

Package ‘BayesSUR’

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Type Package

Title Bayesian Seemingly Unrelated Regression

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Description Bayesian seemingly unrelated regression with general variable selection and dense/sparse covariance matrix. The sparse seemingly unrelated regression is described in Banterle et al. (2018) <doi:10.1101/467019>.

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BayesSUR

Fitting BayesSUR models

Description

Main function of the package. Fits a range of models introduced in the package vignette BayesSUR.pdf. Returns an object of S3 class BayesSUR. There are three options for the prior on the residual covariance matrix (i.e., independent inverse-Gamma, inverse-Wishart and hyper-inverse Wishart) and three options for the prior on the latent indicator variable (i.e., independent Bernoulli, hotspot and Markov random field). So there are nine models in total. See details for their combinations.

Usage

```
BayesSUR(
  data = NULL,
  Y,
  X,
  X_0 = NULL,
```

```

covariancePrior = "HIW",
gammaPrior = "hotspot",
nIter = 10000,
burnin = 5000,
nChains = 2,
outFilePath = "",
gammaSampler = "bandit",
gammaInit = "R",
mrfG = NULL,
standardize = TRUE,
standardize.response = TRUE,
maxThreads = 1,
output_gamma = TRUE,
output_beta = TRUE,
output_Gy = TRUE,
output_sigmaRho = TRUE,
output_pi = TRUE,
output_tail = TRUE,
output_model_size = TRUE,
output_model_visit = FALSE,
output_CPO = FALSE,
output_Y = TRUE,
output_X = TRUE,
hyperpar = list(),
tmpFolder = "tmp/"
)

```

Arguments

data	a numeric matrix with variables on the columns and observations on the rows, if arguments Y and X (and possibly X_0) are vectors. Can be NULL if arguments Y and X (and possibly X_0) are numeric matrices.
Y, X	vectors of indices (with respect to the data matrix) for the outcomes (Y) and the predictors to select (X) respectively; if the data argument is not provided, these needs to be numeric matrices containing the data instead, with variables on the columns and observations on the rows.
X_0	vectors of indices (with respect to the data matrix) for the fixed predictors that are not selected, i.e. always included in the model; if the data argument is not provided, this needs to be a numeric matrix containing the data instead, with variables on the columns and observations on the rows.
covariancePrior	string indicating the prior for the covariance Σ ; it has to be either HIW for the hyper-inverse-Wishart (which will result in a sparse covariance matrix), IW for the inverse-Wishart prior (dense covariance) or IG for independent inverse-Gamma on all the diagonal elements and 0 otherwise. See the details for the model specification
gammaPrior	string indicating the gamma prior to use, either hotspot (default) for the Hotspot prior of Bottolo (2011), MRF for the Markov Random Field prior or hierarchical

	for a simpler hierarchical prior. See the details for the model specification
nIter	number of iterations for the MCMC procedure. Default 10000.
burnin	number of iterations to discard at the start of the chain. Default is 5000.
nChains	number of parallel tempered chains to run (default 2). The temperature is adapted during the burnin phase.
outFilePath	path to where the output files are to be written. The default path is the current working directory.
gammaSampler	string indicating the type of sampler for gamma, either <code>bandit</code> for the Thompson sampling inspired sampler or <code>MC3</code> for the usual MC^3 sampler. See Russo et al.(2018) or Madigan and York (1995) for details.
gammaInit	gamma initialisation to either all-zeros (\emptyset), all ones (1), MLE-informed (MLE) or (default) randomly (R).
mrfG	either a matrix or a path to the file containing the G matrix for the MRF prior on gamma (if necessary)
standardize	logical flag for X variable standardization. Default is <code>standardize=TRUE</code> . The coefficients are returned on the standardized scale.
standardize.response	Standardization for the response variables. Default is <code>standardize.response=TRUE</code> .
maxThreads	maximum threads used for parallelization. Default is 1. Reproducibility of results with <code>set.seed()</code> is only guaranteed if <code>maxThreads=1</code> .
output_gamma	allow (TRUE) or suppress (FALSE) the output for gamma. See the return value below for more information.
output_beta	allow (TRUE) or suppress (FALSE) the output for beta. See the return value below for more information.
output_Gy	allow (TRUE) or suppress (FALSE) the output for Gy. See the return value below for more information.
output_sigmaRho	allow (TRUE) or suppress (FALSE) the output for sigmaRho. See the return value below for more information.
output_pi	allow (TRUE) or suppress (FALSE) the output for pi. See the return value below for more information.
output_tail	allow (TRUE) or suppress (FALSE) the output for tail (hotspot tail probability). See the return value below for more information.
output_model_size	allow (TRUE) or suppress (FALSE) the output for model_size. See the return value below for more information.
output_model_visit	allow (TRUE) or suppress (FALSE) the output for all visited models over the MCMC iterations. Default is FALSE. See the return value below for more information.
output_CPO	allow (TRUE) or suppress (FALSE) the output for (scaled) conditional predictive ordinates (<code>*_CPO_out.txt</code>), CPO with joint posterior predictive of the response variables (<code>*_CPOsumy_out.txt</code>) and widely applicable information criterion (<code>*_WAIC_out.txt</code>). See the return value below for more information.

output_Y	allow (TRUE) or suppress (FALSE) the output for responses dataset Y.
output_X	allow (TRUE) or suppress (FALSE) the output for predictors dataset X.
hyperpar	a list of named hyperparameters to use instead of the default values. Valid names are mrf_d, mrf_e, a_sigma, b_sigma, a_tau, b_tau, nu, a_eta, b_eta, a_o, b_o, a_pi, b_pi, a_w and b_w. Their default values are a_w=2, b_w=5, a_omega=2, b_omega=1, a_o=2, b_o=p-2, a_pi=2, b_pi=1, nu=s+2, a_tau=0.1, b_tau=10, a_eta=0.1, b_eta=1, a_sigma=1, b_sigma=1, mrf_d=-3 and mrf_e=0.03. See the vignette for more information.
tmpFolder	the path to a temporary folder where intermediate data files are stored (will be erased at the end of the chain) default to local tmpFolder

Details

The arguments covariancePrior and gammaPrior specify the model HRR, dSUR or SSUR with different gamma prior. Let γ_{jk} be latent indicator variable of each coefficient and C be covariance matrix of response variables. The nine models specified through the arguments covariancePrior and gammaPrior are as follows.

	$\gamma_{jk} \sim \text{Bernoulli}$	$\gamma_{jk} \sim \text{hotspot}$	$\gamma_{jk} \sim \text{MRF}$
$C \sim \text{indep}$	HRR-B	HRR-H	HRR-M
$C \sim \text{IW}$	dSUR-B	dSUR-H	dSUR-M
$C \sim \text{HIW}$	SSUR-B	SSUR-H	SSUR-M

Value

An object of class BayesSUR is saved as obj_BayesSUR.RData in the output file, including the following components:

- status - the running status
- input - a list of all input parameters by the user
- output - a list of the all output filenames:
 - `"*_logP_out.txt"` - contains each row for the 1000t-th iteration's log-likelihoods of parameters, i.e., Tau, Eta, JunctionTree, SigmaRho, O, Pi, Gamma, W, Beta and data conditional log-likelihood depending on the models.
 - `"*_gamma_out.txt"` - posterior mean of the latent indicator matrix.
 - `"*_pi_out.txt"` - posterior mean of the predictor effects (prospensity) by decomposing the probability of the latent indicator.
 - `"*_hotspot_tail_p_out.txt"` - posterior mean of the hotspot tail probability. Only available for the hotspot prior on the gamma.
 - `"*_beta_out.txt"` - posterior mean of the coefficients matrix.
 - `"*_Gy_out.txt"` - posterior mean of the response graph. Only available for the HIW prior on the covariance.
 - `"*_sigmaRho_out.txt"` - posterior mean of the transformed parameters. Not available for the IG prior on the covariance.
 - `"*_model_size_out.txt"` - contains each row for the 1000t-th iteration's model sizes of the multiple response variables.

- `"*_model_visit_gy_out.txt"` - contains each row for the nonzero indices of the vectorized estimated graph matrix for each iteration.
 - `"*_model_visit_gamma_out.txt"` - contains each row for the nonzero indices of the vectorized estimated gamma matrix for each iteration.
 - `"*_CPO_out.txt"` - the (scaled) conditional predictive ordinates (CPO).
 - `"*_CPOsumy_out.txt"` - the (scaled) conditional predictive ordinates (CPO) with joint posterior predictive of the response variables.
 - `"*_WAIC_out.txt"` - the widely applicable information criterion (WAIC).
 - `"*_Y.txt"` - responses dataset.
 - `"*_X.txt"` - predictors dataset.
 - `"*_X0.txt"` - fixed predictors dataset.
- `call` - the matched call.

