

# Package ‘lgpr’

December 4, 2019

**Title** Longitudinal Gaussian Process Regression

**Version** 0.30.4

**Description** Implements interpretable nonparametric analysis and covariate selection for longitudinal data using additive Gaussian process regression. Includes specialized non-stationary disease effect modeling features for biomedical studies. Bayesian inference for model parameters is performed using Stan.

**License** GPL (>=3)

**Encoding** UTF-8

**LazyData** true

**Biarch** true

**Depends** R (>= 3.4.0),

**Imports** methods,  
Rcpp (>= 0.12.0),  
rstan (>= 2.18.1),  
rstantools (>= 2.0.0),  
bayesplot (>= 1.7.0),  
MASS (>= 7.3-50),  
stats (>= 3.4),  
ggplot2 (>= 3.1.0),  
ggpubr (>= 0.2)

**LinkingTo** BH (>= 1.66.0),  
Rcpp (>= 0.12.0),  
RcppEigen (>= 0.3.3.3.0),  
rstan (>= 2.18.1),  
StanHeaders (>= 2.18.0)

**SystemRequirements** GNU make

**RoxygenNote** 6.1.1

**Suggests** knitr,  
rmarkdown,  
testthat,  
covr

**VignetteBuilder** knitr

**R topics documented:**

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lgpr-package

The 'lgpr' package.

---

## Description

Longitudinal Gaussian Process regression. The package features

- Additive Gaussian process modeling of longitudinal data
- Posterior inference of the model (hyper)parameters using Stan
- Computation of covariate relevances
- Specialized modeling of a non-stationary disease effect
- Functions for visualizing longitudinal data, posterior samples and model predictions
- Gaussian, Poisson, binomial or negative binomial observation models

## Basic usage

- See the main function `lgp` for creating and fitting additive longitudinal GP models.
- See tutorials at <https://jtimonen.github.io/lgpr-usage/>

## Author(s)

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## References

1. Carpenter, B. et al. (2017). *Stan: A probabilistic programming language*. Journal of Statistical Software 76(1).
2. Jonah Gabry, Ben Goodrich and Martin Lysy (2019). *rstantools: Tools for Developing R Packages Interfacing with 'Stan'*. R package version 2.0.0.
3. Gabry, J. and Mahr, T. (2019). *bayesplot: Plotting for Bayesian Models*. R package version 1.7.0, <http://mc-stan.org/bayesplot>.
4. Stan Development Team (2019). *RStan: the R interface to Stan*. R package version 2.19.2. <http://mc-stan.org/>.

---

add\_test\_caseIDs

Add case IDs to test data frame

---

## Description

Add case IDs to test data frame

## Usage

```
add_test_caseIDs(X_test, X_data)
```

**Arguments**

<code>X_test</code>	test data frame
<code>X_data</code>	data frame

**Value**

Updated `X_test` data frame.

---

<code>affected</code>	<i>Select the affected individuals</i>
-----------------------	--

---

**Description**

Select the affected individuals

**Usage**

```
affected(object, medians.return = FALSE, threshold = 0.5)
```

**Arguments**

<code>object</code>	An object of class <code>lgpfit</code> .
<code>medians.return</code>	Should the medians of beta parameters also be returned?
<code>threshold</code>	A value that the median of beta has to exceed

**Value**

A binary vector indicating the individuals for which the disease effect is inferred to exist.

---

<code>assess_convergence</code>	<i>Assess convergence of the chains</i>
---------------------------------	---

---

**Description**

Assess convergence of the chains

**Usage**

```
assess_convergence(fit, skip_F_gen = TRUE)
```

**Arguments**

<code>fit</code>	An (incomplete) object of class <code>lgpfit</code> .
<code>skip_F_gen</code>	Should <code>F_mean</code> , <code>F_var</code> etc. be ignored

**Value**

A data frame with columns `c("Rhat", "Bulk_ESS", "Tail_ESS")`.

---

average_predictions	<i>Average predictions over samples</i>
---------------------	---

---

**Description**

Average predictions over samples

**Usage**

```
average_predictions(LIST)
```

**Arguments**

LIST	a list over samples
------	---------------------

**Value**

a list

---

check_data	<i>Validate the 'data' input to lgp and resolve covariate types</i>
------------	---

---

**Description**

Validate the 'data' input to lgp and resolve covariate types

**Usage**

```
check_data(data, varInfo, verbose)
```

**Arguments**

data	the data frame that was passed to lgp
varInfo	variable type info
verbose	can this print some info?

**Value**

a list

---

check_formula	<i>Validate the formula of lgp</i>
---------------	------------------------------------

---

**Description**

Checks if the input 'formula' to lgp\_model are valid with the given data

**Usage**

```
check_formula(formula, data)
```

**Arguments**

formula	the formula that was passed to lgp_model
data	the data frame that was passed to lgp_model

**Value**

nothing

---

check_hyperparameter_names	<i>An error message for wrong hyperparameter naming</i>
----------------------------	---

---

**Description**

An error message for wrong hyperparameter naming

**Usage**

```
check_hyperparameter_names(dist, correct)
```

**Arguments**

dist	the distribution
correct	the allowed hyperparameter names

**Value**

nothing



---

check_varInfo	<i>Check that variable types make sense</i>
---------------	---

---

**Description**

Check that variable types make sense

**Usage**

```
check_varInfo(varInfo)
```

**Arguments**

varInfo	a named list
---------	--------------

**Value**

nothing

---

component_index_to_covariate_index	<i>Component index to covariate index</i>
------------------------------------	---

---

**Description**

Component index to covariate index

**Usage**

```
component_index_to_covariate_index(D, idx)
```

**Arguments**

D	integer vector of length 6
idx	integer

**Value**

an integer

---

component\_index\_to\_type

*Component index to component type*

---

### Description

Component index to component type

### Usage

component\_index\_to\_type(D, idx)

### Arguments

D	integer vector of length 6
idx	integer

### Value

an integer

---

compute\_kernel\_matrices

*Evaluate kernel matrices for each component*

---

### Description

Used by [compute\\_predictions](#).

### Usage

compute\_kernel\_matrices(X1, X2, kernel\_info)

### Arguments

X1	Covariate matrix of size n1 x sum(D).
X2	Covariate matrix of size n2 x sum(D).
kernel_info	A list of parameters and other kernel info.

### Value

An array of size n1 x n2 x sum(D).

---

compute_K_beta	<i>Compute the multiplier matrix K_beta (to enable heterogeneous disease effect)</i>
----------------	--

---

**Description**

Compute the multiplier matrix K\_beta (to enable heterogeneous disease effect)

**Usage**

```
compute_K_beta(beta, row_to_caseID_1, row_to_caseID_2)
```

**Arguments**

beta	a row vector of length N_cases
row_to_caseID_1	mapping from row index to case ID
row_to_caseID_2	mapping from row index to case ID

**Value**

a matrix

---

compute_K_var_mask	<i>Compute the variance mask kernel matrix</i>
--------------------	--

---

**Description**

Compute the variance mask kernel matrix

**Usage**

```
compute_K_var_mask(disAge1, disAge2, vm_params, stp, nan_replace = 0)
```

**Arguments**

disAge1	disease-related age covariate vector of length n1
disAge2	disease-related age covariate vector of length n2
vm_params	vector of two mask function parameters
stp	input warping steepness
nan_replace	value to replace nans in disAge vectors

**Value**

a matrix of size n1 x n2

---

compute_lppd	<i>Compute log-posterior predictive density at test points</i>
--------------	--

---

**Description**

Compute log-posterior predictive density at test points

**Usage**

```
compute_lppd(PRED, y_test)
```

**Arguments**

PRED	predictions
y_test	values of the response variable at the test points

**Value**

a matrix with size n\_samples x n\_data

---

compute_noise_level	<i>Determine noise level</i>
---------------------	------------------------------

---

**Description**

Determine noise level

**Usage**

```
compute_noise_level(pars, model, noise_method)
```

**Arguments**

pars	A data frame representing one parameter sample, i.e one row of <code>as.data.frame(stanfit)</code> , where <code>stanfit</code> is an object of class <code>stanfit</code>
model	An object of class <code>lgpmodel</code>
noise_method	Noise level estimation method.

**Value**

a value between 0 and 1

---

 compute\_predicted\_components

*Compute component-wise predictions at test points*


---

**Description**

Used by [compute\\_predictions](#).

**Usage**

```
compute_predicted_components(KK, KKs, KKss, y_data, sigma_n, DELTA)
```

**Arguments**

KK	Kernel matrices data vs. data.
KKs	Kernel matrices test vs. data.
KKss	Kernel matrices test vs. test.
y_data	Response variable.
sigma_n	Noise standard deviation parameter.
DELTA	Diagonal jitter that ensures pos. def. kernel.

**Value**

A list containing predicted means and variances.

---

 compute\_predictions

*Compute component-wise predictions at test points*


---

**Description**

Used by [lgp\\_predict](#).

**Usage**

```
compute_predictions(X_data, y_data, X_test, params, D, info, cnames, TSCL,
  handle_extra = "warning")
```

**Arguments**

X_data	Covariate matrix (data points).
y_data	Response variable (data points).
X_test	Covariate matrix (test points).
params	Kernel function and other hyperparameters
D	a vector of length 6
info	other model info
cnames	Names of the model components.
TSCL	time scaling function and its inverse
handle_extra	What to do if test data contains individuals that are not in the training data? Must be 'silent', 'warning' or 'error'.

**Value**

A list.

---

compute_relevances	<i>Compute component relevances and estimate amount of noise (one MCMC sample)</i>
--------------------	--

---

**Description**

Compute component relevances and estimate amount of noise (one MCMC sample)

**Usage**

```
compute_relevances(pars, model, method, noise_method)
```

**Arguments**

pars	A data frame representing one parameter sample, i.e one row of <code>as.data.frame(stanfit)</code> , where <code>stanfit</code> is an object of class <code>stanfit</code>
model	An object of class <code>lgpmodel</code>
method	Relevance determination method. Must be either <code>"f_mean"</code> or <code>"alpha"</code> .
noise_method	Noise level estimation method.

**Value**

a matrix of size 1 x n\_components + 1

---

compute_relevances_alpha	<i>The alpha relevance determination method</i>
--------------------------	---

---

**Description**

The alpha relevance determination method

**Usage**

```
compute_relevances_alpha(pars, model)
```

**Arguments**

pars	A data frame representing one parameter sample, i.e one row of <code>as.data.frame(stanfit)</code> , where <code>stanfit</code> is an object of class <code>stanfit</code>
model	An object of class <code>lgpmodel</code>

**Value**

a vector of length n\_components

---

`compute_relevances_fmean`*The f\_mean relevance determination method*

---

**Description**

The f\_mean relevance determination method

**Usage**

```
compute_relevances_fmean(pars, model)
```

**Arguments**

<code>pars</code>	A data frame representing one parameter sample, i.e one row of <code>as.data.frame(stanfit)</code> , where <code>stanfit</code> is an object of class <code>stanfit</code>
<code>model</code>	An object of class <code>lgpmodel</code>

**Value**

a vector of length `n_components`

---

`create_covariates_stan`*Create the covariate matrix that is given to stan*

---

**Description**

Create the covariate matrix that is given to stan

**Usage**

```
create_covariates_stan(data, varInfo, types, formula, verbose)
```

**Arguments**

<code>data</code>	the data frame that was passed to <code>lgp</code>
<code>varInfo</code>	original variable type info
<code>types</code>	the types returned by <a href="#">check_data</a>
<code>formula</code>	the model formula
<code>verbose</code>	can this print some info?

