot(result)
```

---

SIR-class

*Definition of the SIR model*

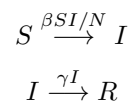
---

**Description**

Class to handle the SIR `SimInf_model`.

**Details**

The SIR model contains three compartments; number of susceptible (S), number of infectious (I), and number of recovered (R). Moreover, it has two state transitions,



where  $\beta$  is the transmission rate,  $\gamma$  is the recovery rate, and  $N = S + I + R$ .



**Examples**

```
## Create an SIR model object.
model <- SIR(u0 = data.frame(S = 99, I = 1, R = 0),
            tspan = 1:100,
            beta = 0.16,
            gamma = 0.077)

## Run the SIR model and plot the result.
set.seed(22)
result <- run(model)
plot(result)
```

SIS

*Create an SIS model***Description**

Create an SIS model to be used by the simulation framework.

**Usage**

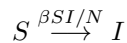
```
SIS(u0, tspan, events = NULL, beta = NULL, gamma = NULL)
```

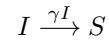
**Arguments**

<code>u0</code>	A data.frame with the initial state in each node (see ‘Details’).
<code>tspan</code>	A vector (length $\geq 1$ ) of increasing time points where the state of each node is to be returned. Can be either an integer or a Date vector. A Date vector is coerced to a numeric vector as days, where <code>tspan[1]</code> becomes the day of the year of the first year of <code>tspan</code> . The dates are added as names to the numeric vector.
<code>events</code>	a data.frame with the scheduled events, see <a href="#">SimInf_model</a> .
<code>beta</code>	A numeric vector with the transmission rate from susceptible to infected where each node can have a different beta value. The vector must have length 1 or <code>nrow(u0)</code> . If the vector has length 1, but the model contains more nodes, the beta value is repeated in all nodes.
<code>gamma</code>	A numeric vector with the recovery rate from infected to recovered where each node can have a different gamma value. The vector must have length 1 or <code>nrow(u0)</code> . If the vector has length 1, but the model contains more nodes, the beta value is repeated in all nodes.

**Details**

The SIS model contains two compartments; number of susceptible (S), and number of infectious (I). Moreover, it has two state transitions,





where  $\beta$  is the transmission rate,  $\gamma$  is the recovery rate, and  $N = S + I$ .

The argument `u0` must be a `data.frame` with one row for each node with the following columns:

**S** The number of susceptible in each node

**I** The number of infected in each node

### Value

A `SimInf_model` of class `SIS`

### Examples

```
## Create an SIS model object.
model <- SIS(u0 = data.frame(S = 99, I = 1),
            tspan = 1:100,
            beta = 0.16,
            gamma = 0.077)

## Run the SIS model and plot the result.
set.seed(22)
result <- run(model)
plot(result)
```

---

SIS-class

*Definition of the SIS model*

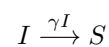
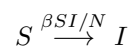
---

### Description

Class to handle the SIS `SimInf_model`.

### Details

The SIS model contains two compartments; number of susceptible (S), and number of infectious (I). Moreover, it has two state transitions,



where  $\beta$  is the transmission rate,  $\gamma$  is the recovery rate, and  $N = S + I$ .

**Examples**

```
## Create an SIS model object.
model <- SIS(u0 = data.frame(S = 99, I = 1),
            tspan = 1:100,
            beta = 0.16,
            gamma = 0.077)

## Run the SIS model and plot the result.
set.seed(22)
result <- run(model)
plot(result)
```

SISe

*Create a SISe model***Description**

Create an ‘SISe’ model to be used by the simulation framework.

**Usage**