References

Russo D, Van Roy B, Kazerouni A, Osband I, Wen Z (2018). *A tutorial on Thompson sampling*. Foundations and Trends in Machine Learning, 11: 1-96.

Madigan D, York J (1995). *Bayesian graphical models for discrete data*. International Statistical Review, 63: 215–232.

Banterle M, Bottolo L, Richardson S, Ala-Korpela M, Jarvelin MR, Lewin A (2018). *Sparse variable and covariance selection for high-dimensional seemingly unrelated Bayesian regression*. bioRxiv: 467019.

Banterle M#, Zhao Z#, Bottolo L, Richardson S, Lewin A, Zucknick M (2019). *BayesSUR: An R package for high-dimensional multivariate Bayesian variable and covariance selection in linear regression*. URL: <https://cran.r-project.org/web/packages/BayesSUR/vignettes/BayesSUR.pdf>

Examples

```
data("exampleEQTL", package = "BayesSUR")
hyperpar <- list( a_w = 2 , b_w = 5 )
set.seed(9173)
fit <- BayesSUR(Y = exampleEQTL[["blockList"]][[1]],
               X = exampleEQTL[["blockList"]][[2]],
               data = exampleEQTL[["data"]], outFilePath = tempdir(),
               nIter = 100, burnin = 50, nChains = 2, gammaPrior = "hotspot",
               hyperpar = hyperpar, tmpFolder = "tmp/", output_CPO=TRUE)

## check output
# show the summary information
summary(fit)

# show the estimated beta, gamma and graph of responses Gy
plot(fit, estimator=c("beta","gamma","Gy"), type="heatmap")

## Not run:
#Set up temporary work directory for saving a pdf figure
td <- tempdir()
oldwd <- getwd()
```

```

setwd(td)

# Produce authentic math formulas in the graph
plot(fit, estimator=c("beta","gamma","Gy"), type="heatmap", fig.tex = TRUE)
system(paste(getOption("pdfviewer"), "ParamEstimator.pdf"))
setwd(oldwd)

## End(Not run)

```

BayesSUR_internal	<i>BayesSUR_internal</i>
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Description

Run a SUR Bayesian sampler – internal function

Arguments

dataFile	path to data file
outFilePath	path to where the output is to be written
nIter	number of iterations
nChains	number of parallel chains to run
	NOTE THAT THIS IS BASICALLY JUST A WRAPPER

coef.BayesSUR	<i>coef method for class BayesSUR</i>
---------------	---------------------------------------

Description

Extract the posterior mean of the coefficients of a BayesSUR class object

Usage

```

## S3 method for class 'BayesSUR'
coef(object, beta.type = "marginal", Pmax = 0, ...)

```

Arguments

object	an object of class BayesSUR
beta.type	type of output beta. Default is marginal, giving marginal beta estimation. If beta.type="conditional", it gives conditional beta estimation
Pmax	valid if beta.type="conditional", giving estimated beta conditioning on gammas>=Pmax. If beta.type="conditional" and Pmax=0.5, it gives median probability model betas. Default is 0.
...	other arguments

Value

Estimated coefficients are from an object of class BayesSUR. If the BayesSUR specified data standardization, the fitted values are base based on standardized data.

Examples

```
data("exampleQTL", package = "BayesSUR")
hyperpar <- list( a_w = 2 , b_w = 5 )

set.seed(9173)
fit <- BayesSUR(Y = exampleEQTL[["blockList"]][[1]],
               X = exampleEQTL[["blockList"]][[2]],
               data = exampleEQTL[["data"]], outFilePath = tempdir(),
               nIter = 100, burnin = 50, nChains = 2, gammaPrior = "hotspot",
               hyperpar = hyperpar, tmpFolder = "tmp/" )

## check prediction
beta.hat <- coef(fit)
```

 elpd

expected log pointwise predictive density

Description

Measure the prediction accuracy by the elpd (expected log pointwise predictive density). The out-of-sample predictive fit can either be estimated by Bayesian leave-one-out cross-validation (LOO) or by widely applicable information criterion (WAIC) (Vehtari et al. 2017).

Usage

```
elpd(object, method = "LOO")
```

Arguments

object	an object of class BayesSUR
method	the name of the prediction accuracy index. Default is the "LOO" (Bayesian LOO estimate of out-of-sample predictive fit). The other index is the "WAIC" (widely applicable information criterion). For the HRR models, both "LOO" and "WAIC" are computed based on the multivariate t-distribution of the posterior predictive rather than approximation of importance sampling.

Value

Return the prediction accuracy measure from an object of class BayesSUR. It is elpd.loo if the argument method="LOO" and elpd.WAIC if method="WAIC".

References

Vehtari, A., Gelman, A., Gabry, J. (2017). *Practical Bayesian model evaluation using leave-one-out cross-validation and WAIC*. *Statistics and Computing*, 27(5): 1413–1432.

Examples

```
data("exampleEQTL", package = "BayesSUR")
hyperpar = list( a_w = 2 , b_w = 5 )

set.seed(9173)
fit <- BayesSUR(Y = exampleEQTL[["blockList"]][[1]],
               X = exampleEQTL[["blockList"]][[2]],
               data = exampleEQTL[["data"]], outFilePath = tempdir(),
               nIter = 100, burnin = 50, nChains = 2, gammaPrior = "hotspot",
               hyperpar = hyperpar, tmpFolder = "tmp/", output_CPO=TRUE)

## check output
# print the prediction accuracy elpd (expected log pointwise predictive density)
# by the Bayesian LOO estimate of out-of-sample predictive fit
elpd(fit, method="LOO")
```

exampleEQTL

Simulated data set to mimic a small expression quantitative trait loci (eQTL) example

Description

Simulated data set to mimic a small expression quantitative trait loci (eQTL) example, with $p=150$ single nucleotide polymorphisms (SNPs) as explanatory variables, $s=10$ gene expression features as response variables and data for $n=100$ observations. Loading the data will load the associated `blockList` object needed to fit the model with `BayesSUR()`. The R code for generating the simulated data is given in the Examples paragraph.

```
#importFrom BDgraph rgwish #importFrom gRbase mcsMAT #importFrom scime simulateSNPs
```

Usage

```
exampleEQTL
```

Format

An object of class `list` of length 4.

Examples

```

# Load the eQTL sample dataset
data("exampleEQTL", package = "BayesSUR")
str(exampleEQTL)

## Not run:
#=====
# The code below is to show how to generate the dataset "exampleEQTL.rda" above
#=====

requireNamespace("BDgraph", quietly = TRUE)
requireNamespace("gRbase", quietly = TRUE)
requireNamespace("scrim", quietly = TRUE)

##### Problem Dimensions
n = 100
p = 150
s = 10

##### Select a set of n x p (SNPs) covariates

## The synthetic data in the paper use a subset of the real SNPs as covariates,
# but as the NFBC66 dataset is confidential we'll use scrim to sample similar data

x = scrim::simulateSNPs(c(n,10), p, c(3, 2),prop.explain = c(0.9, 0.95))$data[1:n,]
x = cbind(rep(1,n),x)

#####

graph_pattern = 2

snr = 25

corr_param = 0.9

### Create the underlying graph
if(graph_pattern==1){
  ### 1) Random but full
  G = matrix(1,s,s)
  Prime = list(c(1:s))
  Res = Prime
  Sep = list()
}else if(graph_pattern==2){

  ### 2) Block Diagonal structure
  Prime = list(c(1:floor(s*2/3)),
              c((floor(s*2/3)+1):(ceiling(s*4/5)-1)),
              c(ceiling(s*4/5):s))

  Res = Prime
  Sep = lapply(Res,function(x) which(x==99))

```

```

G = matrix(0,s,s)
for(i in Prime){
  G[i,i] = 1
}

}else if(graph_pattern==3){

### 3) Decomposable model
Prime = list(c(1:floor(s*5/12),ceiling(s*9/10):s),
             c(floor(s*2/9):(ceiling(s*2/3)-1)),
             c(ceiling(s*2/3):(ceiling(s*4/5)-1)),
             c(ceiling(s*4/5):s))

Sep = list(); H=list()
for( i in 2:length(Prime)){
  H = union(H,Prime[[i-1]])
  Sep[[i-1]] = intersect( H,Prime[[i]])
}

Res = list()
Res[[1]] = Prime[[1]]
for( i in 2:length(Prime)){
  Res[[i]] = setdiff( Prime[[i]],Sep[[i-1]])
}

G = matrix(0,s,s)
for(i in Prime)
  G[i,i] = 1

## decomp check
dimnames(G) = list(1:s,1:s)
length( gRbase::mcsMAT(G - diag(s)) ) > 0

}else if(graph_pattern==4){

### 4) Non-decomposable model
nblocks = 5
nElemPerBlock =c(floor(s/4),floor(s/2)-1-floor(s/4),
                 ceiling(s*2/3)-1-floor(s/2),7)
nElemPerBlock = c(nElemPerBlock , s-sum(nElemPerBlock))
res = 1:s; blockIdx = list()
for(i in 1:nblocks){
  # blockIdx[[i]] = sample(res,nElemPerBlock[i])
  blockIdx[[i]] = res[1:nElemPerBlock[i]]
  res = setdiff(res,blockIdx[[i]])
}

G = matrix(0,s,s)
## add diagonal
for(i in 1:nblocks)
  G[blockIdx[[i]],blockIdx[[i]]] = 1
## add cycle

