# Package 'REBET'

November 29, 2024

**Description** There is an increasing focus to investigate the association between rare variants

Title The subREgion-based BurdEn Test (REBET)

**Version** 1.24.0 **Date** 2018-08-13

and diseases. The REBET package implements the subREgion-based BurdEn Test which is a powerful burden test that simultaneously identifies susceptibility loci and sub-regions.
Imports stats, utils
Depends ASSET
Suggests RUnit, BiocGenerics
License GPL-2
biocViews Software, VariantAnnotation, SNP
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data

Data for the example

## **Description**

Data for the example.

#### **Details**

The data contains a binary phenotype vector response, a genotype matrix genotypes consisting of 20 rare-variant SNPs, and the sub-region annotation vector subRegions for the rebet example.

#### See Also

rebet

### **Examples**

```
data(data, package="REBET")
# Display some of the data
table(response)
dim(genotypes)
subRegions
```

REBET

The REBET package

## **Description**

An R package for the subREgion-based BurdEn Test (REBET).

## **Details**

In rare-variant association studies, aggregating rare and/or low frequency variants, may increase statistical power for detection of the underlying susceptibility gene or region. However, it is unclear which variants, or class of them, in a gene contribute most to the association. This subregion-based burden test (REBET) simultaneously selects susceptibility genes and identifies important underlying sub-regions. The sub-regions are predefined based on shared common biologic characteristics, such as the protein domain or possible functional impact. Based on a subset-based approach considering local correlations between combinations of test statistics of sub-regions, REBET is able to properly control the type I error rate while adjusting for multiple comparisons in a computationally efficient manner. See the reference for the details of this test. The main function in this package is rebet, which performs the REBET test.

## Author(s)

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#### References

Zhu, B., Mirabello, L., Chatterjee, N. (2018) A Subregion-based Burden Test for Simultaneous Identification of Susceptibility Loci and Sub-regions within Genetic Epidemiology. In press. https://doi.org/10.1002/gepi

rebet The subREgion-based BurdEn Test (REBET)
rebet The subREgion-based BurdEn Test (REBET)

## Description

A Subregion-based Burden Test for Simultaneous Identification of Susceptibility Loci and Subregions within

## Usage

## **Arguments**

response	Numerical vector of phenotypes. A binary phenotype must be coded as 0 and 1.
genotypes	Matrix of genotypes with each column as a locus.
subRegions	Sub-region annotation vector with length equal to the number of columns of genotypes. In the returned object, these regions will appear as paste("Region_", subRegions, sep="").
responseType	NULL, "continuous" or "binary". If NULL, then "continuous" or "binary" will be chosen based on Y. The default is NULL.
covariates	NULL or matrix of covariates. The default is NULL.
shape1	The shape1 parameter in the beta distribution. The default is 1.
shape2	The shape2 parameter in the beta distribution. The default is 1.
saveMem	TRUE or FALSE to conserve memory (see details). The default is FALSE.

## Details

See the reference for details of this method.

Missing values in any of response, genotypes or covariates will be removed before the analysis. Setting saveMem to TRUE will allow for the analysis of a much larger number of subjects, but will take more time to compute. When saveMem is FALSE, there needs to be enough memory available to hold two or three NxN matrices, where N is the number of subjects.

This function calls the h. traits function in the ASSET package.

## Value

The object returned from h.traits in the ASSET package.

## Author(s)

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## References

Zhu, B., Mirabello, L., Chatterjee, N. (2018) A Subregion-based Burden Test for Simultaneous Identification of Susceptibility Loci and Sub-regions within Genetic Epidemiology. In press. https://doi.org/10.1002/gepi

## **Examples**

```
data(data, package="REBET")
res <- rebet(response, genotypes, subRegions)
h.summary(res)</pre>
```

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