```
SISe(
  u0,
  tspan,
  events = NULL,
  phi = NULL,
  epsilon = NULL,
  gamma = NULL,
  alpha = NULL,
  beta_t1 = NULL,
  beta_t2 = NULL,
  beta_t3 = NULL,
  beta_t4 = NULL,
  end_t1 = NULL,
  end_t2 = NULL,
  end_t3 = NULL,
  end_t4 = NULL,
  epsilon = NULL
)
```

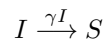
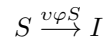
**Arguments**

<code>u0</code>	A data frame with the initial state in each node (see ‘Details’).
<code>tspan</code>	A vector (length $\geq 1$ ) of increasing time points where the state of each node is to be returned. Can be either an integer or a Date vector. A Date vector is coerced to a numeric vector as days, where <code>tspan[1]</code> becomes the day of the year of the first year of <code>tspan</code> . The dates are added as names to the numeric vector.

events	a <code>data.frame</code> with the scheduled events, see <a href="#">SimInf_model</a> .
phi	A numeric vector with the initial environmental infectious pressure in each node. Will be repeated to the length of <code>nrow(u0)</code> . Default is <code>NULL</code> which gives 0 in each node.
upsilon	Indirect transmission rate of the environmental infectious pressure
gamma	The recovery rate from infected to susceptible
alpha	Shed rate from infected individuals
beta_t1	The decay of the environmental infectious pressure in interval 1.
beta_t2	The decay of the environmental infectious pressure in interval 2.
beta_t3	The decay of the environmental infectious pressure in interval 3.
beta_t4	The decay of the environmental infectious pressure in interval 4.
end_t1	vector with the non-inclusive day of the year that ends interval 1 in each node. Will be repeated to the length of <code>nrow(u0)</code> .
end_t2	vector with the non-inclusive day of the year that ends interval 2 in each node. Will be repeated to the length of <code>nrow(u0)</code> .
end_t3	vector with the non-inclusive day of the year that ends interval 3 in each node. Will be repeated to the length of <code>nrow(u0)</code> .
end_t4	vector with the non-inclusive day of the year that ends interval 4 in each node. Will be repeated to the length of <code>nrow(u0)</code> .
epsilon	The background environmental infectious pressure

### Details

The ‘SISe’ model contains two compartments; number of susceptible ( $S$ ) and number of infectious ( $I$ ). Additionally, it contains an environmental compartment to model shedding of a pathogen to the environment. Consequently, the model has two state transitions,



where the transition rate per unit of time from susceptible to infected is proportional to the concentration of the environmental contamination  $\varphi$  in each node. Moreover, the transition rate from infected to susceptible is the recovery rate  $\gamma$ , measured per individual and per unit of time. Finally, the environmental infectious pressure in each node is evolved by,

$$\frac{d\varphi(t)}{dt} = \frac{\alpha I(t)}{N(t)} - \beta(t)\varphi(t) + \epsilon$$

where  $\alpha$  is the average shedding rate of the pathogen to the environment per infected individual and  $N = S + I$  the size of the node. The seasonal decay and removal of the pathogen is captured by  $\beta(t)$ . It is also possible to include a small background infectious pressure  $\epsilon$  to allow for other indirect sources of environmental contamination. The environmental infectious pressure  $\varphi(t)$  in each node is evolved each time unit by the Euler forward method. The value of  $\varphi(t)$  is saved at the time-points specified in `tspan`.

The argument `u0` must be a `data.frame` with one row for each node with the following columns:

**S** The number of susceptible in each node

**I** The number of infected in each node

### Value

SISe

### Beta

The time dependent beta is divided into four intervals of the year

where  $0 \leq \text{day} < 365$

Case 1:  $\text{END}_1 < \text{END}_2 < \text{END}_3 < \text{END}_4$   
 INTERVAL\_1 INTERVAL\_2 INTERVAL\_3 INTERVAL\_4 INTERVAL\_1  
 $[\text{END}_1, \text{END}_2)$   $[\text{END}_2, \text{END}_3)$   $[\text{END}_3, \text{END}_4)$   $[\text{END}_4, 365)$

Case 2:  $\text{END}_3 < \text{END}_4 < \text{END}_1 < \text{END}_2$   
 INTERVAL\_3 INTERVAL\_4 INTERVAL\_1 INTERVAL\_2 INTERVAL\_3  
 $[\text{END}_3, \text{END}_4)$   $[\text{END}_4, \text{END}_1)$   $[\text{END}_1, \text{END}_2)$   $[\text{END}_2, 365)$

Case 3:  $\text{END}_4 < \text{END}_1 < \text{END}_2 < \text{END}_3$   
 INTERVAL\_4 INTERVAL\_1 INTERVAL\_2 INTERVAL\_3 INTERVAL\_4  
 $[\text{END}_4, \text{END}_1)$   $[\text{END}_1, \text{END}_2)$   $[\text{END}_2, \text{END}_3)$   $[\text{END}_3, 365)$

---

SISe-class

*Definition of the SISe model*

---

### Description

Class to handle the SISe [SimInf\\_model](#).

---

SISe3

*Create a SISe3 model*

---

### Description

Create a SISe3 model to be used by the simulation framework.

**Usage**

