METHOD ARTICLE

Fluent genomics with *plyranges* and *tximeta* [version 1; peer review: 1 approved, 2 approved with reservations]

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**Abstract**

We construct a simple workflow for fluent genomics data analysis using the R/Bioconductor ecosystem. This involves three core steps: *import* the data into an appropriate abstraction, *model* the data with respect to the biological questions of interest, and *integrate* the results with respect to their underlying genomic coordinates. Here we show how to implement these steps to integrate published RNA-seq and ATAC-seq experiments on macrophage cell lines. Using *tximeta*, we *import* RNA-seq transcript quantifications into an analysis-ready data structure, called the *SummarizedExperiment*, that contains the ranges of the reference transcripts and metadata on their provenance. Using *SummarizedExperiments* to represent the ATAC-seq and RNA-seq data, we *model* differentially accessible (DA) chromatin peaks and differentially expressed (DE) genes with existing Bioconductor packages. Using *plyranges* we then *integrate* the results to see if there is an enrichment of DA peaks near DE genes by finding overlaps and aggregating over log-fold change thresholds. The combination of these packages and their integration with the Bioconductor ecosystem provide a coherent framework for analysts to iteratively and reproducibly explore their biological data.

**Keywords**

Gene Expression, Chromatin Accessibility, Workflow, Data Integration, Bioconductor, plyranges, tximeta

This article is included in the Bioconductor gateway.
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Author roles: Lee S: Conceptualization, Methodology, Software, Visualization, Writing – Original Draft Preparation, Writing – Review & Editing; Lawrence M: Conceptualization, Software, Supervision, Writing – Review & Editing; Love MI: Conceptualization, Data Curation, Formal Analysis, Methodology, Software, Supervision, Visualization, Writing – Original Draft Preparation, Writing – Review & Editing

Competing interests: No competing interests were disclosed.

Grant information: SL is supported by an Australian Government Research Training Program (RTP) scholarship with a top up scholarship from CSL Limited. MIL is supported by NIH grant R01 HG009937.
The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

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How to cite this article: Lee S, Lawrence M and Love MI. Fluent genomics with plyranges and tximeta [version 1; peer review: 1 approved, 2 approved with reservations] F1000Research 2020, 9:109 https://doi.org/10.12688/f1000research.22259.1

First published: 12 Feb 2020, 9:109 https://doi.org/10.12688/f1000research.22259.1
Introduction

In this workflow, we examine a subset of the RNA-seq and ATAC-seq data from Alasoo et al. (2018), a study that involved treatment of macrophage cell lines from a number of human donors with interferon gamma (IFNg), Salmonella infection, or both treatments combined. Alasoo et al. (2018) examined gene expression and chromatin accessibility in a subset of 86 successfully differentiated induced pluripotent stem cells (iPSC) lines, and compared baseline and response with respect to chromatin accessibility and gene expression at specific quantitative trait loci (QTL). The authors found that many of the stimulus-specific expression QTL were already detectable as chromatin QTL in naive cells, and further hypothesize about the nature and role of transcription factors implicated in the response to stimulus.

We will perform a much simpler analysis than the one found in Alasoo et al. (2018), using their publicly available RNA-seq and ATAC-seq data (ignoring the genotypes). We will examine the effect of IFNg stimulation on gene expression and chromatin accessibility, and look to see if there is an enrichment of differentially accessible (DA) ATAC-seq peaks in the vicinity of differentially expressed (DE) genes. This is plausible, as the transcriptomic response to IFNg stimulation may be mediated through binding of regulatory proteins to accessible regions, and this binding may increase the accessibility of those regions such that it can be detected by ATAC-seq.

Throughout the workflow (Figure 1), we will use existing Bioconductor infrastructure to understand these datasets. In particular, we will emphasize the use of the Bioconductor packages plyranges and tximeta. The plyranges package fluently transforms data tied to genomic ranges using operations like shifting, window construction, overlap detection, etc. It is described by Lee et al. (2019) and leverages underlying core Bioconductor infrastructure (Lawrence et al., 2013; Huber et al., 2015) and the tidyverse design principles Wickham et al. (2019).

The tximeta package described by Love et al. (2019) is used to read RNA-seq quantification data into R/Bioconductor, such that the transcript ranges and their provenance are automatically attached to the object containing expression values and differential expression results.

Figure 1. An overview of the fluent genomics workflow. First, we import data as a SummarizedExperiment object, which enables interoperability with downstream analysis packages. Then we model our assay data, using the existing Bioconductor packages DESeq2 and limma. We take the results of our models for each assay with respect to their genomic coordinates, and integrate them. First, we compute the overlap between the results of each assay, then aggregate over the combined genomic regions, and finally summarize to compare enrichment for differentially expressed genes to non differentially expressed genes. The final output can be used for downstream visualization or further transformation.
Experimental data

The data used in this workflow is available from two packages: the macrophage Bioconductor ExperimentData package and from the workflow package fluentGenomics (Lee & Love, 2020).

The macrophage package contains RNA-seq quantification from 24 RNA-seq samples, a subset of the RNA-seq samples generated and analyzed by Alasoo et al. (2018). The paired-end reads were quantified using Salmon (Patro et al., 2017), using the Gencode 29 human reference transcripts (Frankish et al., 2019). For more details on quantification, and the exact code used, consult the vignette of the macrophage package. The package also contains the Snakemake file that was used to distribute the Salmon quantification jobs on a cluster (Köster & Rahmann, 2012).

The fluentGenomics package (Lee & Love, 2020) contains functionality to download and generate a cached SummarizedExperiment object from the normalized ATAC-seq data provided by Alasoo & Gaffney (2017). This object contains all 145 ATAC-seq samples across all experimental conditions as analyzed by Alasoo et al. (2018). The data can be also be downloaded directly from the Zenodo deposition.

The following code loads the path to the cached data file, or if it is not present, will create the cache and generate a SummarizedExperiment using the the BiocFileCache package (Shepherd & Morgan, 2019).

```r
library(fluentGenomics)
path_to_se <- cache_atac_se()
```

We can then read the cached file and assign it to an object called atac.

```r
atac <- readRDS(path_to_se)
```

A precise description of how we obtained this SummarizedExperiment object can be found in Importing ATAC-seq data as a SummarizedExperiment object.

Import data as a SummarizedExperiment

Using tximeta to import RNA-seq quantification data

First, we specify a directory dir, where the quantification files are stored. You could simply specify this directory with:

```r
dir <- "/path/to/quant/files"
```

where the path is relative to your current R session. However, in this case we have distributed the files in the macrophage package. The relevant directory and associated files can be located using system.file.

```r
dir <- system.file("extdata", package="macrophage")
```

Information about the experiment is contained in the coldata.csv file. We leverage the dplyr and readr packages (as part of the tidyverse) to read this file into R (Wickham et al., 2019). We will see later that plyranges extends these packages to accommodate genomic ranges.

```r
library(dplyr)
library(readr)
colfile <- file.path(dir, "coldata.csv")
coldata <- read_csv(colfile) %>%
          dplyr::select(
            names,
```
```r
id = sample_id,
line = line_id,
condition = condition_name
)

dplyr::mutate(
files = file.path(dir, "quants", names, "quant.sf.gz"),
line = factor(line),
condition = relevel(factor(condition), "naive")
)
```

After we have read the `coldata.csv` file, we select relevant columns from this table, create a new column called `files`, and transform the existing `line` and `condition` columns into factors. In the case of `condition`, we specify the “naive” cell line as the reference level. The `files` column points to the quantifications for each observation – these files have been gzipped, but would typically not have the ‘gz’ ending if used from `Salmon` directly. One other thing to note is the use of the pipe operator, `%>%`, which can be read as “then”, i.e. first read the data, `then` select columns, `then` mutate them.
Now we have a table summarizing the experimental design and the locations of the quantifications. The following lines of code do a lot of work for the analyst: importing the RNA-seq quantification (dropping inferential replicates in this case), locating the relevant reference transcriptome, attaching the transcript ranges to the data, and fetching genome information. Inferential replicates are especially useful for performing transcript-level analysis, but here we will use a point estimate for the per-gene counts and perform gene-level analysis.

