

# Package ‘GxEprsDummy’

June 14, 2023

**Title** What the Package Does (One Line, Title Case)

**Version** 0.0.0.9000

**Description** What the package does (one paragraph).

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## R topics documented:

Bcov_discovery . . . . .	2
Bcov_target . . . . .	3
Bphe_discovery . . . . .	4
Bphe_target . . . . .	4
DummyData . . . . .	5
GWAS_binary . . . . .	5
GWAS_quantitative . . . . .	6
GWEIS_binary . . . . .	7
GWEIS_quantitative . . . . .	8
PRS_binary . . . . .	9
PRS_quantitative . . . . .	10
Qcov_discovery . . . . .	11
Qcov_target . . . . .	12
Qphe_discovery . . . . .	13
Qphe_target . . . . .	14
summary_permuted_binary . . . . .	14
summary_permuted_quantitative . . . . .	15
summary_regular_binary . . . . .	16
summary_regular_quantitative . . . . .	18
<b>Index</b>	<b>20</b>

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Bcov_discovery	<i>Covariate data file of the discovery dataset when the outcome is binary This contains covariate information of the individuals in the discovery dataset following confounders</i>
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**Description**

Covariate data file of the discovery dataset when the outcome is binary This contains covariate information of the individuals in the discovery dataset following confounders

**Usage**

```
Bcov_discovery
```

**Format**

A dataframe with 7916 rows and 18 columns

**Column 1** Family ID

**Column 2** Individual ID

**Column 3** Standardized covariate

**Column 4** Square of the standardized covariate

**Column 5** Confounder 1

**Column 6** Confounder 2

**Column 7** Confounder 3

**Column 8** Confounder 4

**Column 9** Confounder 5

**Column 10** Confounder 6

**Column 11** Confounder 7

**Column 12** Confounder 8

**Column 13** Confounder 9

**Column 14** Confounder 10

**Column 15** Confounder 11

**Column 16** Confounder 12

**Column 17** Confounder 13

**Column 18** Confounder 14

---

Bcov_target	<i>Covariate data file of the target dataset when the outcome is binary This contains covariate information of the individuals in the target dataset following confounders</i>
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### Description

Covariate data file of the target dataset when the outcome is binary This contains covariate information of the individuals in the target dataset following confounders

### Usage

```
Bcov_target
```

### Format

A dataframe with 1939 rows and 18 columns

**Column 1** Family ID

**Column 2** Individual ID

**Column 3** Standardized covariate

**Column 4** Square of the standardized covariate

**Column 5** Confounder 1

**Column 6** Confounder 2

**Column 7** Confounder 3

**Column 8** Confounder 4

**Column 9** Confounder 5

**Column 10** Confounder 6

**Column 11** Confounder 7

**Column 12** Confounder 8

**Column 13** Confounder 9

**Column 14** Confounder 10

**Column 15** Confounder 11

**Column 16** Confounder 12

**Column 17** Confounder 13

**Column 18** Confounder 14

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Bphe_discovery	<i>Phenotype data file of the discovery dataset when the outcome is binary This contains phenotype information of the individuals in the discovery dataset</i>
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### Description

Phenotype data file of the discovery dataset when the outcome is binary This contains phenotype information of the individuals in the discovery dataset

### Usage

Bphe\_discovery

### Format

A dataframe with 7916 rows and 3 columns

**Column 1** Family ID

**Column 2** Individual ID

**Column 3** Phenotype (1=controls, 2=cases)

---

Bphe_target	<i>Phenotype data file of the target dataset when the outcome is binary This contains phenotype information of the individuals in the target dataset</i>
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### Description

Phenotype data file of the target dataset when the outcome is binary This contains phenotype information of the individuals in the target dataset

### Usage

Bphe\_target

### Format

A dataframe with 1939 rows and 3 columns

**Column 1** Family ID

**Column 2** Individual ID

**Column 3** Phenotype (0=controls, 1=cases)

---

DummyData	<i>PLINK binary data files This contains DummyData.fam, DummyData.bim and DummyData.bed files, all in one</i>
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**Description**

