

Package ‘ICGE’

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Title Estimation of Number of Clusters and Identification of Atypical Units

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Depends R (>= 2.0.1), utils, stats, MASS, cluster, fastcluster

Description It is a package that helps to estimate the number of real clusters in data as well as to identify atypical units. The underlying methods are based on distances rather than on unit x variables.

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LazyData yes

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chowdary	<i>Chowdary Database</i>
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Description

The original authors compared pairs of snap-frozen and RNAlater preservative-suspended tissue from lymph node-negative breast tumors (B) and Dukes’ B colon tumors (C). The actual data set, by de Souto et. al (2008), is build with purpose of separating B from C.

Usage

data(chowdary)

Format

Data frame with 183 rows and 104 columns.

Source

Original source from ‘National Center for Biotechnology Information’ from the United States of America, query GSE3726.

References

de Souto MCP, Costa IG, de Araujo DSA, Ludermir TB, and Schliep A (2008). Clustering Cancer Gene Expression Data: a Comparative Study. *BMC Bioinformatics*, **8**, 497–511.

Chowdary D, Lathrop J, Skelton J, Curtin K, Briggs T, Zhang Y, Yu J, Wang X, and Mazumder A (2006). Prognostic gene expression signatures can be measured in tissues collected in RNAlater preservative. *Journal Molecular Diagnosis*, **8**, 31–39.

Examples

```
data(chowdary)

tumor <- as.factor(as.matrix(chowdary[1,]))
x <- as.matrix(chowdary[-1,])
mode(x) <- "numeric"

s <- sample(row.names(x),1)
boxplot( x[s,] ~ tumor , ylab=s)
```

dbhatta	<i>Bhattacharyya Distance</i>
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Description

dbhatta computes and returns the Bhattacharyya distance matrix between the rows of a data matrix. This distance is defined between two units $i = (p_{i1}, \dots, p_{im})$ and $j = (p_{j1}, \dots, p_{jm})$ being p_{kl} frequencies with $p_{kl} \geq 0$ and $p_{k1} + \dots + p_{km} = 1$.

Usage

```
dbhatta(x)
```

Arguments

x a matrix containing, in its rows, the frequencies for each unit. Note: check that each row adds up to 1

Value

A [dist](#) object with distance information.

Author(s)

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References

Bhattacharyya, A. (1946). On a measure of divergence of two multinomial populations. *Sankhya: The Indian Journal of Statistics, Series A*. **14**, 177-136.

See Also

[dist](#), [dmahal](#), [dgower](#), [dcor](#), [dproc2](#)

Examples

```
#5 individuals represented by their relative frequencies of 4 characteristics (M1-M4):
f <- matrix(c(0.36, 0.21, 0.23, 0.20,
              0.66, 0.18, 0.11, 0.05,
              0.01, 0.24, 0.62, 0.13,
              0.43, 0.38, 0.08, 0.11,
              0.16, 0.07, 0.09, 0.68),
            byrow=TRUE, nrow=5, dimnames=list(1:5, paste("M", 1:4, sep="")))

```

```
# Bhattacharyya distances between pairs
d <- dbhatta(f)
```

dcor

Correlation Distance

Description

dcor computes and returns the Correlation distance matrix between the rows of a data matrix. This distance is defined by $d = \sqrt{1 - r}$.

Usage

```
dcor(x)
```

Arguments

x a numeric matrix.

Value

A [dist](#) object with distance information.

Author(s)

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References

Gower, J.C. (1985). Measures of similarity, dissimilarity and distance. In: *Encyclopedia of Statistical Sciences*, volume **5**, 397–405. J. Wiley and Sons.

