

# Package ‘vsclust’

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**Type** Package

**Title** Feature-based variance-sensitive quantitative clustering

**Version** 1.8.0

**Date** 2022-03-23

**Description** Feature-based variance-sensitive clustering of omics data. Optimizes cluster assignment by taking into account individual feature variance. Includes several modules for statistical testing, clustering and enrichment analysis.

**License** GPL-2

**Imports** matrixStats, limma, parallel, shiny, qvalue, grDevices, stats, MultiAssayExperiment, graphics

**Suggests** knitr, yaml, testthat (>= 3.0.0), rmarkdown, BiocStyle, clusterProfiler

**LinkingTo** Rcpp

**biocViews** Clustering, Annotation, PrincipalComponent, DifferentialExpression, Visualization, Proteomics, Metabolomics

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vsclust-package	<i>VSClust provides a powerful method to run variance-sensitive clustering</i>
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## Description

Clustering of high-dimensional quantitative data with data points that come with multiple measurements. In this clustering method, each feature is represented by a) its quantitative profile and b) its variance. Hence, the uncertainty about a measurement enter in the determination of the most common patterns. This methods is both insensitive to noisy measurements and avoids finding clusters in homogeneously distributed data.

## Details

The functions in this package comprise (i) methods to prepare the data for cluster analysis like statistical analysis ('SignAnal' and 'SignPairedAnal'), PCA ('PCAwithVar'), (ii) direct application of the clustering algorithm on a (standardized) data matrix ('vsclust\_algorithm'), (iii) for the further evaluation and visualization (such as 'calcBHI' and 'mfuzz.plot'), and (iv) wrappers for the over workflow including statistical preparation ('statWrapper'), estimation of the cluster number ('estimClustNum'), running the clustering ('runClustWrapper') and functional evaluation ('runFuncEnrich').

## Author(s)

Maintainer: Veit Schw"ammle" <veits@bmb.sdu.dk>

## References

- Schw"ammle V, Hagensen CE. A Tutorial for Variance-Sensitive Clustering and the Quantitative Analysis of Protein Complexes. *Methods Mol Biol.* 2021;2228:433-451. doi: 10.1007/978-1-0716-1024-4\_30. PMID: 33950508.
- Schw"ammle V, Jensen ON. VSCLust: feature-based variance-sensitive clustering of omics data. *Bioinformatics.* 2018 Sep 1;34(17):2965-2972. doi: 10.1093/bioinformatics/bty224. PMID: 29635359.
- Schw"ammle V, Jensen ON. A simple and fast method to determine the parameters for fuzzy c-means cluster analysis. *Bioinformatics.* 2010 Nov 15;26(22):2841-8. doi: 10.1093/bioinformatics/btq534. Epub 2010 Sep 29. PMID: 20880957.

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artificial\_clusters     *Synthetic/artificial data comprising 5 clusters*

---

## Description

10-dimensional data set with 500 simulating features measured over 5 replicates each, comprising a total of 50 samples. The first 250 features were modeled through normal distributions shifted in the 10-dimensional space to form 5 different clusters. The 2nd half of the features were modeled through a normal distribution around the origin and thus should be assigned to any cluster

## Usage

artificial\_clusters

## Format

A data frame consisting of 500 features distributed over 5 clusters and being replicated 5 times each

## Source

Protein Research Group, University of Southern Denmark, Odense

---

averageCond     *Calculate mean over replicates*

---

## Description

Simple method to calculate the means for each feature across its replicates

## Usage

averageCond(data, NumReps, NumCond)

## Arguments

data	Matrix of data frame with numerical values. Columns corresponds to samples
NumReps	Number of replicates per experimental condition
NumCond	Number of different experimental conditions

**Value**

Matrix of data frame with averaged values over replicates for each conditions

**Examples**

```
data <- matrix(rnorm(1000), nrow=100)
av_data <- averageCond(data, NumCond=2, NumReps=5)
```

---

calcBHI

*Calculate "biological homogeneity index"*

---

**Description**

This index is providing a number for the enriched GO terms and pathways to assess the biological content within a set of genes or proteins. The calculation is according to Datta, S. & Datta, S. Methods for evaluating clustering algorithms for gene expression data using a reference set of functional classes. *BMC bioinformatics* 7, 397 (2006).

