Package 'bGWAS'

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Type Package

Title Bayesian Genome-Wide Association Study

Description Package regrouping functions to perform Bayesian Genome-Wide Association Studies (bGWAS). See McDaid et al (2017) for more information about the method.

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URL https://github.com/n-mounier/bGWAS

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Depends R (>= 3.5.0),
dplyr (>= 0.7.8),
magrittr (>= 1.5),

Suggests testthat,
knitr,
remotes,
rmarkdown

Imports calibrate (>= 1.7.2),
data.table (>= 1.12.0),
ggplot2 (>= 2.2.1),
gplots (>= 3.0.1),
qqman (>= 0.1.4),
readr (>= 1.3.1),
Rcpp (>= 0.12.15),
rlang (>= 0.4.0),
R.utils (>= 2.9.0),
stringr (>= 1.4.0),
tibble (>= 2.1.1),
tidyr (>= 0.8.3),
TwoSampleMR (>= 0.3.0)

Remotes MRCIEU/TwoSampleMR

VignetteBuilder knitr

R topics documented:

`all.equal.bGWAS` *Equality test for bGWAS objects*

Description

Equality test for bGWAS objects

Usage

```
## S3 method for class 'equal.bGWAS'
all(obj1, obj2)
```

Arguments

`obj1`	an object of class bGWAS
`obj2`	an object of class bGWAS

Value

all.equal

bGWAS *bGWAS - main function*

Description

Performs a Bayesian GWAS from Summary Statistics, using publicly available results to calculate the prior effects of the SNPs and compare it to observed z-scores

Usage

```
bGWAS(name, GWAS, Z_matrices = "~/ZMatrices/", prior_studies = NULL,
  MR_threshold = 1e-06, MR_ninstruments = 3, MR_pruning_dist = 500,
  MR_pruning_LD = 0, MR_shrinkage = 1, stepwise_threshold = NULL,
  prior_shrinkage = NULL, sign_method = "p", sign_thresh = 5e-08,
  use_permutations = FALSE, res_pruning_dist = 500,
  res_pruning_LD = 0, save_files = FALSE, verbose = TRUE)
```

Arguments

name	The name of the analysis (character)
GWAS	The path to the conventional GWAS of interest, the ID of the GWAS from the list of studies available (prior GWASs), or a data.frame (character, numeric or data.frame)
Z_matrices	The path to the folder containing Z-Matrices, default="~/ZMatrices/" (character)
prior_studies	The IDs of prior GWASs to use for the analysis, default=NULL, will include all the prior GWASs available (numeric vector)
MR_threshold	The threshold used to select strong instruments for MR, should be lower than 1e-5, default=1e-6 (numeric)
MR_ninstruments	The minimum number of strong instruments needed to use a prior GWAS, should be between 2 and 8, default=3 (numeric)
MR_pruning_dist	The distance used for pruning MR instruments (in Kb), should be between 10 and 1000, default=500 (numeric)
MR_pruning_LD	The LD threshold used for pruning MR instruments, should be between 0 and 1 (if 0, distance-based pruning is used), default=0 (numeric)
MR_shrinkage	The p-value threshold used for shrinkage before performing MR, should be between MR_threshold and 1 (no shrinkage), default=1 (numeric)
stepwise_threshold	The p-value threshold used for inclusion/exclusion of Prior GWASs during the stepwise selection approach, should be between 0.05 and 0.0005, default=NULL will use 0.05 divided by the number of Prior GWASs tested (numeric)
prior_shrinkage	The p-value threshold used for shrinkage before calculating the prior, should be between MR_threshold and 1, default=NULL will use MR_shrinkage (numeric)

`sign_method` The method used to identify significant SNPs, should be `"p"` for p-value or `"fdr"` for false discovery rate, `default="p"` (character)

`sign_thresh` The threshold used to identify significant SNPs, `default="5e-8"` (numeric)

`use_permutations`
A logical indicating if BF p-values should be estimated using the permutation approach, `default=FALSE`

`res_pruning_dist`
The distance used for pruning results (in Kb), should be between 10 and 1000, (if set to NULL, no pruning is done), `default=500` (numeric)