See Also

[dist](#), [dmahal](#), [dgower](#), [dbhatta](#), [dproc2](#)

Examples

```
#Generate 10 objects in dimension 8
n <- 10
mu <- sample(1:10, 8, replace=TRUE)
x <- matrix(rnorm(n*8, mean=mu, sd=1), nrow=n, byrow=TRUE)

# Correlation distances between pairs
d <- dcor(x)
```

deltas	<i>Distance Between Groups</i>
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Description

Assume that n units are divided into k groups $C_1, \dots, C_k$. Function `deltas` computes and returns the distance between each pair of groups. It uses the distances between pairs of units.

Usage

```
deltas(d, pert = "onegroup")
```

Arguments

<code>d</code>	a distance matrix or a <code>dist</code> object with distance information between units.
<code>pert</code>	an n -vector that indicates which group each unit belongs to. Note that the expected values of <code>pert</code> are numbers greater than or equal to 1 (for instance 1,2,3,4,..., k). The default value indicates there is only one group in data.

Value

A matrix containing the distances between each pair of groups.

Author(s)

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Conchita Arenas <carenas@ub.edu>; Departament d'Estadística, Universitat de Barcelona, Barcelona, Spain.

References

Arenas, C. and Cuadras, C.M. (2002). Some recent statistical methods based on distances. *Contributions to Science*, **2**, 183–191.

Cuadras, C.M., Fortiana, J. and Oliva, F. (1997). The proximity of an individual to a population with applications in discriminant analysis. *Journal of Classification*, **14**, 117–136.

See Also

[vgeo](#), [proxi](#)

Examples

```
data(iris)
d <- dist(iris[,1:4])
deltas(d,iris[,5])
```

dermatology

*Dermatology Database***Description**

Data from a dermatology study provided by H.A. Guvenir (Dpt. Computer Engineering and Information Science, Bilkent University, Turkey). The data set contains 366 instances presenting 34 different clinical attributes (12 clinical features as age or family history and 22 histopathological features obtained from a biopsy), and a class variable indicating the disease. There are 8 missing values. This data set has been used extensively for classification tasks.

Usage

```
data(dermatology)
```

Format

Matrix with 366 rows.

Details

Attribute information obtained from the UCI KDD data repository:

Clinical Attributes: (they take values 0, 1, 2, 3, unless otherwise indicated)

1: erythema; 2: scaling; 3: definite borders; 4: itching; 5: koebner phenomenon; 6: polygonal papules; 7: follicular papules; 8: oral mucosal involvement; 9: knee and elbow involvement; 10: scalp involvement; 11: family history, (0 or 1); 34: Age.

Histopathological Attributes: (they take values 0, 1, 2, 3)

12: melanin incontinence; 13: eosinophils in the infiltrate; 14: PNL infiltrate; 15: fibrosis of the papillary dermis; 16: exocytosis; 17: acanthosis; 18: hyperkeratosis; 19: parakeratosis; 20: clubbing of the rete ridges; 21: elongation of the rete ridges; 22: thinning of the suprapapillary epidermis; 23: spongiform pustule; 24: munro microabcess; 25: focal hypergranulosis; 26: disappearance of the granular layer; 27: vacuolisation and damage of basal layer; 28: spongiosis; 29: saw-tooth appearance of retes; 30: follicular horn plug; 31: perifollicular parakeratosis; 32: inflammatory mononuclear infiltrate; 33: band-like infiltrate.

The considered diseases are: 1 - psoriasis, 2 - seboric dermatitis, 3- lichen planus, 4 - pityriasis rosea, 5 - chronic dermatitis, 6 - pityriasis rubra pilaris.

Source

The UCI KDD Archive.

References

- Guvenir H, Demiroz G, Ilter N (1998). Learning differential diagnosis of erythemato-squamous diseases using voting feature intervals. *Artificial Intelligence in Medicine*, **13**, 147–165.
- Irigoin I, Arenas C (2008). INCA: New statistic for estimating the number of clusters and identifying atypical units. *Statistics in Medicine*, **27**, 2948–2973.

