

Package ‘HIMA’

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Type Package

Title High-Dimensional Mediation Analysis

Version 1.1.0

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Description Allows to estimate and test high-dimensional mediation effects based on advanced mediator screening and penalized regression techniques. Methods used in the package refer to Haixiang Zhang, Yinan Zheng, Zhou Zhang, Tao Gao, Brian Joyce, Grace Yoon, Wei Zhang, Joel Schwartz, Alan Just, Elena Colicino, Pantel Vokonas, Lihui Zhao, Jinchi Lv, Andrea Baccarelli, Lifang Hou, Lei Liu (2016) <doi:10.1093/bioinformatics/btw351>.

License GPL-3

Depends R (>= 3.4.0), ncvreg, glmnet

Imports utils, stats, MASS, survival, HDMT, iterators, parallel,
foreach, doParallel

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URL <https://github.com/YinanZheng/HIMA/>

BugReports <https://github.com/YinanZheng/HIMA/issues/>

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R topics documented:

HIMA-package	2
hima	2
simHIMA	5
survHIMA	6

Index

9

HIMA-package	<i>High-dimensional Mediation Analysis</i>
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Description

HIMA is an R package for estimating and testing high-dimensional mediation effects in omic studies.

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References

1. Zhang H, Zheng Y, Zhang Z, Gao T, Joyce B, Yoon G, Zhang W, Schwartz J, Just A, Colicino E, Vokonas P, Zhao L, Lv J, Baccarelli A, Hou L, Liu L. Estimating and Testing High-dimensional Mediation Effects in Epigenetic Studies. Bioinformatics. 2016. DOI: 10.1093/bioinformatics/btw351. PubMed PMID: 27357171.
2. Zhang H, Zheng Y, Hou L, Liu L. Mediation Analysis for Survival Data with High-Dimensional Mediators. Bioinformatics. 2021 (under review).

hima	<i>High-dimensional Mediation Analysis</i>
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Description

hima is used to estimate and test high-dimensional mediation effects.

Usage

```
hima(
  X,
  Y,
  M,
  COV.XM = NULL,
  COV.MY = COV.XM,
  family = c("gaussian", "binomial"),
  penalty = c("MCP", "SCAD", "lasso"),
  topN = NULL,
  parallel = FALSE,
  ncore = 1,
  verbose = FALSE,
  ...
)
```

Arguments

X	a vector of exposure.
Y	a vector of outcome. Can be either continuous or binary (0-1).
M	a <code>data.frame</code> or <code>matrix</code> of high-dimensional mediators. Rows represent samples, columns represent variables.
COV.XM	a <code>data.frame</code> or <code>matrix</code> of covariates dataset for testing the association $M \sim X$. Covariates specified here will not participate penalization. Default = <code>NULL</code> . If the covariates contain mixed data types, please make sure all categorical variables are properly formatted as <code>factor</code> type.
COV.MY	a <code>data.frame</code> or <code>matrix</code> of covariates dataset for testing the association $Y \sim M$. Covariates specified here will not participate penalization. If not specified, the same set of covariates for $M \sim X$ will be applied. Using different sets of covariates is allowed but this needs to be handled carefully.
family	either ' <code>gaussian</code> ' or ' <code>binomial</code> ', depending on the data type of outcome (Y). See ncvreg
penalty	the penalty to be applied to the model. Either ' <code>MCP</code> ' (the default), ' <code>SCAD</code> ', or ' <code>lasso</code> '. See ncvreg .
topN	an integer specifying the number of top markers from sure independent screening. Default = <code>NULL</code> . If <code>NULL</code> , <code>topN</code> will be either <code>ceiling(n/log(n))</code> if <code>family = 'gaussian'</code> , or <code>ceiling(n/(2*log(n)))</code> if <code>family = 'binomial'</code> , where <code>n</code> is the sample size. If the sample size is greater than <code>topN</code> (pre-specified or calculated), all mediators will be included in the test (i.e. low-dimensional scenario).
parallel	logical. Enable parallel computing feature? Default = <code>TRUE</code> .
ncore	number of cores to run parallel computing Valid when <code>parallel == TRUE</code> . By default max number of cores available in the machine will be utilized.
verbose	logical. Should the function be verbose? Default = <code>FALSE</code> .
...	other arguments passed to ncvreg .

Value

A data.frame containing mediation testing results of selected mediators.

- alpha: coefficient estimates of exposure (X) → mediators (M).
- beta: coefficient estimates of mediators (M) → outcome (Y) (adjusted for exposure).
- gamma: coefficient estimates of exposure (X) → outcome (Y) (total effect).
- alpha*beta: mediation effect.
- % total effect: alpha*beta / gamma. Percentage of the mediation effect out of the total effect.
- Bonferroni.p: statistical significance of the mediator (Bonferroni procedure).
- BH.FDR: statistical significance of the mediator (Benjamini-Hochberg procedure).

References

Zhang H, Zheng Y, Zhang Z, Gao T, Joyce B, Yoon G, Zhang W, Schwartz J, Just A, Colicino E, Vokonas P, Zhao L, Lv J, Baccarelli A, Hou L, Liu L. Estimating and Testing High-dimensional Mediation Effects in Epigenetic Studies. Bioinformatics. 2016. DOI: 10.1093/bioinformatics/btw351. PubMed PMID: 27357171.