```

```

G[blockIdx[[1]],blockIdx[[2]]] = 1 ; G[blockIdx[[2]],blockIdx[[1]]] = 1
G[blockIdx[[1]],blockIdx[[5]]] = 1 ; G[blockIdx[[5]],blockIdx[[1]]] = 1
G[blockIdx[[2]],blockIdx[[3]]] = 1 ; G[blockIdx[[3]],blockIdx[[2]]] = 1
G[blockIdx[[3]],blockIdx[[5]]] = 1 ; G[blockIdx[[5]],blockIdx[[3]]] = 1

## decomp check
dimnames(G) = list(1:s,1:s)
length( gRbase::mcsMAT(G -diag(s) ) ) > 0

# Prime = blockIdx
Res = blockIdx ## this is not correct but not used in the non-decomp case

}

### Gamma Pattern
gamma = matrix(0,p+1,s)
gamma[1,] = 1

### 2) Extra Patterns

## outcomes (correlated in the decomp model) have some predictors in common
gamma[6:10,6:9] = 1

## outcomes (correlated in the decomp model) have some predictors in common
#gamma[16:20,14:15] = 1

## outcomes (sort-of correlated [pair-wise] in the decomp model)
# have predictors in common 6:15
gamma[26:30,4:8] = 1

## outcomes (NOT correlated in the decomp model) have predictors in common 16:17
gamma[36:40,c(3:5,9:10)] = 1

## these predictors are associated with ALL the outcomes
gamma[46:50,] = 1

combn11 = combn(rep((6:9-1)*p,each=length(6:10-1)) + rep(6:10-1,times=length(6:9)), 2)
combn31 = combn(rep((4:8-1)*p,each=length(26:30-1)) + rep(26:30-1,times=length(4:8)), 2)
combn32 = combn(rep((4:8-1)*p,each=length(46:50-1)) + rep(46:50-1,times=length(4:8)), 2)
combn41 = combn(rep((3:5-1)*p,each=length(36:40-1)) + rep(36:40-1,times=length(3:5)), 2)
combn42 = combn(rep((3:5-1)*p,each=length(46:50-1)) + rep(46:50-1,times=length(3:5)), 2)
combn51 = combn(rep((9:10-1)*p,each=length(36:40-1)) + rep(36:40-1,times=length(9:10)), 2)
combn52 = combn(rep((9:10-1)*p,each=length(46:50-1)) + rep(46:50-1,times=length(9:10)), 2)

Gmrf = rbind(t(combn11), t(combn31), t(combn32), t(combn41), t(combn42), t(combn51), t(combn52))

## get for every correlated bunch in the decomposable model,

if(graph_pattern<4){
  # a different set of predictors
  for(i in 1:length(Prime))
    gamma[6:10 + (i+6) * 10, Prime[[i]]] = 1 ## for each Prime component

```

```

## for every Residual instead
for(i in 1:length(Res))
  gamma[6:10 + (i+10) * 10, Res[[i]]] = 1

}else{

  for(i in 1:length(Prime))
    gamma[6:10 + (i+4) * 10, Prime[[i]]] = 1  ## for each Prime component

  ## for every Residual instead
  for(i in 1:length(Res))
    gamma[6:10 + (i+9) * 10, Res[[i]]] = 1
}

#### Sample the betas
sd_b = 1
b = matrix(rnorm((p+1)*s,0,sd_b),p+1,s)

xb = matrix(NA,n,s)

for(i in 1:s){
  if(sum(gamma[,i])>1){
    xb[,i] = x[,gamma[,i]==1] %*% b[gamma[,i]==1,i]
  }else{
    xb[,i] = rep(1,n) * b[1,i]
  }
}

##Sample the variance
v_r = mean(diag(var(xb))) / snr

nu = s+1

M = matrix(corr_param,s,s)
diag(M) = rep(1,s)

P = BDgraph::rgwish(n=1,adj=G,b=3,D=v_r*M)

var = solve(P)

factor = 10 ; factor_min = 0.01; factor_max = 1000
count = 0 ; maxit = 10000

factor_prev = 1

repeat{

  var = var / factor * factor_prev

  ### Sample the errors and the Ys
  cVar = chol(as.matrix(var))
  #err = matrix(rnorm(n*s),n,s) %*% cVar

```

```

err = matrix(rnorm(n*s,sd=0.5),n,s) %%% cVar
y = xb+err

## Reparametrisation ( assuming PEO is 1:s )
cVar = t(cVar) # make it lower-tri
S = diag(diag(cVar))
sigma = S*S
L = cVar %%% solve(S)
rho = diag(s) - solve(L)

### S/N Ratio
emp_snr = mean( diag( var(xb) %%% solve(sigma) ))
emp_g_snr = mean( diag( var( (err)%%%t(rho) ) %%% solve(sigma) ))

#####

if( abs(emp_snr - snr) < (snr/10) | count > maxit ){
  break
}else{
  if( emp_snr < snr ){ # increase factor
    factor_min = factor
  }else{ # decrease factor
    factor_max = factor
  }
  factor_prev = factor
  factor = (factor_min + factor_max)/2
}
count = count+1
}

#####
colnames(y) <- paste("GEX",1:ncol(y),sep="")
colnames(G) <- colnames(y); Gy <- G
gamma <- gamma[-1,]
mrfG <- Gmrf[!duplicated(Gmrf),]
data = cbind(y,x[-1]) # leave out the intercept because is coded inside already

exampleEQTL = list(data=data, blockList=list(1:s,s+1:p))

## Write data file to the user's directory by save()

## End(Not run)

```

Description

Preprocessed data set to mimic a small pharmacogenetic example from the Genomics of Drug Sensitivity in Cancer (GDSC) database, with $p=850$ gene features as explanatory variables, $s=7$ drugs sensitivity data as response variables and data for $n=498$ cell lines. Gene features include $p_1=343$ gene expression features (GEX), $p_2=426$ by copy number variations (CNV) and $p_3=68$ mutated genes (MUT). Loading the data will load the associated blockList (and mrfG) objects needed to fit the model with BayesSUR(). The R code for generating the simulated data is given in the Examples paragraph.

```
#importFrom plyr mapvalues #importFrom data.table like
```

Usage

```
exampleGDSC
```

Format

An object of class list of length 3.