**Value**

a list

---

create_data_plot_df	Create a plotting data frame for ggplot
---------------------	---

---

**Description**

A helper function for plot\_data.

**Usage**

```
create_data_plot_df(data, hl_1, hl_2, hl_cont)
```

**Arguments**

data	a data frame
hl_1	highlighting by color
hl_2	highlighting by linestyle
hl_cont	highlighting continuous

**Value**

an extended data frame

---

create_example_fit	Create an example fit object
--------------------	------------------------------

---

**Description**

Create an example fit object

**Usage**

```
create_example_fit(N = 4, t = 10 * c(1, 2, 3, 4, 5), iter = 100,  
  chains = 1)
```

**Arguments**

N	number of individuals
t	time points
iter	number of iterations
chains	number of chains

**Value**

an object of class `lgpfit`



create\_F

*Simulate latent function components for longitudinal data analysis***Description**

Simulate latent function components for longitudinal data analysis

**Usage**

```
create_F(X, covariates, relevances, lengthscales, X_affected, dis_fun,
        bin_kernel, steepness, vm_params, force_zeromean)
```

**Arguments**

X	input data matrix (generated by <a href="#">create_X</a> )
covariates	Integer vector that defines the types of covariates (other than id and age). Different integers correspond to the following covariate types: <ul style="list-style-type: none"> <li>• 0 = disease-related age</li> <li>• 1 = other continuous covariate</li> <li>• 2 = a categorical covariate that interacts with age</li> <li>• 3 = a categorical covariate that acts as a group offset</li> <li>• 4 = a categorical covariate that that acts as a group offset AND is restricted to have value 0 for controls and 1 for cases</li> </ul>
relevances	Relative relevance of each component. Must have be a vector so that <code>length(relevances) = 2 + length(covariates)</code> . First two values define the relevance of the individual-specific age and shared age component, respectively.
lengthscales	A vector so that <code>length(lengthscales) = 2 + sum(covariates %in% c(0,1,2))</code> .
X_affected	which individuals are affected by the disease
dis_fun	A function or a string that defines the disease effect. If this is a function, that function is used to generate the effect. If <code>dis_fun</code> is "gp_vm" or "gp_ns", the disease component is drawn from a nonstationary GP prior (vm is the variance masked version of it).
bin_kernel	Should the binary kernel be used for categorical covariates? If this is TRUE, the effect will exist only for group 1.
steepness	Steepness of the input warping function. This is only used if the disease component is in the model.
vm_params	Parameters of the variance mask function. This is only needed if <code>useMaskedVarianceKernel = TRUE</code> .
force_zeromean	Should each component (excluding the disease age component) be forced to have a zero mean?

**Value**

a data frame FFF where one column corresponds to one additive data component

---

```
create_predictions_plot_df1
```

*Create a plotting data frame for ggplot*

---

**Description**

A helper function for plot\_predictions.

**Usage**

```
create_predictions_plot_df1(fit, scale_f = TRUE, n_sds)
```

**Arguments**

fit	An object of class lgpfit.
scale_f	Should the predictions be scaled back to the original data scale?
n_sds	number of standard deviations for the uncertainty band width

**Value**

a data frame

---

```
create_predictions_plot_df2
```

*Create a plotting data frame for ggplot*

---

**Description**

A helper function for plot\_predictions.

**Usage**

```
create_predictions_plot_df2(model, PRED, scale_f = TRUE, mode, n_sds)
```

**Arguments**

model	An object of class lgpmode1.
PRED	Predictions computed using lgp_predict.
scale_f	Should the predictions be scaled back to the original data scale?
mode	mode
n_sds	number of standard deviations for the uncertainty band width

**Value**

a data frame

---

```
create_simdata_plot_df
```

*Create a plotting data frame for ggplot*

---

### Description

A helper function for `plot_simdata_by_component`.

### Usage

```
create_simdata_plot_df(simData)
```

### Arguments

`simData`            An object created using `simulate_data`.

### Value

a data frame

---

```
create_stan_input
```

*Create input for Stan*

---

### Description

Parses the formula and data input to [lgp\\_model](#). Also performs many input checks.

### Usage

```
create_stan_input(formula, data, prior, likelihood, varInfo, standardize,
  uncertain_effect_time, equal_effect, C_hat, DELTA, sample_F, verbose,
  variance_mask, N_trials, skip_gen_quant)
```

### Arguments

<code>formula</code>	A formula of the form $y \sim x_1 + x_2 + x_3$ defining the response variable $y$ and covariates $x_i$ . The formula must contain exactly one tilde ( $\sim$ ), with response variable on the left-hand side and covariates on the right-hand side. Covariates should be separated by a plus (+) sign. All variables that appear in the formula must exist as columns of data. Note that effects of categorical covariates are by default defined as interactions with <code>time_variable</code> . If you wish to change this, see the argument <code>offset_vars</code> . The subject identifier variable cannot currently be included in <code>offset_vars</code> . If you wish to model the effect of <code>id_variable</code> as a constant offset, you can create another covariate with the same values and use it in your formula and <code>offset_vars</code> instead.
<code>data</code>	A data frame containing the variables given in <code>formula</code> as columns.
<code>prior</code>	A named list, defining the prior distribution of model (hyper)parameters. It is recommended to first create this using the function <a href="#">prior_default</a> , and then possibly modify it.

likelihood	Determines the observation model. Must be either "Gaussian" (default), "Poisson", "NB" (negative binomial) or "binomial". To use Bernoulli likelihood, use likelihood="binomial" and set N_trials as a vector of ones.
varInfo	Variable type info.
standardize	Should the response variable be standardized?
uncertain_effect_time	Do we wish to model uncertainty in the disease effect time?
equal_effect	Is the disease effect assumed to be equally strong for all diseased individuals?
C_hat	The constant GP mean. By default this is NULL, and set to <ul style="list-style-type: none"> <li>C_hat = 0, if likelihood is "Gaussian", because with Gaussian likelihood the response variable is by default centered to have zero mean.</li> <li>C_hat = log(mean(y)) if likelihood is "Poisson" or "NB",</li> <li>C_hat = log(p/(1-p)), where p = mean(y/N_trials) if likelihood is "binomial"</li> </ul> <p>Above, y denotes the response variable.</p>
DELTA	the amount of added jitter to ensure positive definiteness of the kernel
sample_F	Determines if the function values are be sampled (must be TRUE if likelihood is not "Gaussian").
verbose	Should more verbose output be printed?
variance_mask	Should a variance mask be used to force disease component variance to zero before disease onset?
N_trials	This argument (number of trials) is only needed when likelihood is binomial. Must have length one or equal to number of data points. Setting N_trials=1 corresponds to Bernoulli observation model.
skip_gen_quant	If this is true, the generated quantities block of Stan is skipped.

**Value**

A list containing the data to be given to `rstan::sampling`, some info about preprocessing and all the information about scaling the inputs and response, and updated variable type info.

---

create_test_points	<i>Create a matrix of test points</i>
--------------------	---------------------------------------

---

**Description**

Create a matrix of test points

**Usage**

```
create_test_points(object, t_test)
```

**Arguments**

object	An object of class <code>lgpmodel</code> or <code>lgpfit</code>
t_test	Test time points (will be same for each individual).

**Value**

A data frame.

---

create_X	<i>Simulate an input data frame X</i>
----------	---------------------------------------

---

## Description

Simulate an input data frame X

## Usage

```
create_X(N, covariates, names, n_cats, t_data, t_jitter, t_effect_range,
         continuous_info, verbose)
```

## Arguments

N	Number of individuals.
covariates	Integer vector that defines the types of covariates (other than id and age). If not given, only the id and age covariates are created. Different integers correspond to the following covariate types: <ul style="list-style-type: none"> <li>• 0 = disease-related age</li> <li>• 1 = other continuous covariate</li> <li>• 2 = a categorical covariate that interacts with age</li> <li>• 3 = a categorical covariate that acts as a group offset</li> <li>• 4 = a categorical covariate that that acts as a group offset AND is restricted to have value 0 for controls and 1 for cases</li> </ul>
names	Covariate names.
n_cats	An integer vector defining the number of categories for each categorical covariate, so that <code>length(n_cats)</code> equals to the number of 2's and 3's in the <code>covariates</code> vector.
t_data	Measurement times.
t_jitter	Standard deviation of the jitter added to the given measurement times.
t_effect_range	Time interval from which the disease effect times are sampled uniformly. Alternatively, This can any function that returns the (possibly randomly generated) real disease effect time for one individual.
continuous_info	Info for generating continuous covariates. Must be a list containing fields <code>lambda</code> and <code>mu</code> , which have length 3. The continuous covariates are generated so that $x < -\sin(a \cdot t + b) + c$ , where <ul style="list-style-type: none"> <li>• <code>t &lt;- seq(0, 2*pi, length.out = k)</code></li> <li>• <code>a &lt;- -mu[1] + lambda[1]*stats::runif(1)</code></li> <li>• <code>b &lt;- -mu[2] + lambda[2]*stats::runif(1)</code></li> <li>• <code>c &lt;- -mu[3] + lambda[3]*stats::runif(1)</code></li> </ul>
verbose	verbosity mode

## Value

`list(X, onsets, par_cont)`

---

create_X_star	<i>Create X_star</i>
---------------	----------------------

---

**Description**

Create X\_star

**Usage**

```
create_X_star(X, D, t_test, SCL, X_notnan)
```

**Arguments**

X	covariate matrix
D	covariate type information
t_test	Test time points (will be same for each individual).
SCL	time scaling function and its inverse
X_notnan	indicates where X_diseaseAge is not NaN

**Value**

A data frame.

---

create_y	<i>Generate noisy observations</i>
----------	------------------------------------

---

**Description**

Generate noisy observations

**Usage**

```
create_y(noise_type, f, snr, phi, N_trials)
```

**Arguments**

noise_type	Either "Gaussian", "Poisson", NB" (negative binomial) or "binomial".
f	The underlying signal.
snr	The desired signal-to-noise ratio. This argument is valid only with noise_type = "Gaussian".
phi	The dispersion parameter for negative binomial data. The variance is $g + g^2/\phi$ .
N_trials	The number of trials parameter for binomial data.

**Value**

A list out, where

- out\$g is f mapped through an inverse link function and
- out\$y is the noisy response variable.