The result is a `SummarizedExperiment` object.

```r
suppressPackageStartupMessages(library(SummarizedExperiment))
library(tximeta)
se <- tximeta(coldata, dropInfReps=TRUE)
## importing quantifications
## reading in files with read_tsv
## 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22 23 24
## found matching linked transcriptome:
## [ GENCODE - Homo sapiens - release 29 ]
## loading existing TxDb created: 2019-11-22 01:02:58
## Loading required package: GenomicFeatures
## Loading required package: AnnotationDbi
## ## Attaching package: 'AnnotationDbi'
## ## The following object is masked from 'package:dplyr':
## ##     select
## ## loading existing transcript ranges created: 2019-11-22 01:06:45
## ## fetching genome info for GENCODE
## se
## ## class: RangedSummarizedExperiment
## ## dim: 205870 24
## ## metadata(6): tximetaInfo quantInfo ... txomeInfo txdbInfo
## ## assays(3): counts abundance length
## ## rownames(205870): ENST00000456328.2 ENST00000450305.2 ...
## ## ENST00000387460.2 ENST00000387461.2
## ## rowData names(3): tx_id gene_id tx_name
## ## colnames(24): SAMEA103885102 SAMEA103885347 ... SAMEA103885308
## ## SAMEA103884949
## ## colData names(4): names id line condition

On a machine with a working internet connection, the above command works without any extra steps, as the `tximeta` function obtains any necessary metadata via FTP, unless it is already cached locally. The `tximeta` package can also be used without an internet connection, in this case the linked transcriptome can be created directly from a `Salmon` index and gtf.

```r
makeLinkedTxome(
  indexDir=file.path(dir, "gencode.v29_salmon_0.12.0"),
  source="Gencode",
  organism="Homo sapiens",
  release="29",
  genome="GRCh38",
  fasta="ftp://ftp.ebi.ac.uk/pub/databases/gencode/Gencode_human/release_29/gencode.v29.transcripts.fa.gz",
  gtf=file.path(dir, "gencode.v29.annotation.gtf.gz"), # local version
  write=FALSE
)
Because tximeta knows the correct reference transcriptome, we can ask tximeta to summarize the transcript-level data to the gene level using the methods of Soneson et al. (2015).

gse <- summarizeToGene(se)
## loading existing TxDb created: 2019-11-22 01:02:58
## obtaining transcript-to-gene mapping from TxDb
## loading existing gene ranges created: 2019-11-23 02:30:13
## summarizing abundance
## summarizing counts
## summarizing length

One final note is that the start of positive strand genes and the end of negative strand genes is now dictated by the genomic extent of the isoforms of the gene (so the start and end of the reduced GRanges). Another alternative would be to either operate on transcript abundance, and perform differential analysis on transcript (and so avoid defining the TSS of a set of isoforms), or to use gene-level summarized expression but to pick the most representative TSS based on isoform expression.

**Importing ATAC-seq data as a SummarizedExperiment object**

The SummarizedExperiment object containing ATAC-seq peaks can be created from the following tab-delimited files from Alasoo & Gaffney (2017):

- The sample metadata: ATAC_sample_metadata.txt.gz (<1M)
- The matrix of normalized read counts: ATAC_cqn_matrix.txt.gz (109M)
- The annotated peaks: ATAC_peak_metadata.txt.gz (5.6M)

To begin, we read in the sample metadata, following similar steps to those we used to generate the coldata table for the RNA-seq experiment:

```r
atac_coldata <- read_tsv("ATAC_sample_metadata.txt.gz") %>%
  select(sample_id, donor, condition = condition_name) %>%
  mutate(condition = relevel(factor(condition), "naive"))
```

The ATAC-seq counts have already been normalized with cqn (Hansen et al., 2012) and log2 transformed. Loading the cqn-normalized matrix of log2 transformed read counts takes ~30 seconds and loads an object of ~370 Mb. We set the column names so that the first column contains the rownames of the matrix, and the remaining columns are the sample identities from the atac_coldata object.

```r
atac_mat <- read_tsv("ATAC_cqn_matrix.txt.gz",
  skip = 1,
  col_names = c("rownames", atac_coldata["sample_id"])))
rownames <- atac_mat[["rownames"]]
atac_mat <- as.matrix(atac_mat[, -1])
rownames(atac_mat) <- rownames
```

We read in the peak metadata (locations in the genome), and convert it to a GRanges object. The as_granges() function automatically converts the data.frame into a GRanges object. From that result, we extract the peak_id column and set the genome information to the build “GRCh38”. We know this from the Zenodo entry.
library(plyranges)

peaks_df <- read_tsv("ATAC_peak_metadata.txt.gz",
                      col_types = c("cidciicdc")
)

peaks_gr <- peaks_df %>%
          as_granges(seqnames = chr) %>%
          select(peak_id=gene_id) %>%
          set_genome_info(genome = "GRCh38")

Finally, we construct a SummarizedExperiment object. We place the matrix into the assays slot as a named list, the annotated peaks into the row-wise ranges slot, and the sample metadata into the column-wise data slot:

atac <- SummarizedExperiment(assays = list(cqndata=atac_mat),
                              rowRanges=peaks_gr,
                              colData=atac_coldata)

**Model assays**

**RNA-seq differential gene expression analysis**

We can easily run a differential expression analysis with DESeq2 using the following code chunks (Love et al., 2014). The design formula indicates that we want to control for the donor baselines (line) and test for differences in gene expression on the condition. For a more comprehensive discussion of DE workflows in Bioconductor see Love et al. (2016) and Law et al. (2018).

library(DESeq2)

dds <- DESeqDataSet(gse, ~line + condition)

## using counts and average transcript lengths from tximeta

# filter out lowly expressed genes
# at least 10 counts in at least 6 samples

keep <- rowSums(counts(dds) >= 10) >= 6

dds <- dds[keep,]

The model is fit with the following line of code:

dds <- DESeq (dds)

## estimating size factors
## using 'avgTxLength' from assays(dds), correcting for library size
## estimating dispersions
## gene-wise dispersion estimates
## mean-dispersion relationship
## final dispersion estimates
## fitting model and testing

Below we set the contrast on the condition variable, indicating we are estimating the log₂ fold change (LFC) of IFNg stimulated cell lines against naive cell lines. We are interested in LFCs greater than 1 at a nominal false discovery rate (FDR) of 1%.

res <- results(dds, 
               contrast=c("condition","IFNg","naive"),
               lfcThreshold=1, alpha=0.01)
To see the results of the expression analysis, we can generate a summary table and an MA plot (Figure 2):

```r
summary(res)
## out of 17806 with nonzero total read count
## adjusted p-value < 0.01
## LFC > 1.00 (up) : 502, 2.8%
## LFC < -1.00 (down): 247, 1.4%
## outliers [1] : 0, 0%
## low counts [2] : 0, 0%
## (mean count < 3)
## [1] see 'cooksCutoff' argument of ?results
## [2] see 'independentFiltering' argument of ?results
DESeq2::plotMA(res, ylim=c(-10,10))
```

![Figure 2. Visualization of DESeq2 results as an “MA plot”. Genes that have an adjusted p-value below 0.01 are colored red.](image)

