

Package ‘cgdv17’

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Title Complete Genomics Diversity Panel, chr17 on 46 individuals

Version 0.0.17

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Description Complete Genomics Diversity Panel, chr17 on 46 individuals

Suggests

Imports Biobase, IRanges

Depends R (>= 2.15), VariantAnnotation, org.Hs.eg.db, methods, GGtools, TxDb.Hsapiens.UCSC.hg19.knownGene, parallel

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License Artistic 2.0

LazyLoad yes

biocViews genetics

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cgdv17-package

Complete Genomics Diversity Panel, chr17 on 46 individuals

Description

Complete Genomics Diversity Panel, chr17 on 46 individuals, illustrating subject-specific variant sets

Details

```

Package:      cgdv17
Version:      0.0.9
Suggests:    
Imports:       Biobase, IRanges
Depends:      R (>= 2.14), VariantAnnotation, org.Hs.eg.db, methods
License:      Artistic 2.0
LazyLoad:     yes
biocViews:   genetics
Built:        R 2.15.0; ; 2012-03-09 12:45:57 UTC; unix

```

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<code>countVariants</code>	count variants in a <code>raggedVariantSet</code> instance
<code>getRVS</code>	acquire data for and construct a ragged variant set instance
<code>padToReference</code>	create a <code>snpStats SnpMatrix</code> instance by padding a ragged variant set to reference alleles wherever a variant is not recorded
<code>raggedVariantSet-class</code>	Class <code>""raggedVariantSet""</code> ,
<code>variantGRanges</code>	acquire a list of GRanges recording variants and locations

see vignette; CY17 is an ExpressionSet on individuals from CEU and YRI overlapping with the diversity set, popvec enumerates source populations, h1 is an exemplar VCF header structure

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<code>countVariants</code>	<i>count variants in a raggedVariantSet instance</i>
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Description

count variants in a `raggedVariantSet` instance

Usage

```
countVariants(rvs, delim, qthresh = 160, applier = lapply)
```

Arguments

<code>rvs</code>	instance of <code>raggedVariantSet</code>
<code>delim</code>	GRanges instance
<code>qthresh</code>	quality threshold for keeping a variant in count
<code>applier</code>	lapply-like function

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getRVS

acquire data for and construct a ragged variant set instance

Description

acquire data for and construct a ragged variant set instance

Usage

```
getRVS(packname, fns2samplenames = function(x)
  gsub(".*(NA.....).*", "\\\1", x))

getrd(x, id)
```

Arguments

packname	string naming package where the resources are found
fns2samplenames	function to transform filenames to sample name tokens
x	instance of raggedVariantSet
id	character to select sample

Details

currently very specialized, as the protocol for managing collections of VCF files with discrepant variant sets per subject is not clear

assumes the package has inst/rowdata where row data of [readVcf](#) results are held

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padToReference

create a snpStats SnpMatrix instance by padding a ragged variant set to reference alleles wherever a variant is not recorded

Description

create a snpStats SnpMatrix instance by padding a ragged variant set to reference alleles wherever a variant is not recorded

Usage

```
padToReference(rv, gr, qthresh = 160, applier = lapply)
```

Arguments

rv	<code>raggedVariantSet</code> instance
gr	<code>GRanges</code> instance
qthresh	quality lower bound for retention of variant
applier	lapply like function

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`raggedVariantSet-class`

Class "raggedVariantSet"

Description

manage information on non-aligned variant sets from multiple VCFs

Objects from the Class

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

Slots

filenames: files will be held in `inst/rowdata`, named here

sampleNames: names of samples managed

Methods

```
[ signature(x = "raggedVariantSet", i = "ANY", j = "ANY", drop = "ANY"): familiar sub-
  setting syntax
sampleNames signature(object = "raggedVariantSet"): getter
show signature(object = "raggedVariantSet"): concise report
variantGRanges signature(rvs = "raggedVariantSet", delim = "GRanges", qthresh = "missing", appli-
  getter
variantGRanges signature(rvs = "raggedVariantSet", delim = "GRanges", qthresh = "numeric", appli-
  getter with quality threshold
```

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Examples

```
showClass("raggedVariantSet")
```

variantGRanges	<i>acquire a list of GRanges recording variants and locations</i>
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Description

acquire a list of GRanges recording variants and locations

Usage

```
variantGRanges(rvs, delim, qthresh = 160, applier = lapply)
```

```
variantNames(rvs, delim, qthresh=160, applier=lapply)
```

Arguments

rvs	raggedVariantSet instance
delim	GRanges instance for confinement
qthresh	lower bound on quality
applier	lapply like function

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