---

disease_effect	<i>Draw disease component from a parameteric form</i>
----------------	---

---

**Description**

Draw disease component from a parameteric form

**Usage**

```
disease_effect(X_id, X_disAge, dis_fun)
```

**Arguments**

X_id	the id covariate
X_disAge	the diseaseAge covariate
dis_fun	the disease age effect function

**Value**

a vector

---

drawCategorical	<i>Independently draw categorical variables for each individual</i>
-----------------	---

---

**Description**

Independently draw categorical variables for each individual

**Usage**

```
drawCategorical(N, k, v)
```

**Arguments**

N	number of individuals
k	number of timepoints
v	vector of numbers of different categories

**Value**

a matrix of size N x D, where  $D \leq \text{length}(v)$

---

drawContinuous	<i>Independently draw continuous variables for each individual</i>
----------------	--

---

**Description**

Independently draw continuous variables for each individual

**Usage**

```
drawContinuous(N, k, D, mu, lambda)
```

**Arguments**

N	number of individuals
k	number of timepoints
D	number of variables
mu	a vector of length 3
lambda	a vector of length 3

**Value**

a matrix of size N x D

---

drawLatentComponents	<i>Draw realizations of multivariate normals</i>
----------------------	--

---

**Description**

Draw realizations of multivariate normals

**Usage**

```
drawLatentComponents(KK)
```

**Arguments**

KK	3D matrix where $KK[, , j]$ is the $j$ th kernel matrix
----	---

**Value**

a matrix FFF



---

drawMeasurementTimes    *Draw the age covariate*

---

**Description**

Draw the age covariate

**Usage**

```
drawMeasurementTimes(N, t_data, t_jitter)
```

**Arguments**

N	number of individuals
t_data	a vector of length k
t_jitter	Standard deviation of the jitter added to the given measurement times.

**Value**

a vector of length N\*k

---

extract\_t\_effect\_samples  
*Extract samples of T\_effect*

---

**Description**

Extract samples of T\_effect

**Usage**

```
extract_t_effect_samples(fit)
```

**Arguments**

fit	an object of class lgpfit
-----	---------------------------

**Value**

a matrix

---

full_model	Create a full model with all covariates included
------------	--

---

**Description**

Create a full model with all covariates included

**Usage**

```
full_model(data, ...)
```

**Arguments**

data	a data frame
...	additional parameters to <a href="#">lgp_model</a>

**Value**

a ggplot object

---

full_model_formula	Get formula of a full model with all covariates included
--------------------	--

---

**Description**

Get formula of a full model with all covariates included

**Usage**

```
full_model_formula(data)
```

**Arguments**

data	a data frame, where the response variable is the last column
------	--

**Value**

a formula

---

get_case_ids	<i>Get case ids in original data</i>
--------------	--------------------------------------

---

**Description**

Get case ids in original data

**Usage**

```
get_case_ids(fit)
```

**Arguments**

fit	an object of class <code>lgpfit</code>
-----	--

**Value**

a character vector

---

get_case_row_mappings	<i>Create case ID to rows and back mappings</i>
-----------------------	---

---

**Description**

Create mappings

- from case ID to data rows (`caseID_to_rows`, `caseID_nrows`)
- from row number to case ID (`row_to_caseID`)

**Usage**

```
get_case_row_mappings(X_notnan, X_id, only_R2C = FALSE)
```

**Arguments**

X_notnan	binary vector indicating if <code>diseaseAge</code> is available for that measurement
X_id	the id covariate in X
only_R2C	should this return only the rows-to-caseID mapping

**Value**

a list

---

get_diseased_info	<i>Get some variables related to diseased individuals</i>
-------------------	---

---

**Description**

Get some variables related to diseased individuals

**Usage**

```
get_diseased_info(D, X, X_notnan, uncertain_effect_time, equal_effect,
                  TSCL)
```

**Arguments**

D	an integer vector of length 6
X	the design matrix
X_notnan	a binary vector of length n
uncertain_effect_time	Boolean value
equal_effect	Boolean value
TSCL	time scaling function and its inverse

**Value**

a list

---

get_function_components_from_df	<i>Get values of function components at data points, for one MCMC sample</i>
---------------------------------	--

---

**Description**

Get values of function components at data points, for one MCMC sample

**Usage**

```
get_function_components_from_df(pars, model)
```

**Arguments**

pars	A data frame representing one parameter sample, i.e one row of <code>as.data.frame(stanfit)</code> , where <code>stanfit</code> is an object of class <code>stanfit</code>
model	An object of class <code>lgpmodel</code>

**Value**

A matrix of size `n_data x n_components+2`

---

`get_function_components_from_df_all`*Get values of function components at data points*

---

**Description**

Get values of function components at data points

**Usage**

```
get_function_components_from_df_all(df, model)
```

**Arguments**

<code>df</code>	A stanfit object as data frame, obtained as <code>as.data.frame(stanfit)</code>
<code>model</code>	An object of class <code>lgpmodel</code>

**Value**

An array of size `n_samples x n_data x n_components+2`

---

`get_g_from_f`*Get signal on data scale from process f*

---

**Description**

Get signal on data scale from process f

**Usage**

```
get_g_from_f(f, model)
```

**Arguments**

<code>f</code>	A vector
<code>model</code>	an object of class <code>lgpmodel</code>

**Value**

A vector g

---

get_model_dims	<i>Get some dimension variables that the Stan model needs as input</i>
----------------	--

---

**Description**

Get some dimension variables that the Stan model needs as input

**Usage**

```
get_model_dims(X, D)
```

**Arguments**

X	the design matrix
D	a vector of length 6

**Value**

a list

---

get_obs_onset_times	<i>Extract observed disease onset times from diseaseAge covariate vector</i>
---------------------	--

---

**Description**

Extract observed disease onset times from diseaseAge covariate vector

**Usage**

```
get_obs_onset_times(id, age, disAge)
```

**Arguments**

id	the id covariate, vector of length n
age	the age covariate, vector of length n
disAge	the observed disease-related age covariate, vector of length n

**Value**

vector of observed onset times

---

get_onset_info	<i>Get disease onset info</i>
----------------	-------------------------------

---

**Description**

This returns

- a vector of observed onsets
- mapping from case ID to average sampling interval before the observed disease onset

**Usage**

```
get_onset_info(D, X, MAPS, TSCL)
```

**Arguments**

D	an integer vector of length 6
X	the design matrix
MAPS	mappings created by <code>get_case_row_mappings</code>
TSCL	time scaling function and its inverse

**Value**

two vectors of length `N_cases`

---

get_pkg_description	<i>Get lgpr version description</i>
---------------------	-------------------------------------

---

**Description**

Get lgpr version description

**Usage**

```
get_pkg_description()
```

**Value**

package description

---

get_predicted	<i>A helper function</i>
---------------	--------------------------

---

**Description**

A helper function

**Usage**

```
get_predicted(fit)
```

**Arguments**

fit	An (incomplete) object of class <code>lgpfit</code> .
-----	---

**Value**

a list

---

get_prior_params	<i>Get prior parameters</i>
------------------	-----------------------------

---

**Description**

Get prior parameters

**Usage**

```
get_prior_params(dist, add_correct)
```

**Arguments**

dist	the distribution
add_correct	additional correct parameter names

**Value**

a hyperparameter vector of length 2



---

get_prior_type	<i>A dictionary from distribution names to integer encoding</i>
----------------	---

---

**Description**

A dictionary from distribution names to integer encoding

**Usage**

```
get_prior_type(type)
```

**Arguments**

type	type of the distribution as a string
------	--------------------------------------

**Value**

an integer

---

get_response	<i>Get the (scaled) response variable</i>
--------------	---

---

**Description**

Gets and possibly scales the response variable.

**Usage**

```
get_response(data, varInfo, standardize, LH)
```

**Arguments**

data	the data frame given as input to lgp
varInfo	variable type info
standardize	should the response be standardized to unit variance and zero mean
LH	likelihood as integer

**Value**

a list with the (scaled) response variable

---

get_runtime	<i>Get average runtime of a chain</i>
-------------	---------------------------------------

---

**Description**

Get average runtime of a chain

**Usage**

```
get_runtime(object)
```

**Arguments**

object	An object of class lgpfit.
--------	----------------------------

**Value**

Average runtimes for warmup and sampling

---

get_stan_model	<i>Get main stan model of the package</i>
----------------	---

---

**Description**

Get main stan model of the package

**Usage**

```
get_stan_model()
```

**Value**

an object of class stanmodel

---

get_transform_type	<i>A dictionary from transform names to integer encoding</i>
--------------------	--

---

**Description**

A dictionary from transform names to integer encoding

**Usage**

```
get_transform_type(type)
```

**Arguments**

type	Type of the transform as a string. Allowed arguments are "none" or "square". If NULL, "none" is used.
------	---

**Value**

an integer (0, 1 or 2)

---

hyperparam_estimate	<i>Get a posterior estimate of model (hyper)parameters</i>
---------------------	--

---

**Description**

Get a posterior estimate of model (hyper)parameters

**Usage**

```
hyperparam_estimate(object, type = "mean")
```

**Arguments**

object	An (incomplete) object of class <code>lgpfit</code> .
type	Must be "mean", "median", or "map".

**Value**

a data frame

---

hyperparam_samples	<i>Get a set of model (hyper)parameter samples</i>
--------------------	--

---

**Description**

Get a set of model (hyper)parameter samples

**Usage**

```
hyperparam_samples(object, samples = NULL)
```

**Arguments**

object	An (incomplete) object of class <code>lgpfit</code> .
samples	Sample indices. If <code>NULL</code> , all samples are taken.

**Value**

a data frame

---

idx_to_cont_index	<i>Component index to how manyth continuous covariate it is</i>
-------------------	---

---

**Description**

Component index to how manyth continuous covariate it is

**Usage**

```
idx_to_cont_index(D, idx)
```

**Arguments**

D	integer vector of length 6
idx	an integer

**Value**

an integer

---

kernel_bin	<i>Compute a binary kernel matrix</i>
------------	---------------------------------------

---

**Description**

Compute a binary kernel matrix

**Usage**

```
kernel_bin(x1, x2 = NULL, alpha = 1, pos_class = 1)
```

**Arguments**

x1	(integer) vector of length n
x2	(integer) vector of length m
alpha	marginal std (default = 1)
pos_class	the positive class label

**Value**

A kernel matrix of size n x m

---

kernel_ns	<i>Compute a nonstationary kernel matrix using input warping</i>
-----------	--

---

**Description**

Compute a nonstationary kernel matrix using input warping

**Usage**

```
kernel_ns(x1, x2 = NULL, alpha = 1, ell, a, b, c, nan_replace = 0)
```

**Arguments**

x1	vector of length n
x2	vector of length m
alpha	marginal std (default = 1)
ell	lengthscale in the warped space
a	steepness of the warping function rise
b	location of the effective time window
c	maximum range
nan_replace	the value to use for replacing NaN values

**Value**

A kernel matrix of size n x m

---

kernel_se	<i>Compute a squared exponential kernel matrix</i>
-----------	--

---

**Description**

Compute a squared exponential kernel matrix

**Usage**

```
kernel_se(x1, x2, alpha = 1, ell = 1)
```

**Arguments**

x1	vector of length n
x2	vector of length m
alpha	marginal std (default = 1)
ell	lengthscale (default = 1)

**Value**

A kernel matrix of size n x m

---

kernel_zerohsum	<i>Compute a zeros-sum kernel matrix</i>
-----------------	--

---

### Description

Compute a zeros-sum kernel matrix

### Usage

```
kernel_zerohsum(x1, x2, M, alpha = 1)
```

### Arguments

x1	(integer) vector of length n
x2	(integer) vector of length m
M	number of categories
alpha	marginal std (default = 1)

### Value

A (binary) kernel matrix of size n x m

---

lgp	<i>The main function of the 'lgpr' package</i>
-----	--

---

### Description

This is a wrapper for both [lgp\\_model](#) and [lgp\\_fit](#). It first creates an `lgpmodel` object and then fits the model, finally returning an `lgpfit` object. Note that the covariate types are automatically inferred from the given data. If you wish to change these, see the arguments

- `id_variable`
- `time_variable`
- `disAge_variable`
- `continuous_vars` and
- `categorical_vars`.

