

# Working with Genomic Ranges

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## Introduction to GenomicRanges<sup>1</sup>

The GenomicRanges serves as the foundation for representing genomic locations within the Bioconductor project. This package lays a foundation for genomic analysis by introducing three classes (GRanges, GPos, and GRangesList), which are used to represent genomic ranges, genomic positions, and groups of genomic ranges.

```
library(GenomicRanges)
z <- GRanges("chr1",IRanges(1000001,1001000),strand="+")
start(z)
end(z)
width(z)
strand(z)
mcols(z) # the 'metadata columns', any information stored alongside each range
ranges(z) # gives the IRanges
seqnames(z) # the chromosomes for each ranges
seqlevels(z) # the possible chromosomes
seqlengths(z) # the lengths for each chromosome
```

## Intra-range methods

Affects ranges independently

function	description
shift	moves left/right
narrow	narrows by relative position within range
resize	resizes to width, fixing start for +, end for -
flank	returns flanking ranges to the left +, or right -
promoters	similar to flank
restrict	restricts ranges to a start and end position
trim	trims out of bound ranges
+/-	expands/contracts by adding/subtracting fixed amount
*	zooms in (positive) or out (negative) by multiples

## Inter-range methods

Affects ranges as a group

function	description
range	one range, leftmost start to rightmost end

<sup>1</sup>Reference card extracted from mikelove/bioc-refcard

function	description
<code>reduce</code>	cover all positions with only one range
<code>gaps</code>	uncovered positions within range
<code>disjoin</code>	breaks into discrete ranges based on original starts/ends

## Nearest methods

Given two sets of ranges, `x` and `subject`, for each range in `x`, returns...

function	description
<code>nearest</code>	index of the nearest neighbor range in subject
<code>precede</code>	index of the range in subject that is directly preceded by the range in <code>x</code>
<code>follow</code>	index of the range in subject that is directly followed by the range in <code>x</code>
<code>distanceToNearest</code>	distances to its nearest neighbor in subject (Hits object)
<code>distance</code>	distances to nearest neighbor (integer vector)

A Hits object can be accessed with `queryHits`, `subjectHits` and `mcols` if a distance is associated.

## set methods

If `y` is a `GRangesList`, then use `punion`, etc. All functions have default `ignore.strand=FALSE`, so are strand specific.

```
union(x,y)
intersect(x,y)
setdiff(x,y)
```

## Overlaps

```
x %over% y # logical vector of which x overlaps any in y
fo <- findOverlaps(x,y) # returns a Hits object
queryHits(fo) # which in x
subjectHits(fo) # which in y
```

## Seqnames and seqlevels

GenomicRanges and GenomeInfoDb

```
gr.sub <- gr[seqlevels(gr) == "chr1"]
seqlevelsStyle(x) <- "UCSC" # convert to 'chr1' style from "NCBI" style '1'
```