

# Package 'BIGL'

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**Type** Package

**Title** Biochemically Intuitive Generalized Loewe Model

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**Description** Response surface methods for drug synergy analysis. Available methods include generalized and classical Loewe formulations as well as Highest Single Agent methodology. Response surfaces can be plotted in an interactive 3-D plot and formal statistical tests for presence of synergistic effects are available. Implemented methods and tests are described in the article "BIGL: Biochemically Intuitive Generalized Loewe null model for prediction of the expected combined effect compatible with partial agonism and antagonism" by Koen Van der Borgh, Annelies Tourny, Rytis Bagdziunas, Olivier Thas, Maxim Nazarov, Heather Turner, Bie Verbist & Hugo Ceulemans (2017) <doi:10.1038/s41598-017-18068-5>.

**License** GPL-3

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Blissindependence	<i>Bliss Independence Model</i>
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### Description

This function returns fractional response levels for when these are based on Bliss Independence Model.

### Usage

```
Blissindependence(doseInput, parmInput, ...)
```

### Arguments

doseInput	Dose-response dataframe containing "d1" and "d2" columns
parmInput	Numeric vector or list with appropriately named parameter inputs. Typically, it will be coefficients from a MarginalFit object.
...	Further arguments that are currently unused

---

bootstrapData	<i>Data generating function used for constructing null distribution of meanR and maxR statistics</i>
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---

### Description

This function uses [simulateNull](#) and simulates all necessary steps to calculate null distribution which will furtherly be used in either [meanR](#) or [maxR](#) functions.

### Usage

```
bootstrapData(data, fitResult, transforms = fitResult$transforms,
  null_model = c("loewe", "hsa", "bliss"), ...)
```

**Arguments**

<code>data</code>	Dose-response dataframe.
<code>fitResult</code>	Monotherapy (on-axis) model fit, e.g. produced by <code>fitMarginals</code> . It has to be a "MarginalFit" object or a list containing <code>df</code> , <code>sigma</code> , <code>coef</code> , <code>shared_asymptote</code> and <code>method</code> elements for, respectively, marginal model degrees of freedom, residual standard deviation, named vector of coefficient estimates, logical value of whether shared asymptote is imposed and method for estimating marginal models during bootstrapping (see <code>fitMarginals</code> ). If biological and power transformations were used in marginal model estimation, <code>fitResult</code> should contain <code>transforms</code> elements with these transformations. Alternatively, these can also be specified via <code>transforms</code> argument.
<code>transforms</code>	Transformation functions. If non-null, <code>transforms</code> is a list containing 5 elements, namely biological and power transformations along with their inverse functions and <code>composeArgs</code> which is a list with argument values shared across the 4 functions. See <code>vignette</code> for more information.
<code>null_model</code>	Specified null model for the expected response surface. Currently, allowed options are "loewe" for generalized Loewe model, and "hsa" for Highest Single Agent model.
<code>...</code>	Further arguments that will be passed to <code>generateData</code> function

---

`boxcox.transformation` *Apply two-parameter Box-Cox transformation*

---

**Description**

Apply two-parameter Box-Cox transformation

**Usage**

```
boxcox.transformation(y, lambda, alpha = 0)
```

**Arguments**

<code>y</code>	Numeric vector
<code>lambda</code>	Power parameter in power transform
<code>alpha</code>	Shift parameter in 2-parameter power transform. Defaults to 0 which implies a 1-parameter Box-Cox transform.

**Value**

Power-transformed data

---

coef.MarginalFit	<i>Coefficients from marginal model estimation</i>
------------------	--

---

**Description**

Coefficients from marginal model estimation

**Usage**

```
## S3 method for class 'MarginalFit'
coef(object, ...)
```

**Arguments**

object	Output of <code>fitMarginals</code> function
...	Further arguments

---

constructFormula	<i>Construct a model formula from parameter constraint matrix</i>
------------------	---

---

**Description**

For parameter names defined in naming vector, formula is constructed so that `consMatrix %*% naming = consVector` is satisfied. Constraint coefficients are normalized and convert into fractions.

**Usage**

```
constructFormula(consMatrix = NULL, consVector = NULL,
  naming = c("h1", "h2", "b", "m1", "m2", "e1", "e2"),
  extraVars = c("d1", "d2"), formulaArgs = c("effect", "fn"))
```

**Arguments**

consMatrix	Constraint matrix
consVector	Constraint vector
naming	Parameter names
extraVars	Non-parameter variables used in the formula and function evaluation. These will be appended to the formula.
formulaArgs	Character vector of length two. First element indicates name for the response variable. Second element indicates name of the function.

**Value**

This function returns a model construct appropriate for `fitMarginals` function. It also separates variables into those that are free and those which are constrained.

**Examples**

```
constM <- rbind(c(0, 0, 1, 0, 0, 0, 0),
               c(0, 0, 0, -1, 1, 0, 0))
constV <- c(0.9, 0)
constructFormula(constM, constV)
```

---

contour.ResponseSurface

*Method for plotting of contours based on maxR statistics*

---

**Description**

Method for plotting of contours based on maxR statistics

**Usage**

```
## S3 method for class 'ResponseSurface'
contour(x, ...)
```

**Arguments**

x	Output of <a href="#">fitSurface</a>
...	Further parameters passed to <a href="#">plot.maxR</a>

---

CPBootstrap

*Estimate CP matrix with bootstrap*

---

**Description**

This function is generally called from within [fitSurface](#).

**Usage**

```
CPBootstrap(data, fitResult, transforms = fitResult$transforms,
            null_model = c("loewe", "hsa", "bliss"), B.CP, ...)
```

**Arguments**

data	Dose-response dataframe.
fitResult	Monotherapy (on-axis) model fit, e.g. produced by <code>fitMarginals</code> . It has to be a "MarginalFit" object or a list containing <code>df</code> , <code>sigma</code> , <code>coef</code> , <code>shared_asymptote</code> and <code>method</code> elements for, respectively, marginal model degrees of freedom, residual standard deviation, named vector of coefficient estimates, logical value of whether shared asymptote is imposed and method for estimating marginal models during bootstrapping (see <code>fitMarginals</code> ). If biological and power transformations were used in marginal model estimation, <code>fitResult</code> should contain <code>transforms</code> elements with these transformations. Alternatively, these can also be specified via <code>transforms</code> argument.
transforms	Transformation functions. If non-null, <code>transforms</code> is a list containing 5 elements, namely biological and power transformations along with their inverse functions and <code>compositeArgs</code> which is a list with argument values shared across the 4 functions. See vignette for more information.
null_model	Specified null model for the expected response surface. Currently, allowed options are "loewe" for generalized Loewe model, and "hsa" for Highest Single Agent model.
B.CP	Number of bootstrap iterations to use for CP matrix estimation
...	Further parameters that will be passed to <code>generateData</code>

**Value**

Estimated CP matrix

**Examples**

```
data <- subset(directAntivirals, experiment == 5)
## Data must contain d1, d2 and effect columns
fitResult <- fitMarginals(data)
CPBootstrap(data, fitResult, null_model = "loewe", B.CP = 5)
```

---

df.residual.MarginalFit

*Residual degrees of freedom in marginal model estimation*

---

**Description**

Residual degrees of freedom in marginal model estimation

**Usage**

```
## S3 method for class 'MarginalFit'
df.residual(object, ...)
```

**Arguments**

object            Output of `fitMarginals` function  
...                Further arguments

---

directAntivirals        *Partial data with combination experiments of direct-acting antivirals*

---

**Description**

A dataset containing 11 combination experiments of direct-acting antivirals.