Examples

```
# Load the GDSC sample dataset
data("exampleGDSC", package = "BayesSUR")
str(exampleGDSC)

## Not run:
#=====
# This code below is to do preprocessing of GDSC data and obtain the complete dataset
# "exampleGDSC.rda" above. The user needs load the datasets from
# ftp://ftp.sanger.ac.uk/pub4/cancerrxgene/releases/release-5.0/.
# But downloading and transforming the three used datasets below to *.csv files first.
#=====

requireNamespace("plyr", quietly = TRUE)
requireNamespace("data.table", quietly = TRUE)

features <- data.frame(read.csv("/gdsc_en_input_w5.csv", head=T))
names.fea <- strsplit(rownames(features), ",")
features <- t(features)
p <- c(13321, 13747-13321, 13818-13747)
Cell.Line <- rownames(features)
features <- data.frame(Cell.Line, features)

ic50_00 <- data.frame(read.csv("gdsc_drug_sensitivity_fitted_data_w5.csv", head=T))
ic50_0 <- ic50_00[,c(1,4,7)]
drug.id <- data.frame(read.csv("gdsc_tissue_output_w5.csv", head=T))[,c(1,3)]
drug.id2 <- drug.id[!duplicated(drug.id$drug.id),]
# delete drug.id=1066 since ID1066 and ID156 both correspond drug AZD6482,
# and no ID1066 in the "suppl.Data1" by Garnett et al. (2012)
drug.id2 <- drug.id2[drug.id2$drug.id!=1066,]
drug.id2$drug.name <- as.character(drug.id2$drug.name)
```

```

drug.id2$drug.name <- substr(drug.id2$drug.name, 1, nchar(drug.id2$drug.name)-6)
drug.id2$drug.name <- gsub(" ", "-", drug.id2$drug.name)

ic50 <- ic50_0
# mapping the drug_id to drug names in drug sensitivity data set
ic50$drug_id <- plyr::mapvalues(ic50$drug_id, from = drug.id2[,2], to = drug.id2[,1])
colnames(ic50) <- c("Cell.Line", "compound", "IC50")

# transform drug sensitivity overall cell lines to a data matrix
y0 <- reshape(ic50, v.names="IC50", timevar="compound", idvar="Cell.Line", direction="wide")
y0$Cell.Line <- gsub("-", ".", y0$Cell.Line)

#=====
# select nonmissing pharmacological data
#=====
y00 <- y0
m0 <- dim(y0)[2]-1
eps <- 0.05
# r1.na is better to be not smaller than r2.na
r1.na <- 0.3
r2.na <- 0.2
k <- 1
while(sum(is.na(y0[,2:(1+m0)]))>0){
  r1.na <- r1.na - eps/k
  r2.na <- r1.na - eps/k
  k <- k + 1
  ## select drugs with <30% (decreasing with k) missing data overall cell lines
  na.y <- apply(y0[,2:(1+m0)], 2, function(xx) sum(is.na(xx))/length(xx))
  while(sum(na.y<r1.na)<m0){
    y0 <- y0[,-c(1+which(na.y>=r1.na))]
    m0 <- sum(na.y<r1.na)
    na.y <- apply(y0[,2:(1+m0)], 2, function(xx) sum(is.na(xx))/length(xx))
  }

  ## select cell lines with treatment of at least 80% (increasing with k) drugs
  na.y0 <- apply(y0[,2:(1+m0)], 1, function(xx) sum(is.na(xx))/length(xx))
  while(sum(na.y0<r2.na)<(dim(y0)[1])){
    y0 <- y0[na.y0<r2.na,]
    na.y0 <- apply(y0[,2:(1+m0)], 1, function(xx) sum(is.na(xx))/length(xx))
  }
  num.na <- sum(is.na(y0[,2:(1+m0)]))
  message("#{NA}=", num.na, "\n", "r1.na =", r1.na, ", r2.na =", r2.na, "\n")
}

#=====
# combine drug sensitivity, tissues and molecular features
#=====
yx <- merge(y0, features, by="Cell.Line")
names.cell.line <- yx$Cell.Line
names.drug <- colnames(yx)[2:(dim(y0)[2])]
names.drug <- substr(names.drug, 6, nchar(names.drug))
# numbers of gene expression features, copy number features and mutation features
p <- c(13321, 13747-13321, 13818-13747)

```



```

num.nonpen <- 13
yx <- data.matrix(yx[,-1])
y <- yx[,1:(dim(y0)[2]-1)]
x <- cbind(yx[,dim(y0)[2]-1+sum(p)+1:num.nonpen], yx[,dim(y0)[2]-1+1:sum(p)])

# delete genes with only one mutated cell line
x <- x[,-c(num.nonpen+p[1]+p[2]+which(colSums(x[,num.nonpen+p[1]+p[2]+1:p[3]])<=1))]
p[3] <- ncol(x) - num.nonpen - p[1] - p[2]

GDSC <- list(y=y, x=x, p=p, num.nonpen=num.nonpen, names.cell.line=names.cell.line,
            names.drug=names.drug)

#####
#####
## select a small set of drugs
#####
#####

name_drugs <- c("Methotrexate", "RDEA119", "PD-0325901", "CI-1040", "AZD6244", "Nilotinib",
               "Axitinib")

# extract the drugs' pharmacological profiling and tissue dummy
YX0 <- cbind(GDSC$y[,colnames(GDSC$y) %in% paste("IC50.", name_drugs, sep="")]
            [,c(1,3,6,4,7,2,5)], GDSC$x[,1:GDSC$num.nonpen])
colnames(YX0) <- c(name_drugs, colnames(GDSC$x)[1:GDSC$num.nonpen])
# extract the genetic information of CNV & MUT
X23 <- GDSC$x[, GDSC$num.nonpen+GDSC$p[1]+1:(p[2]+p[3])]
colnames(X23)[1:p[2]] <- paste(substr(colnames(X23)[1:p[2]], 1,
                                   nchar(colnames(X23)[1:p[2]] )-3), ".CNV", sep="")

# locate all genes with CNV or MUT information
name_genes_duplicate <- c( substr(colnames(X23)[1:p[2]], 1, nchar(colnames(X23)[1:p[2]]-4),
                             substr(colnames(X23)[p[2]+1:p[3]], 1, nchar(colnames(X23)[p[2]+1:p[3]]-4) )
name_genes <- name_genes_duplicate[!duplicated(name_genes_duplicate)]

# select the GEX which have the common genes with CNV or MUT
X1 <- GDSC$x[,GDSC$num.nonpen+which(colnames(GDSC$x)[GDSC$num.nonpen+1:p[1]] %in% name_genes)]

p[1] <- ncol(X1)
X1 <- log(X1)

# summary the data information
exampleGDSC <- list( data=cbind( YX0, X1, X23 ) )
exampleGDSC$blockList <- list(1:length(name_drugs), length(name_drugs)+1:GDSC$num.nonpen,
                             ncol(YX0)+1:sum(p))

#####
# construct the G matrix: edge potentials in the MRF prior
#####

# edges between drugs: Group1 ("RDEA119", "17-AAG", "PD-0325901", "CI-1040" and "AZD6244")
# indexed as (2:5)

```

```

# http://software.broadinstitute.org/gsea/msigdb/cards/KEGG_MAPK_SIGNALING_PATHWAY
pathway_genes <- read.table("MAPK_pathway.txt")[[1]]
Idx_Pathway1 <- which(c(colnames(X1),name_genes_duplicate) %in% pathway_genes)
Gmrf_Group1Pathway1 <- t(combn(rep(Idx_Pathway1,each=length(2:5)) +
                             rep((2:5-1)*sum(p),times=length(Idx_Pathway1)), 2))

# edges between drugs: Group2 ("Nilotinib","Axitinib") indexed as (6:7)
# delete gene ABL2
Idx_Pathway2 <- which( c(colnames(X1),name_genes_duplicate) %like% "BCR" |
                      c(colnames(X1),name_genes_duplicate) %like% "ABL" )[-c(3,5)]
Gmrf_Group2Pathway2 <- t(combn(rep(Idx_Pathway2,each=length(6:7)) +
                              rep((6:7-1)*sum(p),times=length(Idx_Pathway2)), 2))

# edges between the common gene in different data sources
Gmrf_CommonGene <- NULL
list_CommonGene <- list(0)
k <- 1
for(i in 1:length(name_genes)){
  Idx_CommonGene <- which( c(colnames(X1),name_genes_duplicate) == name_genes[i] )
  if(length(Idx_CommonGene) > 1){
    Gmrf_CommonGene <- rbind(Gmrf_CommonGene,t(combn(rep(Idx_CommonGene,each=length(name_drugs))
                                                    + rep((1:length(name_drugs)-1)*sum(p),times=length(Idx_CommonGene)), 2)))
    k <- k+1
  }
}
Gmrf_duplicate <- rbind( Gmrf_Group1Pathway1, Gmrf_Group2Pathway2, Gmrf_CommonGene )
Gmrf <- Gmrf_duplicate[!duplicated(Gmrf_duplicate),]
exampleGDSC$mrfG <- Gmrf

# create the target gene names of the two groups of drugs
targetGenes1 <- matrix(Idx_Pathway1,nrow=1)
colnames(targetGenes1) <- colnames(exampleGDSC$data)[seq_along(targetGene$group1)]
targetGenes2 <- matrix(Idx_Pathway2,nrow=1)
colnames(targetGenes2) <- colnames(exampleGDSC$data)[seq_along(targetGene$group2)]

targetGene <- list(group1=targetGenes1, group2=targetGenes2)

## Write data file exampleGDSC.rda to the user's directory by save()

## End(Not run)

```

fitted.BayesSUR

get fitted responses from a BayesSUR object

Description

Return the fitted response values that correspond to the posterior mean estimates from a BayesSUR class object.

Usage

```
## S3 method for class 'BayesSUR'
fitted(object, Pmax = 0, beta.type = "marginal", ...)
```

Arguments

object	an object of class BayesSUR
Pmax	valid if beta.type="conditional", giving estimated beta conditioning on gamma>=Pmax. If beta.type="conditional" and Pmax=0.5, it gives median probability model betas. Default is 0
beta.type	type of estimated beta for the fitted model. Default is marginal, giving marginal beta estimation. If beta.type="conditional", it gives conditional beta estimation
...	other arguments

Value

Fitted values extracted from an object of class BayesSUR. If the BayesSUR specified data standardization, the fitted values are base based on standardized data.

Examples

```
data("exampleEQTL", package = "BayesSUR")
hyperpar <- list( a_w = 2 , b_w = 5 )

set.seed(9173)
fit <- BayesSUR(Y = exampleEQTL[["blockList"]][[1]],
               X = exampleEQTL[["blockList"]][[2]],
               data = exampleEQTL[["data"]], outFilePath = tempdir(),
               nIter = 100, burnin = 50, nChains = 2, gammaPrior = "hotspot",
               hyperpar = hyperpar, tmpFolder = "tmp/" )

## check fitted values
fitted.val <- fitted(fit)
```

getEstimator

extract the posterior mean of the parameters

Description

Extract the posterior mean of the parameters of a BayesSUR class object.