We now output the results as a GRanges object, and due to the conventions of plyranges, we construct a new column called gene_id from the row names of the results. Each row now contains the genomic region (seqnames, start, end, strand) along with corresponding metadata columns (the gene_id and the results of the test). Note that tximeta has correctly identified the reference genome as “hg38”, and this has also been added to the GRanges along the results columns. This kind of book-keeping is vital once overlap operations are performed to ensure that plyranges is not comparing across incompatible genomes.

```r
suppressPackageStartupMessages(library(plyranges))
de_genes <- results(dds,
  contrast=c("condition","IFNg","naive"),
  lfcThreshold=1,
  format="GRanges") %>%
  names_to_column("gene_id")
de_genes
## GRanges object with 17806 ranges and 7 metadata columns:
## seqnames              ranges strand | gene_id
## <Rle>           <IRanges>  <Rle> |        <character>
## [1] chrX 100627109-100639991      - | ENSG00000000003.14
```
##       | chr20   50934867-50958555      - | ENSG00000000419.12
##       | chr1   169849631-169894267      - | ENSG00000000457.13
##       | chr1   169662007-169854080      + | ENSG00000000460.16
##       | chr1   27612064-27635277      - | ENSG00000000493.12
##       |  ...    ...                 ...    ... .                ...
##   | [17802]    chr10   84167228-84172093      - |  ENSG00000285972.1
##   | [17803]     chr6    63572012-63583587      + |  ENSG00000285976.1
##   | [17804]    chr16    57177349-57181390      + |  ENSG00000285979.1
##   | [17805]     chr8    103398658-103501895      - |  ENSG00000285982.1
##   | [17806]    chr10    12563151-12567351      + |  ENSG00000285994.1

### seqinfo: 25 sequences (1 circular) from hg38 genome

From this, we can restrict the results to those that meet our FDR threshold and select (and rename) the metadata columns we’re interested in:

```r
de_genes <- de_genes %>%
  filter(padj < 0.01) %>%
  select(gene_id, de_log2FC = log2FoldChange, de_padj = padj)
```

We now wish to extract genes for which there is evidence that the LFC is not large. We perform this test by specifying an LFC threshold and an alternative hypothesis (altHypothesis) that the LFC is less than the threshold in absolute value. To visualize the result of this test, you can run `results` without `format="GRanges"`, and pass this object to `plotMA` as before.

We label these genes as `other_genes` and later as “non-DE genes”, for comparison with our `de_genes` set.

```r
other_genes <- results(dds, 
  contrast=c("condition","IFNg","naive"),
  lfcThreshold=1,
  altHypothesis="lessAbs",
  format="GRanges")
```
filter(padj < 0.01) %>%
names_to_column(c("gene_id")) %>%
dplyr::select(gene_id,  
de_log2FC = log2FoldChange,  
de_padj = padj)

**ATAC-seq peak differential abundance analysis**

The following section describes the process we have used for generating a *GRanges* object of differential peaks from the ATAC-seq data in Alasoo et al. (2018).

The code chunks for the remainder of this section are not run.

For assessing differential accessibility, we run *limma* (Smyth, 2004), and generate the a summary of LFCs and adjusted p-values for the peaks:

```r
library(limma)

design <- model.matrix(~donor + condition, colData(atac))
fit <- lmFit(assay(atac), design)
fit <- eBayes(fit)
idx <- which(colnames(fit$coefficients) == "conditionIFNg")

tt <- topTable(fit, coef=idx, sort.by="none", n=nrow(atac))
```

We now take the *rowRanges* of the *SummarizedExperiment* and attach the LFCs and adjusted p-values from *limma*, so that we can consider the overlap with differential expression. Note that we set the genome build to “hg38” and restyle the chromosome information to use the “UCSC” style (e.g. “chr1”, “chr2”, etc.). Again, we know the genome build from the Zenodo entry for the ATAC-seq data.

```r
atac_peaks <- rowRanges(atac) %>%
  remove_names() %>%
  mutate(  
    da_log2FC = tt$logFC,  
    da_padj = tt$adj.P.Val
  ) %>%
  set_genome_info(genome = "hg38")

seqlevelsStyle(atac_peaks) <- "UCSC"
```

The final *GRanges* object containing the DA peaks is included in the workflow package and can be loaded as follows:

```r
library(fluentGenomics)

peaks
## GRanges object with 296220 ranges and 3 metadata columns:
## seqnames ranges strand | peak_id
## <Rle> <IRanges> <Rle> | <character>
## [1] chr1 9979-10668 * | ATAC_peak_1
## [2] chr1 10939-11473 * | ATAC_peak_2
## [3] chr1 15505-15729 * | ATAC_peak_3
## [4] chr1 21148-21481 * | ATAC_peak_4
## [5] chr1 21864-22067 * | ATAC_peak_5
## ... ... ... ... | ...
## [296216] chrX 155896572-155896835 * | ATAC_peak_296216
## [296217] chrX 155958507-155958646 * | ATAC_peak_296217
## [296218] chrX 156016760-156016975 * | ATAC_peak_296218
## [296219] chrX 156028551-156029422 * | ATAC_peak_296219
## [296220] chrX 156030135-156030785 * | ATAC_peak_296220
## da_log2FC da_padj
## <numeric> <numeric>
## [1] 0.266185396736073 9.10672732956434e-05
## Integrate ranges

### Finding overlaps with plyranges

We have already used plyranges a number of times above, to filter, mutate, and select on GRanges objects, as well as ensuring the correct genome annotation and style has been used. The plyranges package provides a grammar for performing transformations of genomic data (Lee et al., 2019). Computations resulting from compositions of plyranges “verbs” are performed using underlying, highly optimized range operations in the GenomicRanges package (Lawrence et al., 2013).

For the overlap analysis, we filter the annotated peaks to have a nominal FDR bound of 1%.

```r
da_peaks <- peaks %>%
  filter(da_padj < 0.01)
```

We now have GRanges objects that contain DE genes, genes without strong signal of DE, and DA peaks. We are ready to answer the question: is there an enrichment of DA ATAC-seq peaks in the vicinity of DE genes compared to genes without sufficient DE signal?

### Down sampling non-differentially expressed genes

As plyranges is built on top of dplyr, it implements methods for many of its verbs for GRanges objects. Here we can use slice to randomly sample the rows of the other_genes. The sample.int function will generate random samples of size equal to the number of DE-genes from the number of rows in other_genes:

```r
size <- length(de_genes)
slice(other_genes, sample.int(n(), size))
```

---

```r
# seqinfo: 23 sequences from hg38 genome; no seqlengths
```

### Example Data

```r
##        [2]  0.32217712436691 2.03434717570469e-05
##        [3] -0.574159538548115 3.41707743345703e-08
##        [4]  -1.14706617895329 8.22298606986521e-26
##        [5] -0.896143162633654 4.79452571676397e-11
##        ...
##   [296216] -0.834628897017445  1.3354605397165e-11
##   [296217] -0.147537281935847    0.313014754316915
##   [296218] -0.609732301631964 3.62338775135558e-09
##   [296219] -0.347678474957794 6.94823191242968e-06
##   [296220]  0.492442459200901 7.07663984067763e-13
##   -------
```
seqinfo: 25 sequences (1 circular) from hg38 genome

We can repeat this many times to create many samples via replicate. By replicating the sub-sampling multiple times, we minimize the variance on the enrichment statistics induced by the sampling process.

```r
# set a seed for the results
set.seed(2019-08-02)
boot_genes <- replicate(10,
  slice(other_genes, sample.int(n(), size)),
  simplify = FALSE)
```

This creates a list of GRanges objects as a list, and we can bind these together using the bind_ranges function. This function creates a new column called “resample” on the result that identifies each of the input GRanges objects:

```r
boot_genes <- bind_ranges(boot_genes, .id = "resample")
```

Similarly, we can then combine the boot_genes GRanges, with the DE GRanges object. As the resample column was not present on the DE GRanges object, this is given a missing value which we recode to a 0 using mutate()

```r
all_genes <- bind_ranges(
  de=de_genes,
  not_de = boot_genes,
  .id="origin"
)%>%
mutate(
  origin = factor(origin, c("not_de", "de")),
  resample = ifelse(is.na(resample), 0L, as.integer(resample))
)
```

