

Package ‘MMGFM’

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Type Package

Title Multi-Study Multi-Modality Generalized Factor Model

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Description We introduce a generalized factor model designed to jointly analyze high-dimensional multi-modality data from multiple studies by extracting study-shared and specified factors. Our factor models account for heterogeneous noises and overdispersion among modality variables with augmented covariates. We propose an efficient and speedy variational estimation procedure for estimating model parameters, along with a novel criterion for selecting the optimal number of factors. More details can be referred to Liu et al. (2024) <[doi:10.48550/arXiv.2408.10542](https://doi.org/10.48550/arXiv.2408.10542)>.

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gendata_mmgfm *Generate simulated data*

Description

Generate simulated data from MMGFM models

Usage

```
gendata_mmgfm(
  seed = 1,
  nvec = c(300, 200),
  pveclist = list(gaussian = c(50, 150), poisson = c(50), binomial = c(100, 60)),
  q = 6,
  d = 3,
  qs = rep(2, length(nvec)),
  rho = rep(1, length(pveclist)),
  rho_z = 1,
  sigmavec = rep(0.5, length(pveclist)),
  n_bin = 1,
  sigma_eps = 1,
  heter_error = FALSE
)
```

Arguments

seed	a positive integer, the random seed for reproducibility of data generation process.
nvec	a vector with positive integers, specify the sample size in each study/source.
pveclist	a named list, specify the number of modalities for each type and variable dimension in each type of modality.
q	a positive integer, specify the number of study-shared factors.
d	a positive integer, specify the dimension of covariate matrix.
qs	a vector with positive integers, specify the number of study-specified factors.
rho	a numeric vector with length(pveclist) and positive elements, specify the signal strength of loading matrices for each modality type.
rho_z	a positive real, specify the signal strength of covariates.
sigmavec	a positive real vector with length(pveclist), specify the variance of study-specified and modality variable-shared factors; default as 0.5 for each element.
n_bin	a positive integer, specify the number of trails when generate Binomial modality matrix; default as 1.
sigma_eps	a positive real, the variance of overdispersion error; default as 1.
heter_error	a logical value, whether to generate the heterogeneous error; default as FALSE.

Value

return a list including the following components:

- hbeta - a M-length list composed by the estimated regression coefficient matrix for each modality;
- hA - a M-length list composed by the loading matrix corresponding to study-shared factors for each modality;
- hB - a S-length list composed by a M-length loading matrix list corresponding to study-specified factors for each study;
- hF - a S-length list composed by the posterior estimation of study-shared factor matrix for each study;
- hH - a S-length list composed by the posterior estimation of study-specified factor matrix for each study;
- hSigma - a S-length list composed by the estimated posterior variance of the study-shared factor;
- hPhi - a S-length list composed by the estimated posterior variance of study-specified factor;
- hv - a S-length list composed by a M-length vector list corresponding to the posterior estimation of study-specified and modality variable-shared factor for each study and modality;
- hzeta - the estimated posterior variance for study-specified and modality variable-shared factor;
- hsigma2 - the estimated variance for study-specified and modality variable-shared factor;
- hinvlambda - a S-length list composed by a M-length vector list corresponding to the inverse of the estimated variances of error;
- S - the approximated posterior covariance for each row of F;
- ELBO - the ELBO value when algorithm stops;
- ELBO_seq - the sequence of ELBO values.
- time_use - the running time in model fitting of SpaCOAP;

Examples

```
q <- 3; qsvec<-rep(2,3)
nvec <- c(100, 120, 100)
pveclist <- list('gaussian'=rep(150, 1), 'poisson'=rep(50, 2), 'binomial'=rep(60, 2))
datlist <- gndata_mmgfm(seed = 1, nvec = nvec, pveclist =pveclist,
                        q = q, d= 3,qs = qsvec, rho = rep(3,length(pveclist)), rho_z=0.5,
                        sigmavec=rep(0.5, length(pveclist)), sigma_eps=1)
```

MMGFM

Fit the high-dimensional multi-study multi-modality covariate-augmented generalized factor model

Description

Fit the high-dimensional multi-study multi-modality covariate-augmented generalized factor model via variational inference.

Usage

```
MMGFM(
  XList,
  ZList,
  numvarmat,
  tauList = NULL,
  q = 15,
  qsvec = rep(2, length(XList)),
  init = c("MSFRVI", "random", "LFM"),
  epsELBO = 1e-12,
  maxIter = 30,
  verbose = TRUE,
  seed = 1
)
```

Arguments

XList	a S-length list with each component a m-length list composed by a combined modality matrix of the same type modalities, which is the observed matrix from each source/study and each modality, where m is the number of modality types.
ZList	a S-length list with each component a matrix that is the covariate matrix from each study.
numvarmat	a m-by-T matrix with rownames modality types that specifies the variable number for each modality of each modality type, where m is the number of modality types, T is the maximum number of modalities for one of modality types .
tauList	an optional S-length list with each component a m-length list corresponding the offset term for each combined modality of each study; default as full-zero matrix.
q	an optional string, specify the number of study-shared factors; default as 15.
qsvec	a integer vector with length S, specify the number of study-specified factors; default as 2.
init	an optional string, specify the initialization method, supporting "MSFRVI", "random" and "LFM", default as "MSFRVI".
epsELBO	an optional positive vlaue, tolerance of relative variation rate of the evidence lower bound value, default as '1e-5'.

maxIter	the maximum iteration of the VEM algorithm. The default is 30.
verbose	a logical value, whether output the information in iteration.
seed	an optional integer, specify the random seed for reproducibility in initialization.