PLINK binary data files This contains DummyData.fam, DummyData.bim and DummyData.bed files, all in one

**Usage**

```
DummyData
```

**Format**

The data files follow the general PLINK format

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GWAS_binary	<i>GWAS_binary function This function performs GWAS using plink2 and outputs the GWAS summary statistics file with additive SNP effects named B_out.trd.sum</i>
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**Description**

GWAS\_binary function This function performs GWAS using plink2 and outputs the GWAS summary statistics file with additive SNP effects named B\_out.trd.sum

**Usage**

```
GWAS_binary(
  plink_path,
  b_file,
  Bphe_discovery,
  Bcov_discovery,
  thread = 20,
  summary_output = "B_out.trd.sum"
)
```

**Arguments**

plink_path	Path to the PLINK executable application
b_file	Prefix of the binary files, where all .fam, .bed and .bim files have a common prefix
Bphe_discovery	Name (with file extension) of the phenotype file containing family ID, individual ID and phenotype of the discovery dataset as columns, without heading
Bcov_discovery	Name (with file extension) of the covariate file containing family ID, individual ID, standardized covariate, square of standardized covariate, and/or confounders of the discovery dataset as columns, without heading

thread            Number of threads used  
summary\_output    Name of the SNP effects of the GWAS summary statistics file specified by the user

### Value

This function will perform GWAS and output  
B\_out.trd.sum

### Examples

```
## Not run:
x <- GWAS_binary(plink_path, DummyData, Bphe_discovery, Bcov_discovery)
head(x)

## End(Not run)
```

---

GWAS\_quantitative    *GWAS\_quantitative function This function performs GWAS using plink2 and outputs the GWAS summary statistics file with additive SNP effects named Q\_out.trd.sum*

---

### Description

GWAS\_quantitative function This function performs GWAS using plink2 and outputs the GWAS summary statistics file with additive SNP effects named Q\_out.trd.sum

### Usage

```
GWAS_quantitative(
  plink_path,
  b_file,
  Qphe_discovery,
  Qcov_discovery,
  thread = 20,
  summary_output = "Q_out.trd.sum"
)
```

### Arguments

plink\_path    Path to the PLINK executable application  
b\_file        Prefix of the binary files, where all .fam, .bed and .bim files have a common prefix  
Qphe\_discovery    Name (with file extension) of the phenotype file containing family ID, individual ID and phenotype of the discovery dataset as columns, without heading  
Qcov\_discovery    Name (with file extension) of the covariate file containing family ID, individual ID, standardized covariate, square of standardized covariate, and/or confounders of the discovery dataset as columns, without heading

thread	Number of threads used
summary_output	Name of the SNP effects of the GWAS summary statistics file specified by the user

### Value

This function will perform GWAS and output

Q\_out.trd.sum

### Examples

```
## Not run:
x <- GWAS_quantitative(plink_path, DummyData, Qphe_discovery, Qcov_discovery)
head(x)

## End(Not run)
```

---

GWEIS_binary	<i>GWEIS_binary function This function performs GWEIS using plink2 and outputs the GWEIS summary statistics files with additive SNP effects named B_out.add.sum and interaction SNP effects named B_out.gxe.sum</i>
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### Description

GWEIS\_binary function This function performs GWEIS using plink2 and outputs the GWEIS summary statistics files with additive SNP effects named B\_out.add.sum and interaction SNP effects named B\_out.gxe.sum

### Usage

```
GWEIS_binary(
  plink_path,
  b_file,
  Bphe_discovery,
  Bcov_discovery,
  thread = 20,
  summary_output = "B_out"
)
```

### Arguments

plink_path	Path to the PLINK executable application
b_file	Prefix of the binary files, where all .fam, .bed and .bim files have a common prefix
Bphe_discovery	Phenotype file containing family ID, individual ID and phenotype of the discovery dataset as columns, without heading