Usage

```
getEstimator(object, estimator = "gamma", Pmax = 0, beta.type = "marginal")
```

Arguments

object	an object of class BayesSUR
estimator	the name of one estimator. Default is the latent indicator estimator "gamma". Other options "beta", "Gy", "CPO" and "logP" correspond the marginal (conditional) coefficient matrix if beta.type="marginal"("conditional"), response graph and conditional predictive ordinate (CPO) respectively
Pmax	threshold that truncate the estimator "gamma" or "Gy". Default is 0
beta.type	the type of output beta. Default is marginal, giving marginal beta estimation. If beta.type="conditional", it gives conditional beta estimation

Value

Return the estimator from an object of class BayesSUR. It is a matrix if the length of argument marginal is greater than 1. Otherwise, it is a list

Examples

```
data("exampleEQTL", package = "BayesSUR")
hyperpar <- list( a_w = 2 , b_w = 5 )

set.seed(9173)
fit <- BayesSUR(Y = exampleEQTL[["blockList"]][[1]],
               X = exampleEQTL[["blockList"]][[2]],
               data = exampleEQTL[["data"]], outFilePath = tempdir(),
               nIter = 100, burnin = 50, nChains = 2, gammaPrior = "hotspot",
               hyperpar = hyperpar, tmpFolder = "tmp/" )

## check output
# extract the posterior mean of the coefficients matrix
beta_hat <- getEstimator(fit, estimator="beta")
```

plot.BayesSUR

create a selection of plots for a BayesSUR class object

Description

plot method for class BayesSUR. This is the main plot function to be called by the user. This function calls one or several of the following functions: plotEstimator(), plotGraph(), plotMCMCdiag(), plotManhattan(), plotNetwork(), plotCPO().

Usage

```
## S3 method for class 'BayesSUR'
plot(x, estimator = NULL, type = NULL, ...)
```

Arguments

x	an object of class BayesSUR
estimator	It is in <code>c(NULL, 'beta', 'gamma', 'Gy', 'logP', 'CPO')</code> and works by combining with argument type. <ul style="list-style-type: none"> • If estimator is in <code>c("beta", "gamma", "Gy")</code> and argument <code>type="heatmap"</code>, it prints heatmaps of the specified estimator in estimator, and refers to function <code>plotEstimator()</code> for more other arguments. • If estimator="Gy" and argument <code>type="graph"</code>, it prints a structure graph of "Gy", and refers to function <code>plotGraph()</code> for more other arguments. • If estimator=<code>c("gamma", "Gy")</code> and argument <code>type="network"</code>, it prints the estimated network between the response variables and predictors with nonzero coefficients, and refers to function <code>plotMCMCdiag()</code> for more other arguments. • If estimator=NULL (default) and <code>type=NULL</code> (default), it interactively prints the plots of estimators (i.e., beta, gamma and (or) Gy), response graph Gy, network, Manhattan and MCMC diagnostics.
type	It is one of <code>NULL, "heatmap", "graph", "network", "Manhattan" and "diagnostics"</code> , and works by combining with argument estimator. <ul style="list-style-type: none"> • If <code>type="Manhattan"</code> and argument estimator="gamma", it prints Manhattan-like plots for marginal posterior inclusion probabilities (mPIP) and numbers of associated response variables for individual predictors, and refers to function <code>plotManhattan()</code> for more other arguments. • If <code>type="diagnostics"</code> and argument estimator="logP" it shows trace plots and diagnostic density plots of a fitted model, and refers to function <code>plotMCMCdiag()</code> for more other arguments. • If <code>type="diagnostics"</code> and argument estimator="CPO", it shows the conditional predictive ordinate (CPO) for each individual of a fitted model, and refers to function <code>plotCPO()</code> for more other arguments.
...	other arguments, see functions <code>plotEstimator()</code> , <code>plotGraph()</code> , <code>plotNetwork()</code> , <code>plotManhattan()</code> , <code>plotMCMCdiag()</code> or <code>plotCPO()</code>

Examples

```

data("exampleEQTL", package = "BayesSUR")
hyperpar = list( a_w = 2 , b_w = 5 )

set.seed(9173)
fit <- BayesSUR(Y = exampleEQTL[["blockList"]][[1]],
               X = exampleEQTL[["blockList"]][[2]],
               data = exampleEQTL[["data"]], outFilePath = tempdir(),
               nIter = 100, burnin = 0, nChains = 2, gammaPrior = "hotspot",
               hyperpar = hyperpar, tmpFolder = "tmp/" )

## check output
## Not run:
# Show the interactive plots. Note that it needs at least 2000*(nbloc+1) iterations
# for the diagnostic plots where nbloc=3 by default

```

```

plot(fit)

## End(Not run)

## plot heatmaps of the estimated beta, gamma and Gy
plot(fit, estimator=c("beta", "gamma", "Gy"), type="heatmap")

## plot estimated graph of responses Gy
plot(fit, estimator="Gy", type="graph")

## plot network between response variables and associated predictors
plot(fit, estimator=c("gamma", "Gy"), type="network")

## print Manhattan-like plots
plot(fit, estimator="gamma", type="Manhattan")

## print MCMC diagnostic plots
plot(fit, estimator="logP", type="diagnostics")

```

plotCPO

plot the conditional predictive ordinate

Description

Plot the conditional predictive ordinate (CPO) for each individual of a fitted model generated by BayesSUR which is a BayesSUR object. CPO is a handy posterior predictive check because it may be used to identify outliers, influential observations, and for hypothesis testing across different non-nested models (Gelfand 1996).

Usage

```

plotCPO(
  x,
  outlier.mark = TRUE,
  outlier.thresh = 0.01,
  scale.CPO = TRUE,
  x.loc = FALSE,
  axis.label = NULL,
  las = 0,
  cex.axis = 1,
  mark.pos = c(0, -0.01),
  mark.color = 2,
  mark.cex = 0.8,
  xlab = "Observations",
  ylab = NULL,
  ...
)

```

Arguments

x	an object of class BayesSUR
outlier.mark	mark the outliers with the response names. The default is FALSE
outlier.thresh	threshold for the CPOs. The default is 0.01.
scale.CPO	scaled CPOs which is divided by their maximum. The default is TRUE
x.loc	a vector of features distance
axis.label	a vector of predictor names which are shown in CPO plot. The default is NULL only showing the indices. The value "auto" show the predictor names from the original data.
las	graphical parameter of plot.default
cex.axis	graphical parameter of plot.default
mark.pos	the location of the marked text relative to the point
mark.color	the color of the marked text. The default color is red
mark.cex	the font size of the marked text. The default font size is 0.8
xlab	a title for the x axis
ylab	a title for the y axis
...	other arguments

Details

The default threshold for the CPOs to detect the outliers is 0.01 by Congdon (2005). It can be tuned by the argument `outlier.thresh`.

References

- Statisticat, LLC (2013). *Bayesian Inference*. Farmington, CT: Statisticat, LLC.
- Gelfand A. (1996). *Model Determination Using Sampling Based Methods*. In Gilks W., Richardson S., Spiegelhalter D. (eds.), *Markov Chain Monte Carlo in Practice*, pp. 145–161. Chapman & Hall, Boca Raton, FL.
- Congdon P. (2005). *Bayesian Models for Categorical Data*. John Wiley & Sons, West Sussex, England.

Examples

```
data("exampleEQTL", package = "BayesSUR")
hyperpar <- list( a_w = 2 , b_w = 5 )

set.seed(9173)
fit <- BayesSUR(Y = exampleEQTL[["blockList"]][[1]],
               X = exampleEQTL[["blockList"]][[2]],
               data = exampleEQTL[["data"]], outFilePath = tempdir(),
               nIter = 100, burnin = 50, nChains = 2, gammaPrior = "hotspot",
               hyperpar = hyperpar, tmpFolder = "tmp/", output_CPO=TRUE)

## check output
```

```
# plot the conditional predictive ordinate (CPO)
plotCPO(fit)
```

```
plotEstimator      plot the estimators
```

Description

Plot the posterior mean estimators from a "BayesSUR" class object, including the coefficients beta, latent indicator variable gamma and graph of responses.