```r
all_genes
## GRanges object with 8239 ranges and 5 metadata columns:
##          seqnames              ranges strand |            gene_id
##             <Rle>           <IRanges>  <Rle> |        <character>
##      [1]     chr1 196651878-196747504      + | ENSG00000000971.15
##      [2]     chr6   46129993-46146699      + | ENSG00000001561.6
##      [3]     chr4   17577192-17607972      + | ENSG00000002549.12
##      [4]     chr7 150800403-150805120      + | ENSG00000002933.8
##      [5]     chr4   15778275-15853230      + | ENSG00000004468.12
##      ...      ...                 ...    ... .                ...
##   [8235]    chr17   43527844-43579620      - | ENSG00000175832.12
##   [8236]    chr17   18260534-18266552      + | ENSG00000177427.12
##   [8237]    chr20   63895182-63936031      + | ENSG00000101152.10
##   [8238]    chr1 39081316-39487177      + | ENSG00000127603.25
##   [8239]    chr8  41577187-41625001      + | ENSG00000158669.11
##                   de_log2FC              de_padj  resample   origin
##                   <numeric>            <numeric> <integer> <factor>
##      [1]   4.98711071930695 1.37057050625117e-13         0       de
##      [2]   1.92721595378787  3.1747750217733e-05         0       de
##      [3]   2.93372501059128  2.0131038573066e-11         0       de
##      [4]   3.16721751137972 1.07359906028984e-08         0       de
##      [5]   5.40894352968188  4.2904694023763e-18         0       de
##      ...                ...                  ...       ...
##   [8235] -0.240918426099239  0.00991611085813261        10   not_de
```
Expanding genomic coordinates around the transcription start site

Now we would like to modify our gene ranges so they contain the 10 kilobases on either side of their transcription start site (TSS). There are many ways one could do this, but we prefer an approach via the anchoring methods in plyranges. Because there is a mutual dependence between the start, end, width, and strand of a GRanges object, we define anchors to fix one of start and end, while modifying the width. As an example, to extract just the TSS, we can anchor by the 5’ end of the range and modify the width of the range to equal 1.

```r
all_genes <- all_genes %>%
  anchor_5p() %>%
  mutate(width = 1)
```

Anchoring by the 5’ end of a range will fix the end of negatively stranded ranges, and fix the start of positively stranded ranges.

We can then repeat the same pattern but this time using anchor_center() to tell plyranges that we are making the TSS the midpoint of a range that has total width of 20 kb, or 10 kb both upstream and downstream of the TSS.

```r
all_genes <- all_genes %>%
  anchor_center() %>%
  mutate(width = 2 * 1e4)
```

Use overlap joins to find relative enrichment

We are now ready to compute overlaps between RNA-seq genes (our DE set and bootstrap sets) and the ATAC-seq peaks. In plyranges, overlaps are defined as joins between two GRanges objects: a left and a right GRanges object. In an overlap join, a match is any range on the left GRanges that is overlapped by the right GRanges. One powerful aspect of the overlap joins is that the result maintains all (metadata) columns from each of the left and right ranges which makes downstream summaries easy to compute.

To combine the DE genes with the DA peaks, we perform a left overlap join. This returns to us the `all_genes` ranges (potentially with duplication), but with the metadata columns from those overlapping DA peaks. For any gene that has no overlaps, the DA peak columns will have NA’s.

```r
genes_olap_peaks <- all_genes %>%
  join_overlap_left(da_peaks)
```

```r
genes_olap_peaks
#> GRanges object with 27766 ranges and 8 metadata columns:
#>  seqnames  ranges strand |    gene_id      
#>     <Rle> <IRanges>  <Rle> |      <character>  
#> [1] chr1 196641878-196661877 + |  ENSG0000000971.15  
#> [2] chr6 46119993-46139992  + |  ENSG00000001561.6   
#> [3] chr4 17567192-17587191  + |  ENSG00000002549.12  
#> [4] chr4 17567192-17587191  + |  ENSG00000002549.12  
#> [5] chr4 17567192-17587191  + |  ENSG00000002549.12  
#> ... ... ... ... ... ... ...  
#> [27762] chr1 39071316-39091315 + |  ENSG00000127603.25  
#> [27763] chr1 39071316-39091315 + |  ENSG00000127603.25  
#> [27764] chr8 41567187-41587186  + |  ENSG00000158669.11  
#> [27765] chr8 41567187-41587186  + |  ENSG00000158669.11  
#> [27766] chr8 41567187-41587186  + |  ENSG00000158669.11
```
Now we can ask, how many DA peaks are near DE genes relative to “other” non-DE genes? A gene may appear more than once in `genes_olap_peaks`, because multiple peaks may overlap a single gene, or because we have re-sampled the same gene more than once, or a combination of these two cases.

For each gene (that is the combination of chromosome, the start, end, and strand), and the “origin” (DE vs not-DE) we can compute the distinct number of peaks for each gene and the maximum peak based on LFC. This is achieved via `reduce_ranges_directed`, which allows an aggregation to result in a `GRanges` object via merging neighboring genomic regions. The use of the directed suffix indicates we’re maintaining strand information. In this case, we are simply merging ranges (genes) via the groups we mentioned above. We also have to account for the number of resamples we have performed when counting if there are any peaks, to ensure we do not double count the same peak:

```r
# seqinfo: 25 sequences (1 circular) from hg38 genome
```

```r
gene_peak_max_lfc <- genes_olap_peaks %>%
group_by(gene_id, origin) %>%
reduce_ranges_directed(
  peak_count = sum(!is.na(da_padj)) / n_distinct(resample),
  peak_max_lfc = max(abs(da_log2FC))
)
```

We can then filter genes if they have any peaks and compare the peak fold changes between non-DE and DE genes using a boxplot (Figure 3):

```r
library(ggplot2)
gene_peak_max_lfc %>%
  filter(peak_count > 0) %>%
as.data.frame() %>%
ggplot(aes(origin, peak_max_lfc)) +
geom_boxplot()
```
In general, the DE genes have larger maximum DA fold changes relative to the non-DE genes.

Next we examine how thresholds on the DA LFC modify the enrichment we observe of DA peaks near DE or non-DE genes. First, we want to know how the number of peaks within DE genes and non-DE genes change as we change threshold values on the peak LFC. As an example, we could compute this by arbitrarily chosen LFC thresholds of 1 or 2 as follows:

```r
origin_peak_lfc <- genes_olap_peaks %>%
group_by(origin) %>%
summarize(
  peak_count = sum(!is.na(da_padj)) / n_distinct(resample),
  lfc1_peak_count = sum(abs(da_log2FC) > 1, na.rm=TRUE) / n_distinct(resample),
  lfc2_peak_count = sum(abs(da_log2FC) > 2, na.rm=TRUE) / n_distinct(resample)
)
```

Here we see that DE genes tend to have more DA peaks near them, and that the number of DA peaks decreases as we increase the DA LFC threshold (as expected). We now show how to compute the ratio of peak counts from DE compared to non-DE genes, so we can see how this ratio changes for various DA LFC thresholds.

![Boxplot of maximum LFCs for DA peaks for DE genes compared to non-DE genes where genes have at least one DA peak.](image)

**Figure 3.** A boxplot of maximum LFCs for DA peaks for DE genes compared to non-DE genes where genes have at least one DA peak.

In general, the DE genes have larger maximum DA fold changes relative to the non-DE genes.
For all variables except for the origin column we divide the first row’s values by the second row, which will be the enrichment of peaks in DE genes compared to other genes. This requires us to reshape the summary table from long form back to wide form using the `tidyr` package. First, we pivot the results of the peak_count columns into name-value pairs, then pivot again to place values into the origin column. Then we create a new column with the relative enrichment:

```r
origin_peak_lfc %>%
as.data.frame() %>%
tidyr::pivot_longer(cols = -origin) %>%
tidyr::pivot_wider(names_from = origin, values_from = value) %>%
mutate(enrichment = de / not_de)
```

## A tibble: 3 x 4

| name      | not_de | de   | enrichment |
|-----------|--------|------|------------|
| peak_count| 2392.  | 3416 | 1.43       |
| lfc1_peak_count | 370. | 1097 | 2.97       |
| lfc2_peak_count  | 32.5 | 234  | 7.2        |

The above table shows that relative enrichment increases for a larger LFC threshold.

Due to the one-to-many mappings of genes to peaks, it is unknown if we have the same number of DE genes participating or less, as we increase the threshold on the DA LFC. We can examine the number of genes with overlapping DA peaks at various thresholds by grouping and aggregating twice. First, the number of peaks that meet the thresholds are computed within each gene, origin, and resample group. Second, within the origin column, we compute the total number of peaks that meet the DA LFC threshold and the number of genes that have more than zero peaks (again averaging over the number of resamples).