Examples

```
data(dermatology)
x <- dermatology[, 1:34]
group <- as.factor(dermatology[,35])

plot(group)
```

dgower

*Gower Distance for Mixed Variables***Description**

dgower computes and returns the Gower distance matrix for mixed variables.

Usage

```
dgower(x, type = list())
```

Arguments

x	data matrix.
type	it is a list with components cuant, bin, nom. Each component indicates the column position of the quantitative, binary or nominal variables, respectively.

Details

The distance between two pairs of objects i and j is obtained as $\sqrt{2(1 - s_{ij})}$ where s_{ij} is the Gower's similarity coefficient for mixed data. This function allows to include missing values (as NA) and therefore calculates distances based on Gower's weighted similarity coefficient.

Value

A `dist` object with distance information.

Note

There is the function `daisy()` in `cluster` package which can perform the Gower distance for mixed variables. The difference is that in `daisy()` the distance is calculated as $d(i, j) = 1 - s_{ij}$ and in `dgower()` it is calculated as $d_{ij} = \sqrt{2(1 - s_{ij})}$.

Author(s)

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References

Gower, J.C. (1971). A general coefficient of similarity and some of its properties. *Biometrics*, **27**, 857–871.

See Also

[dist](#), [dmahal](#), [dbhatta](#), [dcor](#), [dproc2](#)

Examples

```
#Generate 10 objects in dimension 6
# Quantitative variables
mu <- sample(1:10, 2, replace=TRUE)
xc <- matrix(rnorm(10*2, mean = mu, sd = 1), ncol=2, byrow=TRUE)

# Binary variables
xb <- cbind(rbinom(10, 1, 0.1), rbinom(10, 1, 0.5), rbinom(10, 1, 0.9))

# Nominal variables
xn <- matrix(sample(1:3, 10, replace=TRUE), ncol=1)

x <- cbind(xc, xb, xn)

# Distances
d <- dgower(x, type=list(cuant=1:2, bin=3:5, nom=6))
```

dmahal

Mahalanobis Distance

Description

dmahal computes and returns the Mahalanobis distance matrix between the rows of a data matrix.

Usage

```
dmahal(datos, S)
```

Arguments

datos	data matrix.
S	covariance matrix.

Value

A [dist](#) object with distance information.

Note

There is a function `mahalanobis()` in stats package which can perform the Mahalanobis distance. While `mahalanobis()` calculates the Mahalanobis distance with respect to given a center, function `dmahal()` is designed to calculate the distance between each pair of units given a data matrix.

Author(s)

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References

Everitt B. S. and Dunn G. (2001) *Applied Multivariate Data Analysis*. 2 edition, Edward Arnold, London.

See Also

[dist](#), [dbhatta](#), [dgower](#), [dcor](#), [dproc2](#)

Examples

```
#Generate 10 objects in dimension 2
mu <- rep(0, 2)
Sigma <- matrix(c(10,3,3,2),2,2)

x <- mvrnorm(n=10, rep(0, 2), Sigma)

d <- dmahal(x, Sigma)
```

dproc2

Modified Procrustes distance

Description

`dproc2` computes and returns all the pairwise procrustes distances between genes in a time course experiment, using their expression profile.

Usage

```
dproc2(x, timepoints = NULL)
```

Arguments

<code>x</code>	a matrix containing, in its rows, the gene expression values at the T considered time points.
<code>timepoints</code>	a T-vector with the T observed time points. If <code>timepoints=NULL</code> (default), then <code>timepoints=1:T</code> .

Details

Each row i of matrix x is arranged in a two column matrix X_i . In X_i , the first column contains the time points and the second column the observed gene expression values ($x_{i1}...$).

Value

A `dist` object with distance information.

Author(s)

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References

Irigoien, I. , Vives, S. and Arenas, C. (2011). Microarray Time Course Experiments: Finding Profiles. *IEEE/ACM Transactions on Computational Biology and Bioinformatics*, **8**(2), 464–475.

