

# Package ‘FUCHIKOMA’

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**Type** Package

**Title** Detection of Differentially Expressed Genes in one of multiple clusters using BAHSIC and Diffusion Map

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**Depends** R (>= 2.5.0), destiny, doParallel, foreach

**biocViews** Bioinformatics, DifferentialExpression, Clustering, MultipleComparisons, RNAseq, HighThroughputSequencing, Sequencing, Software

**Description** FUCHIKOMA detects differentially expressed genes (DEGs) in one of multiple clusters. To detect DEGs, FUCHIKOMA has two calculation mode; “supervised-mode” and “unsupervised-mode”. In supervised-mode, FUCHIKOMA detects DEGs by using a label vector, in which the cluster of each sample or cell is written. In unsupervised-mode, FUCHIKOMA detects DEGs without the label vector. In this mode, user run the diffusion map, and specify which diffusion components contribute to the difference of such cluster.

**License** Artistic-2.0

**NeedsCompilation** no

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FUCHIKOMA-package	<i>Detection of Differentially Expressed Genes in one of multiple clusters using BAHSIC and Diffusion Map</i>
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## Description

FUCHIKOMA detects differentially expressed genes (DEGs) in one of multiple clusters. To detect DEGs, FUCHIKOMA has two calculation mode; "supervised-mode" and "unsupervised-mode". In supervised-mode, FUCHIKOMA detects DEGs by using a label vector, in which the cluster of each sample or cell is written. In unsupervised-mode, FUCHIKOMA detects DEGs without the label vector. In this mode, user run the diffusion map, and specify which diffusion components contribute to the difference of such cluster.

## Details

The DESCRIPTION file: This package was not yet installed at build time.

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The main function is [FUCHIKOMA](#), which returns an object containing the calculation results.

## Author(s)

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## References

Koki Tsuyuzaki et al. (2015) Fuchikoma: Detection of Differentially Expressed Genes in one of multiple clusters using BAHSIC and Diffusion Map. R package version 1.0.0

Laleh Haghverdi et al. (2015) Diffusion maps for high-dimensional single-cell analysis of differentiation data. *Bioinformatics*, 31(18), 2989-2998

Le Song et al. (2007) Gene selection via the BAHSIC family of algorithms, *Bioinformatics*, 23(13), i490-i498

Y-h Taguchi et al. (2015) Principal component analysis-based unsupervised feature extraction applied to in silico drug discovery for posttraumatic stress disorder-mediated heart disease, *BMC Bioinformatics*, 16(139)

Diego Adhemar Jaitin, Ephraim Kenigsberg, Hadas Keren-Shaul, Naama Elefant, Franziska Paul, Irina Zaretsky, Alexander Mildner, Nadav Cohen, Steffen Jung, Amos Tanay, Ido Amit (2014) Massively Parallel Single-Cell RNA-Seq for Marker-Free Decomposition of Tissues into Cell Types. *Science*, **343** (6172): 776-779

Arthur Gretton et al. (2007) A Kernel Statistical Test of Independence, NIPS 21

## See Also

[DiffusionMap](#)

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**CatKernel***Categorical Kernel function*

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**Description**

CatKernel calculate gram matrix by using label information (e.g., cell label or sample label such as `c(1,1,1,2,2,2,3,3)`).

**Usage**

```
CatKernel(label, type = c("two", "one_vs_rest", "each", "simple"))
```

**Arguments**

<code>label</code>	Label vector descrbing which cells are sample cell type and which cells are different (e.g., <code>c(1,1,1,2,2,2,3,3)</code> ).
<code>type</code>	Categorical kernel functions. simple could be specified even the number of class is two or more. When the number of class is two, only two parameter is accessible. When the number of class is three or more, <code>one_vs_rest</code> and <code>each</code> is accessible.

**Value**

A data frame is returned.

**Author(s)**

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**References**

Le Song et al. (2007) Gene selection via the BAHSIC family of algorithms, *Bioinformatics*, 23(13), i490-i498

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**Examples**

```
data(MARS)
L <- CatKernel(label.MARS, type="one_vs_rest")
dim(L)
```

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custom.DiffusionMap     *Customized Diffusion Map of destiny for extracting Gram Matrix*

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## Description

Inner function of FUCHIKOMA, which returns the gram matrix of diffusion map.

## Usage

```
custom.DiffusionMap(data, sigma = NULL, k = find.dm.k(nrow(data) - 1L), n.eigs = min(20L, nrow(data)))
```

## Arguments

data	Expression data to be analyzed. Provide vars to select specific columns other than the default: all double value columns
sigma	Diffusion scale parameter of the Gaussian kernel. Either a number or a <a href="#">Sigmas</a> object. (Optional. default: will be calculated using <a href="#">find.sigmas</a> ) A larger sigma might be necessary if the eigenvalues can not be found because of a singularity in the matrix
k	Number of nearest neighbors to consider (default: a guess between 100 and $n - 1$ ) NULL or NA are also interpreted as $n - 1$ L.
n.eigs	Number of eigenvectors/values to return (default: 20)
density.norm	logical. If TRUE, use density normalisation
...	All parameter after this are optional and have to be specified by name
distance	Distance measurement method. Euclidean distance (default) or cosine distance ( $1 - \text{corr}(c_1, c_2)$ ).
censor.val	Value regarded as uncertain. Either a single value or one for every dimension (Optional, default: CENSOR.VAL)
censor.range	Uncertainty range for censoring (Optional, default: none). A length-2-vector of certainty range start and end. TODO: also allow $2 \times G$ matrix
missing.range	Whole data range for missing value model. Has to be specified if NAs are in the data
vars	Variables (columns) of the data to use. Specifying NULL will select all columns (default: All floating point value columns)
verbose	Show a progressbar and other progress information (default: do it if censoring is enabled)
.debug.env	If supplied, the rotated transition probaility matrix M and the scaling rotation D.rot is stored in it prior to eigen decomposition.