`res_pruning_LD` The LD threshold used for pruning results, should be between 0 and 1 (if 0, distance-based pruning is used), `default=0` (numeric)

`save_files` A logical indicating if the results should be saved as files, `default=FALSE`

`verbose` A logical indicating if information on progress should be reported, `default=TRUE`

Details

`Name` and `GWAS` are required arguments. If `GWAS` is a path to a file (regular or .gz) or a `data.frame`, it should contain the following columns :
SNPID (rs numbers) should be : `rs`, `rsid`, `snp`, `snpid`, `rnpid`
A1 should be : `a1`, `alt`, `alts`
A2 should be : `a2`, `a0`, `ref`
Z should be : `z`, `Z`, `zscore`
If Z is not present, it can be calculated from BETA and SE.
BETA should be : `b`, `beta`, `beta1`
SE should be : `se`, `std`
Note: in order to get rescaled (prior/posterior/corrected) effects, BETA and SE should be provided.

Z-Matrix files, containing Z-scores for all prior GWASs should be downloaded separately and stored in `"~/ZMatrices"` or in the folder specified with the argument `Z_matrices`.
See [here](https://github.com/n-mounier/bGWAS) for more informations.

Use `list_priorGWASs()` to see all the prior GWASs available. Using one of them as your conventionnal GWAS (argument `GWAS` = numeric ID) will automatically remove it from the list of prior GWASs used to build the prior.

Use select_priorGWASs() to automatically select the prior GWASs to be included/excluded when building the prior (argument `prior_studies`).

Value

`bGWAS()` returns an object of class "bGWAS".
Additionnaly, if `save_files=T`, several files are created in the folder `./name/` :

- "PriorGWASs.tsv" - contains information about all prior GWASs (general info + status (used/excluded) + MR coefficients)
- "CoefficientsByChromosome.csv" - contains the MR estimates when masking the focal chromosome (22 coefficients / prior GWASs used for prior estimation)
- "PriorBFp.csv" - contains BF and p-values, prior, posterior and direct effects estimates for all SNPs
- "SignificantSNPs.csv" - contains BF and p-values, prior, posterior and direct effects estimates for a subset of significant SNPs

Examples

```
# Permorm bGWAS, using a small conventional GWAS included in the package (data.frame)
# and selecting a subset of studies for the prior
## Not run: top
data("SmallGWAS_Timmers2019")
MyStudies = select_priorGWASs(include_traits=c("Blood Pressure", "Education"),
                              include_files=c("cardiogram_gwas_results.txt",
                                   "All_ancestries_SNP_gwas_mc_merge_nogc.tbl.uniq.gz"))
# 6 Prior GWASs used
list_priorGWASs(MyStudies)

 A = bGWAS(name="Test_UsingSmallDataFrame",
           GWAS = SmallGWAS_Timmers2019,
           prior_studies=MyStudies,
           MR_threshold = 1e-6,
           stepwise_threshold=0.05,
           save_files=T)

## End(Not run)

# Permorm bGWAS, using a conventional GWAS from the list of prior GWASs
## Not run: MyGWAS = 3
list_priorGWASs(MyGWAS)
# Coronary Artery Disease GWAS (CARDIoGRAM)
B = bGWAS(name = "Test_UsingGWASfromPriorGWASs",
          GWAS = MyGWAS)

## End(Not run)
```

`coefficients_plot_bGWAS`

Coefficients Plot from bGWAS results

Description

Creates a Coefficients Plot (causal effect of each Prior GWASs)

Usage

```
coefficients_plot_bGWAS(obj, save_file = F, file_name = NULL)
```

Arguments

`obj`	an object of class bGWAS created using `bGWAS()`
`save_file`	A logical indicating if the graphic should be saved, `default=FALSE`, graphic will be displayed on the on-screen device
`file_name`	The name of the file saved (is `save_file` is `TRUE`) `default=NULL`, will used NameOfYourAnalysis_CoefficientsPlot.png

Value

a Coefficients Plot

extract_MRcoeffs_bGWAS

Extract MR coefficients from bGWAS results

Description

Extracts MR coefficients (multivariate genome-wide and per-chromosome estimates)