**Format**

A data frame with 3520 rows and 6 variables:

- experiment: ID of experiment (1-11)
- cpd1: name of the first compound (4 different compounds)
- cpd2: name of the second compound (11 different compounds)
- effect: observed effect (cell count)
- d1: dose of the first compound
- d2: dose of the second compound

---

directAntivirals\_ALL    *Full data with combination experiments of direct-acting antivirals*

---

**Description**

A dataset containing 11 combination experiments of direct-acting antivirals. This dataset is larger than `directAntivirals` dataset as it includes concentrations at levels of  $1e6$  which can render plots visually unappealing.

**Format**

A data frame with 4224 rows and 6 variables:

- experiment: ID of experiment (1-11)
- cpd1: name of the first compound (4 different compounds)
- cpd2: name of the second compound (11 different compounds)
- effect: observed effect (cell count)
- d1: dose of the first compound
- d2: dose of the second compound



---

fitMarginals

*Fit two 4-parameter log-logistic functions for a synergy experiment*


---

## Description

This function uses dose-response data for two compounds and estimates coefficients for monotherapy models of both of these compounds such that they share a common baseline. Currently, these coefficients are estimated by default using a non-linear least squares approximation. Although entire dose-response data can be provided, estimation will subset the part of data where at least one of the compounds is dosed at zero, i.e. on-axis data.

## Usage

```
fitMarginals(data, transforms = NULL, start = NULL,
             constraints = NULL, fixed = NULL, method = c("nls", "nls",
             "optim"), names = NULL, ...)
```

## Arguments

data	Dose-response dataframe. Marginal data will be extracted from it automatically.
transforms	Transformation functions. If non-null, transforms is a list containing 5 elements, namely biological and power transformations along with their inverse functions and compositeArgs which is a list with argument values shared across the 4 functions. See vignette for more information.
start	Starting parameter values. If not specified, they will be obtained from <a href="#">initialMarginal</a> .
constraints	List of constraint matrix and vector which will be passed to <a href="#">constructFormula</a> . If constraints = NULL, no constraints on parameter estimation will be imposed.
fixed	This arguments provides a user-friendly alternative to impose a fixed value for marginal parameters. It must be a named vector with names contained in c("h1", "h2", "b", "m1", "m2"). For example, fixed = c("m1" = 1, "h1" = 1) will automatically generate appropriate constraint matrix and vector to set the maximal response and the Hill coefficient of the first compound to 1. If both constraints and fixed arguments are passed, then only fixed will be used.
method	Which estimation method should be used to obtain the estimates. If method = "nls", simple non-linear least squares <a href="#">nls</a> will be used. If method = "nlsLM" Levenberg-Marquardt non-linear least squares <a href="#">nlsLM</a> is used instead (default). If method = "optim", residual sum of squares will be minimized using general purpose optimization based on Nelder-Mean algorithm in <a href="#">optim</a> . This method can be noticeably slower than the non-linear least squares methods.
names	Compound names to be used on the plot labels.
...	Further arguments that are passed to the optimizer function, such as lower or upper (for the "nlsLM" method), or control.

**Details**

Model formula is specified as `effect ~ fn(h1, h2, ...)` where `fn` is a hard-coded function which fits two 4-parameter log-logistic functions simultaneously so that the baseline can be shared. If transformation functions are provided, `fn` is consequently adjusted to account for them.

**Value**

This function returns a `MarginalFit` object with monotherapy coefficient estimates and diverse information regarding monotherapy estimation. `MarginalFit` object is essentially a list with appropriately named elements.

Among these list elements, `"coef"` is a named vector with parameter estimates. `h1` and `h2` are Hill's slope coefficients for each of the compounds, `m1` and `m2` are their maximal response levels whereas `b` is the shared baseline. Lastly, `e1` and `e2` are log-transformed EC50 values.

`"sigma"` is standard deviation of residuals for the estimated monotherapy model and `"df"` is the degrees of freedom for the residuals. `"vcov"` is the variance-covariance matrix of the estimated parameters.

Return object also contains information regarding data, biological and power transformations used in this estimation as well as model construct and method of estimation.

**Examples**

```
data <- subset(directAntivirals, experiment == 1)
## Data must contain d1, d2 and effect columns
transforms <- getTransformations(data)
fitMarginals(data, transforms)
```

---

fitSurface

---

*Fit response surface model and compute meanR and maxR statistics*


---

**Description**

This function computes predictions for off-axis dose combinations according to the BIGL or HSA null model and, if required, computes appropriate meanR and maxR statistics. Function requires as input dose-response dataframe and output of `fitMarginals` containing estimates for the monotherapy model. If transformation functions were used in monotherapy estimation, these should also be provided.

**Usage**

```
fitSurface(data, fitResult, transforms = fitResult$transforms,
  null_model = c("loewe", "hsa", "bliss"), effect = "effect",
  d1 = "d1", d2 = "d2", statistic = c("none", "meanR", "maxR",
  "both"), CP = NULL, B.CP = 50, B.B = NULL,
  nested_bootstrap = FALSE, error = 4, sampling_errors = NULL,
  wild_bootstrap = FALSE, cutoff = 0.95, parallel = TRUE,
  method = c("equal", "model", "unequal"))
```

**Arguments**

data	Dose-response dataframe.
fitResult	Monotherapy (on-axis) model fit, e.g. produced by <a href="#">fitMarginals</a> . It has to be a "MarginalFit" object or a list containing df, sigma, coef, shared_asymptote and method elements for, respectively, marginal model degrees of freedom, residual standard deviation, named vector of coefficient estimates, logical value of whether shared asymptote is imposed and method for estimating marginal models during bootstrapping (see <a href="#">fitMarginals</a> ). If biological and power transformations were used in marginal model estimation, fitResult should contain transforms elements with these transformations. Alternatively, these can also be specified via transforms argument.
transforms	Transformation functions. If non-null, transforms is a list containing 5 elements, namely biological and power transformations along with their inverse functions and compositeArgs which is a list with argument values shared across the 4 functions. See vignette for more information.
null_model	Specified null model for the expected response surface. Currently, allowed options are "loewe" for generalized Loewe model, and "hsa" for Highest Single Agent model.
effect	Name of the response column in the data ("effect")
d1	Name of the column with doses of the first compound ("d1")
d2	Name of the column with doses of the second compound ("d2")
statistic	Which statistics should be computed. This argument can take one of the values from c("none", "meanR", "maxR", "both").
CP	Prediction covariance matrix. If not specified, it will be estimated by bootstrap using B.CP iterations.
B.CP	Number of bootstrap iterations to use for CP matrix estimation
B.B	Number of iterations to use in bootstrapping null distribution for either meanR or maxR statistics.
nested_bootstrap	When statistics are calculated, if nested_bootstrap = TRUE, CP matrix is recalculated at each bootstrap iteration of B.B using B.CP iterations. Using such nested bootstrap may however significantly increase computational time. If nested_bootstrap = FALSE, CP bootstrapped data reuses CP matrix calculated from the original data.
error	Type of error for resampling in the bootstrapping procedure. This argument will be passed to <a href="#">generateData</a> . If error = 4 (default), the error terms for generating distribution of the null will be resampled from the vector specified in sampling. If error = 1, normal errors are added. If error = 2, errors are sampled from a mixture of two normal distributions. If error = 3, errors are generated from a rescaled chi-square distribution.
sampling_errors	Sampling vector to resample errors from. Used only if error is 4 and is passed as argument to <a href="#">generateData</a> . If sampling_errors = NULL (default), mean residuals at off-axis points between observed and predicted response are taken.