`Bcov_discovery` Covariate file containing family ID, individual ID, standardized covariate, square of standardized covariate, and/or confounders of the discovery dataset as columns, without heading

`thread` Number of threads used

`summary_output` Name (prefix) of the additive or interaction SNP effects of the GWEIS summary statistics file specified by the user

## Value

This function will perform GWEIS and output

`B_out.add.sum`

`B_out.gxe.sum`

## Examples

```
## Not run:
x <- GWEIS_binary(plink_path, DummyData, Bphe_discovery, Bcov_discovery)
head(x[[1]])
head(x[[2]])

## End(Not run)
```

---

`GWEIS_quantitative` *GWEIS\_quantitative function This function performs GWEIS using plink2 and outputs the GWEIS summary statistics files with additive SNP effects named `Q_out.add.sum` and interaction SNP effects named `Q_out.gxe.sum`*

---

## Description

`GWEIS_quantitative` function This function performs GWEIS using plink2 and outputs the GWEIS summary statistics files with additive SNP effects named `Q_out.add.sum` and interaction SNP effects named `Q_out.gxe.sum`

## Usage

```
GWEIS_quantitative(
  plink_path,
  b_file,
  Qphe_discovery,
  Qcov_discovery,
  thread = 20,
  summary_output = "Q_out"
)
```



**Arguments**

<code>plink_path</code>	Path to the PLINK executable application
<code>b_file</code>	Prefix of the binary files, where all .fam, .bed and .bim files have a common prefix
<code>Qphe_discovery</code>	Phenotype file containing family ID, individual ID and phenotype of the discovery dataset as columns, without heading
<code>Qcov_discovery</code>	Covariate file containing family ID, individual ID, standardized covariate, square of standardized covariate, and/or confounders of the discovery dataset as columns, without heading
<code>thread</code>	Number of threads used
<code>summary_output</code>	Name (prefix) of the additive or interaction SNP effects of the GWEIS summary statistics file specified by the user

**Value**

This function will perform GWEIS and output

`Q_out.add.sum`

`Q_out.gxe.sum`

**Examples**

```
## Not run:
x <- GWEIS_quantitative(plink_path, DummyData, Qphe_discovery, Qcov_discovery)
head(x[[1]])
head(x[[2]])

## End(Not run)
```

---

<code>PRs_binary</code>	<i>PRs_binary function This function uses plink2 and outputs PRSs of each individual in the target dataset, using pre-generated GWAS and GWEIS summary statistics files named B_out.trd.sum, B_out.add.sum or B_out.gxe.sum</i>
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**Description**

`PRs_binary` function This function uses plink2 and outputs PRSs of each individual in the target dataset, using pre-generated GWAS and GWEIS summary statistics files named B\_out.trd.sum, B\_out.add.sum or B\_out.gxe.sum

**Usage**

```
PRS_binary(
  plink_path,
  b_file,
  summary_input = "B_out.trd.sum",
  summary_output = "B_trd"
)
```

**Arguments**

plink_path	Path to the PLINK executable application
b_file	Prefix of the binary files, where all .fam, .bed and .bim files have a common prefix
summary_input	Name of the summary statistics file specified by the user
summary_output	Name of the PRS file generated using provided summary statistics file specified by the user

**Value**

This function will output

B\_trd.sscore

**Examples**

```
## Not run:
x <- PRS_binary(plink_path, DummyData)
head(x[[1]])
head(x[[2]])
head(x[[3]])

## End(Not run)
```

---

PRS_quantitative	<i>PRS_binary function This function uses plink2 and outputs PRSs of each individual in the target dataset, using pre-generated GWAS and GWEIS summary statistics files named Q_out.trd.sum, Q_out.add.sum or Q_out.gxe.sum</i>
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---

