

Package: nbfar (via r-universe)

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Type Package

Title Negative Binomial Factor Regression Models ('nbfar')

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Description We developed a negative binomial factor regression model to estimate structured (sparse) associations between a feature matrix X and overdispersed count data Y. With 'nbfar', microbiome count data Y can be used, for example, to associate host or environmental covariates with microbial abundances. Currently, two models are available: a) Negative Binomial reduced rank regression (NB-RRR), b) Negative Binomial co-sparse factor regression (NB-FAR). Please refer the manuscript 'Mishra, A. K., & Müller, C. L. (2021). Negative Binomial factor regression with application to microbiome data analysis. bioRxiv.' for more details.

URL <https://github.com/amishra-stats/nbfar>,
<https://www.biorxiv.org/content/10.1101/2021.11.29.470304v1>

Depends R (>= 3.5.0), stats, utils

Imports Rcpp (>= 0.12.9), MASS, magrittr, rrpck, glmnet,
RcppParallel, mpath

License GPL (>= 3.0)

Encoding UTF-8

LazyData FALSE

LinkingTo Rcpp, RcppArmadillo, RcppParallel

NeedsCompilation yes

VignetteBuilder knitr

Roxygen list(markdown = TRUE)

RoxygenNote 7.1.2

Suggests rmarkdown, knitr, spelling

Language en-US

Repository <https://amishra-stats.r-universe.dev>

RemoteUrl <https://github.com/amishra-stats/nbfar>

RemoteRef HEAD

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nbfar

Negative binomial co-sparse factor regression (NBFAR)

Description

To estimate a low-rank and sparse coefficient matrix in large/high dimensional setting, the approach extracts unit-rank components of required matrix in sequential order. The algorithm automatically stops after extracting sufficient unit rank components.

Usage

```
nbfar(
  Yt,
  X,
  maxrank = 3,
  nlambda = 40,
  cIndex = NULL,
  offset = "CSS",
  control = list(),
  nfold = 5,
  PATH = FALSE,
  nthread = 1,
  trace = FALSE,
  verbose = TRUE
)
```

Arguments

<code>Yt</code>	response matrix
<code>X</code>	design matrix; when <code>X = NULL</code> , we set <code>X</code> as identity matrix and perform generalized sparse PCA.
<code>maxrank</code>	an integer specifying the maximum possible rank of the coefficient matrix or the number of factors
<code>nlambda</code>	number of lambda values to be used along each path
<code>cIndex</code>	specify index of control variables in the design matrix <code>X</code>
<code>offset</code>	offset matrix or microbiome data analysis specific scaling: common sum scaling = CSS (default), total sum scaling = TSS, median-ratio scaling = MRS, centered-log-ratio scaling = CLR
<code>control</code>	a list of internal parameters controlling the model fitting
<code>nfold</code>	number of folds in k-fold crossvalidation
<code>PATH</code>	TRUE/FALSE for generating solution path of sequential estimate after cross-validation step
<code>nthread</code>	number of thread to be used for parallelizing the crossvalidation procedure
<code>trace</code>	TRUE/FALSE checking progress of cross validation error
<code>verbose</code>	TRUE/FALSE checking progress of estimation procedure

Value

<code>C</code>	estimated coefficient matrix; based on GIC
<code>Z</code>	estimated control variable coefficient matrix
<code>Phi</code>	estimted dispersion parameters
<code>U</code>	estimated U matrix (generalize latent factor weights)
<code>D</code>	estimated singular values
<code>V</code>	estimated V matrix (factor loadings)

References

Mishra, A., Müller, C. (2022) *Negative binomial factor regression models with application to microbiome data analysis*. <https://doi.org/10.1101/2021.11.29.470304>

Examples

```
## Load simulated data set:
data('simulate_nbfar')
attach(simulate_nbfar)

# Model with known offset
set.seed(1234)
offset <- log(10)*matrix(1,n,ncol(Y))
control_nbfar <- nbfar_control(initmaxit = 5000, gamma0 = 2, spU = 0.5,
spV = 0.6, lamMinFac = 1e-10, epsilon = 1e-5)
# nbfar_test <- nbfar(Y, X, maxrank = 5, nlambda = 20, cIndex = NULL,
# offset = offset, control = control_nbfar, nfold = 5, PATH = F)
```

<i>nbfar_control</i>	<i>Control parameters for NBFAR and NBRRR</i>
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Description

Default value for a list of control parameters that are used to estimate the parameters of negative binomial co-sparse factor regression (NBFAR) and negative binomial reduced rank regression (NBRRR).