```

SISe3(
  u0,
  tspan,
  events = NULL,
  phi = NULL,
  upsilon_1 = NULL,
  upsilon_2 = NULL,
  upsilon_3 = NULL,
  gamma_1 = NULL,
  gamma_2 = NULL,
  gamma_3 = NULL,
  alpha = NULL,
  beta_t1 = NULL,
  beta_t2 = NULL,
  beta_t3 = NULL,
  beta_t4 = NULL,
  end_t1 = NULL,
  end_t2 = NULL,
  end_t3 = NULL,
  end_t4 = NULL,
  epsilon = NULL
)

```

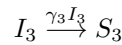
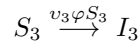
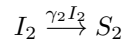
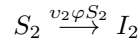
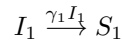
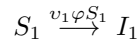
**Arguments**

<code>u0</code>	A data.frame with the initial state in each node (see ‘Details’).
<code>tspan</code>	A vector (length $\geq 1$ ) of increasing time points where the state of each node is to be returned. Can be either an integer or a Date vector. A Date vector is coerced to a numeric vector as days, where <code>tspan[1]</code> becomes the day of the year of the first year of <code>tspan</code> . The dates are added as names to the numeric vector.
<code>events</code>	a data.frame with the scheduled events, see <a href="#">SimInf_model</a> .
<code>phi</code>	A numeric vector with the initial environmental infectious pressure in each node. Will be repeated to the length of <code>nrow(u0)</code> . Default is NULL which gives 0 in each node.
<code>upsilon_1</code>	Indirect transmission rate of the environmental infectious pressure in age category 1
<code>upsilon_2</code>	Indirect transmission rate of the environmental infectious pressure in age category 2
<code>upsilon_3</code>	Indirect transmission rate of the environmental infectious pressure in age category 3
<code>gamma_1</code>	The recovery rate from infected to susceptible for age category 1
<code>gamma_2</code>	The recovery rate from infected to susceptible for age category 2
<code>gamma_3</code>	The recovery rate from infected to susceptible for age category 3

alpha	Shed rate from infected individuals
beta_t1	The decay of the environmental infectious pressure in interval 1.
beta_t2	The decay of the environmental infectious pressure in interval 2.
beta_t3	The decay of the environmental infectious pressure in interval 3.
beta_t4	The decay of the environmental infectious pressure in interval 4.
end_t1	vector with the non-inclusive day of the year that ends interval 1 in each node. Will be repeated to the length of nrow(u0).
end_t2	vector with the non-inclusive day of the year that ends interval 2 in each node. Will be repeated to the length of nrow(u0).
end_t3	vector with the non-inclusive day of the year that ends interval 3 in each node. Will be repeated to the length of nrow(u0).
end_t4	vector with the non-inclusive day of the year that ends interval 4 in each node. Will be repeated to the length of nrow(u0).
epsilon	The background environmental infectious pressure

### Details

The SISe3 model contains two compartments in three age categories; number of susceptible ( $S_1$ ,  $S_2$ ,  $S_3$ ) and number of infectious ( $I_1$ ,  $I_2$ ,  $I_3$ ). Additionally, it contains an environmental compartment to model shedding of a pathogen to the environment. Consequently, the model has six state transitions,



where the transition rate per unit of time from susceptible to infected is proportional to the concentration of the environmental contamination  $\varphi$  in each node. Moreover, the transition rate from infected to susceptible is the recovery rate  $\gamma_1, \gamma_2, \gamma_3$ , measured per individual and per unit of time. Finally, the environmental infectious pressure in each node is evolved by,

$$\frac{d\varphi(t)}{dt} = \frac{\alpha (I_1(t) + I_2(t) + I_3(t))}{N(t)} - \beta(t)\varphi(t) + \epsilon$$

where  $\alpha$  is the average shedding rate of the pathogen to the environment per infected individual and  $N = S_1 + S_2 + S_3 + I_1 + I_2 + I_3$  the size of the node. The seasonal decay and removal of the pathogen is captured by  $\beta(t)$ . It is also possible to include a small background infectious pressure  $\epsilon$  to allow for other indirect sources of environmental contamination. The environmental infectious pressure  $\varphi(t)$  in each node is evolved each time unit by the Euler forward method. The value of  $\varphi(t)$  is saved at the time-points specified in `tspan`.

The argument `u0` must be a `data.frame` with one row for each node with the following columns:

**S\_1** The number of susceptible in age category 1

**I\_1** The number of infected in age category 1

**S\_2** The number of susceptible in age category 2

**I\_2** The number of infected in age category 2

**S\_3** The number of susceptible in age category 3

**I\_3** The number of infected in age category 3

### Value

SISe3

### Beta

The time dependent beta is divided into four intervals of the year

where  $0 \leq \text{day} < 365$

Case 1:  $\text{END}_1 < \text{END}_2 < \text{END}_3 < \text{END}_4$

INTERVAL\_1 INTERVAL\_2 INTERVAL\_3 INTERVAL\_4 INTERVAL\_1  
 $[\emptyset, \text{END}_1)$   $[\text{END}_1, \text{END}_2)$   $[\text{END}_2, \text{END}_3)$   $[\text{END}_3, \text{END}_4)$   $[\text{END}_4, 365)$

Case 2:  $\text{END}_3 < \text{END}_4 < \text{END}_1 < \text{END}_2$

INTERVAL\_3 INTERVAL\_4 INTERVAL\_1 INTERVAL\_2 INTERVAL\_3  
 $[\emptyset, \text{END}_3)$   $[\text{END}_3, \text{END}_4)$   $[\text{END}_4, \text{END}_1)$   $[\text{END}_1, \text{END}_2)$   $[\text{END}_2, 365)$

Case 3:  $\text{END}_4 < \text{END}_1 < \text{END}_2 < \text{END}_3$

INTERVAL\_4 INTERVAL\_1 INTERVAL\_2 INTERVAL\_3 INTERVAL\_4  
 $[\emptyset, \text{END}_4)$   $[\text{END}_4, \text{END}_1)$   $[\text{END}_1, \text{END}_2)$   $[\text{END}_2, \text{END}_3)$   $[\text{END}_3, 365)$

---

SISe3-class

*Definition of the 'SISe3' model*

---

### Description

Class to handle the SISe3 `SimInf_model` model.



---

SISe3\_sp

*Create an SISe3\_sp model*


---

**Description**

Create an SISe3\_sp model to be used by the simulation framework.

**Usage**