Gower, J. C. and Dijksterhuis, G. B. (2004) *Procrustes Problems*. Oxford University Press.

Sibson, R. (1978). Studies in the Robustness of Multidimensional Scaling: Procrustes statistic. *Journal of the Royal Statistical Society, Series B*, **40**, 234–238.

See Also

`dist`, `dmahal`, `dgower`, `dcor` `dbhatta`

Examples

```
# Given 10 hypothetical time course profiles
# over 6 time points at 1, 2, ..., 6 hours.
x <- matrix(c(0.38, 0.39, 0.38, 0.37, 0.385, 0.375,
              0.99, 1.19, 1.50, 1.83, 2.140, 2.770,
              0.38, 0.50, 0.71, 0.72, 0.980, 1.010,
              0.20, 0.40, 0.70, 1.06, 2.000, 2.500,
              0.90, 0.95, 0.97, 1.50, 2.500, 2.990,
              0.64, 2.61, 1.51, 1.34, 1.330, 1.140,
              0.71, 1.82, 2.28, 1.72, 1.490, 1.060,
              0.71, 1.82, 2.28, 1.99, 1.975, 1.965,
              0.49, 0.78, 1.00, 1.27, 0.590, 0.340,
              0.71, 1.00, 1.50, 1.75, 2.090, 1.380), nrow=10, byrow=TRUE)

# Graphical representation
matplot(t(x), type="b")

# Distance matrix between them
d <- dproc2(x)
```

estW

INCA Statistic

Description

Assume that n units are divided into k clusters $C_1, \dots, C_k$, and consider a fixed unit x_0 . Function `estW` calculates the INCA statistic $W(x_0)$ and the related U_i statistics.

Usage

```
estW(d, dx0, pert = "onegroup")
```

Arguments

<code>d</code>	a distance matrix or a <code>dist</code> object with distance information between units.
<code>dx0</code>	an n -vector containing the distances d_{0j} between x_0 and unit j .
<code>pert</code>	an n -vector that indicates which group each unit belongs to. Note that the expected values of <code>pert</code> are consecutive integers bigger or equal than 1 (for instance 1,2,3,4,..., k). The default value indicates the presence of only one group in data.

Value

The function returns an object of class `incaest` which is a list containing the following components:

<code>Wvalue</code>	is the INCA statistic $W(x_0)$.
<code>Uvalue</code>	is a vector containing the statistics U_i .

Note

For a correct geometrical interpretation it is convenient to verify whether the distance matrix `d` is Euclidean.

Author(s)

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References

Arenas, C. and Cuadras, C.M. (2002). Some recent statistical methods based on distances. *Contributions to Science*, **2**, 183–191.

Irigoiien, I. and Arenas, C. (2008). INCA: New statistic for estimating the number of clusters and identifying atypical units. *Statistics in Medicine*, **27**(15), 2948–2973.

See Also

[vgeo](#), [proxi](#), [deltas](#)

Examples

```
data(iris)
d <- dist(iris[,1:4])

# characteristics of a specific flower (likely group 1)
x0 <- c(5.3, 3.6, 1.1, 0.1)
# distances between flower x0 and the rest of flowers in iris
dx0 <- rep(0,150)
for (i in 1:150){
  dif <-x0-iris[i,1:4]
  dx0[i] <- sqrt(sum(dif*dif))
}
estW(d, dx0, iris[,5])
```

INCAindex

INCA index

Description

INCAindex helps to estimate the number of clusters in a dataset.

Usage

```
INCAindex(d, pert_clus)
```

Arguments

d	a distance matrix or a dist object with distance information between units.
pert_clus	an n-vector that indicates which group each unit belongs to. Note that the expected values of pert are numbers greater than or equal to 1 (for instance 1,2,3,4..., k). The default value indicates the presence of only one group in data.

