

Package ‘gwascat’

October 2, 2013

Title representing and modeling data in the NHGRI GWAS catalog

Version 1.4.0

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

Description representing and modeling data in the NHGRI GWAS catalog

Enhances SNPlocs.Hsapiens.dbSNP.20111119, pd.genomewidesnp.6

Depends R (>= 2.14.0), methods, IRanges, GenomicRanges, snpStats, graph, BiocGenerics

Imports Biostrings

Suggests DO.db, Gviz, ggbio, rtracklayer

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

License Artistic-2.0

biocViews genetics

LazyLoad yes

R topics documented:

gwascat-package	2
gwaswloc-class	3
gwcex2gviz	4
gwdf_2012_02_02	5
locon6	6
makeCurrentGwascat	7
obo2graphNEL	8
riskyAlleleCount	9
topTraits	10
traitsManh	11
Index	12

gwascapackage

representing and modeling data in the NHGRI GWAS catalog

Description

representing and modeling data in the NHGRI GWAS catalog, using GRanges and allied infrastructure

Details

Package: gwascap
Version: 0.0.3
Suggests:
Depends: R (>= 2.14.0), methods, IRanges, GenomicRanges
Imports:
License: Artistic-2.0
LazyLoad: yes
Built: R 2.15.0; ; 2012-02-10 21:08:32 UTC; unix

Index:

gwaswloc-class Class "gwaswloc"

Upon attachment, a [GRanges-class](#) structure call gwrngs is formed which can be interrogated by position or through use of element metadata to learn about catalogued GWAS associations.

The data objects

'g17SM' 'gg17N' 'gw6.rs_17' 'low17' 'rules_6.0_1kg_17'

are described in vignettes.

Author(s)

VJ Carey <stvjc@channing.harvard.edu>

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

References

<http://www.genome.gov/gwastudies/>.

Partial support from the Computational Biology Group at Genentech, Inc.

Examples

```
## Not run:  
gwrngs
```

```
## End(Not run)
```

gwaswloc-class	Class "gwaswloc"
----------------	------------------

Description

A container for GRanges instances representing information in the NHGRI GWAS catalog.

Objects from the Class

Objects can be created by calls of the form `new("gwaswloc", ...)`. Any GRanges instance can be supplied.

Slots

`extractDate`: character set manually in `.onAttach` code to indicate date of retrieval of base table
`seqnames`: Object of class "Rle" typically representing chromosome numbers of loci associated with specific traits
`ranges`: Object of class "IRanges" genomic coordinates for locus
`strand`: Object of class "Rle" identifier of chromosome strand
`elementMetadata`: Object of class "DataFrame" general [DataFrame-class](#) instance providing attributes for the locus-trait association
`seqinfo`: Object of class "Seqinfo"
`metadata`: Object of class "list"

Extends

Class "[GRanges](#)", directly. Class "[GenomicRanges](#)", by class "GRanges", distance 2. Class "[Vector](#)", by class "GRanges", distance 3. Class "[GenomicRangesORmissing](#)", by class "GRanges", distance 3. Class "[GenomicRangesORGRangesList](#)", by class "GRanges", distance 3. Class "[Annotated](#)", by class "GRanges", distance 4.

Methods

[`signature(x = "gwaswloc")`]: a character argument to the bracket will be assumed to be a dbSNP identifier for a SNP locus, and records corresponding to this SNP are extracted; numeric indexes are supported as for [GRanges-class](#) instances.

getRsids `signature(x = "gwaswloc")`: extract all dbSNP identifiers as a character vector

getTraits `signature(x = "gwaswloc")`: extract all traits (NHGRI term 'Disease/Trait') as a character vector

subsetByChromosome `signature(x = "gwaswloc")`: select records by chromosome, a vector of chromosomes may be supplied

subsetByTraits `signature(x = "gwaswloc")`: select all records corresponding to a given vector of traits

Note

In gwascats package, the globally accessible gwaswloc instance gwrngs is created upon attachment.

Author(s)

VJ Carey <stvjc@channing.harvard.edu>

References

<http://www.genome.gov/gwastudies/>

Examples

```
showClass("gwaswloc")
```

gwce2gviz

Prepare salient components of GWAS catalog for rendering with Gviz

Description

Prepare salient components of GWAS catalog for rendering with Gviz

Usage

```
gwce2gviz(basegr = gwrngs, contextGR = GRanges(seqnames =
  "chr17", IRanges(start = 37500000, width = 1e+06)),
  txrefpk = "TxDb.Hsapiens.UCSC.hg19.knownGene", genome
  = "hg19", genesympk = "org.Hs.eg.db", plot.it = TRUE,
  maxmlp = 25)
```

Arguments

basegr	gwaswloc instance containing information about GWAS in catalog
contextGR	A GRanges instance delimiting the visualization in genomic coordinates
txrefpk	a TxDb package, typically
genesympk	string naming annotationDbi .db package
genome	character tag like 'hg19'
plot.it	logical, if FALSE, just return list
maxmlp	maximum value of $-10 \log p$ – winsorization of all larger values is performed, modifying the contents of Pvalue_mlogp in the elementMetadata for the call

Examples

```
args(gwce2gviz)
gwascats:::onAttach("", "gwascats")
gwce2gviz()
```

gwdf_2012_02_02 *internal data frame for NHGRI GWAS catalog*

Description

convenience container for imported table from NHGRI GWAS catalog

Usage

data(gwdf_2012_09_22) # or more recent elements available

Format

A data frame with 9000+ observations on the following 34 variables.