```r
genes_olap_peaks %>%
  group_by(gene_id, origin, resample) %>%
  reduce_ranges_directed(
    lfc1 = sum(abs(da_log2FC) > 1, na.rm=TRUE),
    lfc2 = sum(abs(da_log2FC) > 2, na.rm=TRUE)
  ) %>%
  group_by(origin) %>%
  summarize(
    lfc1_gene_count = sum(lfc1 > 0) / n_distinct(resample),
    lfc1_peak_count = sum(lfc1) / n_distinct(resample),
    lfc2_gene_count = sum(lfc2 > 0) / n_distinct(resample),
    lfc2_peak_count = sum(lfc2) / n_distinct(resample)
  )
```

## DataFrame with 2 rows and 5 columns

| origin | lfc1_gene_count | lfc1_peak_count | lfc2_gene_count | lfc2_peak_count |
|--------|-----------------|-----------------|-----------------|-----------------|
| not_de | 271.2           | 369.5           | 30.3            | 32.5            |
| de     | 515             | 1097            | 151             | 234             |

To do this for many thresholds is cumbersome and would create a lot of duplicate code. Instead we create a single function called `count_if_above_threshold` that accepts a variable and a vector of thresholds, and computes the sum of the absolute value of the variable for each element in the thresholds vector.

```r
count_if_above_threshold <- function(var, thresholds) {
  lapply(thresholds, function(.) sum(abs(var) > ., na.rm = TRUE))
}
```
The above function will compute the counts for any arbitrary threshold, so we can apply it over possible LFC thresholds of interest. We choose a grid of one hundred thresholds based on the range of absolute LFC values in the da_peaks GRanges object:

```r
thresholds <- da_peaks %>%
  mutate(abs_lfc = abs(da_log2FC)) %>%
  with(
    seq(min(abs_lfc), max(abs_lfc), length.out = 100)
  )
```

The peak counts for each threshold are computed as a new list-column called value. First, the GRanges object has been grouped by the gene, origin, and the number of resamples columns. Then we aggregate over those columns, so each row will contain the peak counts for all of the thresholds for a gene, origin, and resample. We also maintain another list-column that contains the threshold values.

```r
genes_peak_all_thresholds <- genes_olap_peaks %>%
  group_by(gene_id, origin, resample) %>%
  reduce_ranges_directed(
    value = count_if_above_threshold(da_log2FC, thresholds),
    threshold = list(thresholds)
  )
```

```
# GRanges object with 8239 ranges and 5 metadata columns:
## segnames ranges strand | gene_id origin
## <Rle> <IRanges>  <Rle> | <character> <factor>
## [1] chr1 196641878-196661877 + | ENSG00000000971.15 de
## [2] chr6 46119993-46139992 + | ENSG000000001561.6 de
## [3] chr4 17567192-17587191 + | ENSG000000002549.12 de
## [4] chr7 150790403-150810402 + | ENSG000000002933.8 de
## [5] chr4 15768275-15788274 + | ENSG000000004468.12 de
## ... ... ... ... ...
## [8235] chr17 43569620-43589619 - | ENSG00000175832.12 not_de
## [8236] chr17 18250534-18270533 + | ENSG00000177427.12 not_de
## [8237] chr20 63885182-63905181 + | ENSG00000101152.10 not_de
## [8238] chr1 39071316-39091315 + | ENSG00000127603.25 not_de
## [8239] chr8 41567187-41587186 + | ENSG00000158669.11 not_de
## resample value
## <integer> <IntegerList>
## [1] 0 1,1,1,...
## [2] 0 1,1,1,...
## [3] 0 6,6,6,...
## [4] 0 4,4,4,...
## [5] 0 11,11,11,...
## ... ... ...
## [8235] 10 1,1,1,...
## [8236] 10 3,3,2,...
## [8237] 10 5,5,5,...
## [8238] 10 3,3,3,...
## [8239] 10 3,3,3,...
## threshold <NumericList>
## [1] 0.0658243106359027, 0.118483961449043, 0.171143612262182, ...
## [2] 0.0658243106359027, 0.118483961449043, 0.171143612262182, ...
## [3] 0.0658243106359027, 0.118483961449043, 0.171143612262182, ...
## [4] 0.0658243106359027, 0.118483961449043, 0.171143612262182, ...
## [5] 0.0658243106359027, 0.118483961449043, 0.171143612262182, ...
## ... ...
## [8235] 0.0658243106359027, 0.118483961449043, 0.171143612262182, ...
```
Now we can expand these list-columns into a long `GRanges` object using the `expand_ranges()` function. This function will unlist the `value` and `threshold` columns and lengthen the resulting `GRanges` object. To compute the peak and gene counts for each threshold, we apply the same summarization as before:

```r
origin_peak_all_thresholds <- genes_peak_all_thresholds %>%
  expand_ranges() %>%
  group_by(origin, threshold) %>%
  summarize(
    gene_count = sum(value > 0) / n_distinct(resample),
    peak_count = sum(value) / n_distinct(resample)
  )
```

```
origin_peak_all_thresholds
# DataFrame with 200 rows and 4 columns
#       origin          threshold gene_count  peak_count
#     <factor>          <numeric>  <numeric>  <numeric>
# 1     not_de 0.0658243106359027        708       2391.4
# 2     not_de 0.118483961449043        698       2320.6
# 3     not_de 0.171143612262182        686       2178.6
# 4     not_de 0.223803263075322        672       1989.4
# 5     not_de 0.276462913888462        650       1785.8
# ...     ...                ...        ...        ...
# 196     de   5.06849113788419         2         2
# 197     de   5.12115078869733         0         0
# 198     de   5.17381043951047         0         0
# 199     de   5.22647009032361         0         0
# 200     de   5.27912974113675         0         0
```

Again we can compute the relative enrichment in LFCs in the same manner as before, by pivoting the results to long form then back to wide form to compute the enrichment. We visualize the peak enrichment changes of DE genes relative to other genes as a line chart (**Figure 4**):

```
origin_threshold_counts <- origin_peak_all_thresholds %>%
  as.data.frame() %>%
  tidyr::pivot_longer(cols = -c(origin, threshold),
                     names_to = c("type", "var"),
                     names_sep = "_",
                     values_to = "count") %>%
  select(-var)

origin_threshold_counts %>%
  filter(type == "peak") %>%
  tidyr::pivot_wider(names_from = origin, values_from = count) %>%
  mutate(enrichment = de / not_de) %>%
  ggplot(aes(x = threshold, y = enrichment)) +
  geom_line() +
  labs(x = "logFC threshold", y = "Relative Enrichment")
```

## Warning: Removed 4 row(s) containing missing values (geom_path).
We computed the sum of DA peaks near the DE genes, for increasing LFC thresholds on the accessibility change. As we increased the threshold, the number of total peaks went down (likewise the mean number of DA peaks per gene). It is also likely the number of DE genes with a DA peak nearby with such a large change went down. We can investigate this with a plot (Figure 5) that summarizes many of the aspects underlying the enrichment plot above.

