

Package ‘GenomicScores’

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Type Package

Title Infrastructure to work with genomewide position-specific scores

Description Provide infrastructure to store and access genomewide position-specific scores within R and Bioconductor.

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gscores	<i>Accessing genomic gscores</i>
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Description

Functions to access genomic gscores through GScores objects.

Usage

```
availableGScores()
getGScores(x)
## S4 method for signature 'GScores,GenomicRanges'
gscores(x, ranges, ...)
## S4 method for signature 'GScores,character'
gscores(x, ranges, ...)
## S4 method for signature 'GScores'
score(x, ..., simplify=TRUE)
```

Arguments

- | | |
|--------|--|
| x | For getGScores(), a character vector of length 1 specifying the genomic scores resource to fetch. For gscores() and score(), a GScores object. |
| ranges | A GenomicRanges object with positions from where to retrieve genomic scores, or a character string vector with identifiers associated by the data producer with the genomic scores, e.g., dbSNP 'rs' identifiers in the case of MafDb.* packages. |
| ... | In the call to the gscores() method one can additionally set the following arguments: <ul style="list-style-type: none"> • popCharacter string vector specifying the scores populations to query, when there is more than one. By default, its value is 'defaultPopulation(x)'. Use populations() to find out the available scores populations. • typeCharacter string specifying the type of genomic position being sought, which can be a single nucleotide range (snr), by default, or a nonsnr spanning multiple nucleotides. The latter is the case of indel variants in minor allele frequency data. • scores.onlyFlag setting whether only the scores should be returned as a numeric vector (TRUE), instead of returning them as a metadata column in a GRanges object (FALSE, default). • summaryFunFunction to summarize genomic scores when more than one position is retrieved. By default, this is set to the arithmetic mean, i.e., the mean() function. |

- `quantizedFlag` setting whether the genomic scores should be returned quantized (TRUE) or dequantized (FALSE, default).
- `refVector` of reference alleles in the form of either a character vector, a `DNAStrngSet` object or a `DNAStrngSetList` object. This argument is used only when either there are multiple scores per position or `x` is a `MafDb.*` package.
- `altVector` of alternative alleles in the form of either a character vector, a `DNAStrngSet` object or a `DNAStrngSetList` object. This argument is used only when either there are multiple scores per position or `x` is a `MafDb.*` package.
- `minoverlapInteger` value passed internally to the function `findOverlaps()` from the `IRanges` package, when querying genomic positions associated with multiple-nucleotide ranges (nonSNRs). By default, `minoverlap=1L`, which assumes that the sought nonSNRs are stored as in VCF files, using the nucleotide composition of the reference sequence. This argument is only relevant for genomic scores associated with nonSNRs.
- `cachingFlag` setting whether genomic scores per chromosome should be kept cached in memory (TRUE, default) or not (FALSE). The latter option minimizes the memory footprint but slows down the performance when the `gscores()` method is called multiple times.

`simplify` Flag setting whether the result should be simplified to a vector (TRUE, default) if possible. This happens when scores from a single population are queried.

Details

The function `availableGScores()` shows genomic score sets available as AnnotationHub online resources.

The method `gscores()` takes as first argument a `GScores`-class object that can be loaded from an annotation package or from an AnnotationHub resource. These two possibilities are illustrated in the examples below.

The arguments `ref` and `alt` serve two purposes. One, when there are multiple scores per position, such as with CADD or M-CAP, and we want to select a score matching a specific combination of reference and alternate alleles. The other purpose is when the `GScores` object `x` is a `MafDb.*` package, then by providing `ref` and `alt` alleles we will get separate frequencies for reference and alternate alleles. The current lossy compression of these values yields a correct assignment for biallelic variants in the corresponding `MafDb.*` package and an approximation for multiallelic ones.

Value

The function `availableGScores()` returns a character vector with the names of the AnnotationHub resources corresponding to different available sets of genomic scores. The function `getGScores()` return a `GScores` object. The method `gscores()` returns a `GRanges` object with the genomic scores in a metadata column called `score`. The method `score()` returns a numeric vector with the genomic scores.

Author(s)

R. Castelo

References

Puigdevall, P. and Castelo, R. GenomicScores: seamless access to genomewide position-specific scores from R and Bioconductor. *Bioinformatics*, 18:3208-3210, 2018.