**Description**

PRS\_binary function This function uses plink2 and outputs PRSs of each individual in the target dataset, using pre-generated GWAS and GWEIS summary statistics files named Q\_out.trd.sum, Q\_out.add.sum or Q\_out.gxe.sum

**Usage**

```
PRS_quantitative(
  plink_path,
  b_file,
  summary_input = "Q_out.trd.sum",
  summary_output = "Q_trd"
)
```

**Arguments**

plink_path	Path to the PLINK executable application
b_file	Prefix of the binary files, where all .fam, .bed and .bim files have a common prefix
summary_input	Name of the summary statistics file specified by the user
summary_output	Name of the PRS file generated using provided summary statistics file specified by the user

**Value**

This function will output

```
Q_trd.sscore
```

**Examples**

```
## Not run:
x <- PRS_quantitative(plink_path, DummyData)
head(x[[1]])
head(x[[2]])
head(x[[3]])

## End(Not run)
```

---

Qcov_discovery	<i>Covariate data file of the discovery dataset when the outcome is quantitative This contains covariate information of the individuals in the discovery dataset following confounders</i>
----------------	--

---

**Description**

Covariate data file of the discovery dataset when the outcome is quantitative This contains covariate information of the individuals in the discovery dataset following confounders

**Usage**

```
Qcov_discovery
```

**Format**

A dataframe with 6426 rows and 18 columns

**Column 1** Family ID

**Column 2** Individual ID

**Column 3** Standardized covariate

**Column 4** Square of the standardized covariate

**Column 5** Confounder 1

**Column 6** Confounder 2

**Column 7** Confounder 3

**Column 8** Confounder 4

**Column 9** Confounder 5

**Column 10** Confounder 6

**Column 11** Confounder 7

**Column 12** Confounder 8

**Column 13** Confounder 9

**Column 14** Confounder 10

**Column 15** Confounder 11

**Column 16** Confounder 12

**Column 17** Confounder 13

**Column 18** Confounder 14

---

Qcov_target	<i>Covariate data file of the target dataset when the outcome is quantitative This contains covariate information of the individuals in the target dataset following confounders</i>
-------------	--

---

**Description**

Covariate data file of the target dataset when the outcome is quantitative This contains covariate information of the individuals in the target dataset following confounders

**Usage**

Qcov\_target

**Format**

A dataframe with 1579 rows and 18 columns

**Column 1** Family ID

**Column 2** Individual ID

**Column 3** Standardized covariate

**Column 4** Square of the standardized covariate

**Column 5** Confounder 1

**Column 6** Confounder 2  
**Column 7** Confounder 3  
**Column 8** Confounder 4  
**Column 9** Confounder 5  
**Column 10** Confounder 6  
**Column 11** Confounder 7  
**Column 12** Confounder 8  
**Column 13** Confounder 9  
**Column 14** Confounder 10  
**Column 15** Confounder 11  
**Column 16** Confounder 12  
**Column 17** Confounder 13  
**Column 18** Confounder 14

---

Qphe_discovery	<i>Phenotype data file of the discovery dataset when the outcome is quantitative This contains phenotype information of the individuals in the discovery dataset</i>
----------------	--

---

## Description

Phenotype data file of the discovery dataset when the outcome is quantitative This contains phenotype information of the individuals in the discovery dataset

## Usage

```
Qphe_discovery
```

## Format

A dataframe with 6426 rows and 3 columns

**Column 1** Family ID  
**Column 2** Individual ID  
**Column 3** Phenotype

---

Qphe_target	<i>Phenotype data file of the target dataset when the outcome is quantitative This contains phenotype information of the individuals in the target dataset</i>
-------------	--

---

### Description

Phenotype data file of the target dataset when the outcome is quantitative This contains phenotype information of the individuals in the target dataset

### Usage

```
Qphe_target
```

### Format

A dataframe with 1579 rows and 3 columns

**Column 1** Family ID

**Column 2** Individual ID

**Column 3** Phenotype

---

summary_permuted_binary	<i>summary_permuted_binary function This function uses plink2 and outputs the p value of permuted model in the target dataset, using pre-generated GWAS and GWEIS summary statistics files named B_trd.sum, B_add.sum and B_gxe.sum</i>
-------------------------	---