wild_bootstrap	Whether special bootstrap to correct for heteroskedasticity should be used. If wild_bootstrap = TRUE, errors are generated from sampling_errors multiplied by a random variable following Rademacher distribution. Argument is used only if error = 4.
cutoff	Cut-off to use in maxR procedure for declaring non-additivity (default is 0.95).
parallel	Whether parallel computing should be used for bootstrap. This parameter can take either integer value to specify the number of threads to be used or logical TRUE/FALSE. If parallel = TRUE, then max(1, detectCores()-1) is set to be the number of threads. If parallel = FALSE, then a single thread is used and cluster object is not created.
method	What assumption should be used for the variance of on- and off-axis points. This argument can take one of the values from c("equal", "model", "unequal"). With the value "equal" as the default. "equal" assumes that both on- and off-axis points have the same variance, "unequal" estimates a different parameter for on- and off-axis points and "model" predicts variance based on the average effect of an off-axis point. If no transformations are used the "model" method is recommended. If transformations are used, only the "equal" method can be chosen.

## Details

Please see the example vignette `vignette("analysis", package = "BIGL")` and the report "Lack of fit test for detecting synergy" included in the papers folder for further details on the test statistics used: `system.file("papers", "newStatistics.pdf", package = "BIGL")`

## Value

This function returns a ResponseSurface object with estimates of the predicted surface. ResponseSurface object is essentially a list with appropriately named elements.

Elements of the list include input data, monotherapy model coefficients and transformation functions, null model used to construct the surface as well as estimated CP matrix (see [CPBootstrap](#)), occupancy level at each dose combination according to the generalized Loewe model and "offAxisTable" element which contains observed and predicted effects as well as estimated z-scores for each dose combination.

If statistical testing was done, returned object contains "meanR" and "maxR" elements with output from [meanR](#) and [maxR](#) respectively.

## Examples

```
## Not run:
data <- subset(directAntivirals, experiment == 4)
## Data should contain d1, d2 and effect columns
transforms <- list("PowerT" = function(x, args) with(args, log(x)),
                 "InvPowerT" = function(y, args) with(args, exp(y)),
                 "BioIT" = function(x, args) with(args, N0 * exp(x * time.hours)),
                 "InvBioIT" = function(y, args) with(args, 1/time.hours * log(y/N0)),
                 "compositeArgs" = list(N0 = 1, time.hours = 72))
fitResult <- fitMarginals(data, transforms)
```

```
surf <- fitSurface(data, fitResult, statistic = "meanR")
summary(surf)

## End(Not run)
```

---

fitted.MarginalFit      *Compute fitted values from monotherapy estimation*

---

### Description

Compute fitted values from monotherapy estimation

### Usage

```
## S3 method for class 'MarginalFit'
fitted(object, ...)
```

### Arguments

object	Output of <code>fitMarginals</code> function
...	Further arguments

---

fitted.ResponseSurface  
*Predicted values of the response surface according to the given null model*

---

### Description

Predicted values of the response surface according to the given null model

### Usage

```
## S3 method for class 'ResponseSurface'
fitted(object, ...)
```

### Arguments

object	Output of <code>fitSurface</code>
...	Further parameters

---

generalizedLoewe	<i>Compute combined predicted response from drug doses according to standard or generalized Loewe model.</i>
------------------	--

---

**Description**

Compute combined predicted response from drug doses according to standard or generalized Loewe model.

**Usage**

```
generalizedLoewe(doseInput, parmInput, asymptotes = 2, ...)
```

**Arguments**

doseInput	Dose-response dataframe containing "d1" and "d2" columns
parmInput	Numeric vector or list with appropriately named parameter inputs. Typically, it will be coefficients from a <code>MarginalFit</code> object.
asymptotes	Number of asymptotes. It can be either 1 as in standard Loewe model or 2 as in generalized Loewe model.
...	Further arguments that are currently unused

---

generateData	<i>Generate data from parameters of marginal monotherapy model</i>
--------------	--

---

**Description**

This function is used to generate data for bootstrapping of the null distribution for various estimates. Optional arguments such as specific choice of sampling vector or corrections for heteroskedasticity can be specified in the function arguments.

**Usage**

```
generateData(pars, sigma, data = NULL, transforms = NULL,
  null_model = c("loewe", "hsa", "bliss"), error = 1,
  sampling_errors = NULL, wild_bootstrap = FALSE, ...)
```

**Arguments**

pars	Coefficients of the marginal model along with their appropriate naming scheme. These will typically be estimated using <code>fitMarginals</code> . Furthermore, pars can simply be a <code>MarginalFit</code> object and transforms object will be automatically extracted.
sigma	Standard deviation to use for randomly generated error terms. This argument is unused if <code>error = 4</code> so that sampling error vector is provided.

data	Data frame with dose columns ("d1", "d2") to generate the effect for. Only "d1" and "d2" columns of the dose-response dataframe should be passed to this argument. "effect" column should not be passed and if it is, the column will be replaced by simulated data.
transforms	Transformation functions. If non-null, transforms is a list containing 5 elements, namely biological and power transformations along with their inverse functions and compositeArgs which is a list with argument values shared across the 4 functions. See vignette for more information.
null_model	Specified null model for the expected response surface. Currently, allowed options are "loewe" for generalized Loewe model, and "hsa" for Highest Single Agent model.
error	Type of error for resampling. error = 1 (Default) adds normal errors to the simulated effects, error = 2 adds errors sampled from a mixture of two normal distributions, error = 3 generates errors from a rescaled chi-square distribution. error = 4 will use bootstrap. Choosing this option, the error terms will be resampled from the vector specified in sampling_errors.
sampling_errors	Sampling vector to resample errors from. Used only if error = 4.
wild_bootstrap	Whether special bootstrap to correct for heteroskedasticity should be used. If wild_bootstrap = TRUE, errors are generated from sampling_errors multiplied by a random variable following Rademacher distribution. Argument is used only if error = 4.
...	Further arguments

**Value**

Dose-response dataframe with generated data including "effect" as well as "d1" and "d2" columns.

