

Saying Hello to the Bioconductor Ranges Infrastructure

*Michael Lawrence**

Genentech

*michafla@gene.com

December 5, 2022

Package

HelloRanges 1.24.0

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1 Introduction

The primary purpose of the *HelloRanges* package, from the pedagogical perspective, is to map *bedtools* “recipes” to *R* scripts. The package was born out of the recognition that *Bioconductor* lacks a cookbook that explains how to achieve common, specific tasks using the Ranges infrastructure. And that writing a compiler is more fun (and hopefully more useful) than writing documentation. The goal is to enable those who already use *R/Bioconductor* for modeling and plotting to unify their workflow by integrating their upstream processing.

HelloRanges provides an *R* function corresponding to each *bedtools* command. The output is an *R* language object, and we can print the object to copy and integrate the code into an *R* script. Ideally, the code is first integrated (by copy/paste) into an *R* script. Unlike *bedtools*, the result of evaluation is an *R* object that is suitable for further analysis. There is no automatic output to files, because once we are in *R*, we want to stay there. Assuming that the reader is interested in learning *Bioconductor*, we encourage the reader to inspect the code prior to evaluation by printing the language object. The generated code is meant to be correct, readable, conformant to best practices and performant (in that order). While the code is much more verbose than the corresponding *bedtools* call, we argue that the explicit, low-level Ranges API has the advantage of being self-documenting and more flexible. And, of course, it directly integrates with the rest of *Bioconductor*.

Obviously, performing I/O with each operation will have a negative impact on performance, so it is recommended to import the data once, and perform subsequent operations on in-memory data structures. If memory is exhausted, consider distributing computations.

For the sake of comparison, this tutorial closely follows that of *bedtools* itself (<http://quinlanlab.org/tutorials/bedtools/bedtools.html>). We will analyze the data from Maurano et al [1] assessment of DnaseI hypersensitivity across a range of fetal tissues (20 samples). The *bedtools* tutorial mostly consists of arbitrary range operations on the annotation tracks. Near the end, we will compare samples from the Maurano study in terms of their mutual overlap.

2 Data

The data are provided via the *HelloRangesData* package, which we load presently:

```
> library>HelloRanges)
> library>HelloRangesData)
```

To have convenient paths to the data, we switch our working directory to the one with the data files:

```
> oldwd <- setwd(system.file("extdata", package="HelloRangesData"))
```

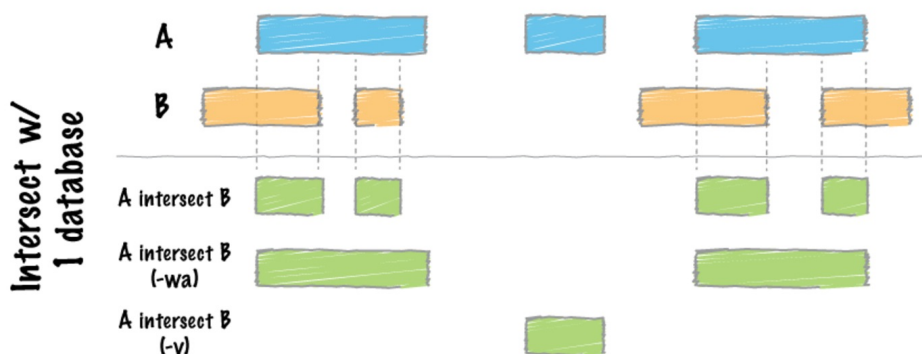
In our working directory are 20 BED files from the DnaseI study, as well as BED files representing the CpG islands ('cpg.bed'), Refseq exons ('exons.bed'), disease-associated SNPs ('gwas.bed'), and functional annotations output by chromHMM given ENCODE human embryonic stem cell ChIP-seq data ('hesc.chromHmm.bed'). There is also a 'hg19.genome' file indicating the chromosome lengths of the hg19 genome build.

One of the advantages of *R*, compared to the shell, is its unified package management system. *R* packages can contain data, and even completed analyses, in addition to libraries of functions. *Bioconductor* provides many annotations and sample datasets through packages.

Packages make obtaining data easy, and they also help with reproducibility and provenance through versioning. Thus, they are more convenient and safer compared to downloading from live URLs. Some packages provide APIs to download data from repositories, and we can use them to programmatically generate data packages in a reproducible way.

For example, the TxDb packages provide transcript annotations for an individual reference genome. *Bioconductor* provides pre-built TxDb packages for the commonly used genome builds, and it is easy to generate a custom TxDb package using annotations from Biomart, UCSC, or just a GFF file. The *rtracklayer* package supports automated retrieval of any dataset from the UCSC table browser, without any pointing-and-clicking involved. We will demonstrate some of these tools later in the tutorial.

3 Overlap and Intersection



One of the most useful ways to compare two tracks is to determine which ranges overlap, and where they intersect (see the above image from the *bedtools* tutorial).

By default, *bedtools* outputs the region of intersection for each overlap between the two tracks. We compile the code for intersecting the CpG islands and the exons.

```
> code <- bedtools_intersect("-a cpg.bed -b exons.bed -g hg19.genome")
> code
{
  genome <- import("hg19.genome")
  gr_a <- import("cpg.bed", genome = genome)
  gr_b <- import("exons.bed", genome = genome)
  pairs <- findOverlapPairs(gr_a, gr_b, ignore.strand = TRUE)
  ans <- pintersect(pairs, ignore.strand = TRUE)
  ans
}
```

This code should be integrated into an R script that implements a larger workflow. For the purposes of this tutorial, we will call `eval` on the language object to yield the result:

```
> ans <- eval(code)
> mcols(ans)$hit <- NULL
> ans
```

GRanges object with 45500 ranges and 1 metadata column:

	seqnames	ranges	strand	name
	<Rle>	<IRanges>	<Rle>	<character>
[1]	chr1	29321-29370	*	CpG:_116
[2]	chr1	135125-135563	*	CpG:_30
[3]	chr1	327791-328229	*	CpG:_29
[4]	chr1	327791-328229	*	CpG:_29
[5]	chr1	327791-328229	*	CpG:_29
...
[45496]	chrY	59213949-59214117	*	CpG:_36
[45497]	chrY	59213949-59214117	*	CpG:_36
[45498]	chrY	59213949-59214117	*	CpG:_36
[45499]	chrY	59213949-59214117	*	CpG:_36
[45500]	chrY	59213949-59214117	*	CpG:_36

seqinfo: 93 sequences from hg19 genome

The result is an instance of *GRanges*, the central data structure in *Bioconductor* for genomic data. A *GRanges* takes the form of a table and resembles a BED file, with a column for the chromosome, start, end, strand. We will see *GRanges* a lot, along with its cousin, *GRangesList*, which stores what *bedtools* calls “split” ranges.