**Usage**

```
calcBHI(Accs, gos)
```

**Arguments**

Accs            list containing gene or protein IDs, such as UniProt accession names  
gos             object from ClusterProfiler

**Value**

Biological Homogeneity Index

**References**

Datta, S. & Datta, S. Methods for evaluating clustering algorithms for gene expression data using a reference set of functional classes. *BMC bioinformatics* 7, 397 (2006).

Schwaemmle V, Jensen ON. VSClust: feature-based variance-sensitive clustering of omics data. *Bioinformatics*. 2018 Sep 1;34(17):2965-2972. doi: 10.1093/bioinformatics/bty224. PMID: 29635359.

Schwaemmle V, Hagensen CE. A Tutorial for Variance-Sensitive Clustering and the Quantitative Analysis of Protein Complexes. *Methods Mol Biol*. 2021;2228:433-451. doi: 10.1007/978-1-0716-1024-4\_30. PMID: 33950508.

Schwaemmle V, Jensen ON. A simple and fast method to determine the parameters for fuzzy c-means cluster analysis. *Bioinformatics*. 2010 Nov 15;26(22):2841-8. doi: 10.1093/bioinformatics/btq534. Epub 2010 Sep 29. PMID: 20880957.

**Examples**

```
# Run enrichment analysis
data(gcSample, package="clusterProfiler")
xx <- clusterProfiler::compareCluster(gcSample, fun="enrichKEGG",
                                     organism="hsa", pvalueCutoff=0.05)
# Generate random list from gcSample
rand_ids <- lapply(gcSample, function(x) sample(unlist(gcSample), 200))
calcBHI(rand_ids, xx)
```

---

ClustComp	<i>Function to run clustering with automatic fuzzifier settings (might become obsolete)</i>
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---

**Description**

Run original fuzzy c-means and vsclust for a number of clusters and the given data set including data pre-processing and automatic setting of the data-dependent parameters like the lower limit of the fuzzifier.

**Usage**

```
ClustComp(
  dat,
  NSs = 10,
  NClust = NClust,
  Sds = Sds,
  cl = parallel::makePSOCKcluster(1),
  verbose = FALSE
)
```

**Arguments**

dat	a numeric data matrix
NSs	number of clusterings runs with different random seeds
NClust	Number of clusters
Sds	Standard deviation of features (either vector of the same length as features numbers in matrix or single value)
cl	object of class 'cluster' or 'SOCKcluster' to specify environment for parallelization
verbose	Show more information during execution

**Value**

List containing the objects

- 'indices' containing minimum centroid distance and Xie-Beni index for both clustering methods
- 'Bestcl' optimal vsclust results (variance-sensitive fcm clustering)
- 'Bestcl2' optimal fuzzy c-means results
- 'm' vector of individual fuzzifier values per feature
- 'withinerror' final optimization score for vsclust
- 'withinerror2' final optimization score for fuzzy c-means clustering

## References

Schwaemmle V, Jensen ON. VSClust: feature-based variance-sensitive clustering of omics data. *Bioinformatics*. 2018 Sep 1;34(17):2965-2972. doi: 10.1093/bioinformatics/bty224. PMID: 29635359.

Schwaemmle V, Hagensen CE. A Tutorial for Variance-Sensitive Clustering and the Quantitative Analysis of Protein Complexes. *Methods Mol Biol*. 2021;2228:433-451. doi: 10.1007/978-1-0716-1024-4\_30. PMID: 33950508.

Schwaemmle V, Jensen ON. A simple and fast method to determine the parameters for fuzzy c-means cluster analysis. *Bioinformatics*. 2010 Nov 15;26(22):2841-8. doi: 10.1093/bioinformatics/btq534. Epub 2010 Sep 29. PMID: 20880957.

## Examples

```
#' # Generate some random data
data <- matrix(rnorm(seq_len(1000)), nrow=100)
# Run clustering
cl <- parallel::makePSOCKcluster(1, nnodes=1)
ClustCompOut <- ClustComp(data, cl=cl, NClust=6, Sds=1)
barplot(ClustCompOut$indices)
```

---

cvalidate.xiebeni

*Xie Beni Index of clustering object*

---

## Description

Calculate the Xie Beni index for validity of the cluster number in clustering results from running fuzzy c-means or vsclust original publication:

## Usage

```
cvalidate.xiebeni(clres, m)
```

## Arguments

clres	Output from clustering. Either fclust object or list containing the objects for 'membership' and cluster 'centers'
m	Fuzzifier value

## Value

Xie Beni index

## References

Xie X.L., Beni G. (1991). A validity measure for fuzzy clustering, *IEEE Transactions on Pattern Analysis and Machine Intelligence*, 13, 841-847.