```
SISe3_sp(
  u0,
  tspan,
  events = NULL,
  phi = NULL,
  epsilon_1 = NULL,
  epsilon_2 = NULL,
  epsilon_3 = NULL,
  gamma_1 = NULL,
  gamma_2 = NULL,
  gamma_3 = NULL,
  alpha = NULL,
  beta_t1 = NULL,
  beta_t2 = NULL,
  beta_t3 = NULL,
  beta_t4 = NULL,
  end_t1 = NULL,
  end_t2 = NULL,
  end_t3 = NULL,
  end_t4 = NULL,
  distance = NULL,
  coupling = NULL
)
```

**Arguments**

<code>u0</code>	A <code>data.frame</code> with the initial state in each node (see ‘Details’).
<code>tspan</code>	A vector (length $\geq 1$ ) of increasing time points where the state of each node is to be returned. Can be either an integer or a Date vector. A Date vector is coerced to a numeric vector as days, where <code>tspan[1]</code> becomes the day of the year of the first year of <code>tspan</code> . The dates are added as names to the numeric vector.
<code>events</code>	a <code>data.frame</code> with the scheduled events, see <a href="#">SimInf_model</a> .
<code>phi</code>	A numeric vector with the initial environmental infectious pressure in each node. Will be repeated to the length of <code>nrow(u0)</code> . Default is NULL which gives 0 in each node.

upsilon_1	Indirect transmission rate of the environmental infectious pressure in age category 1
upsilon_2	Indirect transmission rate of the environmental infectious pressure in age category 2
upsilon_3	Indirect transmission rate of the environmental infectious pressure in age category 3
gamma_1	The recovery rate from infected to susceptible for age category 1
gamma_2	The recovery rate from infected to susceptible for age category 2
gamma_3	The recovery rate from infected to susceptible for age category 3
alpha	Shed rate from infected individuals
beta_t1	The decay of the environmental infectious pressure in interval 1.
beta_t2	The decay of the environmental infectious pressure in interval 2.
beta_t3	The decay of the environmental infectious pressure in interval 3.
beta_t4	The decay of the environmental infectious pressure in interval 4.
end_t1	vector with the non-inclusive day of the year that ends interval 1 in each node. Will be repeated to the length of nrow(u0).
end_t2	vector with the non-inclusive day of the year that ends interval 2 in each node. Will be repeated to the length of nrow(u0).
end_t3	vector with the non-inclusive day of the year that ends interval 3 in each node. Will be repeated to the length of nrow(u0).
end_t4	vector with the non-inclusive day of the year that ends interval 4 in each node. Will be repeated to the length of nrow(u0).
distance	The distance matrix between neighboring nodes
coupling	The coupling between neighboring nodes

### Details

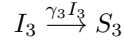
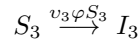
The SISe3\_sp model contains two compartments in three age categories; number of susceptible ( $S_1, S_2, S_3$ ) and number of infectious ( $I_1, I_2, I_3$ ). Additionally, it contains an environmental compartment to model shedding of a pathogen to the environment. Moreover, it also includes a spatial coupling of the environmental contamination among proximal nodes to capture between-node spread unrelated to moving infected individuals. Consequently, the model has six state transitions,

$$S_1 \xrightarrow{v_1 \varphi S_1} I_1$$

$$I_1 \xrightarrow{\gamma_1 I_1} S_1$$

$$S_2 \xrightarrow{v_2 \varphi S_2} I_2$$

$$I_2 \xrightarrow{\gamma_2 I_2} S_2$$



where the transition rate per unit of time from susceptible to infected is proportional to the concentration of the environmental contamination  $\varphi$  in each node. Moreover, the transition rate from infected to susceptible is the recovery rate  $\gamma_1, \gamma_2, \gamma_3$ , measured per individual and per unit of time. Finally, the environmental infectious pressure in each node is evolved by,

$$\frac{d\varphi_i(t)}{dt} = \frac{\alpha (I_{i,1}(t) + I_{i,2}(t) + I_{i,3}(t))}{N_i(t)} + \sum_k \frac{\varphi_k(t) N_k(t) - \varphi_i(t) N_i(t)}{N_i(t)} \cdot \frac{D}{d_{ik}} - \beta(t) \varphi_i(t)$$

where  $\alpha$  is the average shedding rate of the pathogen to the environment per infected individual and  $N = S_1 + S_2 + S_3 + I_1 + I_2 + I_3$  the size of the node. Next comes the spatial coupling among proximal nodes, where  $D$  is the rate of the local spread and  $d_{ik}$  the distance between holdings  $i$  and  $k$ . The seasonal decay and removal of the pathogen is captured by  $\beta(t)$ . The environmental infectious pressure  $\varphi(t)$  in each node is evolved each time unit by the Euler forward method. The value of  $\varphi(t)$  is saved at the time-points specified in `tspan`.

The argument `u0` must be a data.frame with one row for each node with the following columns:

- S\_1** The number of susceptible in age category 1
- I\_1** The number of infected in age category 1
- S\_2** The number of susceptible in age category 2
- I\_2** The number of infected in age category 2
- S\_3** The number of susceptible in age category 3
- I\_3** The number of infected in age category 3

### Value

SISe3\_sp

### Beta

The time dependent beta is divided into four intervals of the year

where  $0 \leq \text{day} < 365$

Case 1:  $\text{END}_1 < \text{END}_2 < \text{END}_3 < \text{END}_4$

INTERVAL\_1 INTERVAL\_2 INTERVAL\_3 INTERVAL\_4 INTERVAL\_1  
 $[\emptyset, \text{END}_1)$   $[\text{END}_1, \text{END}_2)$   $[\text{END}_2, \text{END}_3)$   $[\text{END}_3, \text{END}_4)$   $[\text{END}_4, 365)$

Case 2:  $\text{END}_3 < \text{END}_4 < \text{END}_1 < \text{END}_2$

INTERVAL\_3 INTERVAL\_4 INTERVAL\_1 INTERVAL\_2 INTERVAL\_3  
 $[\emptyset, \text{END}_3)$   $[\text{END}_3, \text{END}_4)$   $[\text{END}_4, \text{END}_1)$   $[\text{END}_1, \text{END}_2)$   $[\text{END}_2, 365)$

Case 3:  $END_4 < END_1 < END_2 < END_3$   
 INTERVAL\_4 INTERVAL\_1 INTERVAL\_2 INTERVAL\_3 INTERVAL\_4  
 $[\emptyset, END_4)$   $[END_4, END_1)$   $[END_1, END_2)$   $[END_2, END_3)$   $[END_3, 365)$