Usage

```
extract_MRcoeffs_bGWAS(obj)
```

Arguments

obj	an object of class bGWAS created using bGWAS()

Value

a tibble containing the MR coefficients (1 estimate using all chromosomes + 22 estimates with 1 chromosome masked)

extract_results_bGWAS *Extract SNPs results from bGWAS results*

Description

Extracts SNPs results from bGWAS results (BFs, p-value, prior, posterior and direct effects, depending on the value of the parameter results)

Usage

```
extract_results_bGWAS(obj, SNPs = "significant", results = "BF")
```

Arguments

obj	an object of class bGWAS created using bGWAS()
SNPs,	"all" / "significant", default="significant"
results,	"BF" / "posterior" / "direct", default="BF"

Details

For all value of results, basic informations about the SNPs will be returned:
rsid : rs number
chrm_UK10K : chromosome (obtained from UK10K data)
pos_UK10K : position (obtained from UK10K data)
alt : alternative (effect) allele
ref : reference allele
z_obs : observed Z-score

In addition, if `results = "BF"` the following information will be returned:
`mu_prior_estimate` : prior effect estimate (z-score scale)
`mu_prior_std_error` : prior effect standard error (z-score scale)
`BF` : Bayes Factor
`BF_p` : Bayes Factor p-value
`BF_fdr` : Bayes Factor FDR (only if FDR used to identify significant SNPs)

Alternatively, if `results = "posterior"` the following information will be returned:
`mu_posterior_estimate` : posterior effect estimate (z-score scale)
`mu_posterior_std_error` : posterior effect standard error (z-score scale)
`z_posterior` : posterior Z-score
`p_posterior` : posterior effect p-value
`fdr_posterior` : posterior effect FDR (only if FDR used to identify significant SNPs)

Alternatively, if `results = "direct"` the following information will be returned:
`mu_direct_estimate` : direct effect estimate (z-score scale)
`mu_direct_std_error` : direct effect standard error (z-score scale)
`z_direct` : direct Z-score
`p_direct` : direct effect p-value
`fdr_direct` : direct effect FDR (only if FDR used to identify significant SNPs)

Value

a `tibble` containing the results for all / significant SNPs

`get_RSquared_bGWAS` *Get squared correlation between observed and prior effects from bGWAS results*

Description

Returns squared correlation between observed and prior effects, for different subsets of SNPs (all, the ones having at least a moderate effects - p-value < 0.001 -, MR instruments)

Usage

```
get_RSquared_bGWAS(obj, SNPs = "all")
```

Arguments

`obj`	an object of class bGWAS created using `bGWAS()`
`SNPs,`	"all" / "moderate" / "instruments"

Value

a squared correlation

heatmap_bGWAS *Heatmap of SNP effects on prior traits from bGWAS results*

Description

Creates a heatmap of SNP effects on prior traits

Usage

```
heatmap_bGWAS(obj, SNPs = NULL, save_file = F, file_name = NULL)
```

Arguments

obj	an object of class bGWAS created using bGWAS()
SNPs	A data.frame containing the SNPs (rsid) to use in the first column, and optionnally the text that should be plotted in addition to rsid in the second column default=NULL.
save_file	A logical indicating if the graphic should be saved, default=FALSE, graphic will be displayed on the on-screen device
file_name	The name of the file saved (is save_file is TRUE) default=NULL, will used NameOfYourAnalysis_Heatmap.png

Value

a Heatmap

list_files *List prior GWASs files*

Description

Lists the filenames of the prior GWASs

Usage

```
list_files(IDs = NULL, Z_matrices = "~/ZMatrices/")
```

Arguments

IDs	the IDs of the studies to print, default="~/ZMatrices/" will list all of them (numeric),
Z_matrices	The path to the folder containing Z-Matrices, default="~/ZMatrices/" (character)

Value

List of files

list_priorGWASs *List prior GWASs*

Description

Lists the studies that can be used as prior GWASs

Usage

```
list_priorGWASs(IDs = NULL, Z_matrices = "~/ZMatrices/")
```

Arguments

IDs	the IDs of the studies to print, default="~/ZMatrices/" will list all of them (numeric),
Z_matrices	The path to the folder containing Z-Matrices, default="~/ZMatrices/" (character)