3.1 Sequence information

Consider the simplest invocation of *bedtools intersect*:

```
> code <- bedtools_intersect("-a cpq.bed -b exons.bed")
> code
{
  genome <- Seqinfo(genome = NA_character_)
  gr_a <- import("cpq.bed", genome = genome)
  gr_b <- import("exons.bed", genome = genome)
  pairs <- findOverlapPairs(gr_a, gr_b, ignore.strand = TRUE)
  ans <- pintersect(pairs, ignore.strand = TRUE)
  ans
}
```

The first line creates an object representing the structure of the genome build:

```
> genome <- eval(code[[2L]])
> genome
```

Seqinfo object with no sequences:

It is an empty object, because the genome identifier is unspecified (*NA_character_*). Having unspecified genome bounds is dangerous and leads to accidents involving incompatible genome builds. Besides error checking, the bounds are useful when computing coverage and finding the gaps (see below). Luckily, *bedtools* lets us specify the genome as an argument:

```
> bedtools_intersect("-a cpq.bed -b exons.bed -g hg19.genome")
{
```

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```
genome <- import("hg19.genome")
gr_a <- import("cpg.bed", genome = genome)
gr_b <- import("exons.bed", genome = genome)
pairs <- findOverlapPairs(gr_a, gr_b, ignore.strand = TRUE)
ans <- pintersect(pairs, ignore.strand = TRUE)
ans
```

We now have a populated genome, since the tutorial had provided the 'hg19.genome' file. However, in general, information on the genome build should be centralized, not sitting somewhere in a file. *Bioconductor* provides [GenomeInfoDb](#) as its central source of sequence information. We can hook into that by passing the genome identifier instead of a file name:

```
> bedtools_intersect("-a cpg.bed -b exons.bed -g hg19")
{
  genome <- Seqinfo(genome = "hg19")
  gr_a <- import("cpg.bed", genome = genome)
  gr_b <- import("exons.bed", genome = genome)
  pairs <- findOverlapPairs(gr_a, gr_b, ignore.strand = TRUE)
  ans <- pintersect(pairs, ignore.strand = TRUE)
  ans
}
```

3.2 Annotations

The next step is the import of the CpG islands and exons using the `import` function from [rtracklayer](#), which can load pretty much any kind of genomic data into the appropriate type of *Bioconductor* object. In this case, we are loading the data as a *GRanges*:

```
> gr_a
GRanges object with 28691 ranges and 1 metadata column:
      seqnames      ranges strand |      name
      <Rle>        <IRanges> <Rle> | <character>
[1]      chr1      28736-29810      * | CpG:_116
[2]      chr1     135125-135563      * | CpG:_30
[3]      chr1     327791-328229      * | CpG:_29
[4]      chr1     437152-438164      * | CpG:_84
[5]      chr1     449274-450544      * | CpG:_99
...      ...      ...      ... | ...
[28687] chrY 27610116-27611088      * | CpG:_76
[28688] chrY 28555536-28555932      * | CpG:_32
[28689] chrY 28773316-28773544      * | CpG:_25
[28690] chrY 59213795-59214183      * | CpG:_36
[28691] chrY 59349267-59349574      * | CpG:_29
-----
seqinfo: 93 sequences from hg19 genome
```

The [rtracklayer](#) package can also download data directly from the UCSC table browser. For example, we could get the CpG islands directly:

```
> ucsc <- browserSession()
> genome(ucsc) <- "hg19"
> cpgs <- ucsc[["CpG Islands"]]
```

Gene annotations, including exon coordinates, should also be stored more formally than in a file, and Bioconductor provides them through its TxDb family of packages:

```
> library(TxDb.Hsapiens.UCSC.hg19.knownGene)
> exons <- exons(TxDb.Hsapiens.UCSC.hg19.knownGene)
```

3.3 Finding Overlaps

The next step is to find all of the overlaps. The workhorse function is `findOverlaps` from the *IRanges* package. Here, we use the variant `findOverlapPairs`, a convenience for creating a *Pairs* object that matches up the overlapping ranges:

```
> pairs

Pairs object with 45500 pairs and 0 metadata columns:
      first second
      <GRanges> <GRanges>
[1]   chr1:28736-29810   chr1:29321-29370:-
[2]   chr1:135125-135563 chr1:134773-139696:-
[3]   chr1:327791-328229 chr1:324439-328581:+
[4]   chr1:327791-328229 chr1:324439-328581:+
[5]   chr1:327791-328229 chr1:327036-328581:+
...
[45496] chrY:59213795-59214183 chrY:59213949-59214117:+
[45497] chrY:59213795-59214183 chrY:59213949-59214117:+
[45498] chrY:59213795-59214183 chrY:59213949-59214117:+
[45499] chrY:59213795-59214183 chrY:59213949-59214117:+
[45500] chrY:59213795-59214183 chrY:59213949-59214117:+
```

Although the ranges are displayed as succinct strings, they data are still represented as *GRanges* objects.

Users of *bedtools* will be familiar with *Pairs* as the analog of the BEDPE file format. We can use *rtracklayer* to export *Pairs* to BEDPE:

```
> export(pairs, "pairs.bedpe")
```

A key parameter to `findOverlapPairs` is `ignore.strand=TRUE`. By default, all operations on *GRanges* take strand into account when determining whether two ranges overlap, and deciding on the orientation of a range. This is surprising to many novice users, particularly to those with *bedtools* experience. Most functions take the `ignore.strand` argument to control this behavior. To avoid confusion, the code generated by *HelloRanges* is always explicit about how it is treating strand. Users are encouraged to follow the same practice.

3.4 Computing intersections

The final step is to find the actual intersecting region between the member of each overlapping pair. We do this with the `pintersect` function, which is the “parallel” or “pairwise” version of the default `intersect` function. If we had just called `intersect(gr_a, gr_b)` instead, the entire set of ranges would have been treated as a set, and overlapping ranges in `gr_a` and `gr_b` would have been merged (this is rarely desirable and requires an extra merge step in *bedtools*).

Notice again the importance of `ignore.strand=TRUE`. Without that, ranges on opposite strands would have zero intersection.

And here is our result:

```
> ans
GRanges object with 45500 ranges and 1 metadata column:
      seqnames      ranges strand |      name
      <Rle>        <IRanges> <Rle> | <character>
[1]    chr1      29321-29370    * |    CpG:_116
[2]    chr1     135125-135563    * |    CpG:_30
[3]    chr1     327791-328229    * |    CpG:_29
[4]    chr1     327791-328229    * |    CpG:_29
[5]    chr1     327791-328229    * |    CpG:_29
...      ...      ...      ... |    ...
[45496] chrY 59213949-59214117    * |    CpG:_36
[45497] chrY 59213949-59214117    * |    CpG:_36
[45498] chrY 59213949-59214117    * |    CpG:_36
[45499] chrY 59213949-59214117    * |    CpG:_36
[45500] chrY 59213949-59214117    * |    CpG:_36
-----
seqinfo: 93 sequences from hg19 genome
```

Again, a *GRanges* object. The `hit` column indicates whether the pair overlapped at all (as opposed to one range being of zero width). It’s useless in this case.

3.5 Keeping the original features

To keep the original form of the overlapping features, the generated code simply neglects to call `pintersect` and ends up with the `pairs` object introduced previously:

```
> bedtools_intersect("-a cpq.bed -b exons.bed -g hg19 -wa -wb")
{
  genome <- Seqinfo(genome = "hg19")
  gr_a <- import("cpq.bed", genome = genome)
  gr_b <- import("exons.bed", genome = genome)
  pairs <- findOverlapPairs(gr_a, gr_b, ignore.strand = TRUE)
  ans <- pairs
  ans
}
```


3.6 Computing the amount of overlap

To compute the width of the overlapping regions, we query the initial result for its width and store as an annotation on the pairs:

```
> bedtools_intersect("-a cpq.bed -b exons.bed -g hg19 -wo")
{
  genome <- Seqinfo(genome = "hg19")
  gr_a <- import("cpq.bed", genome = genome)
  gr_b <- import("exons.bed", genome = genome)
  pairs <- findOverlapPairs(gr_a, gr_b, ignore.strand = TRUE)
  ans <- pairs
  mcols(ans)$overlap_width <- width(pintersect(ans, ignore.strand = TRUE))
  ans
}
```

This code reveals that *GRanges*, along with every other vector-like object in the Ranges infrastructure, is capable of storing tabular annotations, accessible via `mcols`. We actually saw this before with the “name” column on the CpG Islands. Here, we use it to store the overlap width.

3.7 Counting the number of overlaps

A common query, particularly in RNA-seq analysis, is how many ranges in the subject overlap each query range. The `countOverlaps` function serves this particular purpose:

```
> bedtools_intersect("-a cpq.bed -b exons.bed -g hg19 -c")
{
  genome <- Seqinfo(genome = "hg19")
  gr_a <- import("cpq.bed", genome = genome)
  gr_b <- import("exons.bed", genome = genome)
  ans <- gr_a
  mcols(ans)$overlap_count <- countOverlaps(gr_a, gr_b, ignore.strand = TRUE)
  ans
}
```

3.8 Excluding queries with overlaps

We might instead want to exclude all query ranges that overlap any subject range, i.e., any CpG island that overlaps an exon. The `subsetByOverlaps` function is tasked with restricting by overlap. By passing `invert=TRUE`, we exclude ranges with overlaps.

```
> bedtools_intersect("-a cpq.bed -b exons.bed -g hg19 -v")
{
  genome <- Seqinfo(genome = "hg19")
  gr_a <- import("cpq.bed", genome = genome)
  gr_b <- import("exons.bed", genome = genome)
  subsetByOverlaps(gr_a, gr_b, invert = TRUE, ignore.strand = TRUE)
}
```

3.9 Restricting by fraction of overlap

The *bedtools* suite has deep support for restricting overlaps by the fraction of the query/subject range that is overlapped. This is not directly supported by the *Bioconductor* infrastructure, but we can filter post-hoc:

```
> bedtools_intersect("-a cpg.bed -b exons.bed -g hg19 -f 0.5 -wo")
{
  genome <- Seqinfo(genome = "hg19")
  gr_a <- import("cpg.bed", genome = genome)
  gr_b <- import("exons.bed", genome = genome)
  pairs <- findOverlapPairs(gr_a, gr_b, ignore.strand = TRUE)
  olap <- pintersect(pairs, ignore.strand = TRUE)
  keep <- width(olap)/width(first(pairs)) >= 0.5
  pairs <- pairs[keep]
  ans <- pairs
  mcols(ans)$overlap_width <- width(olap)[keep]
  ans
}
```

3.10 Performance

Comparing the performance of *bedtools* and *IRanges* is like comparing apples and oranges. The typical *Bioconductor* workflow imports the data once, paying an upfront cost, and then operates efficiently on in-memory data structures. The BED parser is implemented in R code and will not compete with the parsing performance of special purpose C code. The intersect operation itself is also slower than *bedtools*, but it's still reasonably close for being mostly implemented in R.

```
> a <- import("exons.bed")
> b <- import("hesc.chromHmm.bed")
> system.time(pintersect(findOverlapPairs(a, b, ignore.strand=TRUE),
+                           ignore.strand=TRUE))

   user  system elapsed 
0.424   0.015   0.560
```

3.11 Multiple subjects

Often, we are interested in intersections with multiple annotation tracks, or multiple samples. Note that the command line parser used by *helloRanges* requires that the filenames be comma-separated, instead of space-separated. This is probably more readable anyway.

```
> code <- bedtools_intersect(
+   paste("-a exons.bed",
+         "-b cpg.bed,gwas.bed,hesc.chromHmm.bed -wa -wb -g hg19",
+         "-names cpg,gwas,chromhmm"))
> ans <- eval(code)
> code

{
  genome <- Seqinfo(genome = "hg19")
```

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```
gr_a <- import("exons.bed", genome = genome)
b <- c("cpg.bed", "gwas.bed", "hesc.chromHmm.bed")
names(b) <- c("cpg", "gwas", "chromhmm")
bl <- List(lapply(b, import, genome = genome))
gr_b <- stack(bl, "b")
pairs <- findOverlapPairs(gr_a, gr_b, ignore.strand = TRUE)
ans <- pairs
ans
}
```

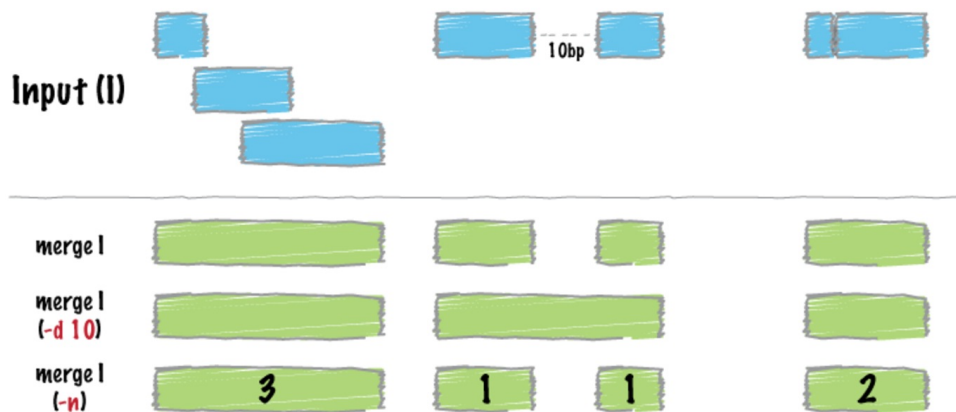
Inspecting the code, we see that we need to loop over the database files and then `stack` them into a single *GRanges* grouped by the column "b":

```
> second(ans)

GRanges object with 541210 ranges and 2 metadata columns:
      seqnames      ranges strand |      b      name
      <Rle>      <IRanges> <Rle> | <Rle> <character>
[1]   chr1      11538-11937      * | chromhmm 11_Weak_Txn
[2]   chr1      11938-12137      * | chromhmm 14_Repetitive/CNV
[3]   chr1      12138-14137      * | chromhmm 11_Weak_Txn
[4]   chr1      12138-14137      * | chromhmm 11_Weak_Txn
[5]   chr1      12138-14137      * | chromhmm 11_Weak_Txn
...     ...             ...   ...   ...
[541206] chrY 59213795-59214183      * |   cpg      CpG:_36
[541207] chrY 59213795-59214183      * |   cpg      CpG:_36
[541208] chrY 59213795-59214183      * |   cpg      CpG:_36
[541209] chrY 59213795-59214183      * |   cpg      CpG:_36
[541210] chrY 59213795-59214183      * |   cpg      CpG:_36
-----
seqinfo: 298 sequences (2 circular) from hg19 genome
```

The "b" column is an *Rle* object, a run-length encoded form of an ordinary R vector, in this case a factor. Since the data are sorted into groups, this encoding is more efficient than a dense representation. The "seqnames" and "strand" columns also benefit from run-length encoding. Not only can we fit more data into memory, many operations become faster.