Usage

```
nbfar_control(
  maxit = 5000,
  epsilon = 1e-07,
  elnetAlpha = 0.95,
  gamma0 = 1,
  spU = 0.5,
  spV = 0.5,
  lamMaxFac = 1,
  lamMinFac = 1e-06,
  initmaxit = 10000,
  initepsilon = 1e-08,
  objI = 0
)
```

Arguments

<code>maxit</code>	maximum iteration for each sequential steps
<code>epsilon</code>	tolerance value required for convergence of inner loop in GCURE
<code>elnetAlpha</code>	elastic net penalty parameter
<code>gamma0</code>	power parameter for generating the adaptive weights
<code>spU</code>	maximum proportion of nonzero elements in each column of U
<code>spV</code>	maximum proportion of nonzero elements in each column of V
<code>lamMaxFac</code>	a multiplier of the computed maximum value (<code>lambda_max</code>) of the tuning parameter
<code>lamMinFac</code>	a multiplier to determine <code>lambda_min</code> as a fraction of <code>lambda_max</code>
<code>initmaxit</code>	maximum iteration for minimizing the objective function while computing the initial estimates of the model parameter
<code>initepsilon</code>	tolerance value required for the convergence of the objective function while computing the initial estimates of the model parameter
<code>objI</code>	1 or 0 to indicate that the convergence will be on the basis of objective function or not

Value

a list of controlling parameter.

References

Mishra, A., Müller, C. (2022) *Negative binomial factor regression models with application to microbiome data analysis*. <https://doi.org/10.1101/2021.11.29.470304>

Examples

```
control <- nbfar_control()
```

nbfar_sim

Simulated data for testing NBFAR and NBRRR model

Description

Simulate response and covariates for multivariate negative binomial regression with a low-rank and sparse coefficient matrix. Coefficient matrix is expressed in terms of U (left singular vector), D (singular values) and V (right singular vector).

Usage

```
nbfar_sim(U, D, V, n, Xsigma, C0, disp, depth)
```

Arguments

U	specified value of U
D	specified value of D
V	specified value of V
n	sample size
Xsigma	covariance matrix used to generate predictors in X
C0	intercept value in the coefficient matrix
disp	dispersion parameter of the generative model
depth	log of the sequencing depth of the microbiome data (used as an offset in the simulated multivariate negative binomial regression model)

Value

Y	Generated response matrix
X	Generated predictor matrix

References

Mishra, A., Müller, C. (2022) *Negative binomial factor regression models with application to microbiome data analysis*. <https://doi.org/10.1101/2021.11.29.470304>

Examples

```

## Model specification:
SD <- 123
set.seed(SD)
p <- 100; n <- 200
pz <- 0
nrank <- 3           # true rank
rank.est <- 5         # estimated rank
nlam <- 20            # number of tuning parameter
s  = 0.5
q <- 30
control <- nbfar_control() # control parameters
#
#
## Generate data
D <- rep(0, nrank)
V <- matrix(0, ncol = nrank, nrow = q)
U <- matrix(0, ncol = nrank, nrow = p)
#
U[, 1] <- c(sample(c(1, -1), 8, replace = TRUE), rep(0, p - 8))
U[, 2] <- c(rep(0, 5), sample(c(1, -1), 9, replace = TRUE), rep(0, p - 14))
U[, 3] <- c(rep(0, 11), sample(c(1, -1), 9, replace = TRUE), rep(0, p - 20))
#
# for similar type response type setting
V[, 1] <- c(rep(0, 8), sample(c(1, -1), 8,
                                replace =
                                TRUE
) * runif(8, 0.3, 1), rep(0, q - 16))
V[, 2] <- c(rep(0, 20), sample(c(1, -1), 8,
                                replace =
                                TRUE
) * runif(8, 0.3, 1), rep(0, q - 28))
V[, 3] <- c(
  sample(c(1, -1), 5, replace = TRUE) * runif(5, 0.3, 1), rep(0, 23),
  sample(c(1, -1), 2, replace = TRUE) * runif(2, 0.3, 1), rep(0, q - 30)
)
U[, 1:3] <- apply(U[, 1:3], 2, function(x) x / sqrt(sum(x^2)))
V[, 1:3] <- apply(V[, 1:3], 2, function(x) x / sqrt(sum(x^2)))
#
D <- s * c(4, 6, 5) # signal strength varries as per the value of s
or <- order(D, decreasing = TRUE)
U <- U[, or]
V <- V[, or]
D <- D[or]
C <- U %*% (D * t(V)) # simulated coefficient matrix
intercept <- rep(0.5, q) # specifying intercept to the model:
C0 <- rbind(intercept, C)
#
Xsigma <- 0.5^abs(outer(1:p, 1:p, FUN = "-"))
# Simulated data
sim.sample <- nbfar_sim(U, D, V, n, Xsigma, C0, disp = 3, depth = 10) # Simulated sample
# Dispersion parameter