---

### Description

summary\_permuted\_binary function This function uses plink2 and outputs the p value of permuted model in the target dataset, using pre-generated GWAS and GWEIS summary statistics files named B\_trd.sum, B\_add.sum and B\_gxe.sum

### Usage

```
summary_permuted_binary(
  Bphe_target,
  Bcov_target,
  iterations = 1000,
  add_score = "B_add.sscore",
  gxe_score = "B_gxe.sscore"
)
```

**Arguments**

Bphe_target	Phenotype file containing family ID, individual ID and phenotype of the target dataset as columns, without heading
Bcov_target	Covariate file containing family ID, individual ID, standardized covariate, square of standardized covariate, and/or confounders of the target dataset as columns, without heading
iterations	Number of iterations used in permutation
add_score	The .sscore file generated using additive SNP effects of GWEIS summary statistics
gxe_score	The .sscore file generated using interaction SNP effects of GWEIS summary statistics

**Value**

This function will output

B\_permuted\_p.txt

**Examples**

```
## Not run:
x <- summary_permuted_binary(Bphe_target, Bcov_target)
x

## End(Not run)
```

---

```
summary_permuted_quantitative
```

*summary\_permuted\_quantitative function This function uses plink2 and outputs the p value of permuted model in the target dataset, using pre-generated GWAS and GWEIS summary statistics files named Q\_trd.sum, Q\_add.sum and Q\_gxe.sum*

---

**Description**

summary\_permuted\_quantitative function This function uses plink2 and outputs the p value of permuted model in the target dataset, using pre-generated GWAS and GWEIS summary statistics files named Q\_trd.sum, Q\_add.sum and Q\_gxe.sum

**Usage**

```
summary_permuted_quantitative(
  Qphe_target,
  Qcov_target,
  iterations = 1000,
  add_score = "Q_add.sscore",
  gxe_score = "Q_gxe.sscore"
)
```

**Arguments**

Qphe_target	Phenotype file containing family ID, individual ID and phenotype of the target dataset as columns, without heading
Qcov_target	Covariate file containing family ID, individual ID, standardized covariate, square of standardized covariate, and/or confounders of the target dataset as columns, without heading
iterations	Number of iterations used in permutation
add_score	The .sscore file generated using additive SNP effects of GWEIS summary statistics
gxe_score	The .sscore file generated using interaction SNP effects of GWEIS summary statistics

**Value**

This function will output

Q\_permuted\_p.txt

**Examples**

```
## Not run:
x <- summary_permuted_quantitative(Qphe_target, Qcov_target)
x

## End(Not run)
```

---

summary\_regular\_binary

*summary\_regular\_binary function This function uses plink2 and outputs the summary of regular model in the target dataset, using pre-generated GWAS and GWEIS summary statistics files named B\_trd.sum, B\_add.sum and B\_gxe.sum*

---

**Description**

summary\_regular\_binary function This function uses plink2 and outputs the summary of regular model in the target dataset, using pre-generated GWAS and GWEIS summary statistics files named B\_trd.sum, B\_add.sum and B\_gxe.sum

**Usage**

```
summary_regular_binary(
  Bphe_target,
  Bcov_target,
  trd_score = "B_trd.sscore",
  add_score = "B_add.sscore",
  gxe_score = "B_gxe.sscore",
  Model,
  summary_output = "Bsummary.txt",
  risk_output = "Individual_risk_values.txt"
)
```