Details

If `init="MSFRVI"`, it will use the results from multi-study linear factor model in MultiCOAP package as initial values; If `init="LFM"`, it will use the results from linear factor model by combing data from all studies as initials.

Value

return a list including the following components:

- `hbeta` - a M-length list composed by the estimated regression coefficient matrix for each modality;
- `hA` - a M-length list composed by the loading matrix corresponding to study-shared factors for each modality;
- `hB` - a S-length list composed by a M-length loading matrix list corresponding to study-specified factors for each study;
- `hF` - a S-length list composed by the posterior estimation of study-shared factor matrix for each study;
- `hH` - a S-length list composed by the posterior estimation of study-specified factor matrix for each study;
- `hSigma` - a S-length list composed by the estimated posterior variance of the study-shared factor;
- `hPhi` - a S-length list composed by the estimated posterior variance of study-specified factor;
- `hv` - a S-length list composed by a M-length vector list corresponding to the posterior estimation of study-specified and modality variable-shared factor for each study and modality;
- `hzeta` - the estimated posterior variance for study-specified and modality variable-shared factor;
- `hsigma2` - the estimated variance for study-specified and modality variable-shared factor;
- `hinvLambda` - a S-length list composed by a M-length vector list corresponding to the inverse of the estimated variances of error;
- `S` - the approximated posterior covariance for each row of F;
- `ELBO` - the ELBO value when algorithm stops;
- `ELBO_seq` - the sequence of ELBO values.
- `time_use` - the running time in model fitting of SpaCOAP;

References

None

See Also

None

Examples

```

q <- 3; qsvec<-rep(2,3)
nvec <- c(100, 120, 100)
pveclist <- list('gaussian'=rep(150, 1),'poisson'=rep(50, 2),'binomial'=rep(60, 2))
datlist <- gendata_mmgfm(seed = 1, nvec = nvec, pveclist =pveclist,
                        q = q, d= 3,qs = qsvec, rho = rep(3,length(pveclist)), rho_z=0.5,
                        sigmavec=rep(0.5, length(pveclist)), sigma_eps=1)

XList <- datlist$XList
ZList <- datlist$ZList
numvarmat <- datlist$numvarmat
### For illustration, we set maxIter=3. Set maxIter=50 when running formally
reslist1 <- MMGFM(XList, ZList=ZList, numvarmat, q=q, qsvec = qsvec, init='MSFRVI',maxIter = 3)
str(reslist1)

```

selectFac.MMGFM	<i>Select the number of study-shared and study-specified factors for MMGFM</i>
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Description

Select the number of study-shared and study-specified factors for the high-dimensional multi-study multi-modality covariate-augmented generalized factor model.

Usage

```

selectFac.MMGFM(
  XList,
  ZList,
  numvarmat,
  q.max = 15,
  qsvec.max = rep(4, length(XList)),
  threshold.vec = c(0.01, 0.001),
  tauList = NULL,
  init = c("MSFRVI", "random", "LFM"),
  epsELBO = 1e-12,
  maxIter = 30,
  verbose = TRUE,
  seed = 1
)

```

Arguments

XList	a S-length list with each component a m-length list composed by a combined modality matrix of the same type modalities, which is the observed matrix from each source/study and each modality, where m is the number of modality types.
ZList	a S-length list with each component a matrix that is the covariate matrix from each study.

numvarmat	a m-by-T matrix with rownames modality types that specifies the variable number for each modality of each modality type, where m is the number of modality types, T is the maximum number of modalities for one of modality types .
q.max	an optional integer, specify the upper bound for the number of study-shared factors; default as 15.
qsvec.max	an optional integer vector with length S, specify the upper bound for the number of study-specified factors; default as 4 for each study.
threshold.vec	an optional real vector with length 2, specify the threshold for the singular values of study-shared loading and study-specified loading matrices, respectively.
tauList	an optional S-length list with each component a m-length list corresponding the offset term for each combined modality of each study; default as full-zero matrix.
init	an optional string, specify the initialization method, supporting "MSFRVI", "random" and "LFM", default as "MSFRVI".
epsELBO	an optional positive vlaue, tolerance of relative variation rate of the evidence lower bound value, default as '1e-5'.
maxIter	the maximum iteration of the VEM algorithm. The default is 30.
verbose	a logical value, whether output the information in iteration.
seed	an optional integer, specify the random seed for reproducibility in initialization.

Value

return a list with two components: q and qs.vec.

Examples

```

q <- 3; qsvec<-rep(2,3)
nvec <- c(100, 120, 100)
pveclist <- list('gaussian'=rep(150, 1),'poisson'=rep(50, 2),'binomial'=rep(60, 2))
datlist <- gendata_mmgfm(seed = 1, nvec = nvec, pveclist =pveclist,
                        q = q, d= 3,qs = qsvec, rho = rep(3,length(pveclist)), rho_z=0.5,
                        sigmavec=rep(0.5, length(pveclist)), sigma_eps=1)
XList <- datlist$XList
ZList <- datlist$ZList
numvarmat <- datlist$numvarmat
### For illustration, we set maxIter=3. Set maxIter=50 when running formally
selectFac.MMGFM(XList, ZList=ZList, numvarmat, q.max=6, qsvec.max = rep(4,3),
init='MSFRVI',maxIter = 3)

```

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