Date Added to Catalog a character vector
PUBMEDID a character vector
First Author a character vector
Date a character vector
Journal a character vector
Link a character vector
Study a character vector
Disease/Trait a character vector
Initial Sample Size a character vector
Replication Sample Size a character vector
Region a character vector
Chr_id a character vector
Chr_pos a character vector
Reported Gene(s) a character vector
Mapped_gene a character vector
Upstream_gene_id a character vector
Downstream_gene_id a character vector
Snp_gene_ids a character vector
Upstream_gene_distance a character vector
Downstream_gene_distance a character vector
Strongest SNP-Risk Allele a character vector
SNPs a character vector
Merged a character vector
Snp_id_current a character vector
Context a character vector

Intergenic a character vector
Risk Allele Frequency a character vector
p-Value a character vector
Pvalue_mlog a character vector
p-Value (text) a character vector
OR or beta a character vector
95% CI (text) a character vector
Platform.. a character vector
CNV a character vector

Note

The .onAttach function specifies which data frame is transformed to GRanges.

Source

<http://www.genome.gov/gwastudies>

Examples

```
## Not run:  
data(gwdf_2012_03_22)  
  
## End(Not run)
```

locon6

location information for 10000 SNPs probed on Affy GW 6.0

Description

location information for 10000 SNPs probed on Affy GW 6.0

Usage

```
data(locon6)
```

Format

A data frame with 10000 observations on the following 3 variables.

dbsnp_rs_id a character vector
chrom a character vector
physical_pos a numeric vector

Details

extracted from pd.genomewidesnp.6 v 1.4.0; for demonstration purposes

Examples

```
data(locon6)
str(locon6)
```

makeCurrentGwascat	<i>read NHGRI GWAS catalog table and construct associated GRanges instance</i>
--------------------	--

Description

read NHGRI table and construct associated GRanges instance

Usage

```
makeCurrentGwascat(table.url = "http://www.genome.gov/admin/gwascatalog.txt", fixNonASCII = TRUE)
```

Arguments

table.url	string identifying the .txt file curated at NHGRI
fixNonASCII	logical, if TRUE, non-ASCII characters as identified by iconv will be replaced by asterisk

Details

records for which clear genomic position cannot be determined are dropped from the ranges instance
an effort is made to use reasonable data types for GRanges metadata, so some qualifying characters such as (EA) in Risk allele frequency field will simply be omitted during coercion of contents of that field to numeric.

Value

a GRanges instance

Author(s)

VJ Carey

Examples

```
## Not run:
# if you have good internet access
newcatr = makeCurrentGwascat()

## End(Not run)
```

obo2graphNEL	<i>convert a typical OBO text file to a graphNEL instance (using Term elements)</i>
--------------	---

Description

convert a typical OBO text file to a graphNEL instance (using Term elements)

Usage

```
obo2graphNEL(obo, kill = "\\[Typedef\\]")
```

Arguments

obo	string naming a file in OBO format
kill	entity types to be excluded from processing – probably this should be in a 'keep' form, but for now this works.

Details

Very rudimentary list and grep operations are used to retain sufficient information to map the DAG to a graphNEL, using formal term identifiers as node names and 'is-a' relationships as edges, and term names and other metadata are assigned to nodeData components.

Value

a graphNEL instance

Note

The OBO for Human Disease ontology is serialized as text with this package.

Author(s)

VJ Carey <stvjc@channing.harvard.edu>

References

For use with human disease ontology, http://www.obofoundry.org/cgi-bin/detail.cgi?id=disease_ontology

riskyAlleleCount	<i>given a matrix of subjects x SNP calls, count number of risky alleles</i>
------------------	--

Description

given a matrix of subjects x SNP calls, count number of risky alleles for various conditions, relative to NHGRI GWAS catalog

Usage

```
riskyAlleleCount(callmat, matIsAB = TRUE, chr,
  gwl = gwrngs, snpap = "SNPlocs.Hsapiens.dbSNP.20111119",
  gencode = c("A/A", "A/B", "B/B"))
```

Arguments

callmat	matrix with subjects as rows, SNPs as columns; entries can be generic A/A, A/B, B/B, or specific nucleotide calls
matIsAB	logical, FALSE if nucleotide codes are present, TRUE if generic call codes are present; in the latter case, gwascats:::ABmat2nuc will be run
chr	code for chromosome, should work with the SNP annotation getSNPlocs function, so likely "ch[nn]"
gwl	an instance of gwaswloc
snpap	name of a Bioconductor SNPlocs.Hsapiens.dbSNP.* package
gencode	codes used for generic SNP call

