

Package ‘ldblock’

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Title data structures for linkage disequilibrium measures in populations

Version 1.6.0

Author VJ Carey <stvjc@channing.harvard.edu>

Description Define data structures for linkage disequilibrium measures in populations.

Suggests RUnit, BiocGenerics, knitr

Imports Matrix, snpStats, erma, VariantAnnotation, GenomeInfoDb, Rsamtools, GO.db, GenomicFiles

Depends R (>= 3.1), methods, Homo.sapiens

Maintainer VJ Carey <stvjc@channing.harvard.edu>

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LazyLoad yes

BiocViews genetics, SNP, GWAS, LinkageDisequilibrium

VignetteBuilder knitr

NeedsCompilation no

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lblockquote-package*data structures for linkage disequilibrium measures in populations*

Description

Define data structures for linkage disequilibrium measures in populations.

Details

The DESCRIPTION file:

| | |
|------------------|---|
| Package: | lblockquote |
| Title: | data structures for linkage disequilibrium measures in populations |
| Version: | 1.6.0 |
| Author: | VJ Carey <stvjc@channing.harvard.edu> |
| Description: | Define data structures for linkage disequilibrium measures in populations. |
| Suggests: | RUnit, BiocGenerics, knitr |
| Imports: | Matrix,.snpStats, erma, VariantAnnotation, GenomeInfoDb, Rsamtools, GO.db, GenomicFiles |
| Depends: | R (>= 3.1), methods, Homo.sapiens |
| Maintainer: | VJ Carey <stvjc@channing.harvard.edu> |
| License: | Artistic-2.0 |
| LazyLoad: | yes |
| BiocViews: | genetics, SNP, GWAS, LinkageDisequilibrium |
| VignetteBuilder: | knitr |

Index of help topics:

| | |
|---------------------|---|
| downloadPopByChr | download hapmap resource with LD estimates |
| expandSnpSet | Given a set of SNP identifiers, use LD to expand the set to include linked loci |
| hmld | import hapmap LD data and create a structure for its management |
| ldByGene | obtain LD statistics in region specified by a gene model |
| lblockquote-package | data structures for linkage disequilibrium measures in populations |
| ldstruct-class | Class '"ldstruct" |
| s3_1kg | Create a URL referencing 1000 genomes content in AWS S3 |

Author(s)

VJ Carey <stvjc@channing.harvard.edu>

Maintainer: VJ Carey <stvjc@channing.harvard.edu>

Examples

```
# see vignette
```

| | |
|------------------|---|
| downloadPopByChr | <i>download hapmap resource with LD estimates</i> |
|------------------|---|

Description

download hapmap resource with LD estimates

Usage

```
downloadPopByChr(chrname = "chr1",
  popname = "CEU",
  urlTemplate = "http://hapmap.ncbi.nlm.nih.gov/downloads/ld_data/2009-02_phaseIII_r2/ld_%%CHRN%%"
  targfolder = Sys.getenv("LDBLOCK_TXTGZ_DIR"))
```

Arguments

| | |
|-------------|---|
| chrname | UCSC format tag for chromosome |
| popname | hapmap three letter code for population, e.g. 'CEU' |
| urlTemplate | pattern for creating URL given chr and pop |
| targfolder | destination |

Details

delivers HapMap LD data to 'targfolder'

Value

just run for side effect of download.file

Examples

```
## Not run:
downloadPopByChr()

## End(Not run)
```

| | |
|--------------|--|
| expandSnpSet | <i>Given a set of SNP identifiers, use LD to expand the set to include linked loci</i> |
|--------------|--|

Description

Given a set of SNP identifiers, use LD to expand the set to include linked loci

Usage

```
expandSnpSet(rsl, lb = 0.8, ldstruct, chrn = "chr17", popn = "CEU",
  txtgfn = dir(system.file("hapmap", package = "lblockquote"), full.names = TRUE))
```

Arguments

| | |
|-----------------------|--|
| <code>rsl</code> | input list – SNPs not found in the LD structure are simply returned along with those found, and the expansion list, all combined in a vector |
| <code>lb</code> | lower bound on statistic used to retrieve loci in LD |
| <code>ldstruct</code> | instance of ldstruct-class |
| <code>chrn</code> | chromosome identifier |
| <code>popn</code> | population identifier (one of 'CEU', 'MEX', ...) |
| <code>txtgfn</code> | path to gzipped hapmap file with LD information |

Details

direct use of elementwise arithmetic comparison

Value

character vector

Note

As of 2015, it appears that locus names are more informative than addresses for determining SNP identity across resources.