See Also

[phastCons100way.UCSC.hg19 MafDb.1Kgenomes.phase1.hs37d5](#)

Examples

```
## one genomic range of width 5
gr1 <- GRanges(seqnames="chr7", IRanges(start=117232380, width=5))
gr1

## five genomic ranges of width 1
gr2 <- GRanges(seqnames="chr7", IRanges(start=117232380:117232384, width=1))
gr2

## accessing genomic gscores from an annotation package
if (require(phastCons100way.UCSC.hg19)) {
  library(GenomicRanges)

  gsco <- phastCons100way.UCSC.hg19
  gsco
  gscores(gsco, gr1)
  score(gsco, gr1)
  gscores(gsco, gr2)
  populations(gsco)
  gscores(gsco, gr2, pop="DP2")
}

if (require(MafDb.1Kgenomes.phase1.hs37d5)) {
  mafdb <- MafDb.1Kgenomes.phase1.hs37d5
  mafdb
  populations(mafdb)

  ## lookup allele frequencies for SNP rs1129038, located at 15:28356859, a
  ## SNP associated to blue and brown eye colors as reported by Eiberg et al.
  ## Blue eye color in humans may be caused by a perfectly associated founder
  ## mutation in a regulatory element located within the HERC2 gene
  ## inhibiting OCA2 expression. Human Genetics, 123(2):177-87, 2008
  ## [http://www.ncbi.nlm.nih.gov/pubmed/18172690]
  gscores(mafdb, GRanges("15:28356859"), pop=populations(mafdb))
  gscores(mafdb, "rs1129038", pop=populations(mafdb))
}

## accessing genomic scores from AnnotationHub resources
## Not run:
availableGScores()
gsco <- getGScores("phastCons100way.UCSC.hg19")
gscores(gsco, gr1)

## End(Not run)
```

Description

The goal of the GenomicScores package is to provide support to store and retrieve genomic scores associated to physical nucleotide positions along a genome. This is achieved through the GScores class of objects, which is a container for genomic score values.

Details

The GScores class attempts to provide a compact storage and efficient retrieval of genomic score values that have been typically processed and stored using some form of lossy compression. This class is currently based on a former version of the SNPlocs class defined in the BSgenome package, with the following slots:

provider (character), the data provider such as UCSC.

provider_version (character), the version of the data as given by the data provider, typically a date in some compact format.

download_url (character), the URL of the data provider from where the original data were downloaded.

download_date (character), the date on which the data were downloaded.

reference_genome (GenomeDescription), object with information about the reference genome whose physical positions have the genomic scores.

data_pkgname (character), name given to the set of genomic scores associated to a particular genome. When the genomic scores are stored within an annotation package, then this corresponds to the name of that package.

data_dirpath (character), absolute path to the local directory where the genomic scores are stored in one file per genome sequence.

data_serialized_objnames (character), named vector of filenames pointing to files containing the genomic scores in one file per genome sequence. The names of this vector correspond to the genome sequence names.

data_group (character), name denoting a category of genomic scores to which the scores stored in the object belong to. Typical values are "Conservation", "MAF", "Pathogenicity", etc.

data_tag (character), name identifying the genomic scores stored in the object and which can be used, for instance, to assign a column name storing these scores.

data_pops (character), vector of character strings storing score population names. The term "default" is reserved to denote a score set that is not associated to a particular population name and is used by default.

data_nonsnrs (logical), flag indicating whether the object stores genomic scores associated with non-single nucleotide ranges.

data_nsites (integer), number of sites in the genome associated with the genomic scores stored in the object.

.data_cache (environment), data structure where objects storing genomic scores are cached into main memory.

The goal of the design behind the GScores class is to load into main memory only the objects associated with the queried sequences to minimize the memory footprint, which may be advantageous in workflows that parallelize the access to genomic scores by genome sequence.

GScores objects are created either from AnnotationHub resources or when loading specific annotation packages that store genomic score values. Two such annotation packages are:

phastCons100way.UCSC.hg19 Nucleotide-level phastCons conservation scores from the UCSC Genome Browser calculated from multiple genome alignments from the human genome version hg19 to 99 vertebrate species.

phastCons100way.UCSC.hg38 Nucleotide-level phastCons conservation scores from the UCSC Genome Browser calculated from multiple genome alignments from the human genome version hg38 to 99 vertebrate species.