Value

matrix with rows corresponding to subjects , columns corresponding to SNP

Examples

```
if (!exists("gwrngs")) gwascats:::onAttach("a", "b")
data(gg17N) # translated from GGdata chr 17 calls using ABmat2nuc
h17 = riskyAlleleCount(gg17N, matIsAB=FALSE, chr="ch17")
h17[1:5,1:5]
table(as.numeric(h17))
```

`topTraits`*operations on GWAS catalog*

Description

operations on GWAS catalog

Usage

```
topTraits (gwwl, n=10, tag="Disease.Trait")  
locs4trait(gwwl, trait, tag="Disease.Trait")  
chklocs(chrtag="20", gwwl=gwrngs)
```

Arguments

<code>gwwl</code>	instance of gwaswloc
<code>n</code>	numeric, number of traits to report
<code>tag</code>	character, name of field to be used for trait enumeration
<code>trait</code>	character, trait to use for filtering
<code>chrtag</code>	character, chromosome identifier

Value

`topTraits` returns a character vector of most frequently occurring traits in the database

`locs4trait` returns a [gwaswloc](#) object with records defining associations to the specified trait

`chklocs` returns a logical that is TRUE when the asserted locations of SNP in the GWAS catalog agree with the locations given in the dbSNP package `SNPlocs.Hsapiens.dbSNP.20110815`

Author(s)

VJ Carey <stvjc@channing.harvard.edu>

Examples

```
if (!exists("gwrngs")) gwascat:::onAttach("a", "b")  
topTraits(gwrngs)
```

traitsManh	<i>use ggbio facilities to display GWAS results for selected traits in genomic coordinates</i>
------------	--

Description

use ggbio facilities to display GWAS results for selected traits in genomic coordinates

Usage

```
traitsManh(gwr, selr = GRanges(seqnames = "chr17", IRanges(3e+07, 5e+07)), traits = c("Asthma", "Parkin
```

Arguments

<code>gwr</code>	GRanges instance as managed by the gwaswloc container design, with Disease.Trait and Pvalue_mlog among elementMetadata columns
<code>selr</code>	A GRanges instance to restrict the gwr for visualization. Not tested for noncontiguous regions.
<code>traits</code>	Character vector of traits to be exhibited; GWAS results with traits not among these will be labeled "other".
<code>truncmlp</code>	Maximum value of $-\log_{10} p$ to be displayed; in the raw data this ranges to the hundreds and can cause bad compression.
<code>...</code>	not currently used

Details

uses a ggbio autoplot

Value

autoplot value

Author(s)

VJ Carey <stvjc@channing.harvard.edu>

Examples

```
# do a p-value truncation if you want to reduce compression
gwascat:::onAttach("A", "gwascat")
traitsManh(gwrngs)
```

Index

- *Topic **classes**
 - gwaswloc-class, 3
- *Topic **datasets**
 - gwdf_2012_02_02, 5
 - locon6, 6
- *Topic **graphics**
 - gwcecx2gviz, 4
 - traitsManh, 11
- *Topic **models**
 - makeCurrentGwascat, 7
 - obo2graphNEL, 8
 - riskyAlleleCount, 9
 - topTraits, 10
 - traitsManh, 11
- *Topic **package**
 - gwascat-package, 2
- [, gwaswloc, ANY, ANY, ANY-method
 - (gwaswloc-class), 3
- [, gwaswloc-method (gwaswloc-class), 3
- Annotated, 3
- chklocs (topTraits), 10
- g17SM (gwascat-package), 2
- GenomicRanges, 3
- GenomicRangesORGRangesList, 3
- GenomicRangesORmissing, 3
- getRsids (gwaswloc-class), 3
- getRsids, gwaswloc-method
 - (gwaswloc-class), 3
- getTraits (gwaswloc-class), 3
- getTraits, gwaswloc-method
 - (gwaswloc-class), 3
- gg17N (gwascat-package), 2
- GRanges, 3
- gw6.rs_17 (gwascat-package), 2
- gwascat (gwascat-package), 2
- gwascat-package, 2
- gwaswloc, 9, 10
- gwaswloc-class, 3
- gwcecx2gviz, 4
- gwdf_2012_02_02, 5
- gwdf_2012_03_20 (gwdf_2012_02_02), 5
- gwdf_2012_09_22 (gwdf_2012_02_02), 5
- locon6, 6
- locs4trait (topTraits), 10
- low17 (gwascat-package), 2
- makeCurrentGwascat, 7
- obo2graphNEL, 8
- riskyAlleleCount, 9
- rules_6.0_1kg_17 (gwascat-package), 2
- subsetByChromosome (gwaswloc-class), 3
- subsetByChromosome, gwaswloc-method
 - (gwaswloc-class), 3
- subsetByTraits (gwaswloc-class), 3
- subsetByTraits, gwaswloc-method
 - (gwaswloc-class), 3
- topTraits, 10
- traitsManh, 11
- Vector, 3