

# Package ‘dQTG.seq’

July 22, 2025

**Type** Package

**Title** A BSA Software for Detecting All Types of QTLs in BC, DH, RIL and F2

**Version** 1.0.2

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**Description** The new (dQTG.seq1 and dQTG.seq2) and existing (SmoothLOD, G', deltaSNP and ED) bulked segregant analysis methods are used to identify various types of quantitative trait loci for complex traits via extreme phenotype individuals in bi-parental segregation populations (F2, backcross, doubled haploid and recombinant inbred line). The numbers of marker alleles in extreme low and high pools are used in existing methods to identify trait-related genes, while the numbers of marker alleles and genotypes in extreme low and high pools are used in the new methods to construct a new statistic Gw for identifying trait-related genes. dQTG.seq2 is feasible to identify extremely over-dominant and small-effect genes in F2. Li P, Li G, Zhang YW, Zuo JF, Liu JY, Zhang YM (2022, <[doi:10.1016/j.xplc.2022.100319](https://doi.org/10.1016/j.xplc.2022.100319)>).

**Depends** R (>= 3.5.0)

**License** GPL (>= 2)

**Imports** BB, data.table, doParallel, openxlsx, qtl, stringr, writexl, vroom, parallel, foreach

**Encoding** UTF-8

**LazyData** true

**RoxygenNote** 7.2.3

**NeedsCompilation** no

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**Repository** CRAN

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|     |                 |
|-----|-----------------|
| BSA | <i>BSA data</i> |
|-----|-----------------|

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Description

BSA format of F2 dataset.

Usage

data(BSA)

Details

Input file for dQTG.seq function.

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|        |                          |
|--------|--------------------------|
| Dodata | <i>To perform method</i> |
|--------|--------------------------|

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Description

To perform method

Usage

Dodata(dir, calculatedata, chr, color1, CL0)

Arguments

- |               |                             |
|---------------|-----------------------------|
| dir           | the path of the output      |
| calculatedata | the inputdata               |
| chr           | chromosome                  |
| color1        | the color of the chromosome |
| CL0           | the numbers of CPU          |

Value

list

Examples

```
data(BSA)
dir<- tempdir()
data.calculatedata<-Readdata1(BSA)
Dodata(dir,calculatedata=data.calculatedata,chr="all",color1="blue",CLO=1)
```

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|          |                                       |
|----------|---------------------------------------|
| dQTG.seq | <i>Title The function of dQTG.seq</i> |
|----------|---------------------------------------|

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Description

Title The function of dQTG.seq

Usage

```
dQTG.seq(dir, filegen, chr, color, CLO)
```

Arguments

- dir                    the path of the output
- filegen                the input data
- chr                    the chromosome
- color                  the color
- CLO                    the numbers of CPU

Value

list

Examples

```
data(BSA)
dQTG.seq(dir=tempdir(),filegen=BSA,chr="all",color="blue",CLO=1)
```

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|           |                                |
|-----------|--------------------------------|
| Readdata1 | <i>Title readdata function</i> |
|-----------|--------------------------------|

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**Description**

Title readdata function  
Title readdata function

**Usage**

Readdata1(File)  
  
Readdata1(File)

**Arguments**

File                    the input file

**Value**

list  
list

**Examples**

data(BSA)  
Readdata1(BSA)  
data(BSA)  
Readdata1(BSA)

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