**Examples**

```

coefs <- c("h1" = 1, "h2" = 1.5, "b" = 0,
          "m1" = 1, "m2" = 2, "e1" = 0.5, "e2" = 0.1)

## Dose levels are set to be integers from 0 to 10
generateData(coefs, sigma = 1)

## Dose levels are taken from existing dataset with d1 and d2 columns
data <- subset(directAntivirals, experiment == 1)
generateData(data = data[, c("d1", "d2")], pars = coefs, sigma = 1)

```

---

get.abs\_tval

*Return absolute t-value, used in optimization call in [optim.bboxcox](#)*


---

**Description**

Return absolute t-value, used in optimization call in [optim.bboxcox](#)

**Usage**

```
get.abs_tval(value, fac, lambda, zero.add2 = 0)
```

**Arguments**

value	data
fac	factor
lambda	box-cox parameter
zero.add2	2nd box-cox parameter

---

get.summ.data	<i>Summarize data by factor</i>
---------------	---------------------------------

---

**Description**

Summarize data by factor

**Usage**

```
get.summ.data(value, fac)
```

**Arguments**

value	data to sumamrize
fac	factor to summarize by

---

GetStartGuess	<i>Estimate initial values for dose-response curve fit</i>
---------------	--

---

**Description**

Estimate initial values for dose-response curve fit

**Usage**

```
GetStartGuess(df, transforms = NULL)
```

**Arguments**

df	Dose-response dataframe containing "dose" and "effect" columns
transforms	Transformation functions. If non-null, transforms is a list containing 5 elements, namely biological and power transformations along with their inverse functions and compositeArgs which is a list with argument values shared across the 4 functions. See vignette for more information.



---

getTransformations      *Return a list with transformation functions*

---

### Description

This function takes in response data from a dose-response model and attempts to find an optimal Box-Cox power transform based on [optim.boxcox](#) function. It then returns a list of transformation functions which contains this power transform and its inverse which can be subsequently used in [fitMarginals](#) and [fitSurface](#).

### Usage

```
getTransformations(data, shift = FALSE, args = list(N0 = 1, time.hours  
= 1))
```

### Arguments

data	Dose-response dataframe.
shift	If TRUE or is a numeric value, then a two-parameter Box-Cox transformation is assumed. This parameter will be passed on to <a href="#">optim.boxcox</a> function.
args	List with elements that are added to the list of transformation function and which can be used by these functions. In particular, this list should be of type <code>args = list("N0" = 1, "time.hours" = 1)</code> where <code>N0</code> and <code>time.hours</code> are arguments used for the biological transform.

### Details

Additionally, returned list contains biological transform and its inverse based on a simple exponential growth model, especially useful when response data is provided in cell counts. User can additionally provide arguments for these biological transforms where `N0` stands for initial cell count and `time.hours` indicates number in hours after which response data was measured.

[getTransformations](#) relies on [optim.boxcox](#) to obtain the optimal Box-Cox transformation parameters. However, [optim.boxcox](#) optimizes for the power parameter only within the interval (0.1, 0.9). Hence, if obtained power parameter is close to 0.1, then a logarithmic transformation is applied instead.

### Value

This function returns a list with transformation functions. These include power transformation ("PowerT") and its inverse ("InvPowerT") as well as biological transformation ("BioLT") and its inverse ("InvBioLT").

Power transformation is a 1-parameter Box-Cox transformation. If `shift = TRUE`, then power transformation is a 2-parameter Box-Cox transformation. Optimal values for power and shift operators are selected by means of [optim.boxcox](#) function.

Biological transformation  $y = N_0 * \exp(x * t)$  where  $N_0$  is the initial cell count and  $t$  is the incubation time. If response/effect variable ( $y$ ) is given in terms of cell counts, biological transformation ensures that modelisation is done for the growth rate instead ( $x$ ).

Returned list also contains "compositeArgs" elements shared by all the transformation functions. These arguments include initial cell count ("N0") and incubation time ("time.hours").

### Examples

```
data <- subset(directAntivirals, experiment == 1)
## Data must contain d1, d2 and effect columns
getTransformations(data)
```

---

hsa	<i>Highest Single Agent model</i>
-----	-----------------------------------

---

### Description

This function returns response levels for when these are based on Highest Single Agent (HSA) model.

### Usage

```
hsa(doseInput, parmInput, ...)
```

### Arguments

doseInput	Dose-response dataframe containing "d1" and "d2" columns
parmInput	Numeric vector or list with appropriately named parameter inputs. Typically, it will be coefficients from a MarginalFit object.
...	Further arguments that are currently unused

---

initialMarginal	<i>Estimate initial values for fitting marginal dose-response curves</i>
-----------------	--

---

### Description

This is a wrapper function which, when a dose-response dataframe is provided, returns start value estimates for both compounds that could be supplied to [fitMarginals](#) function. This function is also used by [fitMarginals](#) if no initials values were supplied.

### Usage

```
initialMarginal(data, transforms = NULL, ...)
```

**Arguments**

data	Dose-response dataframe. Marginal data will be extracted from it automatically.
transforms	Transformation functions. If non-null, transforms is a list containing 5 elements, namely biological and power transformations along with their inverse functions and compositeArgs which is a list with argument values shared across the 4 functions. See vignette for more information.
...	Further parameters that are currently not used

**Details**

Note that this function returns e1 and 2 which are log-transformed inflection points for respective compounds.

**Value**

Named vector with parameter estimates. Parameter names are consistent with parameter names in `fitMarginals`. h1 and h2 are Hill's slope coefficients for each of the compounds, m1 and m2 are their maximal response levels whereas b is the shared baseline. Lastly, e1 and e2 are log-transformed EC50 values.

**Note**

Returns starting value for  $e = \log(\text{EC50})$ .

**Examples**

```
data <- subset(directAntivirals, experiment == 1)
## Data must contain d1, d2 and effect columns
transforms <- getTransformations(data)
initialMarginal(data, transforms)
```

---

isobologram

*Isobologram of the response surface predicted by the null model*


---

**Description**

If transformation functions are used, then the isobologram response levels will be plotted on the transformed scale.

**Usage**

```
isobologram(x, grid.len = 100, logScale = TRUE, ...)
```

**Arguments**

x	Output of <a href="#">fitSurface</a>
grid.len	Number of concentrations to plot for each compound in the contour plot. An evenly spaced grid of doses will be generated for each compound given its respective observed minimum and maximum doses. Note that $\text{grid.len}^2$ computations will be needed later so this number should stay reasonably low.
logScale	If <code>logScale = TRUE</code> , then grid of doses is evenly spaced in the logarithmic scale.
...	Further parameters that are not used at this moment.

---

L4	<i>4-parameter logistic dose-response function</i>
----	--

---

**Description**

4-parameter logistic dose-response function

**Usage**

```
L4(dose, b, L, U, logEC50)
```

**Arguments**

dose	Dose level
b	Hill's coefficient (slope of the curve)
L	Baseline effect (at zero dose)
U	Asymptote effect (at infinite dose)
logEC50	Point of inflection (in logarithmic terms)

---

marginalNLS	<i>Fit two 4-parameter log-logistic functions with non-linear least squares</i>
-------------	---

---

**Description**

This function does not automatically extract marginal data and requires model input obtained from [constructFormula](#).