---

SISe3\_sp-class      *Definition of the 'SISe3\_sp' model*

---

### Description

Class to handle the SISe3\_sp [SimInf\\_model](#) model.

---

SISe\_sp      *Create a SISe\_sp model*

---

### Description

Create a SISe\_sp model to be used by the simulation framework.

### Usage

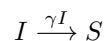
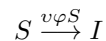
```
SISe_sp(
  u0,
  tspan,
  events = NULL,
  phi = NULL,
  epsilon = NULL,
  gamma = NULL,
  alpha = NULL,
  beta_t1 = NULL,
  beta_t2 = NULL,
  beta_t3 = NULL,
  beta_t4 = NULL,
  end_t1 = NULL,
  end_t2 = NULL,
  end_t3 = NULL,
  end_t4 = NULL,
  coupling = NULL,
  distance = NULL
)
```

**Arguments**

<code>u0</code>	A <code>data.frame</code> with the initial state in each node (see ‘Details’).
<code>tspan</code>	A vector (length $\geq 1$ ) of increasing time points where the state of each node is to be returned. Can be either an <code>integer</code> or a <code>Date</code> vector. A <code>Date</code> vector is coerced to a numeric vector as days, where <code>tspan[1]</code> becomes the day of the year of the first year of <code>tspan</code> . The dates are added as names to the numeric vector.
<code>events</code>	a <code>data.frame</code> with the scheduled events, see <a href="#">SimInf_model</a> .
<code>phi</code>	A numeric vector with the initial environmental infectious pressure in each node. Will be repeated to the length of <code>nrow(u0)</code> . Default is <code>NULL</code> which gives 0 in each node.
<code>upsilon</code>	Indirect transmission rate of the environmental infectious pressure
<code>gamma</code>	The recovery rate from infected to susceptible
<code>alpha</code>	Shed rate from infected individuals
<code>beta_t1</code>	The decay of the environmental infectious pressure in interval 1.
<code>beta_t2</code>	The decay of the environmental infectious pressure in interval 2.
<code>beta_t3</code>	The decay of the environmental infectious pressure in interval 3.
<code>beta_t4</code>	The decay of the environmental infectious pressure in interval 4.
<code>end_t1</code>	vector with the non-inclusive day of the year that ends interval 1 in each node. Will be repeated to the length of <code>nrow(u0)</code> .
<code>end_t2</code>	vector with the non-inclusive day of the year that ends interval 2 in each node. Will be repeated to the length of <code>nrow(u0)</code> .
<code>end_t3</code>	vector with the non-inclusive day of the year that ends interval 3 in each node. Will be repeated to the length of <code>nrow(u0)</code> .
<code>end_t4</code>	vector with the non-inclusive day of the year that ends interval 4 in each node. Will be repeated to the length of <code>nrow(u0)</code> .
<code>coupling</code>	The coupling between neighboring nodes
<code>distance</code>	The distance matrix between neighboring nodes

**Details**

The `SISe_sp` model contains two compartments; number of susceptible ( $S$ ) and number of infectious ( $I$ ). Additionally, it contains an environmental compartment to model shedding of a pathogen to the environment. Moreover, it also includes a spatial coupling of the environmental contamination among proximal nodes to capture between-node spread unrelated to moving infected individuals. Consequently, the model has two state transitions,



where the transition rate per unit of time from susceptible to infected is proportional to the concentration of the environmental contamination  $\varphi$  in each node. Moreover, the transition rate from

infected to susceptible is the recovery rate  $\gamma$ , measured per individual and per unit of time. Finally, the environmental infectious pressure in each node is evolved by,

$$\frac{d\varphi_i(t)}{dt} = \frac{\alpha I_i(t)}{N_i(t)} + \sum_k \frac{\varphi_k(t)N_k(t) - \varphi_i(t)N_i(t)}{N_i(t)} \cdot \frac{D}{d_{ik}} - \beta(t)\varphi_i(t)$$

where  $\alpha$  is the average shedding rate of the pathogen to the environment per infected individual and  $N = S + I$  the size of the node. Next comes the spatial coupling among proximal nodes, where  $D$  is the rate of the local spread and  $d_{ik}$  the distance between holdings  $i$  and  $k$ . The seasonal decay and removal of the pathogen is captured by  $\beta(t)$ . The environmental infectious pressure  $\varphi(t)$  in each node is evolved each time unit by the Euler forward method. The value of  $\varphi(t)$  is saved at the time-points specified in `tspan`.

The argument `u0` must be a data.frame with one row for each node with the following columns:

**S** The number of susceptible

**I** The number of infected

### Value

SISe\_sp

### Beta

The time dependent beta is divided into four intervals of the year

where  $0 \leq \text{day} < 365$

Case 1:  $\text{END}_1 < \text{END}_2 < \text{END}_3 < \text{END}_4$

INTERVAL\_1 INTERVAL\_2 INTERVAL\_3 INTERVAL\_4 INTERVAL\_1  
 $[\emptyset, \text{END}_1)$   $[\text{END}_1, \text{END}_2)$   $[\text{END}_2, \text{END}_3)$   $[\text{END}_3, \text{END}_4)$   $[\text{END}_4, 365)$

Case 2:  $\text{END}_3 < \text{END}_4 < \text{END}_1 < \text{END}_2$

INTERVAL\_3 INTERVAL\_4 INTERVAL\_1 INTERVAL\_2 INTERVAL\_3  
 $[\emptyset, \text{END}_3)$   $[\text{END}_3, \text{END}_4)$   $[\text{END}_4, \text{END}_1)$   $[\text{END}_1, \text{END}_2)$   $[\text{END}_2, 365)$

Case 3:  $\text{END}_4 < \text{END}_1 < \text{END}_2 < \text{END}_3$

INTERVAL\_4 INTERVAL\_1 INTERVAL\_2 INTERVAL\_3 INTERVAL\_4  
 $[\emptyset, \text{END}_4)$   $[\text{END}_4, \text{END}_1)$   $[\text{END}_1, \text{END}_2)$   $[\text{END}_2, \text{END}_3)$   $[\text{END}_3, 365)$

---

SISe\_sp-class

*Definition of the SISe\_sp model*

---

### Description

Class to handle the SISe\_sp [SimInf\\_model](#).

---

summary,SimInf\_abc-method

*Detailed summary of a SimInf\_abc object*

---

**Description**

Detailed summary of a SimInf\_abc object

**Usage**