### Usage

```
lgp(formula, data, likelihood = "Gaussian", prior = prior_default(),
    uncertain_effect_time = FALSE, equal_effect = TRUE,
    id_variable = "id", time_variable = "age", disAge_variable = NULL,
    continuous_vars = NULL, categorical_vars = NULL,
    offset_vars = NULL, C_hat = NULL, DELTA = 1e-08, sample_F = NULL,
    parallel = FALSE, skip_postproc = FALSE, threshold = 0.95,
    variance_mask = TRUE, N_trials = NULL, relevance_method = "f_mean",
    verbose = FALSE, ...)
```

## Arguments

formula	A formula of the form $y \sim x_1 + x_2 + x_3$ defining the response variable $y$ and covariates $x_i$ . The formula must contain exactly one tilde ( $\sim$ ), with response variable on the left-hand side and covariates on the right-hand side. Covariates should be separated by a plus (+) sign. All variables that appear in the formula must exist as columns of data. Note that effects of categorical covariates are by default defined as interactions with <code>time_variable</code> . If you wish to change this, see the argument <code>offset_vars</code> . The subject identifier variable cannot currently be included in <code>offset_vars</code> . If you wish to model the effect of <code>id_variable</code> as a constant offset, you can create another covariate with the same values and use it in your formula and <code>offset_vars</code> instead.
data	A data frame containing the variables given in formula as columns.
likelihood	Determines the observation model. Must be either "Gaussian" (default), "Poisson", "NB" (negative binomial) or "binomial". To use Bernoulli likelihood, use <code>likelihood="binomial"</code> and set <code>N_trials</code> as a vector of ones.
prior	A named list, defining the prior distribution of model (hyper)parameters. It is recommended to first create this using the function <code>prior_default</code> , and then possibly modify it.
uncertain_effect_time	Do we wish to model uncertainty in the disease effect time?
equal_effect	Is the disease effect assumed to be equally strong for all diseased individuals?
id_variable	Name of the unique subject identifier variable (default = "id").
time_variable	Name of the time variable (default = "age").
disAge_variable	Name of the disease-related age variable. If NULL (default), this will be chosen to be "diseaseAge", if such covariate is found in the data.
continuous_vars	Names of other continuous covariates. If NULL, the remaining covariates that have floating point values are interpreted as continuous.
categorical_vars	Names of categorical covariates that interact with the time variable. If NULL (default), the remaining covariates that have integer values are interpreted as categorical.
offset_vars	Names of the categorical covariates that are treated as time-independent group offsets. If NULL (default), no variables are interpreted as such covariates.
C_hat	The constant GP mean. By default this is NULL, and set to <ul style="list-style-type: none"> <li>• <math>C_{\text{hat}} = 0</math>, if likelihood is "Gaussian", because with Gaussian likelihood the response variable is by default centered to have zero mean.</li> <li>• <math>C_{\text{hat}} = \log(\text{mean}(y))</math> if likelihood is "Poisson" or "NB",</li> <li>• <math>C_{\text{hat}} = \log(p/(1-p))</math>, where <math>p = \text{mean}(y/N_{\text{trials}})</math> if likelihood is "binomial"</li> </ul> <p>Above, <math>y</math> denotes the response variable.</p>
DELTA	the amount of added jitter to ensure positive definiteness of the kernel
sample_F	Determines if the function values are to be sampled (must be TRUE if likelihood is not "Gaussian").
parallel	Determines if the chain will be run in parallel (default = FALSE). If TRUE, then Stan is run by first defining <code>options(mc.cores = parallel::detectCores())</code> .

skip_postproc	In this mode the postprocessing after running Stan is skipped.
threshold	Component selection threshold for relevance sum.
variance_mask	Should a variance mask be used to force disease component variance to zero before disease onset?
N_trials	This argument (number of trials) is only needed when likelihood is binomial. Must have length one or equal to number of data points. Setting N_trials=1 corresponds to Bernoulli observation model.
relevance_method	Component relevance determination method. Must be either "f_mean" or "alpha".
verbose	Should more verbose output be printed?
...	Optional arguments passed to <code>rstan::sampling</code> , for example <code>iter</code> , <code>chains</code> or <code>control</code> . See <a href="#">sampling</a> for the possible arguments.

**Value**

An object of class `lgpfit`.

---

<code>lgpfit-class</code>	<i>An S4 class to represent the output of the <code>lgp_fit</code> function</i>
---------------------------	---

---

**Description**

An S4 class to represent the output of the `lgp_fit` function

**Slots**

`stan_fit` The `stanfit` object returned by `rstan::sampling`.  
`model` The `lgpmodel` object returned by `lgp_model`.  
`relevances` Inferred component relevances.  
`selection` Component selection info.  
`pkg_version` Package version number.  
`diagnostics` A data frame with columns `c("Rhat", "Bulk_ESS", "Tail_ESS")`.

---

<code>lgpmodel-class</code>	<i>An S4 class to represent an lgp model</i>
-----------------------------	--

---

**Description**

An S4 class to represent an lgp model

**Slots**

`data` The original unmodified data frame.  
`stan_dat` The data to be given as input to `rstan::sampling`.  
`scalings` Preprocessing scaling functions and their inverse operations.  
`info` Model info.



---

lgp_component_names	<i>Get names of model components</i>
---------------------	--------------------------------------

---

**Description**

Get names of model components

**Usage**

```
lgp_component_names(stan_dat)
```

**Arguments**

stan_dat	The data that was passed to <code>rstan::sampling</code>
----------	--

**Value**

names of model components

---

lgp_covariate_names	<i>Get names of model covariates</i>
---------------------	--------------------------------------

---

**Description**

Get names of model covariates

**Usage**

```
lgp_covariate_names(stan_dat)
```

**Arguments**

stan_dat	The data that was passed to <code>rstan::sampling</code>
----------	--

**Value**

names of model components

---

lgp_fit	<i>Fit an lgp model</i>
---------	-------------------------

---

### Description

Samples the posterior of an additive Gaussian process regression model using [rstan](#).

### Usage

```
lgp_fit(model, threshold = 0.95, parallel = FALSE,
        skip_postproc = FALSE, relevance_method = "f_mean",
        verbose = FALSE, ...)
```

### Arguments

model	An object of class <code>lgpmodel</code> .
threshold	Component selection threshold for relevance sum.
parallel	Determines if the chain will be run in parallel (default = FALSE). If TRUE, then Stan is run by first defining <code>options(mc.cores = parallel::detectCores())</code> .
skip_postproc	In this mode the postprocessing after running Stan is skipped.
relevance_method	Component relevance determination method. Must be either "f_mean" or "alpha".
verbose	should some output be printed?
...	Optional arguments passed to <code>rstan::sampling</code> , for example <code>iter</code> , <code>chains</code> or <code>control</code> . See <a href="#">sampling</a> for the possible arguments.

### Value

An object of class `lgpfit`.

### See Also

For the possible additional arguments, see [sampling](#). For creating the `lgpmodel` input, see [lgp\\_model](#).

---

lgp_model	<i>Create an lgp model</i>
-----------	----------------------------

---

### Description

Creates an object of class `lgpmodel`

### Usage

```
lgp_model(formula, data, likelihood = "Gaussian",
          prior = prior_default(), uncertain_effect_time = FALSE,
          equal_effect = TRUE, C_hat = NULL, DELTA = 1e-08,
          sample_F = NULL, id_variable = "id", time_variable = "age",
          disAge_variable = NULL, continuous_vars = NULL,
          categorical_vars = NULL, offset_vars = NULL, variance_mask = TRUE,
          N_trials = NULL, skip_gen_quant = FALSE, verbose = FALSE)
```

**Arguments**

formula	A formula of the form $y \sim x_1 + x_2 + x_3$ defining the response variable $y$ and covariates $x_i$ . The formula must contain exactly one tilde ( $\sim$ ), with response variable on the left-hand side and covariates on the right-hand side. Covariates should be separated by a plus (+) sign. All variables that appear in the formula must exist as columns of data. Note that effects of categorical covariates are by default defined as interactions with <code>time_variable</code> . If you wish to change this, see the argument <code>offset_vars</code> . The subject identifier variable cannot currently be included in <code>offset_vars</code> . If you wish to model the effect of <code>id_variable</code> as a constant offset, you can create another covariate with the same values and use it in your formula and <code>offset_vars</code> instead.
data	A data frame containing the variables given in formula as columns.
likelihood	Determines the observation model. Must be either "Gaussian" (default), "Poisson", "NB" (negative binomial) or "binomial". To use Bernoulli likelihood, use <code>likelihood="binomial"</code> and set <code>N_trials</code> as a vector of ones.
prior	A named list, defining the prior distribution of model (hyper)parameters. It is recommended to first create this using the function <code>prior_default</code> , and then possibly modify it.
uncertain_effect_time	Do we wish to model uncertainty in the disease effect time?
equal_effect	Is the disease effect assumed to be equally strong for all diseased individuals?
C_hat	The constant GP mean. By default this is NULL, and set to <ul style="list-style-type: none"> <li>• <math>C_{\text{hat}} = 0</math>, if likelihood is "Gaussian", because with Gaussian likelihood the response variable is by default centered to have zero mean.</li> <li>• <math>C_{\text{hat}} = \log(\text{mean}(y))</math> if likelihood is "Poisson" or "NB",</li> <li>• <math>C_{\text{hat}} = \log(p/(1-p))</math>, where <math>p = \text{mean}(y/N_{\text{trials}})</math> if likelihood is "binomial"</li> </ul> <p>Above, <math>y</math> denotes the response variable.</p>
DELTA	the amount of added jitter to ensure positive definiteness of the kernel
sample_F	Determines if the function values are to be sampled (must be TRUE if likelihood is not "Gaussian").
id_variable	Name of the unique subject identifier variable (default = "id").
time_variable	Name of the time variable (default = "age").
disAge_variable	Name of the disease-related age variable. If NULL (default), this will be chosen to be "diseaseAge", if such covariate is found in the data.
continuous_vars	Names of other continuous covariates. If NULL, the remaining covariates that have floating point values are interpreted as continuous.
categorical_vars	Names of categorical covariates that interact with the time variable. If NULL (default), the remaining covariates that have integer values are interpreted as categorical.
offset_vars	Names of the categorical covariates that are treated as time-independent group offsets. If NULL (default), no variables are interpreted as such covariates.
variance_mask	Should a variance mask be used to force disease component variance to zero before disease onset?