Usage

```
plotEstimator(
  x,
  estimator = NULL,
  colorScale.gamma = grey((100:0)/100),
  colorScale.beta = c("blue", "white", "red"),
  legend.cex.axis = 1,
  name.responses = NA,
  name.predictors = NA,
  xlab = "",
  ylab = "",
  fig.tex = FALSE,
  output = "ParamEstimator",
  header = "",
  header.cex = 2,
  tick = FALSE,
  mgp = c(2.5, 1, 0),
  title.beta = paste("Estimator", "$\\hat{\\mathbf{B}}$"),
  title.gamma = paste("Estimator", "$\\hat{\\mathbf{\\Gamma}}$"),
  title.Gy = paste("Estimator", "$\\hat{\\mathcal{G}}$"),
  cex.main = 1.5,
  ...
)
```

Arguments

x	an object of class BayesSUR
estimator	print the heatmap of estimators. The value "beta" is for the estimated coefficients matrix, "gamma" for the latent indicator matrix and "Gy" for the graph of responses
colorScale.gamma	value palette for gamma
colorScale.beta	a vector of three colors for diverging color schemes

legend.cex.axis	magnification of axis annotation relative to cex
name.responses	a vector of the response names. The default is "NA" only to show the locations. The value "auto" show the response names from the original data.
name.predictors	a vector of the predictor names. The default is "NA" only to show the locations. The value "auto" show the predictor names from the original data.
xlab	a title for the x axis
ylab	a title for the y axis
fig.tex	print the figure through LaTeX. Default is "FALSE"
output	the file name of printed figure
header	the main title
header.cex	size of the main title for all estimators
tick	a logical value specifying whether tickmarks and an axis line should be drawn. Default is "FALSE"
mgp	the margin line (in mex units) for the axis title, axis labels and axis line
title.beta	a title for the printed "beta" if fig.tex=TRUE
title.gamma	a title for the printed "gamma" if fig.tex=TRUE
title.Gy	a title for the printed "Gy" if fig.tex=TRUE
cex.main	size of the title for each estimator
...	other arguments

Examples

```

data("exampleEQTL", package = "BayesSUR")
hyperpar <- list( a_w = 2 , b_w = 5 )

set.seed(9173)
fit <- BayesSUR(Y = exampleEQTL[["blockList"]][[1]],
               X = exampleEQTL[["blockList"]][[2]],
               data = exampleEQTL[["data"]], outFilePath = tempdir(),
               nIter = 100, burnin = 50, nChains = 2, gammaPrior = "hotspot",
               hyperpar = hyperpar, tmpFolder = "tmp/" )

## check output
# Plot the estimators from the fitted object
plotEstimator(fit, estimator=c("beta","gamma","Gy"))

## Not run:
#Set up temporary work directory for saving a pdf figure
td <- tempdir()
oldwd <- getwd()
setwd(td)

# Produce authentic math formulas in the graph
plotEstimator(fit, estimator=c("beta","gamma","Gy"), fig.tex = TRUE)

```

```

system(paste(getOption("pdfviewer"), "ParamEstimator.pdf"))
setwd(oldwd)

## End(Not run)

```

plotGraph

plot the estimated graph for multiple response variables

Description

Plot the estimated graph for multiple response variables from a BayesSUR class object.

Usage

```

plotGraph(
  x,
  Pmax = 0.5,
  main = "Estimated graph of responses",
  edge.width = 2,
  edge.weight = FALSE,
  vertex.label = NULL,
  vertex.label.color = "black",
  vertex.size = 30,
  vertex.color = "dodgerblue",
  vertex.frame.color = NA,
  ...
)

```

Arguments

x	either an object of class BayesSUR (default) or a symmetric numeric matrix representing an adjacency matrix for a given graph structure. If x is an adjacency matrix, argument main="Given graph of responses" by default.
Pmax	a value for thresholding the learning structure matrix of multiple response variables. Default is 0.5
main	an overall title for the plot
edge.width	edge width. Default is 2
edge.weight	draw weighted edges after thresholding at 0.5. The default value FALSE is not to draw weighted edges
vertex.label	character vector used to label the nodes
vertex.label.color	label color. Default is "black"
vertex.size	node size. Default is 30
vertex.color	node color. Default is "dodgerblue"
vertex.frame.color	node color. Default is "NA"
...	other arguments

Examples

```

data("exampleEQTL", package = "BayesSUR")
hyperpar <- list( a_w = 2 , b_w = 5 )

set.seed(9173)
fit <- BayesSUR(Y = exampleEQTL[["blockList"]][[1]],
               X = exampleEQTL[["blockList"]][[2]],
               data = exampleEQTL[["data"]], outFilePath = tempdir(),
               nIter = 100, burnin = 50, nChains = 2, gammaPrior = "hotspot",
               hyperpar = hyperpar, tmpFolder = "tmp/" )

## check output
# show the graph relationship between responses
plotGraph(fit, estimator = "Gy")

```

plotManhattan

plot Manhattan-like plots for marginal posterior inclusion probabilities (mPIP) and numbers of responses of association for predictors

Description

Plot Manhattan-like plots for marginal posterior inclusion probabilities (mPIP) and numbers of responses of association for predictors of a "BayesSUR" class object.

Usage

```

plotManhattan(
  x,
  which = c(1, 2),
  x.loc = FALSE,
  axis.label = "auto",
  mark.responses = NULL,
  xlab1 = "Predictors",
  ylab1 = "mPIP",
  xlab2 = "Predictors",
  ylab2 = "No. of responses",
  threshold = 0.5,
  las = 0,
  cex.axis = 1,
  mark.pos = c(0, 0),
  mark.color = 2,
  mark.cex = 0.8,
  header = "",
  ...
)

```

Arguments

x	an object of class BayesSUR
which	if it's value "1" showing the Manhattan-like plot of the marginal posterior inclusion probabilities (mPIP). If it's value "2" showing the Manhattan-like plot of the number of responses. The default is to show both figures.
x.loc	a vector of features distance
axis.label	a vector of predictor names which are shown in the Manhattan-like plot. The value "NULL" only showing the indices. The default "auto" show the predictor names from the original data.
mark.responses	a vector of response names which are shown in the Manhattan-like plot for the mPIP
xlab1	a title for the x axis of Manhattan-like plot for the mPIP
ylab1	a title for the y axis of Manhattan-like plot for the mPIP
xlab2	a title for the x axis of Manhattan-like plot for the numbers of responses
ylab2	a title for the y axis of Manhattan-like plot for the numbers of responses
threshold	threshold for showing number of response variables significantly associated with each feature
las	graphical parameter of plot.default
cex.axis	graphical parameter of plot.default
mark.pos	the location of the marked text relative to the point
mark.color	the color of the marked text. The default color is red.
mark.cex	the fontsize of the marked text. The default fontsize is 0.8.
header	the main title
...	other arguments

Examples

```

data("exampleEQTL", package = "BayesSUR")
hyperpar <- list( a_w = 2 , b_w = 5 )

set.seed(9173)
fit <- BayesSUR(Y = exampleEQTL[["blockList"]][[1]],
               X = exampleEQTL[["blockList"]][[2]],
               data = exampleEQTL[["data"]], outFilePath = tempdir(),
               nIter = 100, burnin = 50, nChains = 2, gammaPrior = "hotspot",
               hyperpar = hyperpar, tmpFolder = "tmp/" )

## check output
# show the Manhattan-like plots
plotManhattan(fit)

```

plotMCMCdiag *plot MCMC diagnostic plots*

Description

Show trace plots and diagnostic density plots of a fitted model object of class "BayesSUR".

Usage

```
plotMCMCdiag(x, nbloc = 3, HIWg = NULL, header = "", ...)
```

Arguments

x	an object of class BayesSUR
nbloc	number of splits for the last half iterations after subtracting burn-in length
HIWg	diagnostic plot of the response graph. Default is NULL. HIW="degree" prints the diagnostic of the degrees of response nodes. HIW="edges" prints the diagnostic of every edge between two responses. HIW="lik" prints the diagnostic of the posterior likelihoods of the hyperparameters related to the response relationships
header	the main title
...	other arguments for the plots of the log-likelihood and model size

Examples

```
data("exampleEQTL", package = "BayesSUR")
hyperpar <- list( a_w = 2 , b_w = 5 )

set.seed(9173)
fit <- BayesSUR(Y = exampleEQTL[["blockList"]][[1]],
               X = exampleEQTL[["blockList"]][[2]],
               data = exampleEQTL[["data"]], outFilePath = tempdir(),
               nIter = 100, burnin = 50, nChains = 2, gammaPrior = "hotspot",
               hyperpar = hyperpar, tmpFolder = "tmp/" )

## check output
plotMCMCdiag(fit)
```

plotNetwork	<i>plot the network representation of the associations between responses and predictors</i>
-------------	---------------------------------------------------------------------------------------------

Description

Plot the network representation of the associations between responses and predictors, based on the estimated gamma matrix and graph of responses from a "BayesSUR" class object.