**Usage**

```
marginalNLS(data, transforms = NULL, start, model, nlsfn = nls, ...)
```

**Arguments**

data	Dose-response dataframe. Marginal data will be extracted from it automatically.
transforms	Transformation functions. If non-null, transforms is a list containing 5 elements, namely biological and power transformations along with their inverse functions and compositeArgs which is a list with argument values shared across the 4 functions. See vignette for more information.
start	Starting parameter values. If not specified, they will be obtained from <a href="#">initialMarginal</a> .
model	List with model parameters. Typically, this is an output from <a href="#">constructFormula</a> .
nlsfn	Non-linear least-squares optimizer function
...	Further arguments that are passed to the optimizer function, such as lower or upper (for the "nlsml" method), or control.

---

marginalOptim	<i>Fit two 4-parameter log-logistic functions with common baseline</i>
---------------	--

---

**Description**

This function is an alternative to non-linear least squares and provides optimization framework with [optim](#) function. It is however noticeably slower than NLS methods and can be especially time consuming in large datasets, in particular if bootstrap statistics are calculated.

**Usage**

```
marginalOptim(data, transforms = NULL, start, model, ...)
```

**Arguments**

data	Dose-response dataframe. Marginal data will be extracted from it automatically.
transforms	Transformation functions. If non-null, transforms is a list containing 5 elements, namely biological and power transformations along with their inverse functions and compositeArgs which is a list with argument values shared across the 4 functions. See vignette for more information.
start	Starting parameter values. If not specified, they will be obtained from <a href="#">initialMarginal</a> .
model	List with model parameters. Typically, this is an output from <a href="#">constructFormula</a> .
...	Further parameters passed to <a href="#">optim</a> function

**Value**

Variance-covariance matrix which is returned by [optim](#) is based on the fact that minimization of sum-of-squared residuals leads essentially to a maximum likelihood estimator and so variance-covariance matrix can be estimated using inverse Hessian evaluated at the optimal parameters. In some cases, so obtained variance-covariance matrix might not be positive-definite which probably means that estimates are unstable because of either a poor choice of initial values or poor properties of the data itself.

maxR

*Compute maxR statistic for each off-axis dose combination***Description**

`maxR` computes maxR statistics for each off-axis dose combination given the data provided. It provides a summary with results indicating whether a given point is estimated to be synergistic or antagonistic. These can be based either on normal approximation or a fully bootstrapped distribution of the statistics.

**Usage**

```
maxR(data, fitResult, transforms = fitResult$transforms,
      null_model = c("loewe", "hsa", "bliss"), Ymean, CP, reps,
      nested_bootstrap = FALSE, B.B = NULL, B.CP = NULL, cutoff = 0.95,
      cl = NULL, method = c("equal", "model", "unequal"), ...)
```

**Arguments**

<code>data</code>	Dose-response dataframe.
<code>fitResult</code>	Monotherapy (on-axis) model fit, e.g. produced by <code>fitMarginals</code> . It has to be a "MarginalFit" object or a list containing <code>df</code> , <code>sigma</code> , <code>coef</code> , <code>shared_asymptote</code> and <code>method</code> elements for, respectively, marginal model degrees of freedom, residual standard deviation, named vector of coefficient estimates, logical value of whether shared asymptote is imposed and method for estimating marginal models during bootstrapping (see <code>fitMarginals</code> ). If biological and power transformations were used in marginal model estimation, <code>fitResult</code> should contain <code>transforms</code> elements with these transformations. Alternatively, these can also be specified via <code>transforms</code> argument.
<code>transforms</code>	Transformation functions. If non-null, <code>transforms</code> is a list containing 5 elements, namely biological and power transformations along with their inverse functions and <code>compositeArgs</code> which is a list with argument values shared across the 4 functions. See vignette for more information.
<code>null_model</code>	Specified null model for the expected response surface. Currently, allowed options are "loewe" for generalized Loewe model, and "hsa" for Highest Single Agent model.
<code>Ymean</code>	Aggregate summary of off-axis predicted responses. In particular, it should contain "effect - predicted" column. If <code>Ymean</code> is missing, it will be calculated automatically from output of <code>predictOffAxis</code> function.
<code>CP</code>	Prediction covariance matrix. If not specified, it will be estimated by bootstrap using B.CP iterations.
<code>reps</code>	Numeric vector containing number of replicates for each off-axis dose combination. If missing, it will be calculated automatically from output of <code>predictOffAxis</code> function.

nested_bootstrap	When statistics are calculated, if <code>nested_bootstrap = TRUE</code> , CP matrix is recalculated at each bootstrap iteration of B.B using B.CP iterations. Using such nested bootstrap may however significantly increase computational time. If <code>nested_bootstrap = FALSE</code> , CP bootstrapped data reuses CP matrix calculated from the original data.
B.B	Number of iterations to use in bootstrapping null distribution for either meanR or maxR statistics.
B.CP	Number of bootstrap iterations to use for CP matrix estimation
cutoff	Cut-off to use in maxR procedure for declaring non-additivity (default is 0.95).
c1	If parallel computations are desired, <code>c1</code> should be a cluster object created by <a href="#">makeCluster</a> . If parallel computing is active, progress reporting messages are not necessarily ordered as it should be expected.
method	What assumption should be used for the variance of on- and off-axis points. This argument can take one of the values from <code>c("equal", "model", "unequal")</code> . With the value "equal" as the default. "equal" assumes that both on- and off-axis points have the same variance, "unequal" estimates a different parameter for on- and off-axis points and "model" predicts variance based on the average effect of an off-axis point. If no transformations are used the "model" method is recommended. If transformations are used, only the "equal" method can be chosen.
...	Further arguments that will be later passed to <a href="#">generateData</a> function during bootstrapping

## Value

This function returns a maxR object with estimates for the maxR statistical test. maxR object is essentially a list with appropriately named elements.

In particular, maxR object contains "Ymean" element which is a summary table of maxR test results for each dose combination. This table contains mean deviation from the predicted surface, normalized deviation ("absR") as well as a statistical call whether this deviation is significant. Distributional information on which these calls are made can be retrieved from the attributes of the "Ymean" dataframe.

Also, maxR object contains "Call" element which indicates the general direction of the deviation of the observed surface from the null. This call is based on the strongest local deviation in the "Ymean" table. 4 values are available here: "Syn", "Ant", "None", "Undefined". If one compound acts as an agonist while another one is an antagonist, then a deviation from the null is classified as "Undefined". If both compounds act in the same direction, then a stronger than individual effect is classified as synergy while a weaker effect would be classified as antagonism.

## Examples

```
data <- subset(directAntivirals, experiment == 2)
## Data must contain d1, d2 and effect columns
fitResult <- fitMarginals(data)
CP <- CPbootstrap(data, fitResult, null_model = "loewe", B.CP = 5)
maxR(data, fitResult, null_model = "loewe", CP = CP)
```

---

 meanR

*Compute meanR statistic for the estimated model*


---

## Description

`meanR` computes the meanR statistic for the provided model and returns the computed F-statistic and the estimated p-value. p-value can be calculated either by assuming an exact distribution or using bootstrapping procedure. In the latter case, null distribution of bootstrapped F-statistics is also returned.