Examples

```
n <- 200 # sample size
p <- 200 # the dimension of covariates

# the regression coefficients alpha (exposure --> mediators)
alpha <- rep(0, p)

# the regression coefficients beta (mediators --> outcome)
beta1 <- rep(0, p) # for continuous outcome
beta2 <- rep(0, p) # for binary outcome

# the first four markers are true mediators
alpha[1:4] <- c(0.45,0.5,0.6,0.7)
beta1[1:4] <- c(0.55,0.6,0.65,0.7)
beta2[1:4] <- c(1.45,1.5,1.55,1.6)

# these are not true mediators
alpha[7:8] <- 0.5
beta1[5:6] <- 0.8
beta2[5:6] <- 1.7

# Generate simulation data
simdat_cont = simHIMA(n, p, alpha, beta1, seed=1029)
simdat_bin = simHIMA(n, p, alpha, beta2, binaryOutcome = TRUE, seed=1029)

# Run HIMA with MCP penalty by default
# When Y is continuous (default)
hima.fit <- hima(simdat_cont$X, simdat_cont$Y, simdat_cont$M, verbose = TRUE)
hima.fit
```

```
# When Y is binary (should specify family)
hima.logistic.fit <- hima(simdat_bin$X, simdat_bin$Y, simdat_bin$M,
family = "binomial", verbose = TRUE)
hima.logistic.fit
```

Description

simHIMA is used to generate simulation data for high-dimensional mediation analysis.

Usage

```
simHIMA(n, p, alpha, beta, binaryOutcome = FALSE, seed)
```

Arguments

n	an integer specifying sample size.
p	an integer specifying the dimension of mediators.
alpha	a numeric vector specifying the regression coefficients alpha (exposure → mediators).
beta	a numeric vector specifying the regression coefficients beta (mediators → outcome).
binaryOutcome	logical. Should the simulated outcome variable be binary?
seed	an integer specifying a seed for random number generation.

See Also

see [hima](#) to run HIMA.

Examples

```
n <- 200 # sample size
p <- 200 # the dimension of covariates

# the regression coefficients alpha (exposure --> mediators)
alpha <- rep(0, p)

# the regression coefficients beta (mediators --> outcome)
beta <- rep(0, p)

# the first four markers are true mediators.
alpha[1:4] <- c(0.45, 0.5, 0.55, 0.6)
beta[1:4] <- c(0.5, 0.45, 0.4, 0.35)

alpha[7:8] <- 0.5
```

```

beta[5:6] <- 0.5

# Generate simulation data
simdat = simHIMA(n, p, alpha, beta, seed=1029)

```

survHIMA

High-dimensional mediation analysis for survival data

Description

survHIMA is used to estimate and test high-dimensional mediation effects for survival data.

Usage

```
survHIMA(X, Z, M, OT, status, FDRcut = 0.05, verbose = FALSE)
```

Arguments

X	a vector of exposure.
Z	a matrix of adjusting covariates. Rows represent samples, columns represent variables. Can be NULL.
M	a data.frame or matrix of high-dimensional mediators. Rows represent samples, columns represent variables.
OT	a vector of observed failure times.
status	a vector of censoring indicator (status = 1: uncensored; status = 0: censored)
FDRcut	FDR cutoff applied to define and select significant mediators. Default = 0.05.
verbose	logical. Should the function be verbose? Default = FALSE.

Value

A data.frame containing mediation testing results of selected mediators (FDR < 0.05).

- ID: index of selected significant mediator.
- alpha: coefficient estimates of exposure (X) → mediators (M).
- alpha_se: standard error for alpha.
- beta: coefficient estimates of mediators (M) → outcome (Y) (adjusted for exposure).
- beta_se: standard error for beta
- p_joint: joint p-value of selected significant mediator.

References

Zhang H, Zheng Y, Hou L, Liu L. Mediation Analysis for Survival Data with High-Dimensional Mediators. *Bioinformatics*. 2021 (under review).

Examples

```

## Generate simulated survival data
set.seed(100)
n <- 100 # sample size
p <- 100 # the dimension of mediators
q <- 1    # the dimension of covariate(s)

sigma_e <- matrix(0.25, p, p)
diag(sigma_e) <- 1
sigma_e[1, 3] <- 0.8
sigma_e[3, 1] <- 0.8
sigma_e[2, 4] <- 0.8
sigma_e[4, 2] <- 0.8

##
beta <- matrix(0, 1, p)
beta[1:5] <- c(0.6, -0.5, 0.4, -0.3, 0.25)

##
alpha <- matrix(0, 1, p)
alpha[1:5] <- c(0.6, -0.5, 0.4, -0.3, 0.25)

##
gamma <- matrix(0.5, 1, q)
eta <- matrix(0.3, p, q)
r <- matrix(0.5, 1, 1)

##
X <- matrix(rnorm(n, mean = 0, sd = 2), n, 1) # exposure
Z <- matrix(rnorm(n * q, mean = 0, sd = 2), n, q) # covariates
mu <- matrix(0, p, 1)
e <- MASS::mvrnorm(n, mu, sigma_e) # the error terms

M <- X%*%(alpha) + Z%*%t(eta) + e
MZ <- cbind(M, Z, X)

beta_gamma <- cbind(beta, gamma, r)

## generate the failure time T
u <- runif(n, 0, 1)
T <- matrix(0, n, 1)
for (i in 1:n)
  T[i] <- -log(1 - u[i])*exp(-sum(beta_gamma*MZ[i,]))

## generate censoring time 0.45 censoring rate
C <- runif(n, min = 0, max = 150)
status <- as.integer(T < C)

## the observed failure time
OT <- apply(cbind(C, T), 1, min)

## Not run:

```

```
survHIMA.fit <- survHIMA(X, Z, M, OT, status)
survHIMA.fit

## End(Not run)
```

Index

* package

HIMA-package, [2](#)

HIMA (HIMA-package), [2](#)

hima, [2](#), [5](#)

HIMA-package, [2](#)

ncvreg, [3](#)

simHIMA, [5](#)

survHIMA, [6](#)