## Value

A DiffusionMap object:

## Author(s)

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## References

Laleh Haghverdi et al. (2015) Diffusion maps for high-dimensional single-cell analysis of differentiation data. *Bioinformatics*, 31(18), 2989-2998

## See Also

[DiffusionMap](#)

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FUCHIKOMA	<i>Detection of Differentially Expressed Genes in one of multiple clusters using BAHSIC and Diffusion Map</i>
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## Description

This package was not yet installed at build time.

## Usage

```
FUCHIKOMA(data, mode = c("Supervised", "Unsupervised"), Comp = FALSE, label = FALSE, cat.type = FALSE)
```

## Arguments

data	A data matrix, in which the row means genes and column means cells or sample.
mode	When Supervised is specified, FUCHIKOMA uses label paramter. When Unsupervised is specified, FUCHIKOMA uses Comp parameter for specifying which diffusion components should be used.
Comp	When mode is specified as Unsupervised, Comp must be specified such as c(1,2).
label	When mode is specified as Supervised, label must be specified such as c(1,1,1,2,2,2,3,3)
cat.type	Type of categorical kernel of CatKernel
n.eigs	Number of eigenvectors/values to return (default: 20)
algorithm	brute means single gene rejection strategies and song means fixed percent of genes rejection strategies in each iteration step of FUCHIKOMA.
per.rej	When algorithm is specified as song, per.rej must be specified such as 20 (default:10)
threshold	In each iteration step, if the difference of HSIC in the step and max value of previous HSICs is lower than threshold, iteration will be halted (default: 0.01).

## Value

DEGs.HSICs	Differentially expressed genes and HSIC values
DEGs.Pvals	Pvalues of HSICs
All.HSICs	HSICs value of all genes
Rej.order	The order of rejecting of genes in each iteration step

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**See Also**

[DiffusionMap](#)

**Examples**

```
data(MARS)
res <- FUCHIKOMA(data=MARS, mode="Supervised", label=label.MARS, type="one_vs_rest", n.eigs=10, algorithm="s
```

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HSIC	<i>Hilbert-Schmidt Independence Criteria (HSIC)</i>
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**Description**

Inner function of FUCHIKOMA. HSIC calculated against two gram-matrix means the independence of two kernel spaces. The higher HSIC value is, the more two space are dependent. When p.value set TRUE, p-value of HSIC is also calculated by moment matching to a gamma distribution.

**Usage**

```
HSIC(K, L, p.value = FALSE)
```

**Arguments**

K	First gram matrix
L	Second gram matrix
p.value	When p.value is specified as TRUE, p-value of HSIC is calculated (default:FALSE)

**Value**

HSIC	The value of HSIC
Pval	P-value of HSIC. The value is accessible only when p.value is specified as TRUE

**Author(s)**

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Le Song et al. (2007) Gene selection via the BAHASIC family of algorithms, *Bioinformatics*, 23(13), i490-i498

Y-h Taguchi et al. (2015) Principal component analysis-based unsupervised feature extraction applied to in silico drug discovery for posttraumatic stress disorder-mediated heart disease, *BMC Bioinformatics*, 16(139)

Arthur Gretton et al. (2007) A Kernel Statistical Test of Independence, *NIPS* 21

**Examples**

```
K <- matrix(runif(100), nrow=10)
L <- matrix(runif(100), nrow=10)
HSIC(K, L, p.value)
```

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MARS

*Count data of MARS-Seq containing four different cell types.*

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**Description**

A data frame with 10801 rows (genes) with following 228 columns (cells).

The data is downloaded from GEO (GSE54006) and genes having no expression are filtered.

The data has four different cell types; B cells (B1-B48), Dendritic cells (DC1-DC89), Monocytes (Mono1-Mono46), and Natural Killer cells (NK1-NK45).

**Usage**

```
data(MARS)
```

**Author(s)**

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**Source**

<https://www.sciencemag.org/content/343/6172/776?related-urls=yes&legid=sci;343/6172/776>

<http://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE54006>

**References**

Diego Adhemar Jaitin, Ephraim Kenigsberg, Hadas Keren-Shaul, Naama Elefant, Franziska Paul, Irina Zaretsky, Alexander Mildner, Nadav Cohen, Steffen Jung, Amos Tanay, Ido Amit (2014) Massively Parallel Single-Cell RNA-Seq for Marker-Free Decomposition of Tissues into Cell Types. *Science*, **343** (6172): 776-779

**Examples**

```
data(MARS)
dim(MARS)
pairs(result.pca.MARS$rotation[,1:10], col=label.MARS, main="MARS-Seq (PCA)")
pairs(result.destiny.MARS@eigenvectors[,1:10], col=label.MARS, main="MARS-Seq (Diffusion Map)")
```



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