

# Package ‘MOSim’

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**Title** Multi-Omics Simulation (MOSim)

**Version** 1.10.0

**Description** MOSim package simulates multi-omic experiments that mimic regulatory mechanisms within the cell, allowing flexible experimental design including time course and multiple groups.

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**License** GPL-3

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**URL** <https://github.com/Neurergus/MOSim>

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**R topics documented:**

MOSim-package . . . . .	2
experimentalDesign . . . . .	2
is.declared . . . . .	3
mosim . . . . .	3
omicData . . . . .	5
omicResults . . . . .	6
omicSettings . . . . .	7
omicSim . . . . .	8
plotProfile . . . . .	9
sampleData . . . . .	9

<b>Index</b>	<b>11</b>
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MOSim-package	<i>MOSim</i>
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**Description**

Multiomics simulation package.

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experimentalDesign	<i>Retrieves the experimental design</i>
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**Description**

Retrieves the experimental design

**Usage**

```
experimentalDesign(simulation)
```

**Arguments**

simulation      A MOSimulation object

**Value**

A data frame containing the experimental design used to simulate the data.

**Examples**

```
omic_list <- c("RNA-seq")
rnaseq_simulation <- mosim(omics = omic_list)
# This will be a data frame with RNA-seq counts

design_matrix <- experimentalDesign(rnaseq_simulation)
```

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is.declared	<i>Check if a variable is declared.</i>
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**Description**

Check if a variable is declared.

**Usage**

```
is.declared(object, key = NULL)
```

**Arguments**

object	Variable name to check
key	Optional key to check inside object.

**Value**

TRUE or FALSE indicating if the variable is initialized & non-empty.

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mosim	<i>mosim</i>
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**Description**

Performs a multiomic simulation by chaining two actions: 1) Creating the "MOSimulation" class with the provided params. 2) Calling "simulate" method on the initialized object.

**Usage**

```
mosim(omics, omicsOptions, diffGenes, numberReps, numberGroups, times,
      depth, profileProbs, minMaxFC, TFtoGene)
```

**Arguments**

omics	Character vector containing the names of the omics to simulate, which can be "RNA-seq", "miRNA-seq", "DNase-seq", "ChIP-seq" or "Methyl-seq" (e.g. c("RNA-seq", "miRNA-seq")). It can also be a list with the omic names as names and their options as values, but we recommend to use the argument omicSim to provide the options to simulated each omic.
omicsOptions	List containing the options to simulate each omic. We recommend to apply the helper method <a href="#">omicSim</a> to create this list in a friendly way, and the function <a href="#">omicData</a> to provide custom data (see the related sections for more information). Each omic may have different configuration parameters, but the common ones are:

	<p><b>simuData/idToGene</b> Seed sample and association tables for regulatory omics. The helper function <code>omicData</code> should be used to provide this information (see the following section).</p> <p><b>regulatorEffect</b> For regulatory omics. List containing the percentage of effect types (repressor, activator or no effect) over the total number of regulators. See vignette for more information.</p> <p><b>totalFeatures</b> Number of features to simulate. By default, the total number of features in the seed dataset.</p> <p><b>depth</b> Sequencing depth in millions of reads. If not provided, it takes the global parameter passed to <code>mosim</code> function.</p> <p><b>replicateParams</b> List with parameters <math>a</math> and <math>b</math> for adjusting the variability in the generation of replicates using the negative binomial. See vignette for more information.</p>
<code>diffGenes</code>	Number of differentially expressed genes to simulate, given in percentage (0 - 1) or in absolute number (> 1). By default 0.15
<code>numberReps</code>	Number of replicates per experimental condition (and time point, if time series are to be generated). By default 3.
<code>numberGroups</code>	Number of experimental groups or conditions to simulate.
<code>times</code>	Vector of time points to consider in the experimental design.
<code>depth</code>	Sequencing depth in millions of reads.
<code>profileProbs</code>	Numeric vector with the probabilities to assign each of the patterns. Defaults to 0.2 for each.
<code>minMaxFC</code>	Numeric vector of length 2 with minimum and maximum fold-change for differentially expressed features, respectively.
<code>TFtoGene</code>	A logical value indicating if default transcription factors data should be used (TRUE) or not (FALSE), or a 3 column data frame containing custom associations. By default FALSE.

## Value

Instance of class "MOSimulation" containing the multiomic simulation data.

## Examples

```

moSimulation <- mosim(
  omics = c("RNA-seq"),
  numberReps = 3,
  times = c(0, 2, 6, 12, 24)
)

# Retrieve simulated count matrix for RNA-seq
dataRNAseq <- omicResults(moSimulation, "RNA-seq")

```

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omicData	<i>Set customized data for an omic.</i>
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### Description

Set customized data for an omic.

### Usage

```
omicData(omic, data = NULL, associationList = NULL)
```

### Arguments

omic	The name of the omic to provide data.
data	Data frame with the omic identifiers as row names and just one column named Counts containing numeric values used as initial sample for the simulation.
associationList	Only for regulatory omics, a data frame with 2 columns, the first called containing the regulator ID and the second called Gene with the gene identifier.

### Value

Initialized simulation object with the given data.

### Examples

```
# Take a subset of the included dataset for illustration
# purposes. We could also load it from a csv file or RData,
# as long as we transform it to have 1 column named "Counts"
# and the identifiers as row names.

data(sampleData)

custom_rnaseq <- head(sampleData$SimRNAseq$data, 100)

# In this case, 'custom_rnaseq' is a data frame with
# the structure:
head(custom_rnaseq)
##           Counts
## ENSMUSG0000000001 6572
## ENSMUSG0000000003     0
## ENSMUSG0000000028 4644
## ENSMUSG0000000031     8
## ENSMUSG0000000037     0
## ENSMUSG0000000049     0

# The helper 'omicData' returns an object with our custom data.
rnaseq_customdata <- omicData("RNA-seq", data = custom_rnaseq)
```

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omicResults	<i>Retrieves the simulated data.</i>
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### Description

Retrieves the simulated data.