```
## S4 method for signature 'SimInf_abc'  
summary(object, ...)
```

**Arguments**

object	The SimInf_abc object
...	Additional arguments affecting the summary produced.

**Value**

None (invisible 'NULL').

---

summary,SimInf\_events-method

*Detailed summary of a SimInf\_events object*

---

**Description**

Shows the number of scheduled events and the number of scheduled events per event type.

**Usage**

```
## S4 method for signature 'SimInf_events'  
summary(object, ...)
```

**Arguments**

object	The SimInf_events object
...	Additional arguments affecting the summary produced.

**Value**

None (invisible 'NULL').

---

summary,SimInf\_model-method

*Detailed summary of a SimInf\_model object*

---

### Description

Detailed summary of a SimInf\_model object

### Usage

```
## S4 method for signature 'SimInf_model'
summary(object, ...)
```

### Arguments

object	The SimInf_model object
...	Additional arguments affecting the summary produced.

### Value

None (invisible 'NULL').

---

summary,SimInf\_pfilter-method

*Detailed summary of a SimInf\_pfilter object*

---

### Description

Detailed summary of a SimInf\_pfilter object

### Usage

```
## S4 method for signature 'SimInf_pfilter'
summary(object, ...)
```

### Arguments

object	The SimInf_pfilter object.
...	Unused additional arguments.

### Value

invisible(NULL).



---

trajectory	<i>Generic function to extract data from a simulated trajectory</i>
------------	---

---

### Description

Generic function to extract data from a simulated trajectory

### Usage

```
trajectory(model, compartments = NULL, index = NULL, ...)
```

### Arguments

model	the object to extract the trajectory from.
compartments	specify the names of the compartments to extract data from. The compartments can be specified as a character vector e.g. <code>compartments = c('S', 'I', 'R')</code> , or as a formula e.g. <code>compartments = ~S+I+R</code> (see 'Examples'). Default ( <code>compartments=NULL</code> ) is to extract the number of individuals in each compartment i.e. the data from all discrete state compartments in the model. In models that also have continuous state variables e.g. the SISE model, they are also included.
index	indices specifying the subset of nodes to include when extracting data. Default ( <code>index = NULL</code> ) is to extract data from all nodes.
...	Additional arguments, see <a href="#">trajectory, SimInf_model-method</a>

---

trajectory, SimInf_model-method	<i>Extract data from a simulated trajectory</i>
---------------------------------	---

---

### Description

Extract the number of individuals in each compartment in every node after generating a single stochastic trajectory with [run](#).

### Usage

```
## S4 method for signature 'SimInf_model'
trajectory(model, compartments, index, format = c("data.frame", "matrix"))
```

**Arguments**

model	the SimInf_model object to extract the result from.
compartments	specify the names of the compartments to extract data from. The compartments can be specified as a character vector e.g. compartments = c('S', 'I', 'R'), or as a formula e.g. compartments = ~S+I+R (see 'Examples'). Default (compartments=NULL) is to extract the number of individuals in each compartment i.e. the data from all discrete state compartments in the model. In models that also have continuous state variables e.g. the SISE model, they are also included.
index	indices specifying the subset of nodes to include when extracting data. Default (index = NULL) is to extract data from all nodes.
format	the default (format = "data.frame") is to generate a data.frame with one row per node and time-step with the number of individuals in each compartment. Using format = "matrix" returns the result as a matrix, which is the internal format (see 'Details').

**Value**

A data.frame if format = "data.frame", else a matrix.

**Internal format of the discrete state variables**

Description of the layout of the internal matrix (U) that is returned if format = "matrix". U[, j] contains the number of individuals in each compartment at tspan[j]. U[1:Nc, j] contains the number of individuals in node 1 at tspan[j]. U[(Nc + 1):(2 \* Nc), j] contains the number of individuals in node 2 at tspan[j] etc, where Nc is the number of compartments in the model. The dimension of the matrix is  $N_n N_c \times \text{length}(\text{tspan})$  where  $N_n$  is the number of nodes.

**Internal format of the continuous state variables**

Description of the layout of the matrix that is returned if format = "matrix". The result matrix for the real-valued continuous state. V[, j] contains the real-valued state of the system at tspan[j]. The dimension of the matrix is  $N_n \text{dim}(\text{ldata})[1] \times \text{length}(\text{tspan})$ .

**Examples**

```
## Create an 'SIR' model with 6 nodes and initialize
## it to run over 10 days.
u0 <- data.frame(S = 100:105, I = 1:6, R = rep(0, 6))
model <- SIR(u0 = u0, tspan = 1:10, beta = 0.16, gamma = 0.077)

## Run the model to generate a single stochastic trajectory.
result <- run(model)

## Extract the number of individuals in each compartment at the
## time-points in 'tspan'.
trajectory(result)

## Extract the number of recovered individuals in the first node
```

```

## at the time-points in 'tspan'.
trajectory(result, compartments = "R", index = 1)

## Extract the number of recovered individuals in the first and
## third node at the time-points in 'tspan'.
trajectory(result, compartments = "R", index = c(1, 3))

