

Package ‘QTL.gCIMapping’

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Type Package

Title QTL Genome-Wide Composite Interval Mapping

Version 1.0

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Description Conduct multi-Quantitative Trait Locus (QTL) mapping under the framework of random-QTL-effect mixed linear model. First, each position on the genome is detected in order to construct a negative logarithm P-value curve against genome position. Then, all the peaks on each effect (additive or dominant) curve are viewed as potential QTL, all the effects of the potential QTL are included in a multi-QTL model, their effects are estimated by empirical Bayes in doubled haploid or by adaptive lasso in F2, and true QTL are identified by likelihood ratio test. Wang S-B, Wen Y-J, Ren W-L, Ni Y-L, Zhang J, Feng J-Y, Zhang Y-M (2016) <doi:10.1038/srep29951>.

Depends MASS, qtl, doParallel, foreach, parallel

Imports methods, openxlsx, stringr, data.table, parcor

License GPL (>= 2)

NeedsCompilation no

Repository CRAN

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f2data	<i>F2 example data</i>
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Description

GCIM format of F2 dataset.

Usage

```
data(f2data)
```

Details

Dataset input of file for QTL.gCIMapping function.

Author(s)

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QTL.gCIMapping	<i>QTL Genome-Wide Composite Interval Mapping</i>
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Description

Conduct multi-QTL mapping under the framework of random-QTL-effect mixed linear model. First, each position on the genome is detected in order to construct a negative logarithm P-value curve against genome position. Then, all the peaks on each effect (additive or dominant) curve are viewed as potential QTL, all the effects of the potential QTL are included in a multi-QTL model, their effects are estimated by empirical Bayes in doubled haploid or by adaptive lasso in F2, and true QTL are identified by likelihood ratio test.

Usage

```
QTL.gCIMapping(file, fileFormat, fileICIMcov, Population, Model, WalkSpeed, CriLOD, Likelihood, flagqrtl, DrawPlot, PlotFormat, Resolution, Trait, dir)
```

Arguments

file	File path and name in your computer.
fileFormat	Format for input file (GCIM, ICIM, Cart).
fileICIMcov	File path and name in your computer.
Population	BC1, BC2, DH, RIL, F2.
Model	Random (random model) or Fixed (fixed model) for QTL effects.

WalkSpeed	Walk speed for Genome-wide Scanning.(WalkSpeed=1)
CriLOD	Critical LOD scores for significant QTL (CriLOD=2.5).
Likelihood	This parameter is only for F2 population, including restricted maximum likelihood (REML) and maximum likelihood (ML).
flagrqtl	This parameter is only for F2 population, flagrqtl="FALSE" in the first running. If the other software detects only one QTL in a neighborhood but this software finds two linked QTLs (one with additive effect and another with dominant effect) in the region, let flagrqtl="TRUE"
DrawPlot	This parameter is for all the populations, including FALSE and TRUE, DrawPlot=FALSE indicates no figure output, DrawPlot=TRUE indicates the output of the figure against genome position.
PlotFormat	This parameter is for all the figure files, including *.jpeg, *.png, *.tiff and *.pdf.
Resolution	This parameter is for all the figure files, including Low and High.
Trait	Trait=1:3 indicates the analysis from the first trait to the third trait.
dir	This parameter is for the save path.

Details

Package: QTL.gCIMapping
 Type: Package
 Version: 1.0
 Date: 2018-4-12
 Depends: MASS,dplyr,parcor,qtl,doParallel
 Imports: methods,openxlsx,stringr
 License: GPL version 2 or newer
 LazyLoad: yes

Author(s)

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References

Mapping small-effect and linked quantitative trait loci for complex traits in backcross or DH populations via a multi-locus GWAS methodology. Wang Shi-Bo, Wen Yang-Jun, Ren Wen-Long, Ni Yuan-Li, Zhang Jin, Feng Jian-Ying, Zhang Yuan-Ming*

Examples

```

G=data(f2data)
QTL.gCIMapping(file=f2data,fileFormat="GCIM",fileICIMcov=NULL,Population="F2",
Model="Random",WalkSpeed=1,CriLOD=2.5,Likelihood="REML",flagrqtl="FALSE",

```

```
DrawPlot="FALSE",PlotFormat="png",Resolution="Low",Trait=1:1,dir=tempdir())
```

WangF

To perform QTL mapping with wang method

Description

Genome-wide Composite Interval Mapping

Usage

```
WangF(pheRaw, genRaw, mapRaw1, yygg1, cov_en, Population, WalkSpeed, CriLOD, dir)
```

Arguments

pheRaw	phenotype matrix.
genRaw	genotype matrix.
mapRaw1	linkage map matrix.
yygg1	the transformed covariate matrix .
cov_en	raw covariate matrix.
Population	population flag.
WalkSpeed	Walk speed for Genome-wide Scanning.(WalkSpeed=1).
CriLOD	Critical LOD scores for significant QTL (CriLOD=2.5).
dir	file path in your computer.

Author(s)

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WenF

To perform QTL mapping with Wen method

Description

An efficient multi-locus mixed model framework for the detection of small and linked QTLs in F2

Usage

```
WenF(pheRaw, genRaw, mapRaw1, yygg1, cov_en, WalkSpeed, CriLOD, dir)
```

Arguments

pheRaw	phenotype matrix.
genRaw	genotype matrix.
mapRaw1	linkage map matrix.
yygg1	the transformed covariate matrix .
cov_en	raw covariate matrix.
WalkSpeed	Walk speed for Genome-wide Scanning.(WalkSpeed=1).
CriLOD	Critical LOD scores for significant QTL (CriLOD=2.5).
dir	file path in your computer.

Author(s)

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