Value

Returns an object of class `incaix` which is a list containing the following components:

well_class	a vector indicating the number of well classified units.
Ni_cluster	a vector indicating each cluster size.
Total	percentage of objects well classified in the partition defined by <code>pert_clus</code> .

Note

For a correct geometrical interpretation it is convenient to verify whether the distance matrix d is Euclidean. It admits the associated methods summary and plot. The first simply returns the percentage of well-classified units and the second offers a barchart with the percentages of well classified units for each group in the given partition.

Author(s)

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References

Arenas, C. and Cuadras, C.M. (2002). Some recent statistical methods based on distances. *Contributions to Science*, **2**, 183–191.

Irigoien, I. and Arenas, C. (2008). INCA: New statistic for estimating the number of clusters and identifying atypical units. *Statistics in Medicine*, **27**(15), 2948–2973.

See Also

[estW](#), [INCAtest](#)

Examples

```
#generate 3 clusters, each of them with 20 objects in dimension 5.
mu1 <- sample(1:10, 5, replace=TRUE)
x1 <- matrix(rnorm(20*5, mean = mu1, sd = 1),ncol=5, byrow=TRUE)
mu2 <- sample(1:10, 5, replace=TRUE)
x2 <- matrix(rnorm(20*5, mean = mu2, sd = 1),ncol=5, byrow=TRUE)
mu3 <- sample(1:10, 5, replace=TRUE)
x3 <- matrix(rnorm(20*5, mean = mu3, sd = 1),ncol=5, byrow=TRUE)
x <- rbind(x1,x2,x3)

# Euclidean distance between units.
d <- dist(x)

# given the right partition, calculate the percentage of well classified objects.
partition <- c(rep(1,20), rep(2,20), rep(3,20))
INCAindex(d, partition)

# In order to estimate the number of cluster in data, try several
# partitions and compare the results
library(cluster)
T <- rep(NA, 5)
for (l in 2:5){
  part <- pam(d,l)$clustering
  T[l] <- INCAindex(d,part)$Total
```

```
}
plot(T, type="b", xlab="Number of clusters", ylab="INCA", xlim=c(1.5, 5.5))
```

INCAnumclu

Estimation of Number of Clusters in Data

Description

INCAnumclu helps to estimate the number of clusters in a dataset. The INCA index associated to different partitions with different number of clusters is calculated.

Usage

```
INCAnumclu(d, K, method = "pam", pert, L= NULL, noise=NULL)
```

Arguments

d	a distance matrix or a dist object with distance information between units.
K	the maximum number of cluster to be considered. For each k value (k=2,...,K) a partition with k clusters is calculated.
method	character string defining the clustering method in order to obtain the partitions. The hierarchical agglomerative clustering methods are performed via hclust function in package fastcluster . Other clustering methods are performed via the functions in package cluster , such as: pam , diana and fanny . The available clustering methods are pam (default method), average (UPGMA), single (single linkage), complete (complete linkage), ward.D2 (Ward's method), ward.D, centroid, median, diana (hierarchical divisive) and fanny (fuzzy clustering). Nevertheless, the user can introduce particular or custom partitions indicating method="partition" and specifying the partitions in argument pert.
pert	only useful when parameter method="partition"; it is a matrix and each column contains a partition of the units. That means that each column is an n-vector that indicates which group each unit belongs to. Note that the expected values of each column of pert are numbers greater than or equal to 1 (for instance 1,2,3,4..., k).
L	default value NULL, but when some units are considered by the user as noise units, L must be specified as follows: (a) L is greater than or equal to 1 and all units in clusters with a cardinal <= L are considered noise units; (b) L="custom" when the user wants to specify which units are considered noise units. These units must be specified in argument noise.
noise	when L="custom", it is a logical vector indicating the units considered by the user as noise units.

Value

Returns an object of class `incanc` which is a numeric vector containing the INCA index associated to each of the k ($k=2,\dots,K$) partitions. When noise is no null, the function returns a list with the INCA index for each partition, which is calculated without noise units as well as with noise units. The associated plot returns INCA index plot, both, with and without noise.