## Usage

```
meanR(data, fitResult, transforms = fitResult$transforms,
      null_model = c("loewe", "hsa", "bliss"), R, CP, reps,
      nested_bootstrap = FALSE, B.B = NULL, B.CP = NULL, c1 = NULL,
      method = c("equal", "model", "unequal"), ...)
```

## Arguments

<code>data</code>	Dose-response dataframe.
<code>fitResult</code>	Monotherapy (on-axis) model fit, e.g. produced by <a href="#">fitMarginals</a> . It has to be a "MarginalFit" object or a list containing <code>df</code> , <code>sigma</code> , <code>coef</code> , <code>shared_asymptote</code> and <code>method</code> elements for, respectively, marginal model degrees of freedom, residual standard deviation, named vector of coefficient estimates, logical value of whether shared asymptote is imposed and method for estimating marginal models during bootstrapping (see <a href="#">fitMarginals</a> ). If biological and power transformations were used in marginal model estimation, <code>fitResult</code> should contain <code>transforms</code> elements with these transformations. Alternatively, these can also be specified via <code>transforms</code> argument.
<code>transforms</code>	Transformation functions. If non-null, <code>transforms</code> is a list containing 5 elements, namely biological and power transformations along with their inverse functions and <code>compositeArgs</code> which is a list with argument values shared across the 4 functions. See vignette for more information.
<code>null_model</code>	Specified null model for the expected response surface. Currently, allowed options are "loewe" for generalized Loewe model, and "hsa" for Highest Single Agent model.
<code>R</code>	Numeric vector containing mean deviation of predicted response surface from the observed one at each of the off-axis points. If missing, it will be calculated automatically from output of <a href="#">predictOffAxis</a> function.
<code>CP</code>	Matrix which is part of covariance matrix for the <code>R</code> argument
<code>reps</code>	Numeric vector containing number of replicates for each off-axis dose combination. If missing, it will be calculated automatically from output of <a href="#">predictOffAxis</a> function.



nested_bootstrap	When statistics are calculated, if <code>nested_bootstrap = TRUE</code> , CP matrix is recalculated at each bootstrap iteration of B.B using B.CP iterations. Using such nested bootstrap may however significantly increase computational time. If <code>nested_bootstrap = FALSE</code> , CP bootstrapped data reuses CP matrix calculated from the original data.
B.B	Number of iterations to use in bootstrapping null distribution for either meanR or maxR statistics.
B.CP	Number of bootstrap iterations to use for CP matrix estimation
c1	If parallel computations are desired, <code>c1</code> should be a cluster object created by <a href="#">makeCluster</a> . If parallel computing is active, progress reporting messages are not necessarily ordered as it should be expected.
method	What assumption should be used for the variance of on- and off-axis points. This argument can take one of the values from <code>c("equal", "model", "unequal")</code> . With the value "equal" as the default. "equal" assumes that both on- and off-axis points have the same variance, "unequal" estimates a different parameter for on- and off-axis points and "model" predicts variance based on the average effect of an off-axis point. If no transformations are used the "model" method is recommended. If transformations are used, only the "equal" method can be chosen.
...	Further arguments that will be later passed to <a href="#">generateData</a> function during bootstrapping

## Value

This function returns a meanR object with estimates for the meanR statistical test. meanR object is essentially a list with appropriately named elements.

meanR object list includes notably the calculated F-statistic, p-value and degrees of freedom ("n1" and "df0" respectively) used to find the critical value of the F-distribution under the null.

If [meanR](#) test is run with bootstrapping, then p-value estimate is based on bootstrapped null distribution of test statistic and an additional element "FDist" (of class "ecdf") is returned.

## Examples

```
data <- subset(directAntivirals, experiment == 2)
## Data must contain d1, d2 and effect columns
fitResult <- fitMarginals(data)
CP <- CPbootstrap(data, fitResult, null_model = "loewe", B.CP = 5)
meanR(data, fitResult, null_model = "loewe", CP = CP)
```

---

optim.boxcox

*Find optimal Box-Cox transformation parameters*

---

## Description

Find optimal Box-Cox transformation parameters

**Usage**

```
optim.boxcox(value, fac, shift = FALSE)
```

**Arguments**

value	Response variable in the data, e.g. "effect" column
fac	Factor indicating groups of replicates, e.g. interaction(d1, d2)
shift	Whether to use 2-parameter Box-Cox transformation. Input may be TRUE/FALSE or a numeric value indicating the shift parameter to use. If FALSE, shift parameter is set to zero.

**Value**

Numeric vector with power and shift parameter in that order.

**Examples**

```
data <- subset(directAntivirals, experiment == 1)
optim.boxcox(data$effect, interaction(data$d1, data$d2))
```

---

outsidePoints	<i>List non-additive points</i>
---------------	---------------------------------

---

**Description**

List all points with corresponding p-values declared non-additive by the maxR statistical test.

**Usage**

```
outsidePoints(maxR, B = 10000)
```

**Arguments**

maxR	maxR statistics table returned by Ymean component from the output of <code>maxR</code> function. This can also be "maxR" element in the output of <code>fitSurface</code> function.
B	Iterations to use for the distribution of the maxR statistic. This is only used if Ymean dataframe does not have a "distr" attribute attached as is normally done when using <code>fitSurface</code> or <code>maxR</code> function.

**Value**

Returns a dataframe listing only dose combinations that exhibit significant deviations from the expected response surface.

**Examples**

```

data <- subset(directAntivirals, experiment == 2)
## Data must contain d1, d2 and effect columns
fitResult <- fitMarginals(data)
CP <- CPBootstrap(data, fitResult, null_model = "loewe", B.CP = 5)
statM <- maxR(data, fitResult, null_model = "loewe", CP = CP)
outsidePoints(statM$Ymean)

```

---

plot.MarginalFit      *Plot monotherapy curve estimates*

---

**Description**

Plot monotherapy curve estimates

**Usage**

```

## S3 method for class 'MarginalFit'
plot(x, ncol = 2, logScale = TRUE, ...)

```

**Arguments**

x	Output of <code>fitMarginals</code> function or a "MarginalFit" object
ncol	Number of plots per row
logScale	Whether x-axis should be plotted on a logarithmic scale
...	Further arguments

**Value**

Returns a ggplot object. It can be consequently modified by using standard operations on ggplot objects (if ggplot2 package is loaded).

---

plot.maxR      *Plot of maxR object*

---

**Description**

Plot of maxR object

**Usage**

```

## S3 method for class 'maxR'
plot(x, main = "Contour plot for maxR",
     xlab = "Dose (Compound 1)", ylab = "Dose (Compound 2)",
     colorPalette = c("blue", "white", "red"), logScale = TRUE,
     zTransform = function(z) { z }, plevels = c(0.7, 0.8, 0.9, 0.95,
     0.99, 0.999), cutoff = max(plevels), maxshow = NULL, ...)