4 Merge



There are many ways to summarize interval data. In the Ranges infrastructure, we call some of them **range** (min start to max end), **reduce** (*bedtools merge*), **disjoin** (involved in *bedtools multiinter*) and **coverage** (counting the number of ranges overlapping each position, *bedtools genomecov*). We are presently concerned with **reduce**, which combines overlapping and adjacent ranges into a single range. The corresponding *bedtools merge* command requires the data to be sorted; however, **reduce** does not have this constraint.

```
> bedtools_merge("-i exons.bed")
{
  genome <- Seqinfo(genome = NA_character_)
  gr_a <- import("exons.bed", genome = genome)
  ans <- reduce(gr_a, ignore.strand = TRUE)
  ans
}
```

4.1 Aggregation

As with any reduction, we often want to simultaneously aggregate associated variables and report the summaries in the result.

We count the number of ranges overlapping each merged range:

```
> code <- bedtools_merge("-i exons.bed -c 1 -o count")
> code
{
  genome <- Seqinfo(genome = NA_character_)
  gr_a <- import("exons.bed", genome = genome)
  ans <- reduce(gr_a, ignore.strand = TRUE, with.revmap = TRUE)
  mcols(ans) <- aggregate(gr_a, mcols(ans)$revmap, seqnames.count = lengths(seqnames),
    drop = FALSE)
  ans
}
```

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The key to aggregation with `reduce` is the `with.revmap=TRUE` argument. That yields a “revmap” column on the result. It is an *IntegerList* holding the subscripts corresponding to each group. We pass it to `aggregate` to indicate the grouping. The named arguments to `aggregate`, in this case `seqnames.count`, are effectively evaluated with respect to each group (although they are actually evaluated only once).

This yields the result:

```
> eval(code)

GRanges object with 229241 ranges and 2 metadata columns:
      seqnames      ranges strand |      grouping
      <Rle>       <IRanges> <Rle> | <ManyToManyGrouping>
[1]   chr1      11874-12227    * |           1
[2]   chr1      12613-12721    * |           2
[3]   chr1      13221-14829    * |          3,4
[4]   chr1      14970-15038    * |           5
[5]   chr1      15796-15947    * |           6
...     ...                ...  ...  .         ...
[229237] chrY 59337091-59337236    * | 459744,459745
[229238] chrY 59337949-59338150    * | 459746,459747
[229239] chrY 59338754-59338859    * | 459748,459749
[229240] chrY 59340194-59340278    * |         459750
[229241] chrY 59342487-59343488    * | 459751,459752

      seqnames.count
      <integer>
[1]             1
[2]             1
[3]             2
[4]             1
[5]             1
...           ...
[229237]         2
[229238]         2
[229239]         2
[229240]         1
[229241]         2
-----
seqinfo: 49 sequences from an unspecified genome; no seqlengths
```

We see that the grouping has been preserved on the object, in case we wish to aggregate further through joins.

Counting the overlaps by counting the “seqnames” is a little circuitous. Instead, we could have just counted the elements in each group:

```
> identical(lengths(ans$grouping), ans$seqnames.count)
[1] TRUE
```

Note that this counting is very fast, because the “revmap” *IntegerList* is not actually a list, but a partitioned vector, and the partitioning already encodes the counts. This is an example of where the flexibility and efficient in-memory representations of *Bioconductor* are particularly effective.

4.2 Merging close features

By default, features are merged if they are overlapping or adjacent, i.e., the `min.gapwidth` (the minimum gap allowed to not be merged) is 1. To merge features that are up to, say, 1000bp away, we need to pass `min.gapwidth=1001`:

```
> bedtools_merge("-i exons.bed -d 1000")
{
  genome <- Seqinfo(genome = NA_character_)
  gr_a <- import("exons.bed", genome = genome)
  ans <- reduce(gr_a, ignore.strand = TRUE, min.gapwidth = 1001L)
  ans
}
```

Here is another example showing how to merge multiple columns at once:

```
> bedtools_merge("-i exons.bed -d 90 -c 1,4 -o count,collapse")
{
  genome <- Seqinfo(genome = NA_character_)
  gr_a <- import("exons.bed", genome = genome)
  ans <- reduce(gr_a, ignore.strand = TRUE, with.revmap = TRUE,
    min.gapwidth = 91L)
  mcols(ans) <- aggregate(gr_a, mcols(ans)$revmap, seqnames.count = lengths(seqnames),
    name.collapse = unstrsplit(name, ","), drop = FALSE)
  ans
}
```

5 Finding the Gaps

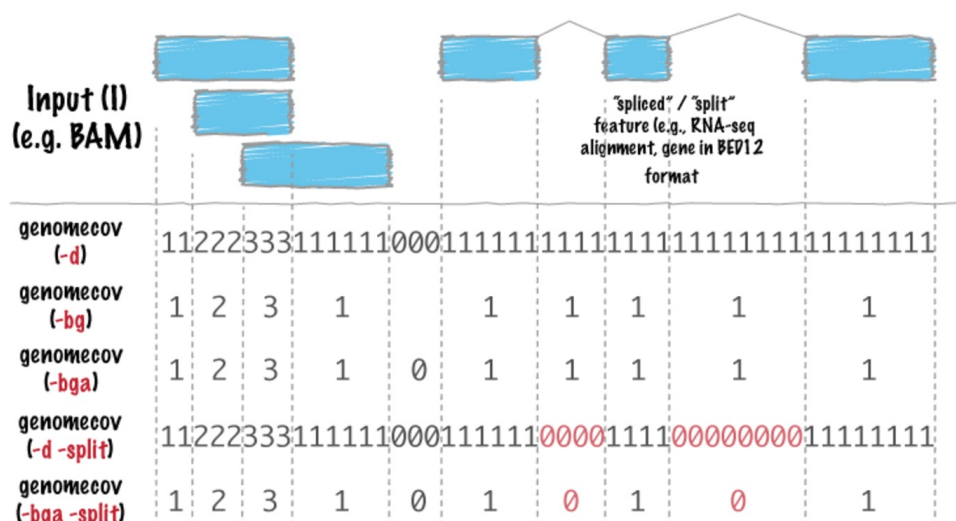


The *bedtools complement* tool finds the gaps in the sequence, i.e., the regions of sequence the track does not cover. This is where having the sequence bounds is critical.

```
> bedtools_complement("-i exons.bed -g hg19.genome")
{
  genome <- import("hg19.genome")
  gr_a <- import("exons.bed", genome = genome)
  ans <- setdiff(as(seqinfo(gr_a), "GRanges"), unstrand(gr_a))
  ans
}
```

The call to `setdiff` is a set operation, along with `intersect` and `union`. Set operations behave a bit surprisingly with respect to strand. The “unstranded” features, those with “*” for their strand, are considered to be in a separate space from the stranded features. If we pass `ignore.strand=TRUE`, both arguments are unstranded and the result is unstranded (strand information is discarded). This makes sense, because there is no obvious way to merge a stranded and unstranded feature. Since we are looking for the gaps, we do not care about the strand, so discarding the strand is acceptable. Best practice is to make this explicit by calling `unstrand` instead of assuming the reader understands the behavior of `ignore.strand=TRUE`.