```

```

X <- sim.sample$X[1:n, ]
Y <- sim.sample$Y[1:n, ]
# disp = 3; depth = 10;
# simulate_nbfar <- list(Y = Y, X = X, U = U, D = D, V = V, n=n,
# Xsigma = Xsigma, C0 = C0, disp = disp, depth = depth)
# save(simulate_nbfar, file = 'data/simulate_nbfar.RData')

```

nbrrr*Negative binomial reduced rank regression (NBRRR)***Description**

In the range of 1 to maxrank, the estimation procedure selects the rank r of the coefficient matrix using a cross-validation approach. For the selected rank, a rank r coefficient matrix is estimated that best fits the observations.

Usage

```

nbrrr(
  Yt,
  X,
  maxrank = 10,
  cIndex = NULL,
  offset = "CSS",
  control = list(),
  nfold = 5,
  trace = FALSE,
  verbose = TRUE
)

```

Arguments

Yt	response matrix
X	design matrix; when X = NULL, we set X as identity matrix and perform generalized PCA.
maxrank	an integer specifying the maximum possible rank of the coefficient matrix or the number of factors
cIndex	specify index of control variable in the design matrix X
offset	offset matrix or microbiome data analysis specific scaling: common sum scaling = CSS (default), total sum scaling = TSS, median-ratio scaling = MRS, centered-log-ratio scaling = CLR
control	a list of internal parameters controlling the model fitting
nfold	number of folds in k-fold crossvalidation
trace	TRUE/FALSE checking progress of cross validation error
verbose	TRUE/FALSE checking progress of estimation procedure

Value

C	estimated coefficient matrix
Z	estimated control variable coefficient matrix
PHI	estimted dispersion parameters
U	estimated U matrix (generalize latent factor weights)
D	estimated singular values
V	estimated V matrix (factor loadings)

References

Mishra, A., Müller, C. (2022) *Negative binomial factor regression models with application to microbiome data analysis*. <https://doi.org/10.1101/2021.11.29.470304>

Examples

```
## Load simulated data set:
data('simulate_nbfar')
attach(simulate_nbfar)

# Model with known offset
set.seed(1234)
offset <- log(10)*matrix(1,n,ncol(Y))
control_nbrr <- nbfar_control(initmaxit = 5000, initepsilon = 1e-4)
# nbrrr_test <- nbrrr(Y, X, maxrank = 5, cIndex = NULL, ofset = offset,
# control = control_nbrr, nfold = 5)
```

offset_sacling	<i>Suitably generates offset matrix for the multivariate regression problem</i>
----------------	---

Description

Suitably generates offset matrix for the multivariate regression problem

Usage

```
offset_sacling(Y, ofset)
```

Arguments

Y	outcome matrix
ofset	offset matrix or microbiome data analysis specific scaling: common sum scaling = CSS (default), total sum scaling = TSS, median-ratio scaling = MRS, centered-log-ratio scaling = CLR

simulate_nbfar	<i>Simulated data for NBFAR</i>
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Description

Simulated data with low-rank and sparse coefficient matrix.

Usage

```
data(simulate_nbfar)
```

Format

A dlist of variables for the analysis using NBFAR and NBRRR:

Y Generated response matrix

X Generated predictor matrix

U specified value of U

V specified value of V

D specified value of D

n sample size

Xsigma covariance matrix used to generate predictors in X

C0 intercept value in the coefficient matrix

disp dispersion parameter of the generative model

depth log of the sequencing depth of the microbiome data (used as an offset in the simulated multivariate negative binomial regression model) Mishra, A., Müller, C. (2022) Negative binomial factor regression models with application to microbiome data analysis. <https://doi.org/10.1101/2021.11.29.470304>

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