Usage

```
plotNetwork(
  x,
  includeResponse = NULL,
  excludeResponse = NULL,
  includePredictor = NULL,
  excludePredictor = NULL,
  MatrixGamma = NULL,
  PmaxPredictor = 0.5,
  PmaxResponse = 0.5,
  nodesizePredictor = 2,
  nodesizeResponse = 15,
  no.isolates = FALSE,
  lineup = 1.2,
  gray.alpha = 0.6,
  edgewith.response = 5,
  edgewith.predictor = 2,
  edge.weight = FALSE,
  label.predictor = NULL,
  label.response = NULL,
  color.predictor = NULL,
  color.response = NULL,
  name.predictors = NULL,
  name.responses = NULL,
  vertex.frame.color = NA,
  layoutInCircle = FALSE,
  header = "",
  ...
)
```

Arguments

x	an object of class BayesSUR
includeResponse	A vector of the response names which are shown in the network
excludeResponse	A vector of the response names which are not shown in the network

includePredictor	A vector of the predictor names which are shown in the network
excludePredictor	A vector of the predictor names which are not shown in the network
MatrixGamma	A matrix or dataframe of the latent indicator variable. Default is NULL and to extrate it from object of class inheriting from an object of class "BayesSUR"
PmaxPredictor	cutpoint for thresholding the estimated latent indicator variable. Default is 0.5
PmaxResponse	cutpoint for thresholding the learning structure matrix of multiple response variables. Default is 0.5
nodesizePredictor	node size of Predictors in the output graph. Default is 15
nodesizeResponse	node size of response variables in the output graph. Default is 25
no.isolates	remove isolated nodes from responses graph and Full graph, may get problem if there are also isolated Predictors
lineup	A ratio of the heights between responses' area and Predictors'
gray.alpha	the opacity. The default is 0.6
edgewith.response	the edge width between response nodes
edgewith.predictor	the edge width between the predictor and response node
edge.weight	draw weighted edges after thresholding at 0.5. The default value FALSE is not to draw weighted edges
label.predictor	A vector of the names of predictors
label.response	A vector of the names of response variables
color.predictor	color of the predictor nodes
color.response	color of the response nodes
name.predictors	A subtitle for the predictors
name.responses	A subtitle for the responses
vertex.frame.color	color of the frame of the vertices. If you don't want vertices to have a frame, supply NA as the color name
layoutInCircle	place vertices on a circle, in the order of their vertex ids. The default is FALSE
header	the main title
...	other arguments

Examples

```

data("exampleEQTL", package = "BayesSUR")
hyperpar <- list( a_w = 2 , b_w = 5 )

set.seed(9173)
fit <- BayesSUR(Y = exampleEQTL[["blockList"]][[1]],
               X = exampleEQTL[["blockList"]][[2]],
               data = exampleEQTL[["data"]], outFilePath = tempdir(),
               nIter = 100, burnin = 50, nChains = 2, gammaPrior = "hotspot",
               hyperpar = hyperpar, tmpFolder = "tmp/" )

## check output
# show the Network representation of the associations between responses and features
plotNetwork(fit)

```

predict.BayesSUR	<i>predict responses corresponding to the posterior mean of the coefficients, return posterior mean of coefficients or indices of nonzero coefficients</i>
------------------	------------------------------------------------------------------------------------------------------------------------------------------------------------

Description

Predict responses corresponding to the posterior mean of the coefficients, return posterior mean of coefficients or indices of nonzero coefficients of a BayesSUR class object.

Usage

```

## S3 method for class 'BayesSUR'
predict(object, newx, type = "response", Pmax = 0, beta.type = "marginal", ...)

```

Arguments

object	an object of class BayesSUR
newx	Matrix of new values for x at which predictions are to be made. Must be a matrix
type	Type of prediction required. type="response" gives the fitted responses; type="coefficients" computes the coefficients truncated the estimated coefficients based on thresholding the estimated latent indicator variable at Pmax; type="nonzero" returns a list of the indices of the nonzero coefficients corresponding to the estimated latent indicator variable thresholding at Pmax
Pmax	threshold that truncates the estimated coefficients based on thresholding the estimated latent indicator variable. Default is 0.
beta.type	the type of estimated coefficients beta for prediction. Default is marginal, giving marginal beta estimation. If beta.type="conditional", it gives conditional beta estimation
...	other arguments

Value

Predicted values extracted from an object of class BayesSUR. If the BayesSUR specified data standardization, the fitted values are base based on standardized data.

Examples

```
data("exampleEQTL", package = "BayesSUR")
hyperpar <- list( a_w = 2 , b_w = 5 )

set.seed(9173)
fit <- BayesSUR(Y = exampleEQTL[["blockList"]][[1]],
               X = exampleEQTL[["blockList"]][[2]],
               data = exampleEQTL[["data"]], outFilePath = tempdir(),
               nIter = 100, burnin = 50, nChains = 2, gammaPrior = "hotspot",
               hyperpar = hyperpar, tmpFolder = "tmp/" )

## check prediction
predict.val <- predict(fit, newx=exampleEQTL[["blockList"]][[2]])
```

print.BayesSUR

print method for class BayesSUR

Description

Print a short summary of a BayesSUR class object. It includes the argument matching information, number of selected predictors based on thresholding the posterior mean of the latent indicator variable at 0.5 by default.

Usage

```
## S3 method for class 'BayesSUR'
print(x, Pmax = 0.5, ...)
```

Arguments

x	an object of class BayesSUR
Pmax	threshold that truncates the estimated coefficients based on thresholding the estimated latent indicator variable. Default is 0.5
...	other arguments

Value

Return a short summary from an object of class BayesSUR, including the number of selected predictors with $mPIP > Pmax$ and the expected log pointwise predictive density estimates (i.e., $elpd.LOO$ and $elpd.WAIC$).

Examples

```

data("exampleEQTL", package = "BayesSUR")
hyperpar = list( a_w = 2 , b_w = 5 )

set.seed(9173)
fit <- BayesSUR(Y = exampleEQTL[["blockList"]][[1]],
               X = exampleEQTL[["blockList"]][[2]],
               data = exampleEQTL[["data"]], outFilePath = tempdir(),
               nIter = 100, burnin = 50, nChains = 2, gammaPrior = "hotspot",
               hyperpar = hyperpar, tmpFolder = "tmp/", output_CPO=TRUE)

## check output
# show the print information
print(fit)

```

summary.BayesSUR	<i>summary method for class BayesSUR</i>
------------------	------------------------------------------

Description

Summary method for class BayesSUR. It includes the argument matching information, Top predictors/responses on average mPIP across all responses/predictors, elpd estimates, MCMC specification, model specification and hyper-parameters. The summarized number of the selected variable corresponds to the posterior mean of the latent indicator variable thresholding at 0.5 by default.

Usage

```

## S3 method for class 'BayesSUR'
summary(object, Pmax = 0.5, ...)

```

Arguments

object	an object of class BayesSUR
Pmax	threshold that truncates the estimated coefficients based on thresholding the estimated latent indicator variable. Default is 0.5
...	other arguments

Value

Return a result summary from an object of class BayesSUR, including the CPOs, number of selected predictors with mPIP>Pmax, top 10 predictors on average mPIP across all responses, top 10 responses on average mPIP across all predictors, Expected log pointwise predictive density (elpd) estimates, MCMC specification, model specification (i.e., covariance prior and gamma prior) and hyper-parameters.

Examples

```

data(exampleEQTL, package = "BayesSUR")
hyperpar = list( a_w = 2 , b_w = 5 )

set.seed(9173)
fit <- BayesSUR(Y = exampleEQTL[["blockList"]][[1]],
               X = exampleEQTL[["blockList"]][[2]],
               data = exampleEQTL[["data"]], outFilePath = tempdir(),
               nIter = 100, burnin = 50, nChains = 2, gammaPrior = "hotspot",
               hyperpar = hyperpar, tmpFolder = "tmp/", output_CPO=TRUE)

## check output
# show the summary information
summary(fit)

```

targetGene

targetGene

Description

Indices list of target genes corresponding the example_GDSC data set. It has two components representing the gene indices of the MAPK/ERK pathway and BCR-ABL gene fusion in the example_GDSC data set.

Usage

```
targetGene
```

Format

An object of class list of length 2.