Examples

```
og = Sys.getenv("LDBLOCK_TXTGZ_DIR")
on.exit( Sys.setenv("LDBLOCK_TXTGZ_DIR" = og) )
Sys.setenv("LDBLOCK_TXTGZ_DIR"=system.file("hapmap", package=" ldblock"))
ld17 = hmld(chr="chr17", pop="CEU")
ee = expandSnpSet( ld17@allrs[1:10], ldstruct = ld17 )
```

`hmld`

import hapmap LD data and create a structure for its management

Description

import hapmap LD data and create a structure for its management

Usage

```
hmld(hmgztxt, poptag, chrom, genome = "hg19", stat = "Dprime")
```

Arguments

| | |
|----------------------|---|
| <code>hmgztxt</code> | name of gzipped text file as distributed at hapmap.ncbi.nlm.nih.gov/downloads/1d_data/2009-02_phaseIII_r2/ . It will be processed by read.delim . |
| <code>poptag</code> | heuristic tag identifying population |
| <code>chrom</code> | heuristic tag for chromosome name |
| <code>genome</code> | genome tag |
| <code>stat</code> | statistic to use, "Dprime", "R2", and "LOD" are options |

Details

generates a sparse matrix representation of pairwise LD statistics and binds metadata on variant name and position

Value

instance of ldstruct class

Examples

```
getClass("ldstruct")
# see vignette
```

ldByGene*obtain LD statistics in region specified by a gene model*

Description

Obtain LD statistics in region specified by a gene model.

Usage

```
ldByGene(sym = "MMP24",
         vcf = system.file("vcf/c20exch.vcf.gz", package = "gQTLstats"),
         flank = 1000,
         vcfSLS = "NCBI",
         genomeSLS = "hg19",
         stats = "D.prime", depth = 10)
```

Arguments

| | |
|-----------|---|
| sym | A standard gene symbol for use with genemodel |
| vcf | Path to a tabix-indexed VCF file |
| flank | number of basepairs to flank gene model for search |
| vcfSLS | seqlevelsStyle (SLS) token for VCF; will be imposed on gene model |
| genomeSLS | character tag for genome, to be used with readVcf |
| stats | passed to ld |
| depth | passed to ld |

Value

sparse matrix representation of selected LD statistic, as returned by [ld](#)

Examples

```
ld1 = ldByGene(depth=150)
image(ld1[1:200,1:200], col.reg=heat.colors(120), colorkey=TRUE,
      main="SNPs in MMP24 (chr20)")
```

ldstruct-class *Class "ldstruct"*

Description

Manage information about LD statistics as reported by HapMap.

Objects from the Class

Objects can be created by calls of the form `new("ldstruct", ...)`.

Slots

ldmat: Object of class "dsCMatrix" sparse representation of statistics
chrom: Object of class "character" chromosome tag in UCSC format
genome: Object of class "character" genome tag
allpos: Object of class "numeric" coordinates
poptag: Object of class "character" hapmap founder population tag, 'CEU', 'MEX' etc.
statInUse: Object of class "character" code for statistic retrieved, one of 'Dprime', 'LOD', 'R2'
allrs: Object of class "character" all SNP identifiers, sometimes in affy format

Methods

ldmat `signature(x = "ldstruct")`: extract sparse matrix

Examples

```
showClass("ldstruct")
```

s3_1kg *Create a URL referencing 1000 genomes content in AWS S3*

Description

Create a URL referencing 1000 genomes content in AWS S3.

Usage

```
s3_1kg(chrnum, tag = "20130502", wrap = function(x) TabixFile(x), tmpl = NULL, dropchr = TRUE)
stack1kg(chrs=as.character(1:22))
```

Arguments

| | |
|---------|--|
| chrnum | a character string denoting a chromosome, such as '22' |
| chrs | a vector of chromosome names for extraction from 1000 genomes VCF collection |
| tag | a character string identifying the version, ignored if <code>tmpl</code> is non-null; valid tag values are the default or "20101123" |
| wrap | The URL is returned after evaluating <code>wrap</code> on it; default is useful when Tabix indexing is to be used |
| tmpl | alternate template for full URL, useful if versions prior to 2010 are of interest |
| dropchr | if TRUE <code>chrnum</code> will have 'chr' removed if present |

Details

`stack1kg` produces a `VcfStack` instance with references to VCF for 1000 genomes autosomal chrs. S3-resident VCF files with version "v5a.20130502" are used.

Value

by default, a `TabixFile` instance

Examples

```
s3_1kg("22")
## Not run:
require(VariantAnnotation)
scanVcfHeader(s3_1kg("22"))

## End(Not run)
```

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