Author(s)

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References

Irigoien, I. and Arenas, C. (2008). INCA: New statistic for estimating the number of clusters and identifying atypical units. *Statistics in Medicine*, **27**(15), 2948–2973.

Arenas, C. and Cuadras, C.M. (2002). Some recent statistical methods based on distances. *Contributions to Science*, **2**, 183–191.

See Also

[INCAindex](#), [estW](#)

Examples

```
#----- Example 1 -----
#generate 3 clusters, each of them with 20 objects in dimension 5.
mu1 <- sample(1:10, 5, replace=TRUE)
x1 <- matrix(rnorm(20*5, mean = mu1, sd = 1),ncol=5, byrow=TRUE)
mu2 <- sample(1:10, 5, replace=TRUE)
x2 <- matrix(rnorm(20*5, mean = mu2, sd = 1),ncol=5, byrow=TRUE)
mu3 <- sample(1:10, 5, replace=TRUE)
x3 <- matrix(rnorm(20*5, mean = mu3, sd = 1),ncol=5, byrow=TRUE)
x <- rbind(x1,x2,x3)

# calculte euclidean distance between them
d <- dist(x)

# calculate the INCA index associated to partitions with k=2, ..., k=5 clusters.
INCAnumclu(d, K=5)
out <- INCAnumclu(d, K=5)
plot(out)

#----- Example 1 cont. -----
# With hypothetical noise elements
noiseunits <- rep(FALSE, 60)
noiseunits[sample(1:60, 20)] <- TRUE
out <- INCAnumclu(d, K=5, L="custom", noise=noiseunits)
plot(out)
```

INCAtest

*INCA Test***Description**

Assume that n units are divided into k groups $C_1, \dots, C_k$. Function `INCAtest` performs the typicality INCA test. Therein, the null hypothesis that a new unit x_0 is a typical unit with respect to a previously fixed partition is tested versus the alternative hypothesis that the unit is atypical.

Usage

```
INCAtest(d, pert, d_test, np = 1000, alpha = 0.05, P = 1)
```

Arguments

<code>d</code>	a distance matrix or a <code>dist</code> object with distance information between units.
<code>pert</code>	an n -vector that indicates which group each unit belongs to. Note that the expected values of <code>pert</code> are numbers greater than or equal to 1 (for instance 1,2,3,4..., k). The default value indicates there is only one group in data.
<code>d_test</code>	an n -vector containing the distances from x_0 to the other units.
<code>np</code>	sample size for the bootstrap sample for the bootstrap procedure.
<code>alpha</code>	fixed level for the test.
<code>P</code>	Number of times the bootstrap procedure is repeated.

Value

A list with class "incat" containing the following components:

<code>StatisticW0</code>	value of the INCA statistic.
<code>ProjectionsU</code>	values of statistics measuring the projection from the specific object to each considered group.
<code>pvalues</code>	p-values obtained in the P times repeated bootstrap procedure. Note: If $P > 1$, it is printed the number of times the p-values were smaller than <code>alpha</code> .
<code>alpha</code>	specified value of the level of the test.

Note

To obtain the INCA statistic distribution, under the null hypothesis, the program can consume long time. For a correct geometrical interpretation it is convenient to verify whether the distance matrix `d` is Euclidean.

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References

Irigoin, I. and Arenas, C. (2008). INCA: New statistic for estimating the number of clusters and identifying atypical units. *Statistics in Medicine*, **27**(15), 2948–2973.

Arenas, C. and Cuadras, C.M. (2002). Some recent statistical methods based on distances. *Contributions to Science*, **2**, 183–191.