N_trials	This argument (number of trials) is only needed when likelihood is binomial. Must have length one or equal to number of data points. Setting N_trials=1 corresponds to Bernoulli observation model.
skip_gen_quant	If this is true, the generated quantities block of Stan is skipped.
verbose	Should more verbose output be printed?

**Value**

An object of class `lgpmodel`.

**See Also**

For fitting the model, see [lgp\\_fit](#).

---

lgp_predict	<i>Compute predictions for a fitted model</i>
-------------	---

---

**Description**

Compute predictions for a fitted model. Only possible for models with Gaussian likelihood.

**Usage**

```
lgp_predict(fit, X_test, samples = "map", print_progress = TRUE,
            print_params = FALSE)
```

**Arguments**

fit	An object of class <code>lgpfit</code> .
X_test	The test points where the predictions should be computed.
samples	The predictions can be computed either by using only the posterior mean ( <code>samples="mean"</code> ), median ( <code>samples="median"</code> ), or MAP ( <code>samples="map"</code> ) parameters, or for all parameter samples ( <code>samples="all"</code> ). This can also be a set of indices, for example <code>samples=c(1:10)</code> gives predictions for the parameter samples 1...10.
print_progress	Should progress be printed (if there is more than one sample)?
print_params	Should the parameter values be printed? (only works if <code>samples</code> is mean or median.)

**Value**

A list.

**See Also**

- For creating an `lgpfit` object, see [lgp\\_fit](#).
- For creating an `lgpmodel` object, see [lgp\\_model](#).

---

lgp_test	<i>Compute predictions and log-posterior predictive density at test points</i>
----------	--

---

### Description

This is a convenience function that wraps `lgp_predict`, `compute_lppd` and `plot_posterior_y`.

### Usage

```
lgp_test(fit, test_data, plot = FALSE, verbose = TRUE,
         samples = "mean")
```

### Arguments

fit	an object of class <code>lgpfit</code>
test_data	a test data matrix
plot	should this return also a plot of the data and predictions?
verbose	Should this print progress?
samples	Sample indices or a keyword "mean", "median", "map", or "all".

### Value

a ggplot object or lppd

---

likelihood_as_int	<i>Convert likelihood string to Stan encoding</i>
-------------------	---

---

### Description

Convert likelihood string to Stan encoding

### Usage

```
likelihood_as_int(likelihood)
```

### Arguments

likelihood	a string
------------	----------

### Value

an integer

---

likelihood_as_str	<i>Convert the Stan likelihood encoding to a string</i>
-------------------	---

---

**Description**

Convert the Stan likelihood encoding to a string

**Usage**

```
likelihood_as_str(LH)
```

**Arguments**

LH	an integer
----	------------

**Value**

a string

---

log_gaussian_density	<i>Compute log-density for gaussian distribution</i>
----------------------	--

---

**Description**

Compute log-density for gaussian distribution

**Usage**

```
log_gaussian_density(x, mu, s2)
```

**Arguments**

x	point x
mu	mean
s2	variance

**Value**

a number

---

`matrix_to_df`*Matrix to data frame without editing column names*

---

**Description**

Matrix to data frame without editing column names

**Usage**

```
matrix_to_df(M)
```

**Arguments**

M                      a matrix

**Value**

a data frame

---

`model_info`*Get model info*

---

**Description**

Get model info

**Usage**

```
model_info(object, print = TRUE)
```

**Arguments**

object                  an object of class lgpmodel or lgpfit  
print                    should this print the info?

**Value**

the info as a string

---

nameComponents	<i>Create names for all components based on covariate names and types</i>
----------------	---

---

**Description**

Create names for all components based on covariate names and types

**Usage**

```
nameComponents(types, names)
```

**Arguments**

types	vector of covariate types
names	names of the covariates

**Value**

a vector of component names

---

onsetsToDiseaseAge	<i>Compute the disease-related ages</i>
--------------------	---

---

**Description**

Compute the disease-related ages

**Usage**

```
onsetsToDiseaseAge(onsets, age, k)
```

**Arguments**

onsets	true disease effect times, a vector of length N
age	the age covariate, a vector of length N*k
k	number of measurements per individual

**Value**

the diseaseAge covariate, a vector of length N\*k



---

```
parse_prior_distribution
```

*Turn a list describing a prior distribution into vectors to be given to Stan*

---

### Description

Turn a list describing a prior distribution into vectors to be given to Stan

### Usage

```
parse_prior_distribution(dist, add_correct = NULL)
```

### Arguments

<code>dist</code>	a list with field type, and possibly others
<code>add_correct</code>	additional correct parameter names

### Value

a list with two vectors to be given to Stan

---

```
parse_prior_t_effect
```

*Turn a list describing an effect time distribution into things to be given to Stan*

---

### Description

Turn a list describing an effect time distribution into things to be given to Stan

### Usage

```
parse_prior_t_effect(dist, N_cases, T_observed, T_last, UNCRT)
```

### Arguments

<code>dist</code>	This is <code>prior\$effect</code> , where <code>prior</code> is an argument of <code>lgp_model</code>
<code>N_cases</code>	number of case individuals
<code>T_observed</code>	observed disease onsets / initiation times
<code>T_last</code>	last time point for each diseased individual
<code>UNCRT</code>	0 or 1

### Value

a list with things to be given to Stan

---

```
plot,lgpfit,ANY-method
```

*Visualize a fitted 'lgpfit' object*

---

### Description

Visualize a fitted 'lgpfit' object

### Usage

```
## S4 method for signature 'lgpfit,ANY'
plot(fit, x = 1, y = 1, color_scheme = "red")
```

### Arguments

fit	an object of class <code>lgpfit</code>
x	does nothing
y	does nothing
color_scheme	bayesplot color scheme

### Value

a ggplot object

---

```
plot_beta
```

*Visualize posterior samples of individual-specific disease effect magnitude parameters*

---

### Description

Can only be used if the disease effect was modeled heterogeneously.

### Usage

```
plot_beta(fit, color_scheme = "red", threshold = 0.5)
```

### Arguments

fit	An object of class <code>lgpfit</code> .
color_scheme	Name of bayesplot color scheme.
threshold	Threshold for median.

### Value

a ggplot object

---

plot_component	<i>Helper function for plotting one component</i>
----------------	---

---

**Description**

Helper function for plotting one component

**Usage**

```
plot_component(MMM, SSS, model, idx, time_is_xvar, linealpha, linetype,
               fill_alpha, X_test, marker, sum_highlight, viridis_option)
```

**Arguments**

MMM	a n array of size n_samples x n_data x n_components
SSS	a n array of size n_samples x n_data x n_components
model	an object of class 'lgpmodel'
idx	Index of component to be plotted.
time_is_xvar	is the time variable the x-axis variable
linealpha	line alpha
linetype	line type
fill_alpha	fill alpha for geom_ribbons
X_test	optional matrix of test points
marker	point type
sum_highlight	name of a categorical covariate to be highlighted
viridis_option	the option argument of ggplot2::scale_colour_viridis_c by colour in the sum plot

**Value**

a ggplot object

---

plot_components	<i>Helper function for plotting components</i>
-----------------	--

---

**Description**

Helper function for plotting components

**Usage**

```
plot_components(MMM, SSS, model, time_is_xvar, X_test = NULL,
                sum_highlight = NULL, linealpha = 1, linetype = 1,
                fill_alpha = 0.3, marker = NULL, ncol = NULL, nrow = NULL,
                legend = NULL, labels = NULL, ylim = NULL, font_size = 9,
                theme = ggplot2::theme_linedraw(), legend_dir = "horizontal",
                xlabel = NULL, ylabel = " ", viridis_option = "viridis",
                return_list = FALSE)
```

**Arguments**

MMM	a n array of size n_samples x n_data x n_components
SSS	a n array of size n_samples x n_data x n_components
model	an object of class 'lgpmodel'
time_is_xvar	is the time variable the x-axis variable
X_test	optional matrix of test points
sum_highlight	name of a categorical covariate to be highlighted
linealpha	line alpha
linetype	line type
fill_alpha	fill alpha for geom_ribbons
marker	point type
ncol	number of plot columns
nrow	number of plot rows
legend	legend argument for ggarrange, use "none" to remove legends
labels	labels argument for ggarrange
ylim	y axis limits
font_size	font size for plots
theme	ggplot theme
legend_dir	direction of legend
xlabel	x-axis label
ylabel	y-axis label
viridis_option	the option argument of ggplot2::scale_colour_viridis_c by colour in the sum plot
return_list	should this return a list of ggplot objects instead of doing ggarrange

**Value**

an object returned by ggpubr::ggarrange or list

---

plot\_components\_posterior  
*Visualize inferred components*

---

**Description**

Visualize inferred components

**Usage**

```
plot_components_posterior(fit, subsamples = NULL, time_is_xvar = TRUE,
  PRED = NULL, marker = NULL, sample_idx = 1, n_sd = 2, ...)
```

**Arguments**

fit	An object of class lgpfit.
subsamples	How many samples to plot. If this is NULL, average over all samples is plotted. If this is "all", all samples are plotted.
time_is_xvar	is the time variable the x-axis variable in all subplots?
PRED	object returned by <a href="#">lgp_predict</a>
marker	point type
sample_idx	Which sample to plot.
n_sd	number of standard deviations (ribbon width)
...	additional arguments for <a href="#">plot_components</a>

**Value**

an object returned by `ggpubr::ggarrange` or a list of `ggplot2` objects

---

plot\_components\_posterior\_sub1

*Helper for [plot\\_components\\_posterior](#)*

---

**Description**

Helper for [plot\\_components\\_posterior](#)

**Usage**

```
plot_components_posterior_sub1(fit, subsamples, time_is_xvar, marker, ...)
```

**Arguments**

fit	An object of class lgpfit.
subsamples	How many samples to plot. If this is NULL, average over all samples is plotted. If this is "all", all samples are plotted.
time_is_xvar	is the time variable the x-axis variable in all subplots?
marker	point type
...	additional arguments for <a href="#">plot_components</a>

**Value**

an object returned by `ggpubr::ggarrange` or a list

---

plot\_components\_posterior\_sub2  
*Helper for [plot\\_components\\_posterior](#)*

---

### Description

Helper for [plot\\_components\\_posterior](#)

### Usage

```
plot_components_posterior_sub2(fit, PRED, sample_idx, time_is_xvar, n_sd,
    ...)
```

### Arguments

fit	An object of class lgpfit.
PRED	object returned by <a href="#">lgp_predict</a>
sample_idx	Which sample to plot.
time_is_xvar	is the time variable the x-axis variable in all subplots?
n_sd	number of standard deviations (ribbon width)
...	additional arguments for <a href="#">plot_components</a>

### Value

an object returned by ggpubr::ggarrange or a list

---

plot\_components\_simdata  
*Visualize the components of a simulated data set*

---

### Description

Visualize the components of a simulated data set

### Usage

```
plot_components_simdata(simData, time_is_xvar = TRUE, marker = 16, ...)
```

### Arguments

simData	simulated data object (list)
time_is_xvar	is the time variable the x-axis variable in all subplots?
marker	point marker
...	additional arguments for <a href="#">plot_components</a>

### Value

an object returned by ggpubr::ggarrange or list

---

plot_data	<i>A spaghetti plot of longitudinal data.</i>
-----------	---

---

**Description**

A spaghetti plot of longitudinal data.

