

# Package ‘oposSOM’

July 2, 2014

**Type** Package

**Title** Package for analysis and visualization of metagenes

**Version** 0.2-3

**Date** 2013-08-13

**Author** Henry Wirth

**Maintainer** Henry Wirth <wirth@izbi.uni-leipzig.de>

**Description** This package translates microarray expression data into metagenes and provides various visualizations and second level analyses.

**License** GPL (>= 2)

**LazyLoad** yes

**URL** <http://som.izbi.uni-leipzig.de>

**Depends** R (>= 2.11.0)

**Imports** som, fastICA, pixmap, igraph, biomaRt

**NeedsCompilation** no

**Repository** CRAN

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oposSOM-package

*Package for analysis and visualization of metagenes*

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## Description

This package translates microarray expression data into metagenes and provides various visualizations and second level analyses. For a given data set, a standardized analysis pipeline is processed and the output is stored in an external folder. Additionally, a HTML summary file is created collating and linking all analysis results.

## Details

Package: oposSOM  
Type: Package  
Version: 0.2-3  
Date: 2013-08-13  
License: GPL (>= 2)  
LazyLoad: yes

## Author(s)

Henry Wirth

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## References

Wirth, Loeffler, v.Bergen, Binder: Expression cartography of human tissues using self organizing maps. (BMC Bioinformatics 2011)

Wirth, v.Bergen, Binder: MALDI-typing of infectious algae of the genus Prototheca using SOM portraits. (Journal of microbiological methods 2012)

## Examples

```
library( oposSOM )

#### Toy example ####

## create data
toy.data = matrix( runif(10000), 1000, 10 )

## run pipeline
run.pipeline( toy.data, dataset.name = "Example", dim.som1 = 20, dim.som2 = 20 )
```

```
#### Real data example - takes some time to process ####

## load data
#data( tissues )

## run pipeline
#run.pipeline( tissue.data, dataset.name="Tissue Example" )
```

---

run.pipeline

*Run the oposSOM pipeline.*


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## Description

This function realizes the complete pipeline functionality: single gene expression values are clustered to metagenes using a self-organizing map. Based on these metagenes, visualizations (e.g. expression profiles) and downstreaming analysis approaches (e.g. hierarchical clustering, ICA or geneset overrepresentation analysis) are performed.

## Usage

```
run.pipeline( indata, dataset.name = "Unnamed", dim.som1 = 20, dim.som2 = 20 )
```

## Arguments

indata	numerical data matrix containing genes as rows and samples as columns.
dataset.name	name of the folder which will be created to contain the results.
dim.som1	size of the primary SOM / resolution of the expression profiles: number of tiles equals $\text{dim.som1}^2$ .
dim.som2	resolution of the second level SOM: number of tiles equals $\text{dim.som2}^2$ .

## Details

If indata is supplied with ENSEMBL-IDs as rownames, geneset overrepresentation analysis is carried out using predefined GO gene sets automatically downloaded from Ensembl database (<http://www.ensembl.org>). The results are contained in summary sheets of the integrated maps, e.g. the overexpression overview map.

## Value

The pipeline will create a folder in the working directory according to dataset.name containing all results. Graphical output is provided in respective files solely to ensure functionality in command line mode.

Additionally, a HTML summary file is created which contains brief descriptions and links to all results:

- Expression profiles for all samples in different color modes (standard, loglog and WAD).

- Supporting maps illustrating properties of the SOM as metagene population, variance and concordance.
- Second level SOM providing sample-centered view on relations and similarities between the samples.
- Metagene based downstream analysis: Hierarchical clustering, pairwise correlation matrix and spanning tree, and independent component analysis.
- Report sheets for the spot summary maps containing various information of identified clusters of metagenes.

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### Examples

```
library( oposSOM )

#### Toy example ####

## create data
toy.data = matrix( runif(10000), 1000, 10 )

## run pipeline
run.pipeline( toy.data, dataset.name = "Example", dim.som1 = 20, dim.som2 = 20 )

#### Real data example - takes some time to process ####

## load data
#data( tissues )

## run pipeline
#run.pipeline( tissue.data, dataset.name="Tissue Example" )
```

---

tissue.data

*Example data set.*

---

### Description

A data set comprising of 12 selected human tissues.

**Usage**

```
data(tissues)
```

**Format**

The data set is stored in RData (binary) format.

**Details**

The data set was downloaded from Gene Expression Omnibus repository (<http://www.ncbi.nlm.nih.gov/geo>, GEO accession no. GSE7307). About 38,000 genes in more than 650 samples were measured using the Affymetrix HGU133-Plus2 microarray. A subset of 12 selected tissues from different categories is used as example data set for the oposSOM-package. ENSEMBL-IDs are used as row-identifier to enable geneset overrepresentation analysis.

**Source**

<http://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE7307>

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