## Examples

```
# Generate some random data
data <- matrix(rnorm(seq_len(1000)), nrow=100)
# Run clustering
clres <- vsclust_algorithm(data, centers=5, m=1.5)
# Calculate Xie-Beni index from results
cvalidate.xiebeni(clres, 1.5)
```

---

determine_fuzz	<i>Determine individual fuzzifier values</i>
----------------	--

---

## Description

This function calculated the values of the fuzzifier from a) the dimensions of the considered data set and b) from the individual feature standard deviations.

## Usage

```
determine_fuzz(dims, NClust, Sds = 1)
```

## Arguments

dims	vector of two integers containing the dimensions of the data matrix for the clustering
NClust	Number of cluster for running vsclust on (does not influence the calculation of 'mm')
Sds	individual standard deviations, set to 1 if not available

## Value

list of 'm': individual fuzzifiers, 'mm': standard fuzzifier for fcm clustering when not using vsclust algorithm

## References

Schwaemmle V, Jensen ON. VSclust: feature-based variance-sensitive clustering of omics data. *Bioinformatics*. 2018 Sep 1;34(17):2965-2972. doi: 10.1093/bioinformatics/bty224. PMID: 29635359.

Schwaemmle V, Hagensen CE. A Tutorial for Variance-Sensitive Clustering and the Quantitative Analysis of Protein Complexes. *Methods Mol Biol*. 2021;2228:433-451. doi: 10.1007/978-1-0716-1024-4\_30. PMID: 33950508.

Schwaemmle V, Jensen ON. A simple and fast method to determine the parameters for fuzzy c-means cluster analysis. *Bioinformatics*. 2010 Nov 15; 26(22):2841-8. doi: 10.1093/bioinformatics/btq534. Epub 2010 Sep 29. PMID: 20880957.

## Examples

```
# Generate some random data
data <- matrix(rnorm(seq_len(1000)), nrow=100)
# Estimate fuzzifiers
fuzz_out <- determine_fuzz(dim(data), 1)
# Run clustering
clres <- vsclust_algorithm(data, centers=5, m=fuzz_out$mm)
```

estimClust.plot      *Plotting results from estimating the cluster number*

---

### Description

This function visualizes the output from estimClustNumber, and there particularly the two validity indices Minimum Centroid Distance and Xie Beni Index.

### Usage

```
estimClust.plot(ClustInd)
```

### Arguments

ClustInd      Matrix with values from validity indices

### Value

Multiple panels showing expression profiles of clustered features passing the minMem threshold

### References

Schwaemmle V, Jensen ON. VSCLust: feature-based variance-sensitive clustering of omics data. *Bioinformatics*. 2018 Sep 1;34(17):2965-2972. doi: 10.1093/bioinformatics/bty224. PMID: 29635359.

Schwaemmle V, Hagensen CE. A Tutorial for Variance-Sensitive Clustering and the Quantitative Analysis of Protein Complexes. *Methods Mol Biol*. '2021;2228:433-451. doi: 10.1007/978-1-0716-1024-4\_30. PMID: 33950508.

Schwaemmle V, Jensen ON. A simple and fast method to determine the parameters for fuzzy c-means cluster analysis. *Bioinformatics*. 2010 Nov 15;26(22):2841-8. doi: 10.1093/bioinformatics/btq534. Epub 2010 Sep 29. PMID: 20880957.

### Examples

```
data("artificial_clusters")
dat <- averageCond(artificial_clusters, 5, 10)
dat <- scale(dat)
dat <- cbind(dat, 1)
ClustInd <- estimClustNum(dat, 6)
estimClust.plot(ClustInd)
```

---

estimClustNum      *Wrapper for estimation of cluster number*

---

### Description

This runs the clustering for different numbers of clusters, and estimates the most suitable numbers from applying the minimum centroid distance and the Xie Beni index. Multi-threading is used to shorten the computation times. Given the hierarchical structure of many data sets, the resulting numbers are suggestions. Inspection of the here plotted indices help to determine alternative cluster numbers, given by a strong decay of the minimum centroid distance and/or a low value of the Xie Beni index.