**Usage**

```
plot_data(data, highlight = NULL, response = "y", id_variable = "id",
  time_variable = "age", psize = 2, lwd = 0.5, title = NULL)
```

**Arguments**

data	A data frame.
highlight	Name of a covariate to be highlighted with color, or id of a subject to be highlighted.
response	Name of the response variable.
id_variable	Name of id variable.
time_variable	Name of time variable.
psize	point size
lwd	line width
title	additional string added to title

**Value**

a ggplot object

---

plot_data_hl_cat	<i>A spaghetti plot of longitudinal data, highlighting a categorical covariate.</i>
------------------	---

---

**Description**

A spaghetti plot of longitudinal data, highlighting a categorical covariate.

**Usage**

```
plot_data_hl_cat(data, highlight = NULL, response = "y",
  id_variable = "id", time_variable = "age", psize = 2, lwd = 0.5)
```

**Arguments**

data	A data frame.
highlight	Name of a categorical covariate to be highlighted with color.
response	Name of the response variable.
id_variable	Name of id variable.
time_variable	Name of time variable.
psize	point size
lwd	line width

**Value**

a ggplot object

---

plot_data_hl_cont	<i>A spaghetti plot of longitudinal data, highlighting a continuous covariate.</i>
-------------------	--

---

**Description**

A spaghetti plot of longitudinal data, highlighting a continuous covariate.

**Usage**

```
plot_data_hl_cont(data, highlight = NULL, response = "y",
  id_variable = "id", time_variable = "age", psize = 2, lwd = 0.5,
  colgrad = ggplot2::scale_colour_gradient2())
```

**Arguments**

data	A data frame.
highlight	Name of a continuous covariate to be highlighted with color.
response	Name of the response variable.
id_variable	Name of id variable.
time_variable	Name of time variable.
psize	point size
lwd	line width
colgrad	color gradient

**Value**

a ggplot object

---

plot_data_hl_disease	<i>A spaghetti plot of longitudinal data, highlighting based on disease group.</i>
----------------------	--

---

**Description**

A spaghetti plot of longitudinal data, highlighting based on disease group.

**Usage**

```
plot_data_hl_disease(data, highlight = "diseaseAge", response = "y",
  id_variable = "id", time_variable = "age", psize = 2, lwd = 0.5)
```



**Arguments**

data	A data frame.
highlight	Name of the disease-related age variable.
response	Name of the response variable.
id_variable	Name of id variable.
time_variable	Name of time variable.
psize	point size
lwd	line width

**Value**

a ggplot object

---

plot\_data\_hl\_individual

*A spaghetti plot of longitudinal data, highlighting one individual.*

---

**Description**

A spaghetti plot of longitudinal data, highlighting one individual.

**Usage**

```
plot_data_hl_individual(data, highlight = 1, response = "y",  
  id_variable = "id", time_variable = "age", psize = 2, lwd = 0.5)
```

**Arguments**

data	A data frame.
highlight	Number indicating the individual to highlight.
response	Name of the response variable.
id_variable	Name of id variable.
time_variable	Name of time variable.
psize	point size
lwd	line width

**Value**

a ggplot object

---

plot_data_plain	<i>A spaghetti plot of longitudinal data without highlighting.</i>
-----------------	--

---

**Description**

A spaghetti plot of longitudinal data without highlighting.

**Usage**

```
plot_data_plain(data, response = "y", id_variable = "id",
  time_variable = "age", psize = 2, lwd = 0.5)
```

**Arguments**

data	A data frame.
response	Name of the response variable.
id_variable	Name of id variable.
time_variable	Name of time variable.
psize	point size
lwd	line width

**Value**

a ggplot object

---

plot_effect_times	<i>Visualize posterior uncertainty in the disease effect times</i>
-------------------	--

---

**Description**

Can only be used if the uncertainty of effect time was modeled.

**Usage**

```
plot_effect_times(fit, color_scheme = "red", prob = 1,
  prob_outer = 1, point_est = "none")
```

**Arguments**

fit	An object of class lgpfitt.
color_scheme	Name of bayesplot color scheme.
prob	Inner interval
prob_outer	Outer interval
point_est	Point estimate type

**Value**

a ggplot object

---

plot_inputwarp	<i>Visualize the input warping function for different parameter samples</i>
----------------	---

---

**Description**

Visualize the input warping function for different parameter samples

**Usage**

```
plot_inputwarp(fit, p = 300, color_scheme = "red", b = 0, c = 1)
```

**Arguments**

fit	An object of class lgpfit.
p	number of plot points
color_scheme	Name of bayesplot color scheme.
b	location of the effective time window (default = 0)
c	maximum range (default = 1)

**Value**

a ggplot object

---

plot_posterior_f	<i>Plot posterior of f</i>
------------------	----------------------------

---

**Description**

This is a wrapper for [plot\\_posterior\\_predictions](#).

**Usage**

```
plot_posterior_f(fit, PRED = NULL, plot_uncertainty = TRUE,
  data_marker = 16, n_sds = 2, ...)
```

**Arguments**

fit	An object of class lgpfit.
PRED	Predictions computed using lgp_predict.
plot_uncertainty	Should an uncertainty ribbon be plotted?
data_marker	pch for data points
n_sds	number of standard deviations for the uncertainty band width
...	additional arguments to <a href="#">plot_posterior_predictions</a>

**Value**

a ggplot object

---

plot\_posterior\_predictions

*Plot posterior of f or predictive distribution for y*


---

## Description

Plot posterior of f or predictive distribution for y

## Usage

```
plot_posterior_predictions(fit, mode, PRED = NULL,
  color_scheme = "red", color_scheme_t_effect = "gray", alpha = 0.5,
  alpha_line = 1, alpha2 = 0.5, plot_uncertainty = TRUE,
  title = NULL, ylim = NULL, plot_obs_onset = FALSE,
  plot_t_effect_samples = FALSE, ypos_dens = NULL, test_data = NULL,
  color_test = "deepskyblue2", pch_test = 21, size_test = 2,
  error_bar = FALSE, n_sds = 2, reference_times = NULL,
  post_t_effect_stat = "none", original_y_scale = TRUE,
  data_color = "black", data_marker = 21, ons_linetypes = c(1, 2, 3),
  ons_linecolors = c("black", "red", "gray50"))
```

## Arguments

fit	An object of class lgpfit.
mode	Must be either "posterior" or "predictive".
PRED	Predictions computed using lgp_predict.
color_scheme	Name of bayesplot color scheme or a list with fieds 'dark' and 'light'.
color_scheme_t_effect	color scheme name for effect time density plotting
alpha	Ribbon fill opacity.
alpha_line	Line opacity.
alpha2	alpha of t_onset density
plot_uncertainty	Should an uncertainty ribbon be plotted?
title	optional prefix to plot title
ylim	y axis limits
plot_obs_onset	should the observed disease onset/initiation time be plotted by a vertical line
plot_t_effect_samples	should a distribution of sampled effect times be plotted
ypos_dens	y-position of the density plot
test_data	Test data frame
color_test	test point color
pch_test	test point marker
size_test	test point size
error_bar	should uncertainty be plotted using error bars instead of a ribbon

n_sds	number of standard deviations for the uncertainty band width
reference_times	reference onset times
post_t_effect_stat	statistic computed from effect time samples (mean or median)
original_y_scale	should the predictions be scaled back to original data scale
data_color	data marker color
data_marker	data marker type
ons_linetypes	onset line types
ons_linecolors	onset line colors

**Value**

a ggplot object

---

plot_posterior_y	<i>Plot posterior predictive distribution</i>
------------------	---

---

**Description**

This is a wrapper for [plot\\_posterior\\_predictions](#).

**Usage**

```
plot_posterior_y(fit, PRED, uncertainty = "ribbon", test_data = NULL,
  data_marker = 16, n_sds = 2, ...)
```

**Arguments**

fit	An object of class lgpfit.
PRED	Predictions computed using lgp_predict.
uncertainty	Either "none", "ribbon" or "errorbar".
test_data	Test data set.
data_marker	pch for data points
n_sds	number of standard deviations for the uncertainty band width
...	additional arguments to <a href="#">plot_posterior_predictions</a>

**Value**

a ggplot object

---

plot\_predictions\_add\_onsets

*Add disease onset / effect times to predictions plot*


---

## Description

NOTE: currently assumes that diseased individuals come first.

## Usage

```
plot_predictions_add_onsets(fit, h, plot_obs_onset, plot_t_effect_samples,
  idvar, timevar, ypos_dens, color_scheme_t_effect, reference_times,
  post_t_effect_stat, linetypes = c(1, 2, 3), linecolors = c("black",
  "red", "gray50"), alpha2 = 1)
```

## Arguments

fit	An object of class lgpfit.
h	a ggplot object
plot_obs_onset	a boolean value
plot_t_effect_samples	a boolean value
idvar	id variable name
timevar	time variable name
ypos_dens	y position of the estimated onset density
color_scheme_t_effect	color scheme
reference_times	reference onset times
post_t_effect_stat	statistic computed from effect time samples
linetypes	onset line types
linecolors	onset line colors
alpha2	alpha parameter

## Value

a modified ggplot object

---

plot\_predictions\_options

*Do input checks and set options for plotting predictions*


---

### Description

Do input checks and set options for plotting predictions

### Usage

```
plot_predictions_options(fit, color_scheme, original_y_scale, PRED,
                        test_data, color_scheme_t_effect, mode, n_sds)
```

### Arguments

fit	An object of class lgpfit.
color_scheme	Name of bayesplot color scheme.
original_y_scale	Boolean value.
PRED	Predictions computed using lgp_predict.
test_data	test data
color_scheme_t_effect	Another color scheme.
mode	mode
n_sds	number of standard deviations for the uncertainty band width

### Value

a list

---

plot\_relevances

*Barplot of covariate relevances*


---

### Description

Barplot of covariate relevances

### Usage

```
plot_relevances(object, violin = FALSE, color_scheme = "red", ...)
```

### Arguments

object	an object of class lgpfit
violin	Should a violin plot be used instead of a boxplot
color_scheme	bayesplot color scheme name
...	Additional arguments to ggplot2::geom_boxplot or ggplot2::geom_violin.

**Value**

a ggplot object

---

plot_samples	<i>Visualize the distribution of the model parameter samples</i>
--------------	--

---

**Description**

This is a wrapper for functions in the bayesplot package.