See Also

[estW](#), [INCAindex](#)

Examples

```
#generate 3 clusters, each of them with 20 objects in dimension 5.
mu1 <- sample(1:10, 5, replace=TRUE)
x1 <- matrix(rnorm(20*5, mean = mu1, sd = 1),ncol=5, byrow=TRUE)
mu2 <- sample(1:10, 5, replace=TRUE)
x2 <- matrix(rnorm(20*5, mean = mu2, sd = 1),ncol=5, byrow=TRUE)
mu3 <- sample(1:10, 5, replace=TRUE)
x3 <- matrix(rnorm(20*5, mean = mu3, sd = 1),ncol=5, byrow=TRUE)
x <- rbind(x1,x2,x3)

# Euclidean distance between units in matrix x.
d <- dist(x)
# given the right partition
partition <- c(rep(1,20), rep(2,20), rep(3,20))

# x0 contains a unit from one group, as for example group 1.
x0 <- matrix(rnorm(1*5, mean = mu1, sd = 1),ncol=5, byrow=TRUE)

# distances between x0 and the other units.
dx0 <- rep(0,60)
for (i in 1:60){
  dif <-x0-x[i,]
  dx0[i] <- sqrt(sum(dif*dif))
}

INCAtest(d, partition, dx0, np=10)

# x0 contains a unit from a new group.
x0 <- matrix(rnorm(1*5, mean = sample(1:10, 5, replace=TRUE),
  sd = 1), ncol=5, byrow=TRUE)

# distances between x0 and the other units in matrix x.
dx0 <- rep(0,60)
for (i in 1:60){
  dif <-x0-x[i,]
  dx0[i] <- sqrt(sum(dif*dif))
}

INCAtest(d, partition, dx0, np=10)
```

 lympa

Lymphatic Database

Description

This lymphography domain was obtained from the University Medical Centre, Institute of Oncology, Ljubljana, Yugoslavia. Thanks go to M. Zwitter and M. Soklic for providing the data. The data are available at the UCI KDD data repository (Hettich S and Bay SD, 1999).

The data set consists of 148 instances presenting 18 different mixed attributes (1 quantitative, 9 binaries and 9 nominals), and a class variable indicating the diagnostic. There are not missing values.

Usage

`data(lympa)`

Format

Data frame with 148 instances and 19 features.

Details

Attribute information:

— NOTE: All attribute values in the database have been entered as numeric values corresponding to their index in the list of attribute values for that attribute domain as given below.

1. class: normal find, metastases, malign lymph, fibrosis
2. lymphatics: normal, arched, deformed, displaced
3. block of affere: no, yes
4. bl. of lymph. c: no, yes
5. bl. of lymph. s: no, yes
6. by pass: no, yes
7. extravasates: no, yes
8. regeneration of: no, yes
9. early uptake in: no, yes
10. lym.nodes dimin: 0-3
11. lym.nodes enlar: 1-4
12. changes in lym.: bean, oval, round
13. defect in node: no, lacunar, lac. marginal, lac. central
14. changes in node: no, lacunar, lac. margin, lac. central
15. changes in stru: no, grainy, drop-like, coarse, diluted, reticular, stripped, faint

- 16. special forms: no, chalices, vesicles
- 17. dislocation of: no, yes
- 18. exclusion of no: no, yes
- 19. no. of nodes in: 0-9, 10-19, 20-29, 30-39, 40-49, 50-59, 60-69, >=70

Source

The UCI KDD Archive.

References

Hettich S and Bay SD (1999). The UCI KDD Archive. Department of Information and Computer Science. University of California at Irvine, Irvine, USA.

Examples

```
data(lympa)
aux <- table(lympa[,1])
barplot(aux, names.arg=c("normal", "metastases", "malign lymph", "fibrosis"))
```

proxi

Proximity Function

Description

Assume that n units are divided into k groups $C_1, \dots, C_k$. The function calculates the proximity function from a specific unit x_0 to the groups C_j .