Examples

```

# Load the indices of gene targets from the GDSC sample dataset
data("targetGene", package = "BayesSUR")
str(targetGene)

## Not run:
#=====
# This code below is to do preprocessing of GDSC data and obtain the complete dataset
# "targetGene.rda" above. The user needs load the datasets from
# ftp://ftp.sanger.ac.uk/pub4/cancerrxgene/releases/release-5.0/.
# But downloading and transforming the three used datasets below to *.csv files first.
#=====

requireNamespace("plyr", quietly = TRUE)
requireNamespace("data.table", quietly = TRUE)

```

```

features <- data.frame(read.csv("/gdsc_en_input_w5.csv", head=T))
names.fea <- strsplit(rownames(features), "")
features <- t(features)
p <- c(13321, 13747-13321, 13818-13747)
Cell.Line <- rownames(features)
features <- data.frame(Cell.Line, features)

ic50_00 <- data.frame(read.csv("gdsc_drug_sensitivity_fitted_data_w5.csv", head=T))
ic50_0 <- ic50_00[,c(1,4,7)]
drug.id <- data.frame(read.csv("gdsc_tissue_output_w5.csv", head=T))[,c(1,3)]
drug.id2 <- drug.id[!duplicated(drug.id$drug.id),]
# delete drug.id=1066 since ID1066 and ID156 both correspond drug AZD6482,
# and no ID1066 in the "suppl.Data1" by Garnett et al. (2012)
drug.id2 <- drug.id2[drug.id2$drug.id!=1066,]
drug.id2$drug.name <- as.character(drug.id2$drug.name)
drug.id2$drug.name <- substr(drug.id2$drug.name, 1, nchar(drug.id2$drug.name)-6)
drug.id2$drug.name <- gsub(" ", "-", drug.id2$drug.name)

ic50 <- ic50_0
# mapping the drug_id to drug names in drug sensitivity data set
ic50$drug_id <- plyr::mapvalues(ic50$drug_id, from = drug.id2[,2], to = drug.id2[,1])
colnames(ic50) <- c("Cell.Line", "compound", "IC50")

# transform drug sensitivity overall cell lines to a data matrix
y0 <- reshape(ic50, v.names="IC50", timevar="compound", idvar="Cell.Line", direction="wide")
y0$Cell.Line <- gsub("-", ".", y0$Cell.Line)

#####
# select nonmissing pharmacological data
#####
y00 <- y0
m0 <- dim(y0)[2]-1
eps <- 0.05
# r1.na is better to be not smaller than r2.na
r1.na <- 0.3
r2.na <- 0.2
k <- 1
while(sum(is.na(y0[,2:(1+m0)]))>0){
  r1.na <- r1.na - eps/k
  r2.na <- r1.na - eps/k
  k <- k + 1
  ## select drugs with <30% (decreasing with k) missing data overall cell lines
  na.y <- apply(y0[,2:(1+m0)], 2, function(xx) sum(is.na(xx))/length(xx))
  while(sum(na.y<r1.na)<m0){
    y0 <- y0[,-c(1+which(na.y>=r1.na))]
    m0 <- sum(na.y<r1.na)
    na.y <- apply(y0[,2:(1+m0)], 2, function(xx) sum(is.na(xx))/length(xx))
  }

  ## select cell lines with treatment of at least 80% (increasing with k) drugs
  na.y0 <- apply(y0[,2:(1+m0)], 1, function(xx) sum(is.na(xx))/length(xx))

```

```

while(sum(na.y0<r2.na)<(dim(y0)[1])){
  y0 <- y0[na.y0<r2.na,]
  na.y0 <- apply(y0[,2:(1+m0)], 1, function(xx) sum(is.na(xx))/length(xx))
}
num.na <- sum(is.na(y0[,2:(1+m0)]))
message("#{NA}=", num.na, "\n", "r1.na =", r1.na, ", r2.na =", r2.na, "\n")
}

#####
# combine drug sensitivity, tissues and molecular features
#####
yx <- merge(y0, features, by="Cell.Line")
names.cell.line <- yx$Cell.Line
names.drug <- colnames(yx)[2:(dim(y0)[2])]
names.drug <- substr(names.drug, 6, nchar(names.drug))
# numbers of gene expression features, copy number features and mutation features
p <- c(13321, 13747-13321, 13818-13747)
num.nonpen <- 13
yx <- data.matrix(yx[,-1])
y <- yx[,1:(dim(y0)[2]-1)]
x <- cbind(yx[,dim(y0)[2]-1+sum(p)+1:num.nonpen], yx[,dim(y0)[2]-1+1:sum(p)])

# delete genes with only one mutated cell line
x <- x[,-c(num.nonpen+p[1]+p[2]+which(colSums(x[,num.nonpen+p[1]+p[2]+1:p[3]])<=1))]
p[3] <- ncol(x) - num.nonpen - p[1] - p[2]

GDSC <- list(y=y, x=x, p=p, num.nonpen=num.nonpen, names.cell.line=names.cell.line,
            names.drug=names.drug)

#####
#####
## select a small set of drugs
#####
#####

name_drugs <- c("Methotrexate","RDEA119","PD-0325901","CI-1040","AZD6244","Nilotinib",
               "Axitinib")

# extract the drugs' pharmacological profiling and tissue dummy
YX0 <- cbind(GDSC$y[,colnames(GDSC$y) %in% paste("IC50.",name_drugs,sep="")]
            [,c(1,3,6,4,7,2,5)], GDSC$x[,1:GDSC$num.nonpen])
colnames(YX0) <- c(name_drugs, colnames(GDSC$x)[1:GDSC$num.nonpen])
# extract the genetic information of CNV & MUT
X23 <- GDSC$x[, GDSC$num.nonpen+GDSC$p[1]+1:(p[2]+p[3])]
colnames(X23)[1:p[2]] <- paste(substr(colnames(X23)[1:p[2]], 1,
                                   nchar(colnames(X23)[1:p[2]])-3), ".CNV", sep="")

# locate all genes with CNV or MUT information
name_genes_duplicate <- c( substr(colnames(X23)[1:p[2]], 1, nchar(colnames(X23)[1:p[2]])-4),
                        substr(colnames(X23)[p[2]+1:p[3]], 1, nchar(colnames(X23)[p[2]+1:p[3]])-4) )
name_genes <- name_genes_duplicate[!duplicated(name_genes_duplicate)]

```

```

# select the GEX which have the common genes with CNV or MUT
X1 <- GDSC$x[,GDSC$num.nonpen+which(colnames(GDSC$x)[GDSC$num.nonpen+1:p[1]] %in% name_genes)]

p[1] <- ncol(X1)
X1 <- log(X1)

# summary the data information
example_GDSC <- list( data=cbind( YX0, X1, X23 ) )
example_GDSC$blockList <- list(1:length(name_drugs), length(name_drugs)+1:GDSC$num.nonpen,
                               ncol(YX0)+1:sum(p))

#=====
# construct the G matrix: edge potentials in the MRF prior
#=====

# edges between drugs: Group1 ("RDEA119","17-AAG","PD-0325901","CI-1040" and "AZD6244")
# indexed as (2:5)
# http://software.broadinstitute.org/gsea/msigdb/cards/KEGG_MAPK_SIGNALING_PATHWAY
pathway_genes <- read.table("MAPK_pathway.txt")[[1]]
Idx_Pathway1 <- which(c(colnames(X1),name_genes_duplicate) %in% pathway_genes)
Gmrf_Group1Pathway1 <- t(combn(rep(Idx_Pathway1,each=length(2:5)) +
                              rep((2:5-1)*sum(p),times=length(Idx_Pathway1)), 2))

# edges between drugs: Group2 ("Nilotinib","Axitinib") indexed as (6:7)
# delete gene ABL2
Idx_Pathway2 <- which( c(colnames(X1),name_genes_duplicate) %like% "BCR" |
                      c(colnames(X1),name_genes_duplicate) %like% "ABL" )[-c(3,5)]
Gmrf_Group2Pathway2 <- t(combn(rep(Idx_Pathway2,each=length(6:7)) +
                              rep((6:7-1)*sum(p),times=length(Idx_Pathway2)), 2))

# edges between the common gene in different data sources
Gmrf_CommonGene <- NULL
list_CommonGene <- list(0)
k <- 1
for(i in 1:length(name_genes)){
  Idx_CommonGene <- which( c(colnames(X1),name_genes_duplicate) == name_genes[i] )
  if(length(Idx_CommonGene) > 1){
    Gmrf_CommonGene <- rbind(Gmrf_CommonGene,t(combn(rep(Idx_CommonGene,each=length(name_drugs))
                                                    + rep((1:length(name_drugs)-1)*sum(p),times=length(Idx_CommonGene)), 2)))
    k <- k+1
  }
}
Gmrf_duplicate <- rbind( Gmrf_Group1Pathway1, Gmrf_Group2Pathway2, Gmrf_CommonGene )
Gmrf <- Gmrf_duplicate[!duplicated(Gmrf_duplicate),]
example_GDSC$mrfG <- Gmrf

# create the target gene names of the two groups of drugs
targetGenes1 <- matrix(Idx_Pathway1,nrow=1)
colnames(targetGenes1) <- colnames(example_GDSC$data)[seq_along(targetGene$group1)]
targetGenes2 <- matrix(Idx_Pathway2,nrow=1)
colnames(targetGenes2) <- colnames(example_GDSC$data)[seq_along(targetGene$group2)]

targetGene <- list(group1=targetGenes1, group2=targetGenes2)

```

```
## Write data file targetGene.rda to the user's directory by save()
```

```
## End(Not run)
```

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