6 Computing Genomic Coverage



One of the useful ways to summarize ranges, particularly alignments, is to count how many times a position is overlapped by a range. This is called the coverage. Unlike *bedtools genomecov*, we do not require the data to be sorted.

6.1 Coverage vector

To compute the coverage, we just call `coverage`. For consistency with *bedtools*, which drops zero runs with the “-bg” option, we convert the coverage vector to a *GRanges* and subset:

```
> bedtools_genomecov("-i exons.bed -g hg19.genome -bg")
{
  genome <- import("hg19.genome")
  gr_a <- import("exons.bed", genome = genome)
  cov <- coverage(gr_a)
  ans <- GRanges(cov)
  ans <- subset(ans, score > 0)
  ans
}
```

6.2 Coverage histogram

The default behavior of *genomecov* is to compute a histogram showing the number and fraction of positions covered at each level. It does this for the individual chromosome, and the entire genome. While computing the coverage vector is as simple as calling `coverage`, forming the histogram is a bit complicated. This is a bit esoteric, but it lets us demonstrate how to aggregate data in R:

```
> code <- bedtools_genomecov("-i exons.bed -g hg19.genome")
> ans <- eval(code)
> code
{
  genome <- import("hg19.genome")
  gr_a <- import("exons.bed", genome = genome)
  cov <- coverage(gr_a)
  tablist <- List(lapply(cov, table))
  mcols(tablist)$len <- lengths(cov, use.names = FALSE)
  covhist <- stack(tablist, "seqnames", "count", "coverage")
  margin <- aggregate(covhist, ~coverage, count = sum(NumericList(count)))[-1L]
  margin <- DataFrame(seqnames = Rle("genome"), margin, len = sum(as.numeric(lengths(cov))))
  covhist <- rbind(covhist, margin)
  ans <- within(covhist, fraction <- count/len)
  ans
}
```

The `cov` object is an *RleList*, with one *Rle* per sequence (chromosome).

```
> cov

RleList of length 93
$chr1
integer-Rle of length 249250621 with 46230 runs
  Lengths: 11873  354  385  109  499 ...  64  2722  677  1868 37276
  Values :    0    1    0    1    0 ...    2    0    1    2    0

$chr10
integer-Rle of length 135534747 with 18962 runs
  Lengths: 92827 1227  500  111  78 ... 1936 1120 3293  260 36289
  Values :    0    1    0    1    0 ...    0    8    0    8    0

$chr11
integer-Rle of length 135006516 with 25138 runs
  Lengths: 126986 4387  93  58  227 ... 104 22067  864 630961
  Values :    0    1    0    1    0 ...    1    0    1    0

$chr11_gl000202_random
integer-Rle of length 40103 with 1 run
  Lengths: 40103
  Values :    0

$chr12
integer-Rle of length 133851895 with 25290 runs
```


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```
Lengths: 87983  34  239  136  177 ... 2463  271 1428  30 39473
Values :    0    1    0    1    0 ...    0    1    0    1    0

...
<88 more elements>
```

We tabulate each coverage vector individually, then stack the tables into an initial histogram. Then, we aggregate over the entire genome and combine the genome histogram with the per-chromosome histogram. The call to `NumericList` is only to avoid integer overflow. Finally, we compute the fraction covered and end up with:

```
> ans

DataFrame with 653 rows and 5 columns
  seqnames coverage    count      len  fraction
   <Rle> <factor> <numeric> <numeric> <numeric>
1   chr1         0 241996316 249250621 0.97089554
2   chr1         1  4276763 249250621 0.01715848
3   chr1         2 1475526 249250621 0.00591985
4   chr1         3  710135 249250621 0.00284908
5   chr1         4  388193 249250621 0.00155744
...      ...      ...      ...      ...      ...
649 genome        73    135 3137161264 4.30325e-08
650 genome        74    263 3137161264 8.38338e-08
651 genome        75   1921 3137161264 6.12337e-07
652 genome        76    705 3137161264 2.24725e-07
653 genome        77   2103 3137161264 6.70351e-07
```

This takes 3 minutes for `bedtools`, but closer to 3 seconds for us, probably because it is working too hard to conserve memory.

7 Combining operations

7.1 Chaining

Most real-world workflows involve multiple operations, chained together. The R objects produced [HelloRanges](#) can be passed directly to existing R functions, and [HelloRanges](#) defines an ordinary R function corresponding to each `bedtools` operation. The arguments of the function correspond to `bedtools` arguments, except they can be R objects, like `GRanges`, in addition to filenames. These functions with ordinary R semantics are prefixed by `R_`, so the analog to `bedtools_intersect` is `R_bedtools_intersect`.

Consider a use case similar to the one mentioned in the `bedtools` tutorial: find the regions of the CpG islands that are not covered by exons. We could do this directly with `bedtools_subtract`, but let us instead compute the coverage of the exons, find the regions of zero coverage, and intersect those with the CpG islands.

First, we generate the code for the coverage operation (and ideally copy it to a script). The result of evaluating that code is a `GRanges`, which we subset for the regions with zero score.

```
> code <- bedtools_genomecov("-i exons.bed -g hg19.genome -bga")
> gr0 <- subset(eval(code), score == 0L) # compare to: awk '$4==0'
```

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```
> gr0
```

GRanges object with 229334 ranges and 1 metadata column:

	seqnames	ranges	strand	score
	<Rle>	<IRanges>	<Rle>	<integer>
[1]	chr1	1-11873	*	0
[2]	chr1	12228-12612	*	0
[3]	chr1	12722-13220	*	0
[4]	chr1	14830-14969	*	0
[5]	chr1	15039-15795	*	0
...
[229330]	chrY	59337237-59337948	*	0
[229331]	chrY	59338151-59338753	*	0
[229332]	chrY	59338860-59340193	*	0
[229333]	chrY	59340279-59342486	*	0
[229334]	chrY	59343489-59373566	*	0

seqinfo: 93 sequences from an unspecified genome

Next, we pass `gr0` directly to the R analog of *intersect*, `R_bedtools_intersect`:

```
> code <- R_bedtools_intersect("cpg.bed", gr0)
> code

{
  genome <- Seqinfo(genome = NA_character_)
  gr_a <- import("cpg.bed", genome = genome)
  gr_b <- gr0
  pairs <- findOverlapPairs(gr_a, gr_b, ignore.strand = TRUE)
  ans <- pintersect(pairs, ignore.strand = TRUE)
  ans
}
```

The generated code already refers to `gr0` explicitly, so it is easy to copy this into the script.