Value

a tibble containing prior GWASs information

list_traits *List prior GWASs traits Lists the traits of the prior GWASs*

Description

List prior GWASs traits Lists the traits of the prior GWASs

Usage

```
list_traits(Z_matrices = "~/ZMatrices/")
```

Arguments

Z_matrices	The path to the folder containing Z-Matrices, default="~/ZMatrices/" (character)

Value

List of traits

manhattan_plot_bGWAS *Manhattan Plot from bGWAS results*

Description

Creates a Manhattan Plot from bGWAS results (for performance, only SNPs with p-value or FDR < 0.05 are plotted)

Usage

```
manhattan_plot_bGWAS(obj, save_file = F, file_name = NULL,
  annotate = T, SNPs = NULL, results = "BF")
```

Arguments

obj	an object of class bGWAS created using bGWAS()
save_file	A logical indicating if the graphic should be saved, default=FALSE, graphic will be displayed on the on-screen device
file_name	The name of the file saved (is save_file is TRUE) default=NULL, will used NameOfYourAnalysis_ManhattanPlot.png
annotate	A logical indicating if the significant SNPs identified in the analysis should be annotated on the plot, default=TRUE If your results are not pruned or if you have a high number of significant SNPs, be aware that annotate=TRUE might decrease readability of the figure. You could define a set of SNPs to annotate using SNPs.
SNPs	A data.frame containing the SNPs (rsid) to annotate in the first column, and optionnally the text that should be plotted in the second column, and the color in the third column, default=NULL, only evaluated if annotate=TRUE.
results,	"BF" / "posterior" / "direct", default="BF"

Details

If results = "BF", BF p-values / fdr-values will be used.
If results = "direct", direct effect p-values / fdr-values will be used.
If results = "posterior", posterior effect p-values / fdr-values will be used.

Value

a Manhattan Plot

print.bGWAS *Print a bGWAS object*

Description

Print a bGWAS object

Usage

```
## S3 method for class 'bGWAS'
print(x, ...)
```

Arguments

x	an object of class bGWAS
...	further arguments

Value

print

print_log_bGWAS *Print log from bGWAS results*

Description

Prints the log (everything that is printed during a bGWAS analysis) with verbose=TRUE)

Usage

```
print_log_bGWAS(obj)
```

Arguments

obj	an object of class bGWAS created using bGWAS()

select_priorGWASs *Select prior GWASs*

Description

Allow the quick selection of a subset of prior GWASs based on 2 criteria. First, include all the files specified (if all including parameters are NULL, include all studies), and then remove all the files specified (if all excluding parameters are NULL, keep all studies included at the step before)

Usage

```
select_priorGWASs(include_files = NULL, include_traits = NULL,
  exclude_files = NULL, exclude_traits = NULL,
  Z_matrices = "~/ZMatrices/", verbose = F)
```

Arguments

include_files	list of file names (see list_files()) (character)
include_traits	list of trait (see list_traits()) (character)
exclude_files	list of file names (see list_files()) (character)
exclude_traits	list of trait (see list_traits()) (character)
Z_matrices	The path to the folder containing Z-Matrices, default="~/ZMatrices/" (character)
verbose	boolean, default = FALSE

Value

IDs (numeric) of studies that meet the criteria

Examples

```
  ## Not run:
  AllStudies = list_priorGWASs()
  list_traits()
 MyStudies = select_priorGWASs(include_traits=c("Heart Rate", "Body Mass Index", "Smoking"))
  AllStudies[AllStudies$ID %in% MyStudies, c("ID", "Name", "Trait", "File")]
## End(Not run)
```

SmallGWAS_Timmers2019 *Assocation results between genotypes and parental lifespan (LifeGen Consortium).*

Description

Subset of the original dataset containing the estimated effect of SNPs on parental survival

Usage

```
SmallGWAS_Timmers2019
```

Format

A data frame with 100000 rows and 5 variables:

rsid rsid of the SNP

a1 effect allele for the SNP

a0 reference allele for the SNP

beta estimated effect size for the SNP

se standard error of the estimated effect size for the SNP

Source

https://datashare.is.ed.ac.uk/handle/10283/3209

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