```

**Arguments**

x	Output of <code>maxR</code> . This can also be "maxR" element in the output of <code>fitSurface</code> .
main	Fixed non-moving title for the 3D plot
xlab	X axis label using font, size and color <code>par(c("font.lab", "cex.lab", "col.lab"))</code> .
ylab	Y axis label, same font attributes as xlab.
colorPalette	Vector of color names for surface
logScale	Draw doses on log-scale (setting zeroes to be finite constant)
zTransform	Optional transformation function for z-axis. By default, identity function is used.
plevels	Probability levels used to generate a color scale
cutoff	Probability cutoff to use for range of colors
maxshow	Forced value for range of colors
...	Further arguments that are passed to <code>format</code> function for formatting of axis labels

---

<code>plot.meanR</code>	<i>Plot bootstrapped cumulative distribution function of meanR null distribution</i>
-------------------------	--

---

**Description**

Plot bootstrapped cumulative distribution function of meanR null distribution

**Usage**

```
## S3 method for class 'meanR'
plot(x, ...)
```

**Arguments**

x	Output from <code>meanR</code>
...	Further arguments

---

plot.ResponseSurface *Method for plotting response surface objects*

---

### Description

Method for plotting response surface objects

### Usage

```
## S3 method for class 'ResponseSurface'
plot(x, color = c("z-score", "maxR",
  "occupancy"), ...)
```

### Arguments

x	Output of <a href="#">fitSurface</a>
color	Character indicating on what values surface coloring will be based. If color = "z-score", surface coloring will be based on median of standartized off-axis Z-scores. Median function can be replaced by other function using an optional colorfun argument which will be passed to plotResponseSurface. Color breaks are determined here by standard deviation of off-axis Z-scores. For color = "maxR", coloring will be based on values of maxR statistic and the quantile of its distribution (bootstrapped or not). If color = "occupancy", coloring will be based on calculated occupancy rate for the respective dose combination.
...	Further parameters passed to <a href="#">plotResponseSurface</a> . colorBy argument in this method is computed automatically and thus cannot be passed to <a href="#">plotResponseSurface</a> .

---

plotResponseSurface *Plot response surface*

---

### Description

Plot the 3-dimensional response surface predicted by one of the null models. This plot allows for a visual comparison between the null model prediction and observed points. This function is mainly used as the workhorse of [plot.ResponseSurface](#) method.

### Usage

```
plotResponseSurface(data, fitResult = NULL,
  transforms = fitResult$transforms, predSurface = NULL,
  null_model = c("loewe", "hsa", "bliss"), colorPalette = c("blue",
  "grey70", "red"), colorBy = "none", colorPoints = c("black",
  "sandybrown", "brown", "white"), breaks = c(-Inf, 0, Inf),
  radius = NULL, logScale = TRUE, colorfun = median,
```

```
zTransform = function(x) x, add = FALSE, main = "",
legend = TRUE, xat = "actual", yat = "actual", plotfun = NULL,
...)
```

### Arguments

data	Dose-response dataframe.
fitResult	Monotherapy (on-axis) model fit, e.g. produced by <a href="#">fitMarginals</a> . It has to be a "MarginalFit" object or a list containing df, sigma, coef, shared_asymptote and method elements for, respectively, marginal model degrees of freedom, residual standard deviation, named vector of coefficient estimates, logical value of whether shared asymptote is imposed and method for estimating marginal models during bootstrapping (see <a href="#">fitMarginals</a> ). If biological and power transformations were used in marginal model estimation, fitResult should contain transforms elements with these transformations. Alternatively, these can also be specified via transforms argument.
transforms	Transformation functions. If non-null, transforms is a list containing 5 elements, namely biological and power transformations along with their inverse functions and compositeArgs which is a list with argument values shared across the 4 functions. See vignette for more information.
predSurface	Vector of all predicted responses based on <code>expand.grid(uniqueDoses)</code> . If not supplied, it will be computed with <a href="#">predictOffAxis</a> function.
null_model	If predSurface is not supplied, it is computed using one of the available null models, i.e. "loewe" or "hsa". See also <a href="#">fitSurface</a> .
colorPalette	Vector of color names for surface
colorBy	This parameter determines values on which coloring is based for the 3-dimensional surface. If matrix or a data frame with d1 and d2 columns is supplied, dose combinations from colorBy will be matched automatically to the appropriate dose combinations in data. Unmatched dose combinations will be set to 0. This is especially useful for plotting results for off-axis estimates only, e.g. off-axis Z-scores or maxR test statistics. If colorBy = "colors", surface will be colored using colors in colorPalette argument.
colorPoints	Colors for off-axis and on-axis points. Character vector of length four with colors for 1) off-axis points; 2) on-axis points of the first drug (i.e. second drug is dosed at zero); 3) on-axis points of the second drug; 4) on-axis points where both drugs are dosed at zero.
breaks	Numeric vector with numerical breaks. To be used in conjunction with colorPalette argument.
radius	Radius of spheres. If missing, an educated guess based on number of digits in average effect will be made.
logScale	Draw doses on log-scale (setting zeroes to be finite constant)
colorfun	If replicates in colorBy variable are present, these will be aggregated using colorfun function. This can also be a custom function returning a scalar.
zTransform	Optional transformation function for z-axis. By default, identity function is used.

add	Add the predicted response surface to an existing plot. Will not draw any points, just the surface. Must be called after another call to <code>plotResponseSurface</code> .
main	Fixed non-moving title for the 3D plot
legend	Whether legend should be added
xat	x-axis ticks: "pretty", "actual" or a numeric vector
yat	y-axis ticks: "pretty", "actual" or a numeric vector
plotfun	If replicates for dose combinations in data are available, points can be aggregated using <code>plotfun</code> function. Typically, it will be <code>mean</code> , <code>median</code> , <code>min</code> or <code>max</code> but a custom-defined function returning a scalar from a vector is also possible.
...	Further arguments to format axis labels

### Details

Title for the plot and legend are drawn as bitmaps and do not rotate with the rest of the plot. Since they are bitmaps, they do not scale properly, hence resizing window will result in unappealing visuals. For them to look properly, it suffices to set the appropriate RGL window size and rerun the plotting command.

### Value

Plot is shown on a rgl device.

### Examples

```
## Not run:
data <- subset(directAntivirals, experiment == 1)
## Data must contain d1, d2 and effect columns
fitResult <- fitMarginals(data)
data_mean <- aggregate(effect ~ d1 + d2, data = data[, c("d1", "d2", "effect")],
                       FUN = mean)

## Construct the surface from marginal fit estimates based on HSA
## model and color it by mean effect level
plotResponseSurface(data, fitResult, null_model = "hsa",
                   colorBy = data_mean, breaks = 10^(c(0, 3, 4, 6)),
                   colorPalette = c("grey", "blue", "green"))

## Response surface based on Loewe additivity model and colored with
## rainbow colors. Legend will not be displayed in any case.
plotResponseSurface(data, fitResult, null_model = "loewe",
                   colorBy = "colors", colorPalette = rainbow(6))

## End(Not run)
```

---

predict.MarginalFit     *Predict values on the dose-response curve*

---

### Description

Predict values on the dose-response curve

### Usage

```
## S3 method for class 'MarginalFit'
predict(object, newdata, ...)
```

### Arguments

object	Output of <a href="#">fitMarginals</a> function
newdata	An optional data frame in which to look for d1 and d2 variables with which to predict. If omitted, the fitted values are used. Doses that are passed to this function must correspond to marginal data, i.e. at least one of the doses must be zero.
...	Further arguments

---

predictOffAxis     *Compute off-axis predictions*

---

### Description

Given a dataframe with dose-response data, this function uses coefficient estimates from the marginal (on-axis) monotherapy model to compute the expected values of response at off-axis dose combinations using a provided null model.

