Introduction to R and Bioconductor

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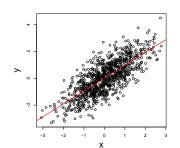
R: Statistical Computing Environment

- Vectors logical, integer, numeric, character, . . .
 - ▶ list() contains other vectors (recursive)
 - ► factor(), NA statistical concepts
 - ► Can be named c(Germany=1, Argentina=0)
- ▶ matrix(), array() a vector with a 'dim' attribute.
- ▶ data.frame() like spreadsheets; list of equal length vectors.
 - Homogenous types within a column, heterogenous types across columns.
 - An example of an R class.
- Other classes more complicated arrangement of vectors.
 - Examples: the value returned by lm(); the DNAStringSet class used to hold DNA sequences.
 - plain, 'accessor', 'generic', and 'method' functions
- Packages base, recommended, contributed.

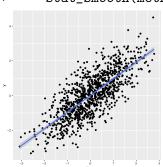
R: Statistical Computing Environment

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```
> plot(y ~ x, df, cex.lab=2)
> abline(fit, col="red",
+ lwd=2)
```



```
> library(ggplot2)
> ggplot(df, aes(x, y)) +
+         geom_point() +
+         stat_smooth(method="lm")
```



R: programming concepts

- Functions built-in (e.g., rnorm()); user-defined
- Subsetting logical, numeric, character; df [df\$x > 0,];
- lteration over vector elements, lapply(), mapply(),
 apply(), ...; e.g., lapply(df, mean)

R: help!

- ?data.frame
- methods(lm), methods(class=class(fit))
- ?"plot<tab>"
- help(package="Biostrings")
- vignette(package="GenomicRanges")
- StackOverflow; R-help mailing list

"Hey, can you help me with this? I tried..."

Bioconductor

Analysis & comprehension of high-throughput genomic data

- > 12 years old; 1024 packages; widely used
- Sequencing (RNAseq, ChIPseq, variants, copy number, ...),
 microarrays, flow cytometery, proteomics, ...
- http://bioconductor.org, https://support.bioconductor.org

Themes

- ► Interoperable classes to work with genome-scale data, shared (where possible!) across packages
- Usable package vignettes, man pages, examples, . . .
- Reproducible 'release' and 'devel' versions, updated every 6 months

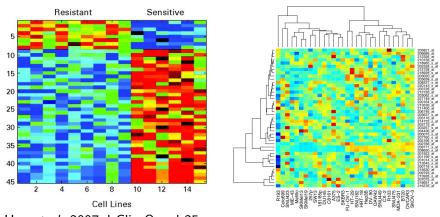
Bioconductor: GenomicRanges

```
GRanges
> gr = exons(TxDb.Hsapiens.UCSC.hg19.knownGene); gr
                                                                    length(gr); gr[1:5]
GRanges with 289969 ranges and 1 metadata column:
                                                                    segnames(gr)
          segnames
                                  ranges strand
                                                      exon_id
                                                                    start(gr)
              <Rle>
                               <IRanges> <Rle>
                                                  | <integer>
                                                                    end(gr)
       [1]
               chr1
                          [11874, 12227]
                                                                    width(gr)
       Γ27
               chr1
                         Γ12595, 127217
                                                                    strand(gr)
       [3]
              chr1
                         [12613, 12721]
                                                                  DataFrame
  [289967]
              chrY [59358329, 59359508]
                                                       277748
                                                                    mcols(gr)
  F2899687
              chrY [59360007, 59360115]
                                                       277749
                                                                    gr$exon id
  Γ2899697
               chrY [59360501, 59360854]
                                                       277750
                                                                  Seginfo
  seginfo: 93 seguences (1 circular) from hg19 genome
                                                                    seglevels(gr)
                                                                    seqlengths(gr)
                                                                    genome(gr)
```

- ▶ Data: aligned reads, called peaks, SNP locations, CNVs, . . .
- ▶ Annotation: gene models, variants, regulatory regions, . . .
- findOverlaps(), nearest(), and many other useful range-based operations.

Bioconductor: SummarizedExperiment motivation

Cisplatin-resistant non-small-cell lung cancer gene sets



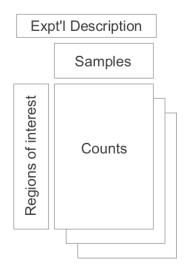
Hsu *et al.* 2007 J Clin Oncol 25: 4350-4357 (retracted)

Baggerly & Coombes 2009 Ann Appl Stat 3: 1309-1334

Coordinated, programmatic manipulation of feature, sample, and assay data



Bioconductor: SummarizedExperiment



Regions of interest \times samples

- assay() matrix, e.g., counts of reads overlapping regions of interest.
- rowData() regions of interest as GRanges or GRangesList
- colData() DataFrame describing samples.

> assay(se)[,se\$Treatment == "Control"] # Control counts

Bioconductor: a fun demo of GRanges interoperability

GenomicFeatures And 'annotation' packages to represent gene models as GRanges.

GenomicAlignments To input aligned reads as GRanges.

Gviz For visualization.

shiny For interactivity.

Bioconductor: Resources

http://bioconductor.org

- ▶ Packages biocViews, landing pages (e.g., AnnotationHub)
- Course & conference material; work flows; publications
- Developer resources

https://support.bioconductor.org

- Queston & answer forum for users; usually fast, expert, friendly responses
- Contributed tutorials, news

Citations

- ► Huber et al. (2015) Orchestrating high-throughput genomic analysis with *Bioconductor*. Nature Methods 12:115-121.
- Lawrence et al. (2013) Software for Computing and Annotating Genomic Ranges. PLoS Comput Biol 9(8): e1003118.

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https://bioconductor.org,
https://support.bioconductor.org
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