To generalize, the chaining workflow is:

1. Generate code for first operation,
2. Integrate and evaluate the code,
3. Interactively inspect the result of evaluation,
4. Perform intermediate operations, while inspecting results,
5. Call `R_` analog to generate second stage code.

Generating and integrating *R* code is the best way to learn, and the best way to produce a readable, flexible and performant script. However, there are probably those who are tempted to evaluate the code directly, as we have done in this vignette. Further, there are those who wish to chain these operations together with the so-called “pipe” operator, because it would come so tantalizingly close to the syntax of the shell. Thus, we created a third family of functions, prefixed by `do_`, which provide the same interface as the `R_` family, except they evaluate the generated code:

```
> do_bedtools_genomecov("exons.bed", g="hg19.genome", bga=TRUE) %>%
+   subset(score > 0L) %>%
+   do_bedtools_intersect("cpg.bed", .)
```

7.2 Coalescence

In the previous section, we chained together independent operations. Having access to the underlying code gives us the flexibility to merge operations so that they are faster than the sum of their parts. We call this coalescence.

Consider a use case cited by the *bedtools* tutorial: compute the distribution of coverage over all exons. To integrate better with this tutorial, we adapt that to finding the distribution of exon coverage over all CpG islands.

We could mimic the example by computing the coverage complete histogram and extracting only the margin:

```
> bedtools_coverage("-a cpg.bed -b exons.bed -hist -g hg19.genome")
{
  genome <- import("hg19.genome")
  gr_a <- import("cpg.bed", genome = genome)
  gr_b <- import("exons.bed", genome = genome)
  cov <- unname(coverage(gr_b)[gr_a])
  tab <- t(table(cov))
  tab <- cbind(tab, all = rowSums(tab))
  covhist <- DataFrame(as.table(tab))
  colnames(covhist) <- c("coverage", "a", "count")
  len <- c(lengths(cov, use.names = FALSE), sum(lengths(cov)))
  covhist$len <- rep(len, each = nrow(tab))
  covhist <- subset(covhist, count > 0L)
  covhist$fraction <- with(covhist, count/len)
  ans <- gr_a
  covhistList <- split(covhist, ~a)[, -2L]
  mcols(ans)$coverage <- head(covhistList, -1L)
  metadata(ans)$coverage <- covhistList$all
  ans
}
```

The code is quite complex, because the Ranges infrastructure does not attempt to generate high-level summaries of the data. The rationale, which is validated in this case, is that the desired summary depends on the specific question, and the number of questions is effectively infinite. In this case, we only care about the margin, i.e., `metadata(ans)$coverage`.

Thus, we can simplify the code. We begin with the same lines:

```
> genome <- import("hg19.genome")
> gr_a <- import("cpg.bed", genome = genome)
> gr_b <- import("exons.bed", genome = genome)
> cov <- unname(coverage(gr_b)[gr_a])
```

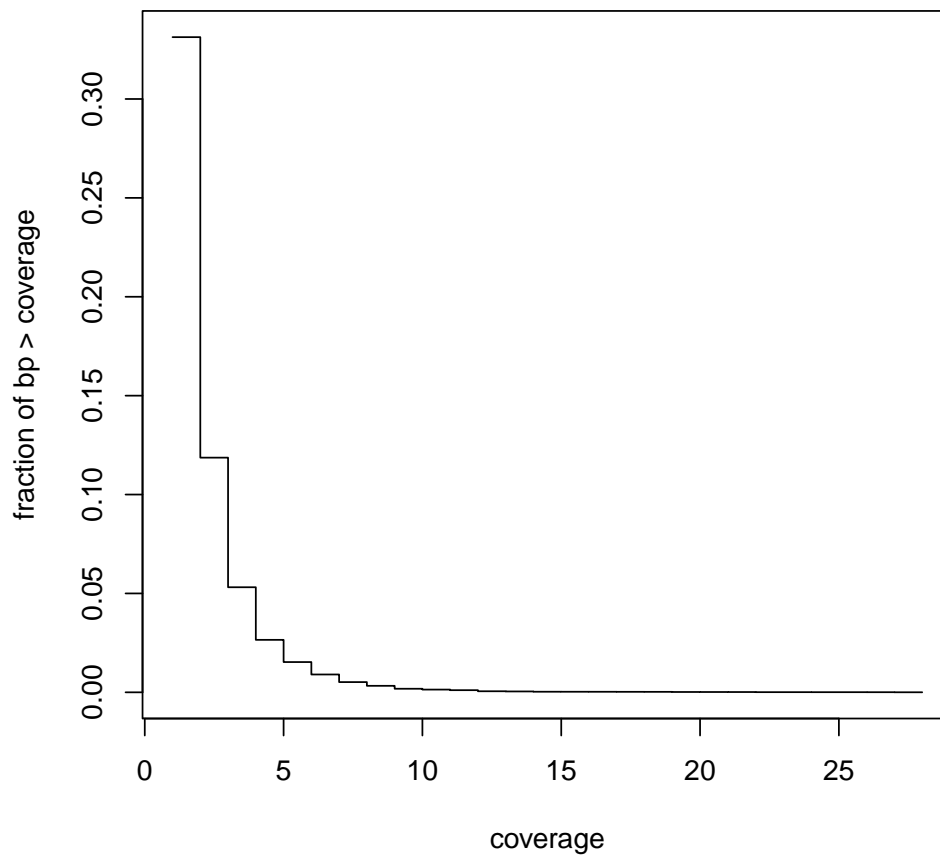
And summarize all of the coverage at once:

```
> all_cov <- unlist(cov)
> df <- as.data.frame(table(coverage=all_cov))
> df$fraction <- df$Freq / length(all_cov)
```

This is much faster, because we are only computing one table, not 30,000, and the `table` method for `Rle` is very efficient.

We now have a simple `data.frame` that we can plot as an inverted ECDF:

```
> plot((1-cumsum(fraction)) ~ as.integer(coverage), df, type="s",
+      ylab = "fraction of bp > coverage", xlab="coverage")
```



8 Jaccard Statistic

In order to compare the Dnasel hypersensitivity across tissues, we will employ the *bedtools jaccard* statistic, a measure of similarity between two tracks. It is defined as the total width of their intersection over the total width of their union.