**Usage**

```
plot_samples(object, pars = character(), regex_pars = character(),
  type = "intervals", prob = 0.5, prob_outer = 0.9,
  color_scheme = "red", point_est = "median", binwidth = NULL,
  transformations = list(), off_diag_args = list(size = 1),
  facet_args = list())
```

**Arguments**

object	An object of class lgpfit.
pars	parameter names
regex_pars	regex for parameter names
type	Visualization type. Must be either "dens", "areas", "intervals"(default) or "hist".
prob	inner interval
prob_outer	outer interval
color_scheme	See different color schemes in the bayesplot package.
point_est	the point estimate type
binwidth	width of histogram bins if type = "hist"
transformations	the parameter transformations
off_diag_args	Additional argument list for the pairs plot.
facet_args	additional facetting arguments

**Value**

a ggplot object



---

plot_simdata	<i>Plot a simulated longitudinal data set for each individual separately</i>
--------------	--

---

**Description**

Plot a simulated longitudinal data set for each individual separately

**Usage**

```
plot_simdata(simData, linecolor = "gray50", nrow = NULL, ncol = NULL,
             i_test = NULL, color_point = "black", color_test = "steelblue2",
             signal_name = "signal", y_transform = function(x) { x })
```

**Arguments**

simData	a list returned by <a href="#">simulate_data</a>
linecolor	line color
nrow	an argument for <code>ggplot2::facet_wrap</code>
ncol	an argument for <code>ggplot2::facet_wrap</code>
i_test	test point indices
color_point	data point color
color_test	test point color
signal_name	name of signal
y_transform	function to transform the data y

**Value**

a ggplot object

**See Also**

For plotting each component separately, see [plot\\_components\\_simdata](#)

---

postproc	<i>Finalize the lgpfit object after sampling</i>
----------	--

---

**Description**

Finalize the lgpfit object after sampling

**Usage**

```
postproc(fit, threshold = 0.95, relevance_method = "f_mean",
         verbose = FALSE)
```

**Arguments**

fit	An (incomplete) object of class <code>lgpfit</code> .
threshold	Threshold for relevance sum. Must be a value between 0 and 1.
relevance_method	Component relevance determination method. Must be either "f_mean" or "alpha".
verbose	Should some output be printed?

**Value**

An updated object of class `lgpfit`.

---

postproc_relevances	<i>Compute component relevances and estimate amount of noise (one MCMC sample)</i>
---------------------	--

---

**Description**

Compute component relevances and estimate amount of noise (one MCMC sample)

**Usage**

```
postproc_relevances(fit, relevance_method = "f_mean",
  noise_method = "SSE", verbose = FALSE)
```

**Arguments**

fit	An (incomplete) object of class <code>lgpfit</code> .
relevance_method	Component relevance determination method. Must be either "f_mean" or "alpha".
noise_method	Noise level determination method. Currently must be "SSE".
verbose	Should some output be printed?

**Value**

An updated object of class `lgpfit`.

---

predict_preproc	<i>Preprocess some things before computing predictions</i>
-----------------	--

---

### Description

This is a helper function for [lgp\\_predict](#).

### Usage

```
predict_preproc(fit, X_test, samples)
```

### Arguments

fit	An object of class <code>lgpfit</code> .
X_test	The test points where the predictions should be computed.
samples	The samples argument to <a href="#">lgp_predict</a>

---

PRED_to_arrays	<i>PRED object to arrays</i>
----------------	------------------------------

---

### Description

PRED object to arrays

### Usage

```
PRED_to_arrays(PRED)
```

### Arguments

PRED	an object returned by <a href="#">lgp_predict</a>
------	---

### Value

a list containing two arrays

---

print_prior	<i>Human-readable description of a specified prior</i>
-------------	--

---

**Description**

Print human-readable info about the prior specification that was used or will be used

**Usage**

```
print_prior(object)
```

**Arguments**

object            An object of class lgpfit or a valid prior argument for the ‘lgp’ function.

**Value**

nothing

---

prior_default	<i>Create the default prior</i>
---------------	---------------------------------

---

**Description**

Create the default prior

**Usage**

```
prior_default(sigma_alpha = 1)
```

**Arguments**

sigma\_alpha      Sigma parameter of the student-t distribution for all alpha.

**Value**

A list defining a valid prior argument for the lgp function.

---

prior_LonGP	<i>Create a similar default prior as in LonGP (Cheng et. al, 2019)</i>
-------------	--

---

**Description**

Not recommended, because a lengthscale close to 0 is possible.

**Usage**

```
prior_LonGP()
```

**Value**

A list defining a valid prior argument for the lgp\_model function.

---

prior\_stan\_to\_readable

*Human-readable information about the priors in the Stan data object*

---

### Description

Human-readable information about the priors in the Stan data object

### Usage

```
prior_stan_to_readable(stan_dat)
```

### Arguments

stan\_dat            The list that is passed as data to `rstan::sampling`.

### Value

Info as a string.

---

prior\_statement

*Human-readable prior statement*

---

### Description

Human-readable prior statement

### Usage

```
prior_statement(parname, TYP, P, dist, row_change = TRUE)
```

### Arguments

parname	parameter name
TYP	two integers
P	three real numbers
dist	list of distribution names
row_change	should a newline be last character?

### Value

Sampling statement as a string.

---

prior_to_stan	<i>Get priors as a format that can be input to Stan</i>
---------------	---

---

**Description**

Get priors as a format that can be input to Stan

**Usage**

```
prior_to_stan(D, prior, HMGNS, UNCRT, N_cases, T_observed, T_last)
```

**Arguments**

D	an integer vector of length 6
prior	The prior argument supplied to lgp().
HMGNS	Is diseaseAge assumed to have a homogenous effect (1) or not (0)?
UNCRT	Boolean value, is uncertainty of disease onset modeled?
N_cases	number of case individuals
T_observed	observed disease onsets
T_last	last time point for each diseased individual

**Value**

a list with all things related to priors that Stan needs

---

repvec	<i>Repeat a vector as a rows of an array</i>
--------	--

---

**Description**

Repeat a vector as a rows of an array

**Usage**

```
repvec(v, n)
```

**Arguments**

v	a vector of length m
n	number of times to repeat

**Value**

returns an array of size n x m

---

rtgeom	<i>Sample from the 'truncated geometric' distribution</i>
--------	---

---

**Description**

Sample from the 'truncated geometric' distribution

**Usage**

```
rtgeom(s, p, n = 1)
```

**Arguments**

s	an integer
p	a number between 0 and 1
n	number of samples

**Value**

an integer from the interval 1...n

---

scaleRelevances	<i>Scale the effect sizes</i>
-----------------	-------------------------------

---

**Description**

Scale the effect sizes

**Usage**

```
scaleRelevances(FFF, relevances, force_zeromean, i_skip)
```

**Arguments**

FFF	matrix where one column corresponds to one additive data component
relevances	the desired variance of each component (column)
force_zeromean	Should each component (excluding the disease age component) be forced to have a zero mean.
i_skip	induces of components for which the zero-mean forcing is skipped

**Value**

a new matrix FFF

---

selection	<i>Selection of relevant components</i>
-----------	---

---

**Description**

Selection of relevant components

**Usage**

```
selection(object, threshold = 0.95)
```

**Arguments**

object	An object of class lgpfit.
threshold	Threshold for relevance sum. Must be a value between 0 and 1.

**Value**

A named list

---

selection_fixed_threshold	<i>Select relevant components</i>
---------------------------	-----------------------------------

---

**Description**

Select relevant components

**Usage**

```
selection_fixed_threshold(rel, threshold)
```

**Arguments**

rel	a named vector of component relevances
threshold	value between 0 and 1

**Value**

indices of selected components (including "noise" always)



---

selection_prob	<i>Probabilistic selection of relevant components</i>
----------------	---

---

**Description**

Probabilistic selection of relevant components

**Usage**

```
selection_prob(object, p = function(x) { stats::dbeta(x, 100, 5) },
  h = 0.01, show_progbar = FALSE)
```

**Arguments**

object	An object of class lgpfit.
p	a function defining a density over interval [0,1]
h	discretization parameter for computing a quadrature
show_progbar	Should this show a progress bar?

**Value**

Selection probabilities for each component

---

selection_prob_fixed_threshold	<i>Selection probabilities using a fixed threshold</i>
--------------------------------	--

---

**Description**

Selection probabilities using a fixed threshold

**Usage**

```
selection_prob_fixed_threshold(relevances, threshold)
```

**Arguments**

relevances	The relevances\$samples slot of an lgpfit object.
threshold	value between 0 and 1

**Value**

proportion of times each component was selected

---

selection_prob_plot	<i>Plot of probabilistic selection of relevant components</i>
---------------------	---

---

**Description**

Plot of probabilistic selection of relevant components

**Usage**

```
selection_prob_plot(PROB, H, P)
```

**Arguments**

PROB	computed probabilities at points H
H	a grid on interval [0,1]
P	threshold probability distribution evaluated at H

**Value**

a ggplot object

---

set_C_hat	<i>Set C_hat (Non-gaussian observation models)</i>
-----------	--

---

**Description**

Set C\_hat (Non-gaussian observation models)

**Usage**

```
set_C_hat(C_hat, response, LH, N_trials)
```

**Arguments**

C_hat	the C_hat argument given as input to lgp
response	response variable
LH	likelihood as int
N_trials	the N_trials data (binomial likelihood)

**Value**

a real number

---

`set_N_cat`*Count numbers of different categories for each categorical variable*

---

**Description**

Count numbers of different categories for each categorical variable

**Usage**

```
set_N_cat(X, D)
```

**Arguments**

X	the design matrix
D	a vector of length 6

**Value**

a numeric vector

---

`set_N_trials`*Set N\_trials (binomial and Bernoulli observation models)*

---

**Description**

Set N\_trials (binomial and Bernoulli observation models)

**Usage**

```
set_N_trials(N_trials, response, LH)
```

**Arguments**

N_trials	the N_trials argument given as input to lgp
response	response variable
LH	likelihood as int

**Value**

a numeric vector

---

show,lgpfit-method	<i>Show a summary of results of the lgp function</i>
--------------------	--

---

**Description**

Show a summary of results of the lgp function

**Usage**

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

**Arguments**

object                    an object of class lgpfit

**Value**

nothing

---

show,lgpmodel-method	<i>Show a summary of an lgpmodel</i>
----------------------	--------------------------------------

---

**Description**

Show a summary of an lgpmodel

**Usage**

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

**Arguments**

object                    an object of class lgpmodel

**Value**

nothing

---

simdata\_colnames\_pretty

*Simulated data column names in a prettier form*


---

**Description**

Simulated data column names in a prettier form

**Usage**

```
simdata_colnames_pretty(cn)
```

**Arguments**

cn                      column names

**Value**

names of model components

---

simulate\_data

*Generate an artificial longitudinal data set*


---

**Description**

Generate an artificial longitudinal data set.