```r
origin_threshold_counts %>%
ggplot(aes(x = threshold,
y = count + 1,
    color = origin,
    linetype = type)) +
geom_line() +
scale_y_log10()
```

**Figure 4.** A line chart displaying how relative enrichment of DA peaks change between DE genes compared to non-DE genes as the absolute DA LFC threshold increases.

**Figure 5.** A line chart displaying how gene and peak counts change as the absolute DA LFC threshold increases. Lines are colored according to whether they represent a gene that is DE or not. Note the x-axis is on a log_{10} scale.
Discussion
We have shown that by using plyranges and tximeta (with support of Bioconductor and tidyverse ecosystems) we can fluently iterate through the biological data science workflow: from import, through to modeling, and data integration.

There are several further steps that would be interesting to perform in this analysis; for example, we could modify window size around the TSS to see how it affects enrichment, and vary the FDR cut-offs for both the DE gene and DA peak sets. We could also have computed variance in addition to the mean of the bootstrap set, and so drawn an interval around the enrichment line.

Finally, our workflow illustrates the benefits of using appropriate data abstractions provided by Bioconductor such as the SummarizedExperiment and GRanges. These abstractions provide users with a mental model of their experimental data and are the building blocks for constructing the modular and iterative analyses we have shown here. Consequently, we have been able to interoperate many decoupled R packages (from both Bioconductor and the tidyverse) to construct a seamless end-to-end workflow that is far too specialized for a single monolithic tool.

Data availability
All data underlying the results are available as part of the article and no additional source data are required.

Software availability
plyranges is available from Bioconductor: https://doi.org/doi:10.18129/B9.bioc.plyranges.

tximeta is available from Bioconductor: https://doi.org/doi:10.18129/B9.bioc.tximeta.

Source code and all workflow materials are available at: https://github.com/sa-lee/fluentGenomics.

Archived source code at time of publication: https://doi.org/10.5281/zenodo.3633505 (Lee & Love, 2020).

License: MIT License.

The development version of the workflow and all downstream dependencies can be installed using the BiocManager package by running:

```r
# development version from Github
BiocManager::install("sa-lee/fluentGenomics")
# version available from Bioconductor
BiocManager::install("fluentGenomics")
```

This article and the analyses were performed with R (R Core Team, 2019) using the rmarkdown (Allaire et al., 2019), and knitr (Xie, 2019; Xie, 2015) packages.

Session Info
```
sessionInfo()
```