Usage

```
proxi(d, dx0, pert = "onegroup")
```

Arguments

<code>d</code>	a distance matrix or a <code>dist</code> object with distance information between units.
<code>dx0</code>	an n -vector containing the distances from x_0 to the other units.
<code>pert</code>	an n -vector that indicates which group each unit belongs to. Note that the expected values of <code>pert</code> are numbers greater than or equal to 1 (for instance 1,2,3,4..., k). The default value indicates there is only one group in data.

Value

k -vector containing the proximity function value from x_0 to each group.

Author(s)

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References

Arenas, C. and Cuadras, C.M. (2002). Some recent statistical methods based on distances. *Contributions to Science*, **2**, 183–191.

Cuadras, C.M., Fortiana, J. and Oliva, F. (1997). The proximity of an individual to a population with applications in discriminant analysis. *Journal of Classification*, **14**, 117–136.

See Also

[vgeo](#), [deltas](#)

Examples

```
data(iris)
d <- dist(iris[,1:4])

# x0 contains a unit from one group, as for example group 1.
x0 <- c(5.3, 3.6, 1.1, 0.1)
# distances between x0 and the other units.
dx0 <- rep(0,150)
for (i in 1:150){
  dif <- x0-iris[i,1:4]
  dx0[i] <- sqrt(sum(dif*dif))
}

proxi(d, dx0, iris[,5])

# x0 contains a unit from one group, as for example group 2.
x0 <- c(6.4, 3.0, 4.8, 1.3)
# distances between x0 and the other units.
dx0 <- rep(0,150)
for (i in 1:150){
  dif <- x0-iris[i,1:4]
  dx0[i] <- sqrt(sum(dif*dif))
}

proxi(d, dx0, iris[,5])
```

SyntheticTimeCourse	<i>Synthetic Time Course data</i>
---------------------	-----------------------------------

Description

Synthetic time course data where 210 genes profiles along 6 time points are reported and where the genes are drawn from 8 different populations.

Usage

```
data(SyntheticTimeCourse)
```

Format

Data frame with 120 rows and 7 columns.

Details

Attribute information: Column cl: the class that the gen belongs to. Columns t1 - t6: gene's expression along the t1, ..., t6 time points considered.

Examples

```
data(SyntheticTimeCourse)
x <- SyntheticTimeCourse[, 2:7]
cl <- SyntheticTimeCourse[, 1]
par(mfrow=c(3,3))
for (g in 1:8){
  xx <- t(x[cl==g,] )
  yy <- matrix(c(1:6 ), nrow=6, ncol=15, byrow=FALSE)
  matplot(yy,xx, pch=21, type="b", axes=FALSE,
    ylim=c(0,3.5), xlim=c(0.5,6.5), xlab="", ylab="", col="black", main=paste("G",g))
  abline(h=0)
  abline(v=0.5)
  mtext("Time", side=1)
  mtext("Expression", side=2)
}
```

vgeo

Geometric Variability

Description

Assume that n units are divided into k groups C1,...,Ck. The function calculates the geometrical variability for each group in data.

Usage

```
vgeo(d, pert = "onegroup")
```

Arguments

d	a distance matrix or a dist object with distance information between units.
pert	an n-vector that indicates which group each unit belongs to. Note that the expected values of pert are numbers greater than or equal to 1 (for instance 1,2,3,4..., k). The default value indicates there is only one group in data.

Value

It is a matrix containing the geometric variability for each group.

Author(s)

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Conchita Arenas <carenas@ub.edu>; Departament d'Estadística, Universitat de Barcelona, Barcelona, Spain.

References

Arenas, C. and Cuadras, C.M. (2002). Some recent statistical methods based on distances. *Contributions to Science*, **2**, 183–191.

Cuadras, C.M. (1992). Some examples of distance based discrimination. *Biometrical Letters*, **29**(1), 3–20.

See Also

[deltas](#), [proxi](#)

Examples

```
data(iris)
d <- dist(iris[,1:4])
vgeo(d,iris[,5])
```

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