
gUtils Documentation

Release 1.0.0

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August 04, 2015

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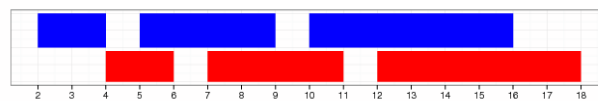
Introduction

Range Operations

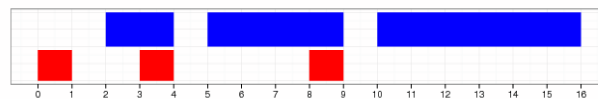
This section will describe additional GRanges operations provided by gUtils.

```
## make some example data sets
refl9 <- readRDS(system.file("extdata", "refGene.hg19.gr.rds", package="gUtils"))
gr <- GRanges(1, IRanges(c(2,5,10), c(4,9,16)), seqinfo=Seqinfo("1", 20))
gr2 <- c(gr, GRanges(1, IRanges(c(1,9), c(6,14)), seqinfo=Seqinfo("1", 20)))
dt <- data.table(seqnames=1, start=c(2,5,10), end=c(3,8,15))
```

```
shift(gr, 2)
```



```
flank(gr, width=2)
```



```
gr.start(gr, width=3)
```

```
gr.end(gr, width=3)
```

```
gr.mid(gr) + 2
```

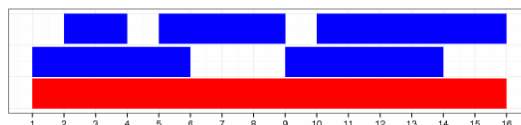
```
grbind
```

```
## add metadata to one field
mcols(gr)$score = 3
## try to concatenate
c(gr, gr2) ## ERROR
## with grbind
grbind(gr, gr2) ## SUCCESS. Adds NA for missing fields
## GenomicRanges::c does this already for GRangesList
```

```
streduce(gr2)
gr.sample(gr2, 2, len=2, replace=TRUE)
```

```
## output GRanges
GRanges object with 3 ranges and 1 metadata column:
      seqnames      ranges strand |   query.id
      <Rle> <IRanges>  <Rle> | <integer>
[1]          1 [ 8,  9]      * |         2
[2]          1 [ 5,  6]      * |         2
[3]          1 [11, 12]      * |         3
```

```
gr.rand(w=c(2,5,3), seqinfo(gr))
gr.simplify
gr.tile(GRanges(1, IRanges(1,9)), w=3) + 1
gr.refactor
gr.tile.map
gr.round
```



Data manipulation

R provides a number of data structures for storing genomic data, each with its advantages and drawbacks.

The most useful structures for this purpose are:

GRanges Store ranges along with metadata, sequences and the coordinates of the reference genome.

GRangesList Store groups of ranges, with additional metadata belonging to the group.

data.table Fast and efficient general-purpose container similar to `data.frame`, but with significant performance improvements.

In `gUtils` functions, we often manipulate the data to move between these data structures where one is more useful than another. A key example is in `gr.findoverlaps`, which converts the input `GRanges` into `data.table` objects to take advantage of the blazing fast `foverlaps` util. For the most part, these conversions should be invisible to the user.

However, often there are data structures conversions that may be useful to the end user. This includes unlisting `GRangesList` objects into `GRanges`, making `data.table` objects from `GRanges`, and binding together multiple `GRanges` or `GRangesList` objects, among others. This section will describe and demonstrate the functionality `gUtils` provides for manipulating these data structures.

```
refl9 <- readRDS(system.file("extdata", "refGene.hg19.gr.rds", package="gUtils"))
gr <- GRanges(1, IRanges(c(2,5,10), c(4,9,16)), seqinfo=Seqinfo("1", 20))
dt <- data.table(seqnames=1, start=c(2,5,10), end=c(3,8,15))
```

`grbind`

`grlbind`

`dtgr`

`grdt`

`si2gr`

`gr2gatk`

`gr.flatten`

`gr.flatmap`

`grl.split`

`grl.stripnames`

`grl.unlist`

`grl.span`

grl.pivot

rrbind

Additional utilities

`gr.string` and `grl.string`

`parse.grl` and `parse.gr`

`gr.gatk`

`gr.chr` and `gr.nochr`

`gr.fix`

`gr.tostring`

`affine.map`

`gr.findoverlaps`

`gr.match`

`gr.in`

`gr.duplicated`

`gr.val`

`gr.dist`

`alpha`

`grl.in`

`chunk`

`import.ucsc`

BAM/SAM operations

This section will describe utility functions for manipulating BAM files, using functions based on the [Rsamtools](#) package.

`read.bam`

`bam.cov.gr`

`bam.cov.tile.st`

`bam.pair.cov`

`counts2rpkm`

`get.pairs.grl`

`count.clips`

`varbase`

`varcount`

`mafcount`

`splice.cigar`

`gc_content`

`bamtag`

Indices and tables

- `genindex`
- `modindex`
- `search`