

Package ‘NPLStoolbox’

June 10, 2025

Title N-Way Partial Least Squares Modelling of Multi-Way Data

Version 1.0.0

Description Creation and selection of N-way Partial Least Squares (NPLS) models. Selection of the optimal number of components can be done using `ncrossreg()`. NPLS was originally described by Rasmus Bro, see <[doi:10.1002/%28SICI%291099-128X%28199601%2910%3A1%3C47%3A%3AAID-CEM400%3E3.0.CO%3B2-C](https://doi.org/10.1002/%28SICI%291099-128X%28199601%2910%3A1%3C47%3A%3AAID-CEM400%3E3.0.CO%3B2-C)>.

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Encoding UTF-8

RoxygenNote 7.3.2

Imports dplyr, parafac4microbiome, pracma, rTensor, stats

Depends R (>= 2.10)

LazyData true

Suggests knitr, rmarkdown, testthat (>= 3.0.0)

Config/testthat/edition 3

URL <https://github.com/GRvanderPloeg/NPLStoolbox>,
<https://grvanderploeg.com/NPLStoolbox/>

BugReports <https://github.com/GRvanderPloeg/NPLStoolbox/issues>

VignetteBuilder knitr

NeedsCompilation no

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Cornejo2025	<i>Cornejo2025 longitudinal dataset measured in transgender persons</i>
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Description

The Cornejo2025 longitudinal dataset as three-dimensional arrays, with subjects in mode 1, features in mode 2 and time in mode 3.

Usage

Cornejo2025

Format

'Cornejo2025':

A list object with three elements:

Tongue_microbiome List object of the tongue longitudinal microbiota data.

Salivary_microbiome List object of the saliva longitudinal microbiota data.

Salivary_cytokines List object of the longitudinal salivary cytokine data.

Salivary_biochemistry List object of the longitudinal salivary biochemistry data.

Circulatory_hormones List object of the longitudinal circulatory hormone data.

Clinical_measurements List object of the longitudinal clinical outcome data.

Subject_metadata Matrix with subject metadata.

Source

TBD

`ncrossreg`*Cross-validation of NPLS by classical K-fold CV.*

Description

This function runs ACMTF-R with cross-validation. A deterministic K-fold partition is used: the subjects are split in order into `cvFolds` groups. For each fold the training set consists of the other folds and the test set is the current fold.

Usage

```
ncrossreg(X, y, maxNumComponents = 5, maxIter = 120, cvFolds = dim(X)[1])
```

Arguments

<code>X</code>	Centered tensor of independent data
<code>y</code>	Centered dependent variable
<code>maxNumComponents</code>	Maximum number of components to investigate (default 5).
<code>maxIter</code>	Maximum number of iterations (default 100).
<code>cvFolds</code>	Number of folds to use in the cross-validation. For example, if <code>cvFolds</code> is 5, then the subjects are deterministically partitioned into 5 groups (each CV iteration uses 4/5 for training and 1/5 for testing). Default: equal to the number of subjects (i.e. jack-knifing).

Value

A list with two elements: - **varExp**: a tibble with the variance-explained (for X and Y) per number of components. - **RMSE**: a tibble with the RMSE (computed over the unified CV prediction vector) per number of components.

Examples

```
set.seed(123)
X <- array(rnorm(25 * 5 * 4), dim = c(25, 5, 4))
y <- rnorm(25) # Random response variable
result = ncrossreg(X, y, cvFolds=2, maxNumComponents=2)
```

npred	<i>Predict Y for new data by projecting the data onto the latent space defined by an NPLS model.</i>
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Description

Predict Y for new data by projecting the data onto the latent space defined by an NPLS model.

Usage

```
npred(model, newX)
```

Arguments

model	NPLS model
newX	New data organized in a matrix of (Inew x J x K) with Inew new subjects

Value

Ypred: vector of the predicted value(s) of Y for the new data

Examples

```
Y = as.numeric(as.factor(Cornejo2025$Tongue$mode1$GenderID))
Ycnt = Y - mean(Y)
model = triPLS1(Cornejo2025$Tongue$data, Ycnt, numComponents=1)
npred(model, Cornejo2025$Tongue$data[1, ,])
```

triPLS1	<i>Tri-PLS1: three-way PLS regressed onto a y vector</i>
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Description

Tri-PLS1: three-way PLS regressed onto a y vector

Usage

```
triPLS1(X, y, numComponents, tol = 1e-10, maxIter = 100)
```

Arguments

X	Centered tensor of independent data
y	Centered dependent variable
numComponents	Number of components to fit
tol	Relative change in loss for the model to converge (default 1e-10).
maxIter	Maximum number of iterations (default 100).

Value

Model

Examples

```
set.seed(123)
X <- array(rnorm(100 * 5 * 4), dim = c(100, 5, 4)) # Random tensor (100 samples, 5 vars, 4 vars)
y <- rnorm(100) # Random response variable
model <- triPLS1(X, y, numComponents = 2)
```

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* datasets

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