**Usage**

```
estimClustNum(dat, maxClust = 25, cores = 1)
```

**Arguments**

dat	matrix of features averaged over replicates. The last column contains their standard deviation
maxClust	Maximal number of cluster. The minimum is 3
cores	The number of threads to be used for parallelisation

**Value**

list with the items 'ClustInd': list of clustering objects for each number of clusters, 'p' plot object with plots for validity indices, 'numclust' optimal cluster number according to "minimum centroid distance"

**Examples**

```
data <- matrix(rnorm(1000), nrow=100)
estim_out <- estimClustNum(data, maxClust=10)
best_number <- max(estim_out[1])
```

---

mfuzz.plot

*Plotting vsclust results*

---

**Description**

This function visualizes the clustered quantitative profiles in multiple figure panels. The parameters allow specifying the main items like axes labels and color maps. The code is adopted from the MFuzz package.

**Usage**

```
mfuzz.plot(
  dat,
  cl,
  mfrow = c(1, 1),
  colo,
  minMem = 0,
  timeLabels,
  filename = NA,
  xlab = "Time",
  ylab = "Expression changes"
)
```

**Arguments**

dat	a numeric data matrix containing the values used in the clustering
cl	clustering results from vsclust_algorithm or Bestcl object from clustComp function
mfrow	vector of two numbers for the number of rows and columns, figure panels are distributed in the plot
colo	color map to be used (can be missing)
minMem	filter for showing only features with a higher membership values than this value
timeLabels	alternative labels for different conditions
filename	for writing into pdf. Will write on screen when using NA
xlab	Label of x-axis
ylab	Label of y-axis

**Value**

Multiple panels showing expression profiles of clustered features passing the minMem threshold

**References**

Schwaemmle V, Jensen ON. VSclust: feature-based variance-sensitive clustering of omics data. *Bioinformatics*. 2018 Sep 1;34(17):2965-2972. doi: 10.1093/bioinformatics/bty224. PMID: 29635359.

Schwaemmle V, Hagensen CE. A Tutorial for Variance-Sensitive Clustering and the Quantitative Analysis of Protein Complexes. *Methods Mol Biol*. 2021;2228:433-451. doi: 10.1007/978-1-0716-1024-4\_30. PMID: 33950508.

Schwaemmle V, Jensen ON. A simple and fast method to determine the parameters for fuzzy c-means cluster analysis. *Bioinformatics*. 2010 Nov 15;26(22):2841-8. doi: 10.1093/bioinformatics/btq534. Epub 2010 Sep 29. PMID: 20880957.

**Examples**

```
#' # Generate some random data
data <- matrix(rnorm(seq_len(5000)), nrow=500)
# Run clustering
clres <- vsclust_algorithm(data, centers=2, m=1.5)
mfuzz.plot(data, clres, mfrow=c(2,3), minMem=0.0)
```

---

optimalClustNum

*Determine optimal cluster number from validity index*

---

**Description**

Calculated the optimal number from expected behavior of the indices. This would be a large decay for the Minimum Centroid Distance and a minimum for the Xie Beni index

**Usage**

```
optimalClustNum(ClustInd, index = "MinCentroidDist", method = "VSclust")
```

**Arguments**

ClustInd	Output from estimClustNum providing the calculated cluster validity indices
index	Either "MinCentroidDist" or "XieBeni"
method	Either "VSClust" or "FCM" for standard fuzzy c-means clustering

**Value**

optimal cluster number

**References**

- Schwaemmle V, Jensen ON. VSClust: feature-based variance-sensitive clustering of omics data. *Bioinformatics*. 2018 Sep 1;34(17):2965-2972. doi: 10.1093/bioinformatics/bty224. PMID: 29635359.
- Schwaemmle V, Hagensen CE. A Tutorial for Variance-Sensitive Clustering and the Quantitative Analysis of Protein Complexes. *Methods Mol Biol*. 2021;2228:433-451. doi: 10.1007/978-1-0716-1024-4\_30. PMID: 33950508.
- Schwaemmle V, Jensen ON. A simple and fast method to determine the parameters for fuzzy c-means cluster analysis. *Bioinformatics*. 2010 Nov 15;26(22):2841-8. doi: 10.1093/bioinformatics/btq534. Epub 2010 Sep 29. PMID: 20880957.