### Usage

```
predictOffAxis(data, fitResult, transforms = fitResult$transforms,
  null_model = c("loewe", "hsa", "bliss"), ...)
```

### Arguments

data	Dose-response dataframe.
fitResult	Monotherapy (on-axis) model fit, e.g. produced by <a href="#">fitMarginals</a> . It has to be a "MarginalFit" object or a list containing df, sigma, coef, shared_asymptote and method elements for, respectively, marginal model degrees of freedom, residual standard deviation, named vector of coefficient estimates, logical value of whether shared asymptote is imposed and method for estimating marginal models during bootstrapping (see <a href="#">fitMarginals</a> ). If biological and power transformations were used in marginal model estimation, fitResult should contain transforms elements with these transformations. Alternatively, these can also be specified via transforms argument.



transforms	Transformation functions. If non-null, transforms is a list containing 5 elements, namely biological and power transformations along with their inverse functions and compositeArgs which is a list with argument values shared across the 4 functions. See vignette for more information.
null_model	Specified null model for the expected response surface. Currently, allowed options are "loewe" for generalized Loewe model, and "hsa" for Highest Single Agent model.
...	Further arguments that are currently unused

**Value**

This functions returns a list with 3 elements.

"offaxisZTable" is a dataframe containing dose levels, observed effects and effects predicted according to the specified null model. This dataframe also contains replicates, if there are any.

"predSurface" are the predicted effects (without replicates) according to the specified null model. These effects are arranged in a matrix form so that each direction of the matrix rightward or downward corresponds to increasing dose of one of the compounds.

"occupancy" contains occupancy levels at each dose combination as (always) computed by generalized Loewe model.

**Examples**

```
data <- subset(directAntivirals, experiment == 1)
## Data must contain d1, d2 and effect columns
transforms <- getTransformations(data)
fitResult <- fitMarginals(data, transforms)
predictOffAxis(data, fitResult, null_model = "hsa")
```

---

```
print.summary.MarginalFit
```

*Print method for summary of MarginalFit object*

---

**Description**

Print method for summary of MarginalFit object

**Usage**

```
## S3 method for class 'summary.MarginalFit'
print(x, ...)
```

**Arguments**

x	Summary of MarginalFit object
...	Further arguments

---

`print.summary.maxR`     *Print summary of maxR object*

---

**Description**

Print summary of maxR object

**Usage**

```
## S3 method for class 'summary.maxR'  
print(x, ...)
```

**Arguments**

<code>x</code>	Summary of "maxR" object
<code>...</code>	Further arguments

---

`print.summary.meanR`     *Print summary of meanR object*

---

**Description**

Print summary of meanR object

**Usage**

```
## S3 method for class 'summary.meanR'  
print(x, ...)
```

**Arguments**

<code>x</code>	Summary of meanR object
<code>...</code>	Further arguments

---

```
print.summary.ResponseSurface
```

*Print method for the summary function of ResponseSurface object*

---

### Description

Print method for the summary function of ResponseSurface object

### Usage

```
## S3 method for class 'summary.ResponseSurface'  
print(x, ...)
```

### Arguments

x	Summary of ResponseSurface object
...	Further parameters

---

```
residuals.MarginalFit Residuals from marginal model estimation
```

---

### Description

Residuals from marginal model estimation

### Usage

```
## S3 method for class 'MarginalFit'  
residuals(object, ...)
```

### Arguments

object	Output of <code>fitMarginals</code> function
...	Further arguments

---

runBIGL	<i>Run the BIGL application for demonstrating response surfaces</i>
---------	---

---

**Description**

Run the BIGL application for demonstrating response surfaces

**Usage**

```
runBIGL(...)
```

**Arguments**

... Pass parameters to [runApp](#)

**Examples**

```
## Not run:
runBIGL()

## End(Not run)
```

---

simulateNull	<i>Simulate data from a given null model and monotherapy coefficients</i>
--------------	---

---

**Description**

Simulate data from a given null model and monotherapy coefficients

**Usage**

```
simulateNull(data, fitResult, transforms = fitResult$transforms,
  null_model = c("loewe", "hsa", "bliss"), ...)
```

**Arguments**

data	Dose-response dataframe.
fitResult	Monotherapy (on-axis) model fit, e.g. produced by <a href="#">fitMarginals</a> . It has to be a "MarginalFit" object or a list containing df, sigma, coef, shared_asymptote and method elements for, respectively, marginal model degrees of freedom, residual standard deviation, named vector of coefficient estimates, logical value of whether shared asymptote is imposed and method for estimating marginal models during bootstrapping (see <a href="#">fitMarginals</a> ). If biological and power transformations were used in marginal model estimation, fitResult should contain transforms elements with these transformations. Alternatively, these can also be specified via transforms argument.

transforms	Transformation functions. If non-null, transforms is a list containing 5 elements, namely biological and power transformations along with their inverse functions and compositeArgs which is a list with argument values shared across the 4 functions. See vignette for more information.
null_model	Specified null model for the expected response surface. Currently, allowed options are "loewe" for generalized Loewe model, and "hsa" for Highest Single Agent model.
...	Further parameters that will be passed to <a href="#">generateData</a>

**Value**

List with data element containing simulated data and fitResult element containing marginal fit on the simulated data.

**Examples**

```
data <- subset(directAntivirals, experiment == 1)
## Data must contain d1, d2 and effect columns
fitResult <- fitMarginals(data)
simulateNull(data, fitResult, null_model = "hsa")
```

---

summary.MarginalFit    *Summary of MarginalFit object*

---

**Description**

Summary of MarginalFit object

**Usage**

```
## S3 method for class 'MarginalFit'
summary(object, ...)
```

**Arguments**

object	Output of <a href="#">fitMarginals</a> function
...	Further arguments

---

summary.maxR	<i>Summary of maxR object</i>
--------------	-------------------------------

---

**Description**

Summary of maxR object

**Usage**

```
## S3 method for class 'maxR'  
summary(object, ...)
```

**Arguments**

object	Object of "maxR" class
...	Further arguments

---

summary.meanR	<i>Summary of meanR object</i>
---------------	--------------------------------

---

**Description**

Summary of meanR object

**Usage**

```
## S3 method for class 'meanR'  
summary(object, ...)
```

**Arguments**

object	Output from <a href="#">meanR</a>
...	Further arguments

---

summary.ResponseSurface  
*Summary of ResponseSurface object*

---

**Description**

Summary of ResponseSurface object

**Usage**

```
## S3 method for class 'ResponseSurface'  
summary(object, ...)
```

**Arguments**

object	Output of <a href="#">fitSurface</a>
...	Further parameters

---

vcov.MarginalFit      *Estimate of coefficient variance-covariance matrix*

---

**Description**

Estimate of coefficient variance-covariance matrix

**Usage**

```
## S3 method for class 'MarginalFit'  
vcov(object, ...)
```

**Arguments**

object	Output of <a href="#">fitMarginals</a> function
...	Further arguments

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