## Create an 'SISe' model with 6 nodes and initialize
## it to run over 10 days.
u0 <- data.frame(S = 100:105, I = 1:6)
model <- SISe(u0 = u0, tspan = 1:10, phi = rep(0, 6),
  upsi = 0.02, gamma = 0.1, alpha = 1, epsilon = 1.1e-5,
  beta_t1 = 0.15, beta_t2 = 0.15, beta_t3 = 0.15, beta_t4 = 0.15,
  end_t1 = 91, end_t2 = 182, end_t3 = 273, end_t4 = 365)

## Run the model
result <- run(model)

## Extract the continuous state variable 'phi' which represents
## the environmental infectious pressure.
trajectory(result, "phi")

```

---

```
trajectory, SimInf_pfilter-method
```

*Extract filtered trajectory from running a particle filter*

---

## Description

Extract filtered trajectory from running a particle filter

## Usage

```

## S4 method for signature 'SimInf_pfilter'
trajectory(model, compartments, index, format = c("data.frame", "matrix"))

```

## Arguments

model	the SimInf_pfilter object to extract the result from.
compartments	specify the names of the compartments to extract data from. The compartments can be specified as a character vector e.g. compartments = c('S', 'I', 'R'), or as a formula e.g. compartments = ~S+I+R (see 'Examples'). Default (compartments=NULL) is to extract the number of individuals in each compartment i.e. the data from all discrete state compartments in the model. In models that also have continuous state variables e.g. the SISe model, they are also included.
index	indices specifying the subset of nodes to include when extracting data. Default (index = NULL) is to extract data from all nodes.

`format` the default (`format = "data.frame"`) is to generate a `data.frame` with one row per node and time-step with the number of individuals in each compartment. Using `format = "matrix"` returns the result as a matrix, which is the internal format (see ‘Details’ in [trajectory, SimInf\\_model-method](#)).

### Value

A `data.frame` if `format = "data.frame"`, else a matrix.

---

u0\_SEIR

*Example data to initialize the ‘SEIR’ model*

---

### Description

Example data to initialize a population of 1600 nodes and demonstrate the [SEIR](#) model.

### Usage

```
u0_SEIR()
```

### Details

A `data.frame` with the number of individuals in the ‘S’, ‘E’, ‘I’ and ‘R’ compartments in 1600 nodes. Note that the ‘E’, ‘I’ and ‘R’ compartments are zero.

### Value

A `data.frame`

### Examples

```
## Create an 'SEIR' model with 1600 nodes and initialize it to
## run over 4*365 days and record data at weekly time-points.
## Add ten infected individuals to the first node.
u0 <- u0_SEIR()
u0$I[1] <- 10
tspan <- seq(from = 1, to = 4*365, by = 7)
model <- SEIR(u0      = u0,
              tspan   = tspan,
              events  = events_SEIR(),
              beta    = 0.16,
              epsilon = 0.25,
              gamma   = 0.01)

## Run the model to generate a single stochastic trajectory.
result <- run(model)
plot(result)

## Summarize trajectory
summary(result)
```

---

`u0_SIR`*Example data to initialize the 'SIR' model*

---

## Description

Example data to initialize a population of 1600 nodes and demonstrate the [SIR](#) model.

## Usage

```
u0_SIR()
```

## Details

A data.frame with the number of individuals in the 'S', 'I' and 'R' compartments in 1600 nodes. Note that the 'I' and 'R' compartments are zero.

## Value

A data.frame

## Examples

```
## Create an 'SIR' model with 1600 nodes and initialize
## it to run over 4*365 days. Add one infected individual
## to the first node.
u0 <- u0_SIR()
u0$I[1] <- 1
tspan <- seq(from = 1, to = 4*365, by = 1)
model <- SIR(u0      = u0,
             tspan  = tspan,
             events = events_SIR(),
             beta   = 0.16,
             gamma  = 0.01)

## Run the model to generate a single stochastic trajectory.
result <- run(model)
plot(result)

## Summarize trajectory
summary(result)
```

---

`u0_SIS`*Example data to initialize the 'SIS' model*

---

## Description

Example data to initialize a population of 1600 nodes and demonstrate the [SIS](#) model.

## Usage

```
u0_SIS()
```

## Details

A `data.frame` with the number of individuals in the 'S', and 'I' compartments in 1600 nodes. Note that the 'I' compartment is zero.

## Value

A `data.frame`

## Examples

```
## Create an 'SIS' model with 1600 nodes and initialize
## it to run over 4*365 days. Add one infected individual
## to the first node.
u0 <- u0_SIS()
u0$I[1] <- 1
tspan <- seq(from = 1, to = 4*365, by = 1)
model <- SIS(u0      = u0,
            tspan   = tspan,
            events  = events_SIS(),
            beta    = 0.16,
            gamma   = 0.01)

## Run the model to generate a single stochastic trajectory.
result <- run(model)
plot(result)

## Summarize trajectory
summary(result)
```

---

u0\_SISe

*Example data to initialize the 'SISe' model*


---

### Description

Example data to initialize a population of 1600 nodes and demonstrate the [SISe](#) model.

### Usage

```
u0_SISe()
```

### Details

A data.frame with the number of individuals in the 'S' and 'I' compartments in 1600 nodes. Note that the 'I' compartment is zero.

### Value

A data.frame

### Examples

```
## Create an 'SISe' model with 1600 nodes and initialize it to
## run over 4*365 days and record data at weekly time-points.

