## Package 'bGWAS'

August 30, 2019

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

VignetteBuilder knitr

2 all.equal.bGWAS

## R topics documented:

	all.equal.bGWAS	2
	bGWAS	3
	coefficients_plot_bGWAS	5
	extract_MRcoeffs_bGWAS	6
	extract_results_bGWAS	6
	get_RSquared_bGWAS	7
	heatmap_bGWAS	8
	list_files	8
	list_priorGWASs	9
	list_traits	9
	manhattan_plot_bGWAS	10
	print.bGWAS	10
	print_log_bGWAS	11
	select_priorGWASs	11
	SmallGWAS_Timmers2019	12
Index		13

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 3

bGWAS	bGWAS - main function	
	J.	

#### **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 = 1, 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) #' @param prior_shrinkage The p-value threshold used for shrinkage before calculating the prior, should be between MR_threshold and 1, default=1e-5 (numeric)
stepwise_threshold		
		The p-value threshold used for inclusion/exclusion of Prior GWASs during the

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=1e-5 (numeric)

4 bGWAS

"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,
          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)

#### **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

 $\label{local_chrm_UK10K} \begin{subarray}{ll} chrm\_UK10K : chromosome (obtained from UK10K data) \\ pos\_UK10K : position (obtained from UK10K data) \\ \end{subarray}$ 

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 bG-WAS 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

8 list\_files

heatmap_bGWAS	Heatmap of SNP effects on prior traits from bGWAS results
rica ciliap_boti/to	Treatmap of Sivi effects on prior transfront of Wils results

#### **Description**

Creates a heatmap of SNP effects on prior traits

#### Usage

```
heatmap_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

 $Name Of Your Analysis\_Heatmap.png$ 

#### Value

a Heatmap

## 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/" (char-

acter)

#### Value

List of files

list\_priorGWASs 9

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/" (char-

acter)

#### 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

 $Z\_matrices \qquad \qquad The \ path \ to \ the \ folder \ containing \ Z-Matrices, \ default="\sim/ZMatrices/" \ (characteristic of the \ path) \ default="\simZMatrices/" \ (characteristic of the \ path)$ 

acter)

## Value

List of traits

10 print.bGWAS

### **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, 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.
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, ...)
```

print\_log\_bGWAS 11

#### 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

# **Index**

```
*Topic datasets
    SmallGWAS_Timmers2019, 12
all.equal.bGWAS, 2
bGWAS, 3, 5-8, 10, 11
coefficients_plot_bGWAS, 5
extract_MRcoeffs_bGWAS, 6
extract_results_bGWAS, 6
get_RSquared_bGWAS, 7
heatmap_bGWAS, 8
list_files, 8, 11
list_priorGWASs, 4, 9
list_traits, 9, 11
manhattan_plot_bGWAS, 10
print.bGWAS, 10
print_log_bGWAS, 11
select\_priorGWASs, 4, 11
SmallGWAS_Timmers2019, 12
```