**Examples**

```
data("artificial_clusters")
dat <- averageCond(artificial_clusters, 5, 10)
dat <- scale(dat)
dat <- cbind(dat, 1)
ClustInd <- estimClustNum(dat, 6)
optimalClustNum
```

---

pcaWithVar	<i>Visualize using principal component analysis (both loadings and scoring) including the variance from the replicates</i>
------------	--

---

**Description**

The loading plot shows all features and their scaled variance. This provides an idea of the intrinsic noise in the data.

**Usage**

```
pcaWithVar(data, NumReps, NumCond, Sds = 1)
```

**Arguments**

data	Matrix of data frame with numerical values. Columns corresponds to samples
NumReps	Number of replicates per experimental condition
NumCond	Number of different experimental conditions
Sds	Standard deviation for each features. Usually using the one from LIMMA

**Value**

Loading and scoring plots that include feature variance

**References**

Schwaemmle V, Jensen ON. VSCLust: feature-based variance-sensitive clustering of omics data. *Bioinformatics*. 2018 Sep 1;34(17):2965-2972. doi: 10.1093/bioinformatics/bty224. PMID: 29635359.

Schwaemmle V, Hagensen CE. A Tutorial for Variance-Sensitive Clustering and the Quantitative Analysis of Protein Complexes. *Methods Mol Biol*. 2021;2228:433-451. doi: 10.1007/978-1-0716-1024-4\_30. PMID: 33950508.

Schwaemmle V, Jensen ON. A simple and fast method to determine the parameters for fuzzy c-means cluster analysis. *Bioinformatics*. 2010 Nov 15;26(22):2841-8. doi: 10.1093/bioinformatics/btq534. Epub 2010 Sep 29. PMID: 20880957.

**Examples**

```
data <- matrix(rnorm(1000), nrow=100)
pcaWithVar(data, NumCond=2, NumReps=5, Sds=1)
```

---

PrepareForVSClust      *Wrapper for statistical analysis*

---

**Description**

Prepare data for running vsclust clustering. This includes visualization running the functions for the principal component analysis and its visualization, statistical testing with LIMMA, as well as scaling and filtering of missing values

**Usage**

```
PrepareForVSClust(dat, NumReps, NumCond, isPaired = FALSE, isStat)
```

**Arguments**

dat	matrix or data frame of numerical data. Columns are samples. Replicates are grouped (i.e. A1, B1, C1, A2, B2, C2) when letters denote conditions and numbers the replicates. In case of 'isStat=FALSE', you need a last column for the standard deviations
NumReps	Number replicates in the data
NumCond	Number of different experimental conditions. The total number of columns needs to be NumReps*NumCond
isPaired	Boolean for running paired or unpaired statistical tests
isStat	Boolean for whether to run statistical test or each column corresponds to a different experimental conditions. Then this function reads feature standard deviations from data frame from the last column

**Value**

list with the items 'dat' (data matrix of features averaged over replicates and last column with their standard deviations), 'qvals' FDRs from the statistical tests (each conditions versus the first), 'StatFileOut' all of before for saving in file

## References

- Schwaemmle V, Jensen ON. VSCLust: feature-based variance-sensitive clustering of omics data. *Bioinformatics*. 2018 Sep 1;34(17):2965-2972. doi: 10.1093/bioinformatics/bty224. PMID: 29635359.
- Schwaemmle V, Hagensen CE. A Tutorial for Variance-Sensitive Clustering and the Quantitative Analysis of Protein Complexes. *Methods Mol Biol*. 2021;2228:433-451. doi: 10.1007/978-1-0716-1024-4\_30. PMID: 33950508.
- Schwaemmle V, Jensen ON. A simple and fast method to determine the parameters for fuzzy c-means cluster analysis. *Bioinformatics*. 2010 Nov 15;26(22):2841-8. doi: 10.1093/bioinformatics/btq534. Epub 2010 Sep 29. PMID: 20880957.