**Arguments**

<code>Bphe_target</code>	Phenotype file containing family ID, individual ID and phenotype of the target dataset as columns, without heading
<code>Bcov_target</code>	Covariate file containing family ID, individual ID, standardized covariate, square of standardized covariate, and/or confounders of the target dataset as columns, without heading
<code>trd_score</code>	The .sscore file generated using additive SNP effects of GWAS summary statistics
<code>add_score</code>	The .sscore file generated using additive SNP effects of GWEIS summary statistics
<code>gxe_score</code>	The .sscore file generated using interaction SNP effects of GWEIS summary statistics
<code>Model</code>	Specify the model number (1: $y = \text{PRS\_trd} + E + \text{PRS\_trd} \times E + \text{confounders}$ , 2: $y = \text{PRS\_add} + E + \text{PRS\_add} \times E + \text{confounders}$ , 3: $y = \text{PRS\_add} + E + \text{PRS\_gxe} \times E + \text{confounders}$ , 4: $y = \text{PRS\_add} + E + \text{PRS\_gxe} + \text{PRS\_gxe} \times E + \text{confounders}$ , 5: $y = \text{PRS\_add} + E + E^2 + \text{PRS\_gxe} + \text{PRS\_gxe} \times E + \text{confounders}$ , where $y$ is the outcome variable, $E$ is the covariate of interest, $\text{PRS\_trd}$ and $\text{PRS\_add}$ are the polygenic risk scores computed using additive SNP effects of GWAS and GWEIS summary statistics respectively, and $\text{PRS\_gxe}$ is the polygenic risk scores computed using GxE interaction SNP effects of GWEIS summary statistics.)
<code>summary_output</code>	Name of the model summary file specified by the user
<code>risk_output</code>	Name of the file containing risk scores of the target individuals specified by the user

**Value**

This function will output

`Bsummary.txt`  
`Individual_risk_values.txt`

**Examples**

```
## Not run:
x <- summary_regular_binary(Bphe_target, Bcov_target, Model = 5)
x[[1]][[1]]
x[[1]][[2]]
x[[1]][[3]]
x[[1]][[4]]
x[[1]][[5]]
x[[1]][[6]]
x[[1]][[7]]
x[[1]][[8]]
x[[1]][[9]]
x[[1]][[10]]
x[[1]][[11]]
x[[1]][[12]]
x[[1]][[13]]
x[[1]][[14]]
```

```
x[[1]][[15]]
x[[1]][[16]]
x[[1]][[17]]
x[[1]][[18]]
head(x[[2]])

## End(Not run)
```

---

```
summary_regular_quantitative
```

*summary\_regular\_quantitative function This function uses plink2 and outputs the summary of regular model in the target dataset, using pre-generated GWAS and GWEIS summary statistics files named Q\_trd.sum, Q\_add.sum and Q\_gxe.sum*

---

## Description

summary\_regular\_quantitative function This function uses plink2 and outputs the summary of regular model in the target dataset, using pre-generated GWAS and GWEIS summary statistics files named Q\_trd.sum, Q\_add.sum and Q\_gxe.sum

## Usage

```
summary_regular_quantitative(
  Qphe_target,
  Qcov_target,
  trd_score = "Q_trd.sscore",
  add_score = "Q_add.sscore",
  gxe_score = "Q_gxe.sscore",
  Model,
  summary_output = "Qsummary.txt",
  risk_output = "Individual_risk_values.txt"
)
```

## Arguments

Qphe_target	Phenotype file containing family ID, individual ID and phenotype of the target dataset as columns, without heading
Qcov_target	Covariate file containing family ID, individual ID, standardized covariate, square of standardized covariate, and/or confounders of the target dataset as columns, without heading
trd_score	The .sscore file generated using additive SNP effects of GWAS summary statistics
add_score	The .sscore file generated using additive SNP effects of GWEIS summary statistics
gxe_score	The .sscore file generated using interaction SNP effects of GWEIS summary statistics
Model	Specify the model number (1: $y = \text{PRS\_trd} + E + \text{PRS\_trd} \times E + \text{confounders}$ , 2: $y = \text{PRS\_add} + E + \text{PRS\_add} \times E + \text{confounders}$ , 3: $y = \text{PRS\_add} + E + \text{PRS\_gxe} \times E + \text{confounders}$ , 4: $y = \text{PRS\_add} + E + \text{PRS\_gxe} + \text{PRS\_gxe} \times$