**Usage**

```
simulate_data(N, t_data, covariates = c(), names = NULL,
  relevances = c(1, 1, rep(1, length(covariates))), n_cats = rep(2,
  sum(covariates %in% c(2, 3))), t_jitter = 0, lengthscales = rep(12,
  2 + sum(covariates %in% c(0, 1, 2))), f_var = 1,
  noise_type = "Gaussian", snr = 3, phi = 1,
  N_affected = round(N/2), t_effect_range = "auto",
  t_observed = "after_0", C_hat = 0, dis_fun = "gp_vm",
  bin_kernel = FALSE, steepness = 0.5, vm_params = c(0.025, 1),
  continuous_info = list(mu = c(pi/8, pi, -0.5), lambda = c(pi/8, pi,
  1)), N_trials = 1, verbose = FALSE, force_zeromean = TRUE)
```

**Arguments**

N                      Number of individuals.

t\_data                Measurement times.

covariates            Integer vector that defines the types of covariates (other than id and age). If not given, only the id and age covariates are created. Different integers correspond to the following covariate types:

- 0 = disease-related age
- 1 = other continuous covariate

	<ul style="list-style-type: none"> <li>• 2 = a categorical covariate that interacts with age</li> <li>• 3 = a categorical covariate that acts as a group offset</li> <li>• 4 = a categorical covariate that that acts as a group offset AND is restricted to have value 0 for controls and 1 for cases</li> </ul>
names	Covariate names.
relevances	Relative relevance of each component. Must have be a vector so that $\text{length}(\text{relevances}) = 2 + \text{length}(\text{covariates})$ . First two values define the relevance of the individual-specific age and shared age component, respectively.
n_cats	An integer vector defining the number of categories for each categorical covariate, so that $\text{length}(\text{n_cats})$ equals to the number of 2's and 3's in the covariates vector.
t_jitter	Standard deviation of the jitter added to the given measurement times.
lengthscales	A vector so that $\text{length}(\text{lengthscales}) = 2 + \text{sum}(\text{covariates} \%in\% c(0,1,2))$ .
f_var	variance of f
noise_type	Either "Gaussian", "Poisson", "NB" (negative binomial) or "binomial".
snr	The desired signal-to-noise ratio. This argument is valid only with <code>noise_type = "Gaussian"</code> .
phi	The dispersion parameter for negative binomial data. The variance is $g + g^2/\phi$ .
N_affected	Number of diseased individuals that are affected by the disease. This defaults to the number of diseased individuals. This argument can only be given if covariates contains a zero.
t_effect_range	Time interval from which the disease effect times are sampled uniformly. Alternatively, This can any function that returns the (possibly randomly generated) real disease effect time for one individual.
t_observed	Determines how the disease effect time is observed. This can be any function that takes the real disease effect time as an argument and returns the (possibly randomly generated) observed onset/initiation time. Alternatively, this can be a string of the form "after_n" or "random_p" or "exact".
C_hat	A constant added to f
dis_fun	A function or a string that defines the disease effect. If this is a function, that function is used to generate the effect. If <code>dis_fun</code> is "gp_vm" or "gp_ns", the disease component is drawn from a nonstationary GP prior (vm is the variance masked version of it).
bin_kernel	Should the binary kernel be used for categorical covariates? If this is TRUE, the effect will exist only for group 1.
steepness	Steepness of the input warping function. This is only used if the disease component is in the model.
vm_params	Parameters of the variance mask function. This is only needed if <code>useMaskedVarianceKernel = TRUE</code> .
continuous_info	<p>Info for generating continuous covariates. Must be a list containing fields <code>lambda</code> and <code>mu</code>, which have length 3. The continuous covariates are generated so that <math>x &lt; -\sin(a*t + b) + c</math>, where</p> <ul style="list-style-type: none"> <li>• <math>t \leftarrow \text{seq}(0, 2*\pi, \text{length.out} = k)</math></li> </ul>

```

      • a <- mu[1] + lambda[1]*stats::runif(1)
      • b <- mu[2] + lambda[2]*stats::runif(1)
      • c <- mu[3] + lambda[3]*stats::runif(1)
N_trials      The number of trials parameter for binomial data.
verbose        verbosity mode
force_zeromean Should each component (excluding the disease age component) be forced to
                have a zero mean?

```

## Value

A list out, where

- `out$data` is a data frame containing the actual data and
- `out$components` contains more points for smoother visualizations of the generating process.
- `out$onsets` contains the real disease effect times
- `out$p_signal` proportion of variance explained by signal

## Examples

```

# Generate Gaussian data
dat <- simulate_data(N = 4, t_data = c(6,12,24,36,48), snr = 3)

# Generate negative binomially distributed count data
dat <- simulate_data(N = 6, t_data = seq(2, 10, by = 2), noise_type = "NB", phi = 2)

```

---

<code>simulate_kernels</code>	<i>Compute all kernel matrices when simulating data</i>
-------------------------------	---

---

## Description

Compute all kernel matrices when simulating data

## Usage

```
simulate_kernels(X, types, lengthscales, X_affected, bin_kernel,
  useMaskedVarianceKernel, steepness, vm_params)
```

## Arguments

<code>X</code>	covariates
<code>types</code>	vector of covariate types, so that <ul style="list-style-type: none"> <li>• 1 = ID</li> <li>• 2 = age</li> <li>• 3 = diseaseAge</li> <li>• 4 = other continuous covariate</li> <li>• 5 = a categorical covariate that interacts with age</li> <li>• 6 = a categorical covariate that acts as an offset</li> </ul>
<code>lengthscales</code>	vector of lengthscales

<code>X_affected</code>	which individuals are affected by the disease
<code>bin_kernel</code>	whether or not binary (mask) kernel should be used for categorical covariates
<code>useMaskedVarianceKernel</code>	should the masked variance kernel be used for drawing the disease component
<code>steepness</code>	steepness of the input warping function
<code>vm_params</code>	parameters of the variance mask function

**Value**

a 3D array

---

`sim_check_covariates` *Input check for the covariates-related arguments of `simulate_data`*

---

**Description**

Input check for the covariates-related arguments of `simulate_data`

**Usage**

```
sim_check_covariates(covariates, relevances, names, n_cat)
```

**Arguments**

<code>covariates</code>	argument to <code>simulate_data</code>
<code>relevances</code>	argument to <code>simulate_data</code>
<code>names</code>	argument to <code>simulate_data</code>
<code>n_cat</code>	the <code>n_categs</code> argument to <code>simulate_data</code>

**Value**

the covariate names

---

`sim_data_to_observed` *Real generated disease ages to observed ones*

---

**Description**

Real generated disease ages to observed ones

**Usage**

```
sim_data_to_observed(dat, t_observed)
```

**Arguments**

<code>dat</code>	data frame
<code>t_observed</code>	Determines how the disease onset is observed. See documentation of <a href="#">simulate_data</a> .

**Value**

a new data frame and observed onsets



---

sim_generate_names	<i>Generate names for covariates</i>
--------------------	--------------------------------------

---

**Description**

Generate names for covariates

**Usage**

```
sim_generate_names(covariates)
```

**Arguments**

covariates      vector of covariate types

**Value**

covariate names

---

sim_parse_t_obs	<i>Parse the t_observed argument of simulate_data</i>
-----------------	---

---

**Description**

Parse the t\_observed argument of simulate\_data

**Usage**

```
sim_parse_t_obs(t_observed)
```

**Arguments**

t\_observed      a string

**Value**

a list with a name and number

---

split_data	<i>Split data into training and test data according to given row indices</i>
------------	--

---

**Description**

Split data into training and test data according to given row indices

**Usage**

```
split_data(data, i_test, sort_ids = TRUE)
```

**Arguments**

data	a data frame
i_test	test data row indices
sort_ids	should the test indices be sorted into increasing order

**Value**

a list(train, test)

---

split_data_by_id	<i>Split data into training and test data according to given individuals</i>
------------------	--

---

**Description**

Split data into training and test data according to given individuals

**Usage**

```
split_data_by_id(data, test_ids, id_variable = "id")
```

**Arguments**

data	a data frame
test_ids	test data individual identifiers
id_variable	name of id variable

**Value**

a list(train, test)

---

`split_data_by_timepoint`*Split data into training and test data according to time point indices*

---

**Description**

Split data into training and test data according to time point indices

**Usage**

```
split_data_by_timepoint(data, test_idx, id_variable = "id",  
  time_variable = "age")
```

**Arguments**

<code>data</code>	a data frame
<code>test_idx</code>	indices of test time points
<code>id_variable</code>	name of id variable
<code>time_variable</code>	name of time variable

**Value**

a list(train, test)

---

`split_data_random`*Split data into training and test data randomly*

---

**Description**

Split data into training and test data randomly

**Usage**

```
split_data_random(data, p_test = 0.1, n_test = NULL)
```

**Arguments**

<code>data</code>	a data frame
<code>p_test</code>	desired proportion of test data
<code>n_test</code>	desired number of test data points (if NULL, p_test is used to compute this)

**Value**

a list(train, test)

---

```
split_data_random_each
```

*Split data into training and test data by selecting randomly  $k$  points from each individual*

---

### Description

Split data into training and test data by selecting randomly  $k$  points from each individual

### Usage

```
split_data_random_each(data, n_test = 1, id_variable = "id",
  time_variable = "age")
```

### Arguments

<code>data</code>	a data frame
<code>n_test</code>	desired number of test data points per individual
<code>id_variable</code>	name of id variable
<code>time_variable</code>	name of time variable

### Value

a list(train, test)

---

```
standardize_inputs
```

*Standardize continuous input variables in  $X$*

---

### Description

Standardize continuous input variables in  $X$

### Usage

```
standardize_inputs(X, D)
```

### Arguments

<code>X</code>	the design matrix
<code>D</code>	the covariate types, a vector of length 6

### Value

updated  $X$  and info about scaling

---

stan_input_X_and_D	<i>Predictor covariates and types to Stan input</i>
--------------------	---

---

**Description**

Reorders covariates and takes only those that are needed

**Usage**

```
stan_input_X_and_D(data, varInfo, types, formula, verbose)
```

**Arguments**

data	a data frame containing the covariates
varInfo	original variable type info
types	types of the covariates
formula	model formula
verbose	can this print some info?

**Value**

X and needed types and updated varInfo

---

validate_prior	<i>Validate prior by sampling the signal and noise from it</i>
----------------	--

---

**Description**

Validate prior by sampling the signal and noise from it

**Usage**

```
validate_prior(model, chains = 4, iter = 1000, parallel = FALSE)
```

**Arguments**

model	An object of class <code>lgpmodel</code> .
chains	how many chains are used to sample from the prior
iter	for how many iterations are the chains run
parallel	should the chains be run in parallel?

**Value**

An object of class `lgpfit` and random samples of both 'f' and 'y'.

---

warp_input	<i>Warp inputs</i>
------------	--------------------

---

**Description**

Warp inputs

**Usage**

```
warp_input(t, a, b, c)
```

**Arguments**

t	a vector
a	steepness of the rise
b	location of the effective time window
c	maximum range

**Value**

a vector of warped inputs  $w(t)$

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