## Examples

```
data <- matrix(rnorm(2000), nrow=200)
stats <- PrepareForVSCLust(data, 5, 2, isStat=TRUE)
```

---

PrepareSEForVSCLust      *Wrapper for statistical analysis for SummarizedExperiment object*

---

## Description

Prepare data for running vsclust clustering. This includes visualization running the functions for the principal component analysis and its visualization, statistical testing with LIMMA, as well as scaling and filtering of missing values

## Usage

```
PrepareSEForVSCLust(
  se,
  assayname = 1,
  coldatname = NULL,
  isPaired = FALSE,
  isStat
)
```

## Arguments

se	SummarizedExperiment object
assayname	Sample in SummarizedExperiment object
coldatname	Column in colData for extracting replicates
isPaired	Boolean for running paired or unpaired statistical tests
isStat	Boolean for whether to run statistical test or each column corresponds to a different experimental conditions. Then this function reads feature standard deviations from data frame from the last column

## Value

list with the items 'dat' (data matrix of features averaged over replicates and last column with their standard deviations), 'qvals' FDRs from the statistical tests (each conditions versus the first), 'StatFileOut' all of before for saving in file, 'NumReps' number of replicates and 'NumCond' number of different experimental conditions

## References

- Schwaemmle V, Jensen ON. VSCLust: feature-based variance-sensitive clustering of omics data. *Bioinformatics*. 2018 Sep 1;34(17):2965-2972. doi: 10.1093/bioinformatics/bty224. PMID: 29635359.
- Schwaemmle V, Hagensen CE. A Tutorial for Variance-Sensitive Clustering and the Quantitative Analysis of Protein Complexes. *Methods Mol Biol*. 2021;2228:433-451. doi: 10.1007/978-1-0716-1024-4\_30. PMID: 33950508.
- Schwaemmle V, Jensen ON. A simple and fast method to determine the parameters for fuzzy c-means cluster analysis. *Bioinformatics*. 2010 Nov 15;26(22):2841-8. doi: 10.1093/bioinformatics/btq534. Epub 2010 Sep 29. PMID: 20880957.

## Examples

```
data(miniACC, package="MultiAssayExperiment")

stats <- PrepareSEForVSCLust(miniACC, coldatname="COC", isStat=TRUE)
```

---

protein\_expressions     *Data from a typical proteomics experiment*

---

## Description

There are 12 samples coming from mouse fed with the four different diets, measured in three replicates each. Relative protein abundances were obtained using iTRAQ labelling. The given numbers are log<sub>2</sub>-transformed. Protein names as UniProt accession numbers are given as rownames.

## Usage

```
protein_expressions
```

## Format

A data frame consisting of 574 proteins measured in 12 samples:

**HF.Rep.1** Mice fed with a high fat diet, replicate 1

**HF.Rep.2** Mice fed with a high fat diet, replicate 2

**HF.Rep.3** Mice fed with a high fat diet, replicate 3

**TTA.Rep.1** Mice fed with a diet containing TTA (Tetradecylthioacetic Acid) high fat diet, replicate 1

**TTA.Rep.2** Mice fed with a diet containing TTA (Tetradecylthioacetic Acid) high fat diet, replicate 2

**TTA.Rep.3** Mice fed with a diet containing TTA (Tetradecylthioacetic Acid) high fat diet, replicate 3

**FO.Rep.1** Mice fed with a fish oil diet, replicate 1

**FO.Rep.2** Mice fed with a fish oil diet, replicate 2

**FO.Rep.3** Mice fed with a fish oil diet, replicate 3

**TTA.FO.Rep.1** Mice fed with a diet containing fish oil and TTA, replicate 1

**TTA.FO.Rep.2** Mice fed with a diet containing fish oil and TTA, replicate 2

**TTA.FO.Rep.3** Mice fed with a diet containing fish oil and TTA, replicate 3

**Source**

Protein Research Group, University of Southern Denmark, Odense

---

runClustWrapper      *Wrapper for running cluster analysis*

---

**Description**

This function runs the clustering and visualizes the results.

**Usage**

```
runClustWrapper(  
  dat,  
  NClust,  
  proteins = NULL,  
  VSclust = TRUE,  
  cores,  
  verbose = FALSE  
)
```

**Arguments**

dat	matrix or data frame with feature values for different conditions
NClust	Number of cluster for running the clustering
proteins	vector with additional feature information (default is NULL) to be added to the results
VSclust	boolean. TRUE for running the variance-sensitive clustering. Otherwise, the function will call standard fuzzy c-means clustering
cores	Number of threads for the parallelization
verbose	Show more information during execution