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We might expect, for example, that the similarity within a tissue is higher than that between two tissues, and this is indeed the case:

```
> code <- bedtools_jaccard(
+   paste("-a fHeart-DS16621.hotspot.twopass.fdr0.05.merge.bed",
+         "-b fHeart-DS15839.hotspot.twopass.fdr0.05.merge.bed"))
> heart_heart <- eval(code)
> code <- bedtools_jaccard(
+   paste("-a fHeart-DS16621.hotspot.twopass.fdr0.05.merge.bed",
+         "-b fSkin_fibro_bicep_R-DS19745.hg19.hotspot.twopass.fdr0.05.merge.bed"))
> heart_skin <- eval(code)
> mstack(heart_heart=heart_heart, heart_skin=heart_skin)

DataFrame with 2 rows and 5 columns
      name intersection   union   jaccard n_intersections
    <Rle>    <integer> <integer> <numeric>      <integer>
1 heart_heart      81269248 160493950  0.506370         130852
2 heart_skin       28076951 164197278  0.170995          73261
```

The generated code makes the statistic self-documenting:

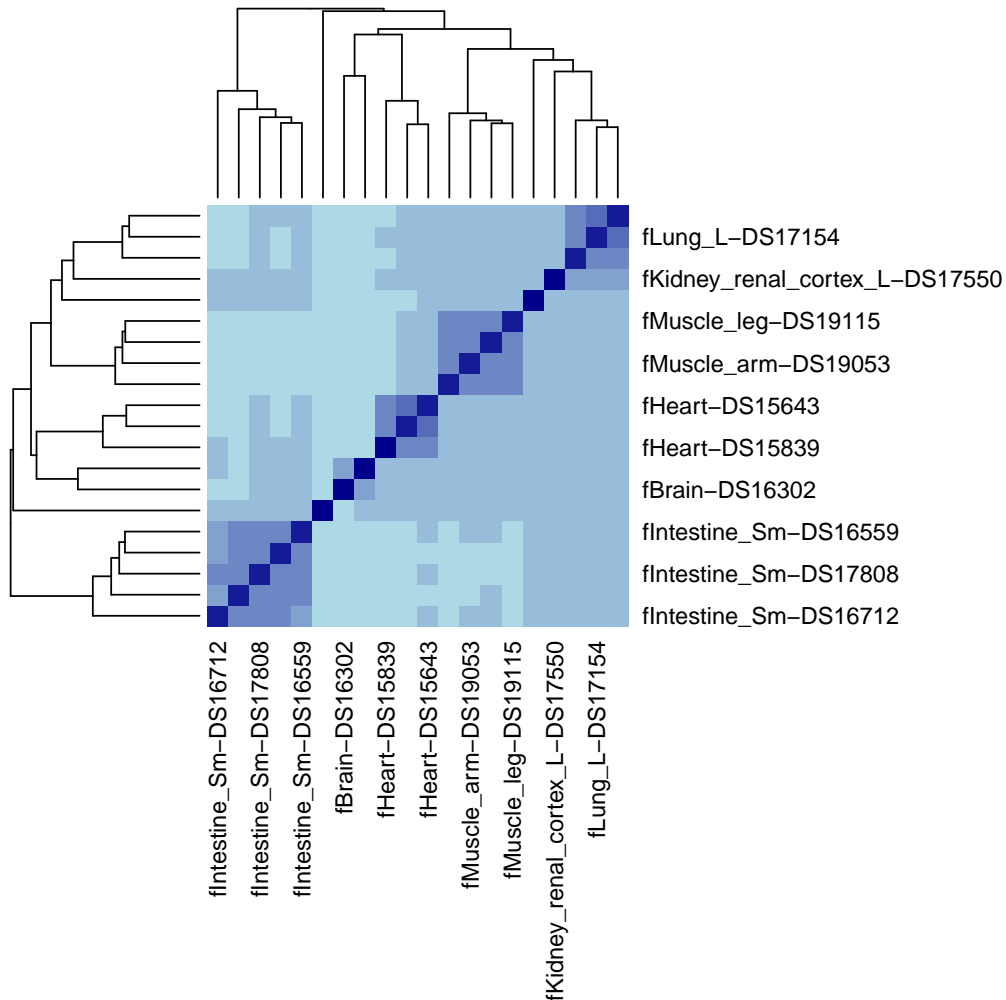
```
> code
{
  genome <- NA_character_
  gr_a <- import("fHeart-DS16621.hotspot.twopass.fdr0.05.merge.bed",
    genome = genome)
  gr_b <- import("fSkin_fibro_bicep_R-DS19745.hg19.hotspot.twopass.fdr0.05.merge.bed",
    genome = genome)
  intersects <- intersect(gr_a, gr_b, ignore.strand = TRUE)
  intersection <- sum(width(intersects))
  union <- sum(width(union(gr_a, gr_b, ignore.strand = TRUE)))
  ans <- DataFrame(intersection, union, jaccard = intersection/union,
    n_intersections = length(intersects))
  ans
}
```

We can compute the statistic over all pairs of samples using functionality included with R, through the *parallel* package. There is no need to learn yet another syntax, such as that of the *parallel* UNIX utility. Nor do we need to download a custom python script, and repeatedly call perl and awk.

```
> files <- Sys.glob("*.merge.bed")
> names(files) <- sub("\\\\.*", "", files)
> ncores <- if (.Platform$OS.type == "windows") 1L else 4L
> library(parallel)
> ans <- outer(files, files,
+   function(a, b) mcmapply(do_bedtools_jaccard, a, b,
+   mc.cores=ncores))
> jaccard <- apply(ans, 1:2, function(x) x[[1]]$jaccard)
```

Since we are already in R, it is easy to create a simple plot:

```
> palette <- colorRampPalette(c("lightblue", "darkblue"))(9)
> heatmap(jaccard, col=palette, margin=c(14, 14))
```



9 Exercises

These were adapted from the *bedtools* tutorial. Try to complete these exercises using *Bioconductor* directly.

1. Create a *GRanges* containing the non-exonic regions of the genome.
2. Compute the average distance from the GWAS SNPs to the closest exon (Hint: `?bedtools_closest` and `?distanceToNearest`).
3. Compute the exon coverage in 500kb windows across the genome (Hint: `?bedtools_makewindows` and `?tileGenome`).
4. How many exons are completely overlapped by an enhancer (from 'hesc.chromHmm.bed') (Hint: `?%within%`)?

5. What fraction of the disease-associated SNPs are exonic (Hint: (Hint: `2%over%`))?
6. Create intervals representing the canonical 2bp splice sites on either side of each exon (bonus: exclude splice sites at the first and last exons) (Hint: `?bedtools_flank`, `?intransByTranscript`).
7. Which hESC ChromHMM state represents the most number of base pairs in the genome? (Hint: `?xtabs`).

9.1 Answers

Below, we give the *bedtools*-style answer first, followed by the essential call against the *Bioconductor* API.