## Load the initial population and add ten infected individuals to
## the first node.
u0 <- u0_SISe()
u0$I[1] <- 10

## Define 'tspan' to run the simulation over 4*365 and record the
## state of the system at weekly time-points.
tspan <- seq(from = 1, to = 4*365, by = 7)

## Load scheduled events for the population of nodes with births,
## deaths and between-node movements of individuals.
events <- events_SISe()

## Create an 'SISe' model
model <- SISe(u0 = u0, tspan = tspan, events = events_SISe(),
             phi = 0, epsilon = 1.8e-2, gamma = 0.1, alpha = 1,
             beta_t1 = 1.0e-1, beta_t2 = 1.0e-1, beta_t3 = 1.25e-1,
             beta_t4 = 1.25e-1, end_t1 = 91, end_t2 = 182,
             end_t3 = 273, end_t4 = 365, epsilon = 0)

## Run the model to generate a single stochastic trajectory.
result <- run(model)

## Summarize trajectory
summary(result)
```

```
## Plot the proportion of nodes with at least one infected
## individual.
plot(result, I~S+I, level = 2, type = "l")
```

---

u0\_SISe3

*Example data to initialize the 'SISe3' model*


---

## Description

Example data to initialize a population of 1600 nodes and demonstrate the [SISe3](#) model.

## Usage

```
data(u0_SISe3)
```

## Format

A data.frame

## Details

A data.frame with the number of individuals in the 'S\_1', 'S\_2', 'S\_3', 'I\_1', 'I\_2' and 'I\_3' compartments in 1600 nodes. Note that the 'I\_1', 'I\_2' and 'I\_3' compartments are zero.

## Examples

```
## Create an 'SISe3' model with 1600 nodes and initialize it to
## run over 4*365 days and record data at weekly time-points.

## Load the initial population and add ten infected individuals to
## I_1 in the first node.
u0 <- u0_SISe3
u0$I_1[1] <- 10

## Define 'tspan' to run the simulation over 4*365 and record the
## state of the system at weekly time-points.
tspan <- seq(from = 1, to = 4*365, by = 7)

## Load scheduled events for the population of nodes with births,
## deaths and between-node movements of individuals.
events <- events_SISe3

## Create a 'SISe3' model
model <- SISe3(u0 = u0, tspan = tspan, events = events,
              phi = rep(0, nrow(u0)), epsilon_1 = 1.8e-2,
              epsilon_2 = 1.8e-2, epsilon_3 = 1.8e-2,
              gamma_1 = 0.1, gamma_2 = 0.1, gamma_3 = 0.1,
              alpha = 1, beta_t1 = 1.0e-1, beta_t2 = 1.0e-1,
              beta_t3 = 1.25e-1, beta_t4 = 1.25e-1, end_t1 = 91,
```



```

end_t2 = 182, end_t3 = 273, end_t4 = 365, epsilon = 0)

## Run the model to generate a single stochastic trajectory.
result <- run(model)

## Summarize trajectory
summary(result)

## Plot the proportion of nodes with at least one infected
## individual.
plot(result, I_1 + I_2 + I_3 ~ ., level = 2, type = "l")

```

---

update\_u0

*Update the initial compartment state u0 in each node*


---

### Description

Update the initial compartment state  $u_0$  in each node

### Usage

```

update_u0(model, u0)

## S4 method for signature 'SimInf_model'
update_u0(model, u0)

```

### Arguments

model	The model to update the initial compartment state $u_0$ .
u0	A data.frame with the initial state in each node. Each row is one node, and the number of rows in $u_0$ must match the number of nodes in model. Only the columns in $u_0$ with a name that matches a compartment in the model will be used.

### Value

a SimInf\_model with the updated initial compartment state  $u_0$ .

### Examples

```

## Create an SIR model object.
model <- SIR(u0 = data.frame(S = 99, I = 1, R = 0),
            tspan = 1:100,
            beta = 0.16,
            gamma = 0.077)

## Run the SIR model and plot the result.
set.seed(22)
result <- run(model)

```

```

plot(result)

## Update u0 and run the model again
model <- update_u0(model, data.frame(S = 990, I = 10, R = 0))
result <- run(model)
plot(result)

```

---

update\_v0

*Update the initial continuous state v0 in each node*


---

### Description

Update the initial continuous state v0 in each node

### Usage

```

update_v0(model, v0)

## S4 method for signature 'SimInf_model'
update_v0(model, v0)

```

### Arguments

model	The model to update the initial continuous state v0.
v0	A data.frame with the initial continuous state in each node. Each row is one node, and the number of rows in v0 must match the number of nodes in model. Only the columns in v0 with a name that matches a continuous state in v0 in the model will be used.

### Value

a SimInf\_model with the updated initial continuous state v0.

### Examples

```

## Create an 'SISe' model with no infected individuals and no
## infectious pressure (phi = 0, epsilon = 0).
model <- SISe(u0 = data.frame(S = 100, I = 0), tspan = 1:100,
             phi = 0, upsilon = 0.02, gamma = 0.1, alpha = 1,
             epsilon = 0, beta_t1 = 0.15, beta_t2 = 0.15,
             beta_t3 = 0.15, beta_t4 = 0.15, end_t1 = 91,
             end_t2 = 182, end_t3 = 273, end_t4 = 365)

## Run the 'SISe' model and plot the result.
set.seed(22)
result <- run(model)
plot(result)

## Update the infectious pressure 'phi' in 'v0' and run

```

```
## the model again.  
model <- update_v0(model, data.frame(phi = 1))  
result <- run(model)  
plot(result)
```

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