```r
# R version 3.6.1 (2019-07-05)
# Platform: x86_64-apple-darwin15.6.0 (64-bit)
# Running under: macOS Mojave 10.14.6
# Matrix products: default
# BLAS: /System/Library/Frameworks/Accelerate.framework/Versions/A/Frameworks/vecLib.framework/Versions/A/libBLAS.
# LAPACK: /Library/Frameworks/R.framework/Versions/3.6/Resources/lib/libRlapack.dylib
# locale:
# [1] en_AU.UTF-8/en_AU.UTF-8/en_AU.UTF-8/C/en_AU.UTF-8/en_AU.UTF-8
# attached base packages:
# [1] parallel stats4 stats graphics grDevices utils datasets
# [8] methods base
```
Author contributions
All authors contributed to the writing and development of the workflow.

Acknowledgements
The authors would like to thank all participants of the Bioconductor 2019 and BiocAsia 2019 conferences who attended and provided feedback on early versions of this workflow paper.
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Current Peer Review Status: ✔️ ⬜️ ⬜️

Version 1

Reviewer Report 21 May 2020

https://doi.org/10.5256/f1000research.24553.r59959

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The authors describe a workflow to perform exploratory analysis of biological data. This paper also highlights the importance of using tools that facilitate the application of essential concepts of exploratory data analysis: reproducibility and code readability.

Major comments:

1. I was not able to successfully complete the protocol myself, failing at the second step. I tried three different environments:

   - R 3.4.4 on Ubuntu 18.04.4 LTS/Windows Subsystem for Linux WSL1/Windows 10 1909 18363.836
     (BiocManager refuses to install fluentGenomics on the system R for what is currently the latest version of Ubuntu available for general availability Windows users).
   - R 4.0.0 x86 on Windows 10 1909 18363.836
     - BiocManager::install("fluentGenomics")
     - BiocManager::install("sa-lee/fluentGenomics")

R version 4.0.0 (2020-04-24) -- "Arbor Day"
Copyright (C) 2020 The R Foundation for Statistical Computing
Platform: i386-w64-mingw32/i386 (32-bit)

R is free software and comes with ABSOLUTELY NO WARRANTY.
You are welcome to redistribute it under certain conditions.
Type 'license()' or 'licence()' for distribution details.
Natural language support but running in an English locale

R is a collaborative project with many contributors. Type 'contributors()' for more information and 'citation()' on how to cite R or R packages in publications.

Type 'demo()' for some demos, 'help()' for on-line help, or 'help.start()' for an HTML browser interface to help. Type 'q()' to quit R.

> library(fluentGenomics)
> path_to_se <- cache_atac_se()
Error in retrieve_cache() :
  Please install rappdirs to set cache directory
> detach("package:fluentGenomics", unload=TRUE)
> BiocManager::install("sa-lee/fluentGenomics")
Bioconductor version 3.11 (BiocManager 1.30.10), R 4.0.0 (2020-04-24)
Installing github package(s) 'sa-lee/fluentGenomics'
Error: package 'remotes' not installed in library path(s)
  C:/Users/mhoffman/R/win-library/4.0
  C:/Program Files/R/R-4.0.0/library
install with 'install("remotes")'
> install.packages("remotes")
[...]
> BiocManager::install("sa-lee/fluentGenomics")
> path_to_se <- cache_atac_se()
Error in retrieve_cache() :
  Please install rappdirs to set cache directory
> sessionInfo()
R version 4.0.0 (2020-04-24)
Platform: i386-w64-mingw32/i386 (32-bit)
Running under: Windows 10 x64 (build 18363)

Matrix products: default
locale:
[1] LC_COLLATE=English_United States.1252  LC_CTYPE=English_United States.1252
[4] LC_MONETARY=English_United States.1252
[7] LC_NUMERIC=C                         LC_TIME=English_United States.1252

attached base packages:
[1] stats  graphics  grDevices  utils  datasets  methods  base

other attached packages:
[1] fluentGenomics_0.99.3

loaded via a namespace (and not attached):
Minor comments:

2. It is unclear who the audience for this paper is. As written it seems to presume a lot of technical knowledge about Bioconductor, R, and specific Bioconductor packages. It would not make a good starting point for someone who doesn’t have large familiarity with all of the above. This is too bad because it has the bones of something that could be useful to people with less expertise if some additional explanatory text were provided and clearer choices made about naming variables. There is also a danger that this will be used as a cookbook without understanding.

I have marked some examples of things that could use more explanation but this is non-exhaustive. It would be worthwhile for the authors to examine the manuscript again and ask whether all of the large number of concepts, jargon, and names introduced are explained.

3. The goal of the paper appeared to be clear after reading the paper. The introduction, figure 1, and following sections, however, leave the impression that the authors tried to reproduce results from another publication which is not the case. The paper would benefit from clarifying this toward the end of the introduction.

4. Abstract: it is not clear which problem does the workflow solve and why is it important to solve it. For example, how does the import, model, and integrate structure increases result reproducibility? Is this structure new? Why would someone use it? We think that the abstract would benefit from more justification and moving the text specific to the tools used (tximeta, SummarizedExperiment) to the introduction.

5. Abstract: The sentence “Using tximeta, ...“ and the following can be confusing. Why is it only using of RNA-seq in the first sentence, and both RNA-seq and ATAC-seq in the following one?

6. p3: Figure 1: The meaning of some of the objects is not immediately apparent. For example: coldata, se, dds, boot_genes. This figure does not make a great introduction for someone who is not already familiar with this workflow because so many of the details are mystifying. You should either simplify or explain more.

7. p3: Figure 1: Text in boxes is blurry.
8. p3: meaning of “fluent” should be described briefly here and not assumed

9. Introduction: This section would benefit from a more thorough description of the tools used in the workflow and explain why they are important. For example: What is Bioconductor? Why is plyranges relevant for this analysis? What are tidyverse design principles and why are they relevant here?

10. Introduction: Authors should also explain in the introduction why the “import, model, integrate” structure benefits users.

11. The authors should consistently use the same tense (we examine ... we will ...).

12. Why ignoring genotypes?

13. It would be useful to note that "genomic ranges" are often referred to as intervals elsewhere such as in BEDTools.

14. Please cite Tximeta at the first occurrence.

15. Experimental data: Here and in the following sections, the authors describe alternatively what the workflow does and what the workflow can do. For example: “The following code loads the path to the cached data file, or if it is not present, will create the cache and generate a SummarizedExperiment using”. We think that it would be clearer if these sections with code examples focus on what exactly the code does, like in a protocol paper. Describing what the tools can do should appear either in the introduction or in a separate paragraph. For the example above, the paragraph would explain how users would benefit from this behavior: save time?

16. “A subset of the RNA-seq ...”, Why use only a subset?

17. Typo: “be also be downloaded”

18. p4: “GENCODE” not “Gencode”

19. p4: Specify whether basic or comprehensive gene set used

20. p4: Unclear what “Importing ATAC-seq data as a SummarizedExperiment object” refers to

21. Import data as SummarizedExperiment: Authors should briefly describe what methods Soneson et al. use to summarize transcripts at gene level, and justify why they are doing it.

22. p4: Unclear where the `coldata.csv` file is or what that name refers to. “file” should not be in monospace text

23. p4: Placement of “Wickham et al., 2019” makes it appear to be a citation for R rather than tidyverse.

24. p5: Many things here unclear to the uninitiated: what is the scary message when you load
dplyr? What is a factor and relevel? What is a tibble? What do the ~ mean?

25. p5: Meaning of “but would typically not have the ‘gz’ ending if used from Salmon directly” unclear

26. p5: Should note the “files” column is going to be different for different users

27. p6: What is suppressPackageStartupMessages?

28. p6: What is a Salmon gtf?

29. p6: Lots of unexplained details in the two transcripts on this page.

30. p7: “tximeta knows the correct reference transcription”. How?

31. p7: GRanges not described or cited

32. p7: Unclear why only the start of + genes and not also their end is affected

33. p7: The sentence starting “Another alternative would be“ is complex and its meaning unclear (alternative to what?)

34. Last paragraph of "Import data as SummarizedExperiment": authors describe alternative ways of summarizing genes and using their representative TSS. Is the point to explain that GRanges make these operations easier? We think a potential user would like to know how to perform precisely these operations. Otherwise move this to an earlier paragraph describing what the tool is capable of doing.

35. p7: What does “109M” refer to? Please use SI or IEC units here and elsewhere

36. p7: Should be “370 MB” or “370 MiB” not “~370 Mb”

37. p7: It's not clear how to actually get the tab-delimited files, a departure from what is otherwise detailed step-by-step instructions

38. p7: Why skip=1? Why [-1]?

39. p7: It would be better if instead of just saying “we know this from the Zenodo entry” you instruct the workflow user to *check* the Zenodo entry to find out what genome assembly is used. A screenshot of that part of the Zenodo entry might be helpful.

40. p7: Citation for Genome Reference Consortium needed

41. p8: what is col_types? How do you know that this is what the col_types should be?

42. p8: This is the first instance of a convention of using comments in monospace text for explanatory notes. I would avoid this and stick to notes in proportional text.
43. p8: what is a “nominal” FDR?

44. p8: greater than 1 or greater-than/equal? Suggest using > or >= for greater clarity

45. Why 1%?

46. p9: “MA” needs expansion, citation, and ideally brief explanation

47. p9: Figure 2 caption has insufficient detail. Should describe what each point represents, and how many points there are. Should describe what method is used to adjust p-values. Please describe triangles versus circles. Explain the red horizontal line. Please indicate that “log fold change” means log2. Describe what “normalized counts are”

48. p9: “due to the convention of plyranges”—meaning unclear

49. p9: “This kind of book-keeping” = ?

50. p9: Not everyone reading this will work on human genomics and know that hg38 = GRCh38

51. p9: The authors should consistently put explanations of transcripts either below the transcript or above and not switch back and forth. (I think below is probably more helpful.)

52. p10: please make clear that the altHypothesis here gets ALL of the other genes, that there are no problems with the edge case at LFC = threshold

53. p11: Lots unexplained in the limma transcript

54. p12: What is a “seqlength”?

55. “genes without strong signal of DE”, are these “non-DE” genes? Would be better if used consistently.

56. p12: Down sampling -> “Downsampling”

57. p12: Unclear why one would want to downsample. The description only mentions that one can.

58. p13: Please explain why one would want to set a seed.

59. p13: “This creates a list of _GRanges_ objects as a list”. It would be clearer to say that this creates a _list_ of 10 _GRanges_ objects.

60. What are “verbs” for GRanges

61. p13: The importance of doing the bootstrap procedure here is really unclear

62. p14: Why is there a circular sequence? I assume this is the mitochondrial chromosome? Does its presence in the background disturb the analysis at all?
63. p14: "Now we would like to modify our gene ranges so they contain..." Why?

64. p14: should be 5 prime not 5 apostrophe

65. p15: the significance of the clarification that each “gene” is just a tuple of chromosome, start, end, strand is unclear

66. p15: A lot of na.rm or is.na from here outward. Would be worth explaining why this is.

67. p16: Figure 3: This figure needs descriptive labels instead of references to variables used in the code.

68. p19: Explain why there are missing values

69. p20: Should comment on why Figure 4 is non-monotonic

70. p20: Saying the x-axis is on a log10 scale is a bit confusing because the y-axis is also on a log10 scale. It is better to make log scales more obvious by having non-uniform tick marks and grid lines. Also I don't think the x-axis is on a log10 scale, log fold change here is measured with log2. (This would be more obvious if in the text you called it \log2 fold change consistently instead of the ambiguous “LFC”.)

71. p21: It is unfortunate that the main point of this paper is to show that you can do something “fluently”, while what that means is never discussed or justified in the paper.