First, we load the files into *R* objects for convenience:

```
> genome <- import("hg19.genome")
> exons <- import("exons.bed", genome=genome)
> gwas <- import("gwas.bed", genome=genome)
> hesc.chromHmm <- import("hesc.chromHmm.bed", genome=genome)
```

Here are the numbered answers:

```
1. > bedtools_complement("-i exons.bed -g hg19.genome")
{
  genome <- import("hg19.genome")
  gr_a <- import("exons.bed", genome = genome)
  ans <- setdiff(as(seqinfo(gr_a), "GRanges"), unstrand(gr_a))
  ans
}
```

```
> ## or without HelloRanges:
> setdiff(as(seqinfo(exons), "GRanges"), unstrand(exons))
```

GRanges object with 229334 ranges and 0 metadata columns:

	seqnames	ranges	strand
	<Rle>	<IRanges>	<Rle>
[1]	chr1	1-11873	*
[2]	chr1	12228-12612	*
[3]	chr1	12722-13220	*
[4]	chr1	14830-14969	*
[5]	chr1	15039-15795	*
...
[229330]	chrY	59337237-59337948	*
[229331]	chrY	59338151-59338753	*
[229332]	chrY	59338860-59340193	*
[229333]	chrY	59340279-59342486	*
[229334]	chrY	59343489-59373566	*

seqinfo: 93 sequences from hg19 genome

```
2. > bedtools_closest("-a gwas.bed -b exons.bed -d")
```

```
{
  genome <- Seqinfo(genome = NA_character_)
  gr_a <- import("gwas.bed", genome = genome)
  gr_b <- import("exons.bed", genome = genome)
  hits <- nearest(gr_a, gr_b, ignore.strand = TRUE, select = "all")
  ans <- pair(gr_a, gr_b, hits, all.x = TRUE)
  mcols(ans)$distance <- distance(ans)
  ans
}
```

> ## or

> distanceToNearest(gwas, exons)

Hits object with 17674 hits and 1 metadata column:

	queryHits	subjectHits	distance
	<integer>	<integer>	<integer>
[1]	1	235	1319
[2]	2	249	0
[3]	3	426	0
[4]	4	1163	2385
[5]	5	1163	2894
...
[17670]	17676	455399	371
[17671]	17677	455399	371
[17672]	17678	455464	5898
[17673]	17679	455674	324706
[17674]	17680	456097	135628

queryLength: 17680 / subjectLength: 459752

3. > code <- bedtools_makewindows("-g hg19.genome -w 500000")

> code

```
{
  genome <- import("hg19.genome")
  ans <- tile(as(genome, "GRanges"), width = 500000L)
  ans
}
```

> windows <- unlist(eval(code))

> R_bedtools_intersect(windows, exons, c=TRUE)

```
{
  genome <- Seqinfo(genome = NA_character_)
  gr_a <- windows
  gr_b <- exons
  ans <- gr_a
  mcols(ans)$overlap_count <- countOverlaps(gr_a, gr_b, ignore.strand = TRUE)
  ans
}
```

> ## or

> str(countOverlaps(tileGenome(seqinfo(exons), tilewidth=500000),


```
+               exons))
int [1:6275] 37 197 477 445 209 96 83 271 9 12 ...
```

```
4. > bedtools_intersect(
+   paste("-a exons.bed -b <\"grep Enhancer hesc.chromHmm.bed\"",
+       "-f 1.0 -wa -u"))
{
  genome <- Seqinfo(genome = NA_character_)
  gr_a <- import("exons.bed", genome = genome)
  gr_b <- import(BEDFile(pipe("\"grep Enhancer hesc.chromHmm.bed\"")),
    genome = genome)
  hits <- findOverlaps(gr_a, gr_b, ignore.strand = TRUE, type = "within")
  gr_a[countQueryHits(hits) > 0L]
}
> quote(length(ans))
length(ans)
> ## or
> sum(exons %within%
+   subset(hesc.chromHmm, grepl("Enhancer", name)))
[1] 13746
```

```
5. > bedtools_intersect("-a gwas.bed -b exons.bed -u")
{
  genome <- Seqinfo(genome = NA_character_)
  gr_a <- import("gwas.bed", genome = genome)
  gr_b <- import("exons.bed", genome = genome)
  subsetByOverlaps(gr_a, gr_b, ignore.strand = TRUE)
}
> quote(length(gr_a)/length(ans))
length(gr_a)/length(ans)
> ## or
> mean(gwas %over% exons)
[1] 0.09191176
```

```
6. > bedtools_flank("-l 2 -r 2 -i exons.bed -g hg19.genome")
{
  genome <- import("hg19.genome")
  gr_a <- import("exons.bed", genome = genome)
  left <- flank(gr_a, 2, ignore.strand = TRUE)
  right <- flank(gr_a, 2, start = FALSE, ignore.strand = TRUE)
  ans <- zipup(Pairs(left, right))
  ans
}
```

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```
> ## or, bonus:
> txid <- sub("_exon.*", "", exons$name)
> tx <- split(exons, txid)
> bounds <- range(tx)
> transpliced <- lengths(bounds) > 1
> introns <- unlist(psetdiff(unlist(bounds[!transpliced]),
+                           tx[!transpliced]))
> Pairs(resize(introns, 2L), resize(introns, 2L, fix="end"))

Pairs object with 386125 pairs and 0 metadata columns:
      first second
      <GRanges> <GRanges>
[1] chr12:9220777-9220778:- chr12:9220436-9220437:-
[2] chr12:9221334-9221335:- chr12:9220821-9220822:-
[3] chr12:9222339-9222340:- chr12:9221439-9221440:-
[4] chr12:9223082-9223083:- chr12:9222410-9222411:-
[5] chr12:9224953-9224954:- chr12:9223175-9223176:-
...
[386121] chr9:123626325-123626326:- chr9:123625028-123625029:-
[386122] chr9:123627986-123627987:- chr9:123626395-123626396:-
[386123] chr9:123628304-123628305:- chr9:123628109-123628110:-
[386124] chr9:123629146-123629147:- chr9:123628375-123628376:-
[386125] chr9:123631084-123631085:- chr9:123629244-123629245:-

> ## better way to get introns:
> library(TxDb.Hsapiens.UCSC.hg19.knownGene)
> txdb <- TxDb.Hsapiens.UCSC.hg19.knownGene
> introns <- unlist(intronsByTranscript(txdb))
```

```
7. > system.time(names(which.max(xtabs(width ~ name,
+                                     hesc.chromHmm))))

      user system elapsed
0.091    0.006    0.118

> ## or
> names(which.max(sum(with(hesc.chromHmm,
+                           splitAsList(width, name)))))

[1] "13_Heterochrom/lo"

> ## or
> df <- aggregate(hesc.chromHmm, ~ name, totalWidth=sum(width))
> df$name[which.max(df$totalWidth)]

[1] 13_Heterochrom/lo
15 Levels: 10_Txn_Elongation 11_Weak_Txn 12_Repressed ... 9_Txn_Transition
```

References

- [1] Matthew T Maurano, Richard Humbert, Eric Rynes, Robert E Thurman, Eric Haugen, Hao Wang, Alex P Reynolds, Richard Sandstrom, Hongzhu Qu, Jennifer Brody, et al. Systematic localization of common disease-associated variation in regulatory dna. *Science*, 337(6099):1190–1195, 2012.