**Value**

list with the items 'dat'(the original data), 'Bestcl' clustering results (same as from vsclust\_algorithm), 'p' (plot object with mfuzz plots), 'outFileClust'(suitable matrix with complete information) , 'ClustInd' (information about being member of any cluster, feature needs on membership values > 0.5)

**Examples**

```
data(iris)  
data <- cbind(iris[,seq_len(4)],1)  
clust_out <- runClustWrapper(data, NClust=3, cores=1)  
clust_out$p
```

---

<code>runVSCLustApp</code>	<i>Run VSCLust as Shiny app</i>
----------------------------	---------------------------------

---

**Description**

You will get the full functionality of the VSCLust workflow with multiple visualizations and downloads

**Usage**

```
runVSCLustApp()
```

**Value**

The shiny app should open in a browser or in RStudio.

**References**

- Schwaemmle V, Jensen ON. VSCLust: feature-based variance-sensitive clustering of omics data. *Bioinformatics*. 2018 Sep 1;34(17):2965-2972. doi: 10.1093/bioinformatics/bty224. PMID: 29635359.
- Schwaemmle V, Hagensen CE. A Tutorial for Variance-Sensitive Clustering and the Quantitative Analysis of Protein Complexes. *Methods Mol Biol*. 2021;2228:433-451. doi: 10.1007/978-1-0716-1024-4\_30. PMID: 33950508.
- Schwaemmle V, Jensen ON. A simple and fast method to determine the parameters for fuzzy c-means cluster analysis. *Bioinformatics*. 2010 Nov 15;26(22):2841-8. doi: 10.1093/bioinformatics/btq534. Epub 2010 Sep 29. PMID: 20880957.

**Examples**

```
runVSCLustApp()
```

---

<code>SignAnalysis</code>	<i>Unpaired statistical testing</i>
---------------------------	-------------------------------------

---

**Description**

Statistical testing and variance estimation in multi-dimensional data set. given by a matrix. This functions runs LIMMA paired tests and calculated the shrunken variance estimates.

**Usage**

```
SignAnalysis(Data, NumCond, NumReps)
```

**Arguments**

- |                      |  |
|----------------------|--|
| <code>Data</code>    | a numeric data matrix with columns as samples. Different experimental conditions are grouped together in their replicates. The number of samples per group needs to be identical |
| <code>NumCond</code> | Number of different experimental conditions  |
| <code>NumReps</code> | Number of replicates per experimental condition  |



**Value**

List containing the objects

‘pvalues’ p-values before correction for multiple testing

‘qvalues’ false discovery rates after correction for multiple testing (‘qvalue’ method from ‘qvalue’ library)

‘Sds’ General standard deviation within replicates after using shrinkage by LIMMA

**References**

Schwaemmle V, Jensen ON. VSClust: feature-based variance-sensitive clustering of omics data. *Bioinformatics*. 2018 Sep 1;34(17):2965-2972. doi: 10.1093/bioinformatics/bty224. PMID: 29635359.

Schwaemmle V, Hagensen CE. A Tutorial for Variance-Sensitive Clustering and the Quantitative Analysis of Protein Complexes. *Methods Mol Biol*. 2021;2228:433-451. doi: 10.1007/978-1-0716-1024-4\_30. PMID: 33950508.

Schwaemmle V, Jensen ON. A simple and fast method to determine the parameters for fuzzy c-means cluster analysis. *Bioinformatics*. 2010 Nov 15;26(22):2841-8. doi: 10.1093/bioinformatics/btq534. Epub 2010 Sep 29. PMID: 20880957.

**Examples**

```
#' # Generate some random data
data <- matrix(rnorm(seq_len(1000)), nrow=100)
# Run statistical testing
stat_out <- SignAnalysis(data, 2, 5)
# Histogram of qvalues (no significant events)
hist(stat_out$qvalues, 50, xlab="q-values")
```

---

SignAnalysisPaired      *Paired statistical testing*

---

**Description**

Statistical testing and variance estimation in multi-dimensional data set. given by a matrix. This functions runs LIMMA paired tests and calculated the shrunken variance estimates.