Constructor

GScores(provider, provider_version, download_url, download_date, reference_genome, data_pkgname, data_dirpath) Creates a GScores object. In principle, the end-user needs not to call this function.

provider character string, containing the data provider.

provider_version character string, containing the version of the data as given by the data provider.

download_url character string, containing the URL of the data provider from where the original data were downloaded.

reference_genome GenomeDescription, storing the information about the associated reference genome.

data_pkgname character string, name given to the set of genomic scores stored through this object.

data_dirpath character string, absolute path to the local directory where the genomic scores are stored.

data_serialized_objname character string vector, containing filenames where the genomic scores are stored.

default_pop character string, containing the name of the default scores population.

data_group character string, containing a name that indicates a category of genomic scores to which the scores in the object belong to. Typical names could be "Conservation", "MAF", etc.

data_tag character string, containing a tag that succinctly labels genomic scores from a particular source. This can be used to automatically give, for instance, a name to a column storing genomic scores in data frame object. Its default value takes the prefix of the package name.

Accessors

name(x): get the name of the set of genomic scores.

type(x): get the substring of the name of the set of genomic scores comprised between the first character until the first period. This should typically match the type of genomic scores such as, phastCons, phyloP, etc.

provider(x): get the data provider.

providerVersion(x): get the provider version.

organism(x): get the organism associated with the genomic scores.

seqlevelsStyle(x): get the genome sequence style.

seqinfo(x): get the genome sequence information.

seqnames(x): get the genome sequence names.

seqlengths(x): get the genome sequence lengths.

populations(x): get the identifiers of the available scores populations. If only one scores population is available, then it shows only the term default.

defaultPopulation(x): get or set the default population of scores.
 gscoresGroup(x): get or set the genomic scores group label.
 gscoresTag(x): get or set the genomic scores tag label.
 gscoresNonSNRs(x): get whether there are genomic scores associated with non-single nucleotide ranges.
 nsites(x): get the number of sites in the genome with genomic scores.
 qfun(x): get the quantizer function.
 dqfun(x): get the dequantizer function.
 citation(x): get citation information for the genomic scores data in the form of a bibentry object.

Author(s)

R. Castelo

References

Puigdevall, P. and Castelo, R. GenomicScores: seamless access to genomewide position-specific scores from R and Bioconductor. *Bioinformatics*, 18:3208-3210, 2018.

See Also

[gscores\(\)](#) [score\(\)](#) [phastCons100way.UCSC.hg19](#)

Examples

```

## one genomic range of width 5
gr1 <- GRanges(seqnames="chr7", IRanges(start=117232380, width=5))
gr1

## five genomic ranges of width 1
gr2 <- GRanges(seqnames="chr7", IRanges(start=117232380:117232384, width=1))
gr2

## supporting annotation packages with genomic scores
if (require(phastCons100way.UCSC.hg19)) {
  library(GenomicRanges)

  gsco <- phastCons100way.UCSC.hg19
  gsco
  gscores(gsco, gr1)
  score(gsco, gr1)
  gscores(gsco, gr2)
  populations(gsco)
  gscores(gsco, gr2, pop="DP2")
}

## supporting AnnotationHub resources
## Not run:
availableGScores()
gsco <- getGScores("phastCons100way.UCSC.hg19")
gsco
gscores(gsco, gr1)

```

```
## End(Not run)

## metadata from a GScores object
name(gsco)
type(gsco)
provider(gsco)
providerVersion(gsco)
organism(gsco)
seqlevelsStyle(gsco)
seqinfo(gsco)
head(seqnames(gsco))
head(seqlengths(gsco))
gscoresTag(gsco)
populations(gsco)
defaultPopulation(gsco)
qfun(gsco)
dqfun(gsco)
citation(gsco)
```

igscores

GenomicScores shiny app

Description

Starts an interactive GenomicScores shiny web app.

Usage

```
igscores()
```

Details

The goal of the GenomicScores package is to provide support to store and retrieve genomic scores associated to physical nucleotide positions along a genome.