### Usage

```
omicResults(simulation, omics = NULL, format = "data.frame")
```

### Arguments

simulation	A MOSimulation object.
omics	List of the omics to retrieve the simulated data.
format	Type of object to use for returning the results

### Value

A list containing an element for every omic specified, with the simulation data in the format indicated, or a numeric matrix with simulated data if the omic name is directly provided.

### Examples

```
omic_list <- c("RNA-seq")
rnaseq_simulation <- mosim(omics = omic_list)
#' # This will be a data frame with RNA-seq counts
rnaseq_simulated <- omicResults(rnaseq_simulation, "RNA-seq")

#           Group1.Time0.Rep1 Group1.Time0.Rep2 Group1.Time0.Rep3 ...
# ENSMUSG00000073155          4539             5374          5808 ...
# ENSMUSG00000026251              0              0              0 ...
# ENSMUSG00000040472          2742             2714          2912 ...
# ENSMUSG00000021598          5256             4640          5130 ...
# ENSMUSG00000032348           421             348           492 ...
# ENSMUSG00000097226            16              14            9 ...
# ENSMUSG00000027857              0              0              0 ...
# ENSMUSG00000032081              1              0              0 ...
# ENSMUSG00000097164           794             822           965 ...
# ENSMUSG00000097871              0              0              0 ...
```

---

omicSettings	<i>Retrieves the settings used in a simulation</i>
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### Description

Retrieves the settings used in a simulation

### Usage

```
omicSettings(simulation, omics = NULL, association = FALSE,  
             reverse = FALSE, only.linked = FALSE, prefix = FALSE,  
             include.lagged = TRUE)
```

### Arguments

simulation	A MOSimulation object.
omics	List of omics to retrieve the settings.
association	A boolean indicating if the association must also be returned for the regulators.
reverse	A boolean, swap the column order in the association list in case we want to use the output directly and the program requires a different ordering.
only.linked	Return only the interactions that have an effect.
prefix	Logical indicating if the name of the omic should prefix the name of the regulator.
include.lagged	Logical indicating if interactions with transitory profile and different minimum/maximum time point between gene and regulator should be included or not.

### Value

A list containing a data frame with the settings used to simulate each of the indicated omics. If association is TRUE, it will be a list with 3 keys: 'associations', 'settings' and 'regulators', with the first two keys being a list containing the information for the selected omics and the last one a global data frame giving the merged information.

### Examples

```
omic_list <- c("RNA-seq", "miRNA-seq")  
multi_simulation <- mosim(omics = omic_list)  
  
# This will be a data frame with RNA-seq settings (DE flag, profiles)  
rnaseq_settings <- omicSettings(multi_simulation, "RNA-seq")  
  
# This will be a list containing all the simulated omics (RNA-seq  
# and DNase-seq in this case)  
all_settings <- omicSettings(multi_simulation)
```





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plotProfile	<i>Generate a plot of a feature's profile for one or two omics.</i>
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**Description**

Generate a plot of a feature's profile for one or two omics.

**Usage**

```
plotProfile(simulation, omics, featureIDS, drawReps = FALSE,
            groups = NULL)
```

**Arguments**

simulation	A MOSimulation object
omics	Character vector of the omics to simulate.
featureIDS	List containing the feature to show per omic. Must have the omics as the list names and the features as values.
drawReps	Logical to enable/disable the representation of the replicates inside the plot.
groups	Character vector indicating the groups to plot in the form "GroupX" (i.e. Group1)

**Value**

A ggplot2 object.

**Examples**

```
omic_list <- c("RNA-seq", "miRNA-seq")
rnaseq_simulation <- mosim(omics = omic_list)

plotProfile(rnaseq_simulation,
            omics = c("RNA-seq", "miRNA-seq"),
            featureIDS = list("RNA-seq"="ENSMUSG00000007682", "miRNA-seq"="mmu-miR-320-3p")
            )
```

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sampleData	<i>Default data</i>
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**Description**

Dataset with base counts and id-gene tables.

**Usage**

```
sampleData
```

**Format**

An object of class `list` of length 6.

**Details**

List with 6 elements:

**SimRNAseq data** Dataframe with base counts with gene id as rownames.

**geneLength** Length of every gene.

**SimChIPseq data** Dataframe with base counts with regions as rownames.

**idToGene** Dataframe with region as "ID" column and gene name on "Gene" column.

**SimDNaseseq data** Dataframe with base counts with regions as rownames.

**idToGene** Dataframe with region as "ID" column and gene name on "Gene" column.

**SimMiRNAseq data** Dataframe with base counts with miRNA id as rownames.

**idToGene** Dataframe with miRNA as "ID" column and gene name on "Gene" column.

**SimMethylseq idToGene** Dataframe with region as "ID" column and gene name on "Gene" column.

**CpGisland** Dataframe of CpG to be used as initialization data, located on "Region" column

# Index

## \* datasets

- sampleData, 9
- experimentalDesign, 2
- is.declared, 3
- mosim, 3, 4
- MOSim-package, 2
- omicData, 3, 4, 5
- omicResults, 6
- omicSettings, 7
- omicSim, 3, 8
- plotProfile, 9
- sampleData, 9