

# Package ‘cgdv17’

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**Title** Complete Genomics Diversity Panel, chr17 on 46 individuals

**Version** 0.26.0

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**Description** Complete Genomics Diversity Panel, chr17 on 46 individuals

**Depends** R (>= 2.15), methods, VariantAnnotation (>= 1.15.15)

**Imports** BiocGenerics, S4Vectors, IRanges, GenomicRanges, Biobase

**Suggests** parallel, GGtools, TxDb.Hsapiens.UCSC.hg19.knownGene,  
org.Hs.eg.db, illuminaHumanv1.db

**License** Artistic-2.0

**LazyLoad** yes

**biocViews** SequencingData, SNPData, BiocViews

**git\_url** <https://git.bioconductor.org/packages/cgdv17>

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**git\_last\_commit\_date** 2020-04-27

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 cgdv17-package

*Complete Genomics Diversity Panel, chr17 on 46 individuals*


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## 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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| countVariants          | count variants in a raggedVariantSet instance  |
| getRVS                 | acquire data for and construct a ragged variant set instance   |
| padToReference         | create a snpStats SnpMatrix instance by padding a ragged variant set to reference alleles wherever a variant is not recorded |
| raggedVariantSet-class | Class "raggedVariantSet"   |
| variantGRanges         | 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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 countVariants

*count variants in a raggedVariantSet instance*


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## Description

count variants in a raggedVariantSet instance

**Usage**

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

**Arguments**

|         |  |
|---------|--|
| rvs     | instance of <a href="#">raggedVariantSet</a>     |
| delim   | GRanges instance                                 |
| qthresh | quality threshold for keeping a variant in count |
| applier | 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 <a href="#">raggedVariantSet</a>          |
| 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/rowranges where row ranges of [readVcf](#) results are held

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|                |   |
|----------------|---|
| padToReference | <i>create a snpStats SnpMatrix instance by padding a ragged variant set to reference alleles wherever a variant is not recorded</i> |
|----------------|---|

---

### 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      | raggedVariantSet instance                    |
| gr      | GRanges instance                             |
| qthresh | quality lower bound for retention of variant |
| applier | lapply like function                         |

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|                        |                                 |
|------------------------|---------------------------------|
| raggedVariantSet-class | <i>Class "raggedVariantSet"</i> |
|------------------------|---------------------------------|

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### 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/rowranges, named here  
**sampleNames:** names of samples managed

### Methods

[ signature(x = "raggedVariantSet", i = "ANY", j = "ANY", drop = "ANY"): familiar subsetting syntax  
**sampleNames** signature(object = "raggedVariantSet"): getter  
**show** signature(object = "raggedVariantSet"): concise report  
**variantGRanges** signature(rvs = "raggedVariantSet", delim = "GRanges", qthresh = "missing", applier = "missing"): getter  
**variantGRanges** signature(rvs = "raggedVariantSet", delim = "GRanges", qthresh = "numeric", applier = "function"): getter with quality threshold

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**Examples**

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

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|                |   |
|----------------|---|
| 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     | <a href="#">raggedVariantSet</a> instance |
| delim   | GRanges instance for confinement          |
| qthresh | lower bound on quality                    |
| applier | lapply like function                      |

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