The `igscores()` function starts an interactive shiny web app that allows the user to query annotation packages storing genomic scores. Internally, it calls to the function `gscores()`; see its manual page for a description of the arguments and their default and alternative values.

Value

None.

Author(s)

P. Rodríguez and R. Castelo

References

Puigdevall, P. and Castelo, R. GenomicScores: seamless access to genomewide position-specific scores from R and Bioconductor. *Bioinformatics*, 18:3208-3210, 2018.

See Also

[gscores](#)

Examples

```
## Not run:  
igscores() ## this will open your browser with the GenomicScores shiny web app  
  
## End(Not run)
```

makeGScoresPackage *Building genomic gscores packages*

Description

Build a genomic gscores packages from a GScores object.

Usage

```
makeGScoresPackage(gsco, version, maintainer, author,  
                  destDir=".", license="Artistic-2.0")
```

Arguments

gsco	GScores. The GScores object from which the package will be created.
version	Character. Version of the package.
maintainer	Character. Maintainer of the package, including email.
author	Character. Author of the package.
destDir	Character. The path to the directory where the package will be created.
license	Character. The license of the package.

Details

This function allows one to create an R package from a GScores object. This may be useful if one wants to have a tar-ball package version of genomic scores available only through the Annotation-Hub; see the vignette.

Value

It returns invisibly the package directory.

Author(s)

R. Castelo

References

Puigdevall, P. and Castelo, R. GenomicScores: seamless access to genomewide position-specific scores from R and Bioconductor. *Bioinformatics*, 18:3208-3210, 2018.

See Also

[availableGScores\(\)](#) [getGScores\(\)](#)

Examples

```
## accessing genomic scores from AnnotationHub resources
## and building a package from them
## Not run:
availableGScores()
gsco <- getGScores("fitCons.UCSC.hg19")
makeGScoresPackage(gsco, version="1.0", maintainer="me <me@example.com>", author="me")

## End(Not run)
```

 rgscores

Sampling genomic gscores

Description

Function for randomly sampling genomic gscores from GScores objects.

Usage

```
## S4 method for signature 'GScores,missing'
rgscores(n, object, ...)
## S4 method for signature 'missing,GScores'
rgscores(n, object, ...)
## S4 method for signature 'numeric,GScores'
rgscores(n, object, ...)
## S4 method for signature 'integer,GScores'
rgscores(n, object, ...)
```

Arguments

n	Number of scores to sample.
object	A GScores object.
...	In the call to the rgscores() method one can additionally set the following arguments: <ul style="list-style-type: none"> • popCharacter string vector of length one, specifying the scores population from which we want to sample scores. By default, its value is 'default-Population(object)'. Use populations() to find out the available scores populations. • scores.onlyFlag setting whether only the scores should be returned as a numeric vector (TRUE), instead of returning them as a metadata column in a GRanges object (FALSE, default). • rangesEither a GRanges object or a character string vector of sequence names. Scores will be sampled from the given genomic regions, which by default correspond to the entire genomic space of object. Currently, only entire chromosomes are considered.

Details

The method rgscores() samples scores randomly from a GScores object.

Value

A GRanges object with the sampled genomic positions and scores. When `scores.only=TRUE` then a numeric vector is returned with the sampled scores.

Author(s)

R. Castelo

References

Puigdevall, P. and Castelo, R. GenomicScores: seamless access to genomewide position-specific scores from R and Bioconductor. *Bioinformatics*, 18:3208-3210, 2018.

See Also

[phastCons100way.UCSC.hg19 MafDb.1Kgenomes.phase1.hs37d5](#)

Examples

```
## accessing genomic gscores from an annotation package
if (require(phastCons100way.UCSC.hg19)) {
  library(GenomicRanges)

  phast <- phastCons100way.UCSC.hg19
  set.seed(123)
  rgcores(10L, phast, ranges=c("chr22", "chrY"))
}

if (require(MafDb.1Kgenomes.phase1.hs37d5)) {
  mafdb <- MafDb.1Kgenomes.phase1.hs37d5
  set.seed(123)
  rgcores(10L, mafdb, ranges=c("21", "22"))
}
```

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