72. p21: “There are several further steps that would be interesting to perform”, Are these steps easy to implement with the tools you described? If so, can you explain how easier it is with your approach?

73. p21: “Finally, our workflow illustrates the benefits of using appropriate data abstractions”, This is probably obvious for R power users but not for other readers. Can you point at a specific section that shows that a user will benefit from using your approach?

74. p21: The data availability statement is confusing. Clearly not all data are part of the article, they are part of external packages or articles (some specified without a particular URL or accession number, just reference to a paper).

75. p21: instructions for installing BiocManager would be helpful, as would listing the *minimum* version of R or any other relevant packages. Also should mention that you need to library(BiocManager) first

76. p21: It is unclear why you would want the development version, especially since it seems to be an older version number than what's on Bioconductor

77. p23: Items in the bibliography are not consistently referenced with specificity in a visible way. DOI should be provided for all Zenodo entries not just some. Some other documents that are not referenced via traditional publishing (journal volume and page number, or monograph from a publisher) should probably have DOI or URL specified also.
Discretionary recommendations:

78. p4: `path_to_se` is a poor name for a variable. It is not orthogonal to the other variable names used here and has little explanatory power. I recommend `atac_filename`.

79. p4: `dir` also is not descriptive. I recommend `quantdirname`.

80. p4: Would suggest more descriptive names than colfile/coldata, orthogonal to whatever naming scheme you use (which means either changes to names here or to use variations of the names I used above).

81. p5: This transcript is a lot to digest, and it's hard to know which parts of the explanatory text refer to which part of the transcript, which has multiple commands that go back a page previous. I suggest breaking it up more with explanatory text to the extent possible. Piping is convenient for a programmer but not for explanation. This comment applies to many of the subsequent transcripts too.

82. p5: Get the display of coldata so that it doesn't exceed one line per row.

83. p6: It's not a good idea to call your experiment object merely `se` when it is one of multiple SummarizedExperiments here.

84. p6: Please use a consistent style for spacing around `=` in R code.

85. p7: I suggest not giving the same name to different objects (`atac_mat` is first something like a data.frame and then a matrix.

86. p7: Suggest not shadowing built-in functions with variable names like `rownames`.

87. p7: “We know this” antecedent unclear.

88. p8: I suggest running all the R code in here through a linter. This will ensure you have consistent style.

89. p8: res is not a specific variable name.

90. p10: Why switch between “other genes” and “non DE genes”? Just call them `not_de_genes` from the start. (Or switch to `genes_de` and `genes_not_de`.)

91. p11: Why is library(fluentGenomics) loaded again?

92. p11: Using library(fluentGenomics) adds a data object called `peaks` to your environment? This seems undesirable.

93. p12: The text would be a bit more readable if you eschewed the abbreviations “DA”, “DE”, and “LFC” in the main text.
94. p13: here and elsewhere: integer literals (e.g. 10L) should be used instead of float literals for integers

95. p14: why `2*1e4`? 20000L would be clearer

96. p16: Figure 3: Please specify what the components of the box plot mean. There is no universal box plot.

Is the rationale for developing the new method (or application) clearly explained?
Partly

Is the description of the method technically sound?
Partly

Are sufficient details provided to allow replication of the method development and its use by others?
No

If any results are presented, are all the source data underlying the results available to ensure full reproducibility?
No

Are the conclusions about the method and its performance adequately supported by the findings presented in the article?
Partly

Competing Interests: No competing interests were disclosed.

Reviewer Expertise: computational genomics

We confirm that we have read this submission and believe that we have an appropriate level of expertise to confirm that it is of an acceptable scientific standard, however we have significant reservations, as outlined above.
Lee, Lawrence and Love present a workflow article about using a group of well-constructed Bioconductor packages for fluent genomic analysis that integrates both RNA-seq and ATAC-seq analysis results from the same samples (baseline and stimulus).

The workflow provides instructions for obtaining differentially accessible (DA; peaks from ATAC-seq) and differential expression (DE; from RNA-seq) data objects, and more importantly a convenient way of integrating the ranged genomic features produced, while taking care of matching reference genome builds and other crucial “bookkeeping” tasks in the process.

The authors have demonstrated analysis can be performed to check whether DA peaks are enriched in the vicinity of DE genes guided by the principle that transcriptomic response to IFNg stimulation may be mediated through binding of regulatory proteins to accessible regions, and these bindings can be detected by ATAC-seq if they increase the accessibility of these regions.

The following steps are involved in the workflow:

1. Downloading peaks data and RNA-seq data
2. Perform DE and DA analysis
3. Integrate ranges of DE and DA
4. Test enrichment of DA in DE gene vicinity versus non-DE genes with various thresholds

**Major comments:**

1. The process of ‘resampling’ subsets of non-DE genes from the pool of non-DE genes should be clarified to explain the purpose of doing such ‘resampling’. In particular, whether it can be considered ‘bootstrapping’ should be addressed. Our understanding is that the resampling here is done *without replacement* from DE results, whereas for bootstrapping we expect to sample data records *with replacement* and recompute statistics of interest from them.
2. Interpretation of the enrichment results or more discussion on the caveats of doing enrichment analysis like this are required.

**Minor comments:**

1. The workflow was built with R-3.6.1 which results in a discrepancy with the current package’s requirement of R >= 4.0.0. However, this is understandable since we are near the new release of Bioconductor but be reminded to re-compile the workflow with released version.
2. A code chunk for checking that all required packages have been installed prior to other analysis in the environment would make the execution smoother for a user wanting to run the workflow.
3. The downloading function `cache_atac_se()` raised an error when running on Windows. It can be fixed by setting mode = “wb” explicitly in the `download_file()`.
4. Figure 1 which summaries the workflow could be improved by incorporating colored blocks for each package module and more descriptive elements companying the actual function.
calls or data objects. The caption also needs to provide more complete description of the elements of the figure for it to be comprehensible. These changes will give readers who are not familiar with all these packages more information.

5. The optional code chunks (such as the Importing ATAC-seq data as a SummarizedExperiment object) could be moved down to appendix to make it clearer to the readers which things are core parts of the workflow and which are included for full transparency/reproducibility/data provenance.

6. As for calculating the relative enrichment of peaks in DE versus non-DE, the description "For all variables except for the origin column we divide the first row's values by the second row," was inaccurate and could corrected to "divide the DE row with the non-DE row"

7. It might worth emphasizing more on the contribution of this work in the abstract and introduction, maybe by stating explicitly the most attractive aspect of this work. For example, a guideline on integrating the ranged genomic features.

8. Make the sub-headings consistent. Importing ATAC-seq, Import data.

9. Some minor comments: library(fluentGenomics) appeared three times and library(plyranges) twice in the workflow. If this was to avoid namespace clashes, the function could be called explicitly plyranges:: instead of loading the packages multiple times?

10. You could just use limma for the DE analysis as well and save yourself the loading of the DESeq2 package (joking!)

Is the description of the method technically sound?
Partially. The “resampling” procedure needs a bit more explanation such as clarifying how and whether this is considered a bootstrapping method.

Is the rationale for developing the new method (or application) clearly explained?
Yes

Is the description of the method technically sound?
Partly

Are sufficient details provided to allow replication of the method development and its use by others?
Yes

If any results are presented, are all the source data underlying the results available to ensure full reproducibility?
Yes

Are the conclusions about the method and its performance adequately supported by the findings presented in the article?
Yes

Competing Interests: No competing interests were disclosed.

Reviewer Expertise: Statistics, bioinformatics, genomics, transcriptomics

We confirm that we have read this submission and believe that we have an appropriate level
of expertise to confirm that it is of an acceptable scientific standard, however we have significant reservations, as outlined above.

Mark Dunning
Sheffield Bioinformatics Core, University of Sheffield, Sheffield, UK

Lee et al. present a case study of using Bioconductor packages to recapitulate an existing analysis. The analysis is performed without the need to worry too much about genome versions and using tools from tidyverse which were previously incompatible with standard Bioconductor methodologies. I see this being a useful reference for experienced users on how to run a modern RNA-seq analysis and perform data integration.

I am pleased to say that I could re-run the code and reproduce the results presented. I can think of a few potential changes to increase its utility for beginner Bioconductor users.

The article assumes quite a lot of knowledge of tidyverse and GenomicRanges to understand the code chunks. Some pointers to where people can get more help on these packages would be useful. Some code chunks explicitly call methods from dplyr (dplyr::mutate), but not all. It would be good to maintain consistency when calling dplyr methods, as the user would know where to get help.

The authors go straight from creating the DESeq dataset to running the differential analysis. In reality, one would never do this without performing adequate QC checks. I'm sure the authors must have done this (as indeed the original study authors must have). But showing the code to generate a PCA would be useful; along with interpretation of the results.

The code is shown to remove genes with low counts prior to the DESeq analysis. I thought this was no longer necessary as DESeq2 incorporates it's own filtering. Or is this a memory-saving exercise? Some justification of why 10 counts and 6 samples was chosen would be beneficial to those new to Bioconductor and RNA-seq.

The authors use Genomic ranges to facilitate the integration of different assay types. Whilst it is critical to keep the data in this form for analysis, ultimately collaborators will want the results in spreadsheet format with recognisable gene names. It would be useful if the authors could include an example code snippet to annotate the overlapping regions (e.g. with gene symbols) and exporting as a csv.

It is my understanding that the authors present a simpler approach than that of Alasoo et al (2018) in order to illustrate the utility of Bioconductor infrastructure. Nevertheless, some
commentary on whether the same results and biological insights are revealed by their approach would be beneficial.

References
1. Alasoo K, Rodrigues J, Mukhopadhyay S, Knights AJ, et al.: Shared genetic effects on chromatin and gene expression indicate a role for enhancer priming in immune response. *Nat Genet.* **50** (3): 424-431 PubMed Abstract | Publisher Full Text

Is the rationale for developing the new method (or application) clearly explained?
Partly

Is the description of the method technically sound?
Yes

Are sufficient details provided to allow replication of the method development and its use by others?
Yes

If any results are presented, are all the source data underlying the results available to ensure full reproducibility?
Yes

Are the conclusions about the method and its performance adequately supported by the findings presented in the article?
Yes

*Competing Interests:* No competing interests were disclosed.

*Reviewer Expertise:* Bioinformatics, Bioconductor, Functional Genomics

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard.
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