**Usage**

```
SignAnalysisPaired(Data, NumCond, NumReps)
```

**Arguments**

Data	a numeric data matrix with columns as samples. Different experimental conditions are grouped together in their replicates. The number of samples per group needs to be identical
NumCond	Number of different experimental conditions
NumReps	Number of replicates per experimental condition

**Value**

List containing the objects

'qvalues' false discovery rates after correction for multiple testing ('qvalue' method from 'qvalue' library)

'Sds' General standard deviation within replicates after using shrinkage (eBayes) by LIMMA

**References**

Schwaemmle V, Jensen ON. VSClust: feature-based variance-sensitive clustering of omics data. *Bioinformatics*. 2018 Sep 1;34(17):2965-2972. doi: 10.1093/bioinformatics/bty224. PMID: 29635359.

Schwaemmle V, Hagensen CE. A Tutorial for Variance-Sensitive Clustering and the Quantitative Analysis of Protein Complexes. *Methods Mol Biol*. 2021;2228:433-451. doi: 10.1007/978-1-0716-1024-4\_30. PMID: 33950508.

Schwaemmle V, Jensen ON. A simple and fast method to determine the parameters for fuzzy c-means cluster analysis. *Bioinformatics*. 2010 Nov 15;26(22):2841-8. doi: 10.1093/bioinformatics/btq534. Epub 2010 Sep 29. PMID: 20880957.

**Examples**

```
#' # Generate some random data with three different experimental conditions
data <- matrix(rnorm(seq_len(1500)), nrow=100)
# Run statistical testing
stat_out <- SignAnalysisPaired(data, 3, 5)
# Histogram of qvalues comparing the second to the first condition
hist(stat_out$qvalues[,1], 50, xlab="q-values")
```

---

SwitchOrder

*arrange cluster member numbers from largest to smallest*


---

**Description**

arrange cluster member numbers from largest to smallest

**Usage**

```
SwitchOrder(Bestcl, NClust)
```

**Arguments**

Bestcl            fclust object

NClust            Number of clusters

**Value**

fclust object with reorder clusters

**Examples**

```
# Generate some random data
data <- matrix(rnorm(seq_len(1000)), nrow=100)
# Run clustering
clres <- vsclust_algorithm(data, centers=5, m=1.5)
clres <- SwitchOrder(clres, 5)
```

---

vsclust\_algorithm      *Run the vsclust clustering algorithm*

---

**Description**

This function calls the c++ implementation of the vsclust algorithm, being an extension of fuzzy c-means clustering with additional variance control and capability to run on data with missing values

**Usage**

```
vsclust_algorithm(
  x,
  centers,
  iterMax = 100,
  verbose = FALSE,
  dist = "euclidean",
  m = 2,
  ratePar = NULL,
  weights = 1,
  control = list()
)
```

**Arguments**

x	a numeric data matrix
centers	Either numeric for number of clusters or numeric matrix with center coordinates
iterMax	Numeric for maximum number of iterations
verbose	Verbose information
dist	Distance to use for the calculation. We prefer "euclidean" (default)
m	Fuzzifier value: numeric or vector of length equal to number of rows of x
ratePar	(experimental) numeric value for punishing missing values
weights	numeric or vector of length equal to number of rows of x
control	list with arguments to vsclust algorithms (now only cutoff for relative tolerance: reltol)

**Value**

list with details about clustering having the objects 'centers' (positions of centroids), 'size' (feature number per cluster), 'cluster' (nearest cluster of each feature), 'membership' matrix of membership values, 'iter' (number of carried out iterations), 'withinerror' (final error from optimization), 'call' (call of function)

## References

Schwaemmle V, Jensen ON. VSclust: feature-based variance-sensitive clustering of omics data. *Bioinformatics*. 2018 Sep 1;34(17):2965-2972. doi: 10.1093/bioinformatics/bty224. PMID: 29635359.

Schwaemmle V, Hagensen CE. A Tutorial for Variance-Sensitive Clustering and the Quantitative Analysis of Protein Complexes. *Methods Mol Biol*. 2021;2228:433-451. doi: 10.1007/978-1-0716-1024-4\_30. PMID: 33950508.

Schwaemmle V, Jensen ON. A simple and fast method to determine the parameters for fuzzy c-means cluster analysis. *Bioinformatics*. 2010 Nov 15;26(22):2841-8. doi: 10.1093/bioinformatics/btq534. Epub 2010 Sep 29. PMID: 20880957.

## Examples

```
#' # Generate some random data
data <- matrix(rnorm(seq_len(1000)), nrow=100)
# Run clustering
clres <- vsclust_algorithm(data, centers=5, m=1.5)
head(clres$membership)
```

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