E + confounders, where y is the outcome variable, E is the covariate of interest, PRS\_trd and PRS\_add are the polygenic risk scores computed using additive SNP effects of GWAS and GWEIS summary statistics respectively, and PRS\_gxe is the polygenic risk scores computed using GxE interaction SNP effects of GWEIS summary statistics.)

summary\_output

Name of the model summary file specified by the user

risk\_output

Name of the file containing risk scores of the target individuals specified by the user

## Value

This function will output

Qsummary.txt

Individual\_risk\_values.txt

## Examples

```
## Not run:
x <- summary_regular_quantitative(Qphe_target, Qcov_target, Model = 4)
x[[1]][[1]]
x[[1]][[2]]
x[[1]][[3]]
x[[1]][[4]]
x[[1]][[5]]
x[[1]][[6]]
x[[1]][[7]]
x[[1]][[8]]
x[[1]][[9]]
x[[1]][[10]]
x[[1]][[11]]
x[[1]][[12]]
head(x[[2]])

## End(Not run)
```

# Index

- \* **datasets**
  - Bcov\_discovery, [2](#)
  - Bcov\_target, [3](#)
  - Bphe\_discovery, [4](#)
  - Bphe\_target, [4](#)
  - DummyData, [5](#)
  - Qcov\_discovery, [11](#)
  - Qcov\_target, [12](#)
  - Qphe\_discovery, [13](#)
  - Qphe\_target, [14](#)
- \* **gwas**
  - GWAS\_binary, [5](#)
  - GWAS\_quantitative, [6](#)
- \* **gwies,**
  - GWEIS\_binary, [7](#)
  - GWEIS\_quantitative, [8](#)
- \* **gxe**
  - GWEIS\_binary, [7](#)
  - GWEIS\_quantitative, [8](#)
- \* **interaction,**
  - GWEIS\_binary, [7](#)
  - GWEIS\_quantitative, [8](#)
- \* **profile**
  - PRS\_binary, [9](#)
  - PRS\_quantitative, [10](#)
- \* **prs,**
  - PRS\_binary, [9](#)
  - PRS\_quantitative, [10](#)
- \* **regression**
  - summary\_permuted\_binary, [14](#)
  - summary\_permuted\_quantitative, [15](#)
  - summary\_regular\_binary, [16](#)
  - summary\_regular\_quantitative, [18](#)
- \* **risk**
  - summary\_regular\_binary, [16](#)
  - summary\_regular\_quantitative, [18](#)
- \* **scores**
  - PRS\_binary, [9](#)
  - PRS\_quantitative, [10](#)
  - summary\_regular\_binary, [16](#)
  - summary\_regular\_quantitative, [18](#)
- \* **summary**
  - summary\_permuted\_binary, [14](#)
  - summary\_permuted\_quantitative, [15](#)
  - summary\_regular\_binary, [16](#)
  - summary\_regular\_quantitative, [18](#)
- Bcov\_discovery, [2](#)
- Bcov\_target, [3](#)
- Bphe\_discovery, [4](#)
- Bphe\_target, [4](#)
- DummyData, [5](#)
- GWAS\_binary, [5](#)
- GWAS\_quantitative, [6](#)
- GWEIS\_binary, [7](#)
- GWEIS\_quantitative, [8](#)
- PRS\_binary, [9](#)
- PRS\_quantitative, [10](#)
- Qcov\_discovery, [11](#)
- Qcov\_target, [12](#)
- Qphe\_discovery, [13](#)
- Qphe\_target, [14](#)
- summary\_permuted\_binary, [14](#)
- summary\_permuted\_quantitative, [15](#)
- summary\_regular\_binary, [16](#)
- summary\_regular\_quantitative, [18](#)