

Preparing Data from Nagalakshmi et al.

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This data set is based on Nagalakshmi et al. [1]. The data was retrieved from the [sequence read archive](#), aligned with [bwa](#), and converted to BAM format with [samtools](#).

Data were downloaded from the sequence read archive by visiting <http://trace.ncbi.nlm.nih.gov/Traces/sra/sra.cgi?study=SRP000227> and downloading reads for the entire study. This results in a directory structure

```
SRP000227
|---SRR002051
|   |-- SRR002051.sra
|---SRR002058
|   |-- SRR002058.sra
|---SRR002059
|   |-- SRR002059.sra
|---SRR002061
|   |-- SRR002061.sra
|--- SRR002062
|   |-- SRR002062.sra
|---SRR002064
|   |-- SRR002064.sra
```

Fastq files were extracted using [NCBI SRA SDK](#).

```
#!/usr/bin/env sh
SRA=/home/mtmorgan/bin/sratoolkit.2.0b4-3-suse_linux32
for f in `ls */*.sra`
do
    ${SRA}/sratoolkit.2.0b4-3-suse_linux32/fastq-dump $f
done
```

Reference sequences were retrieved from [UCSC](#) golden path, unzipped, and catenated into a single file. The file was indexed for use by the [bwa](#) aligner with `bwa index sacCer2.fa`. Reads were aligned and converted to BAM as

```

#!/usr/bin/env sh
BWA=/home/mtmorgan/bin/bwa/bwa
SAMTOOLS=/home/mtmorgan/bin/samtools/samtools
for f in `ls ./SRP000227/*fastq`
do
    echo "processing: $f"
    g=`basename $f`
    ${BWA} aln -t 2 Sacc2r2.fa $f > aln/$g.sai
    ${BWA} samse Sacc2r2.fa aln/$g.sai $f > aln/$g.sam
    ${SAMTOOLS} view -Sb aln/$g.sam > aln/$g.bam
    rm aln/$g.sai aln/$g.sam
done

```

Descriptive information about samples (e.g., protocol and replicate) were extracted by hand from web pages at the SRA.

Objects used in the lab were processed using the following *R* script. This script requires files and a directory structure that are not distributed with this package.

```

> library(SeattleIntro2010)
> readScript("create-objects.R")

[1] ## http://trace.ncbi.nlm.nih.gov/Traces/sra/sra.cgi?study=SRP000227
[2] library(SeattleIntro2010)
[3] datasrc <- "/home/mtmorgan/SeattleIntro2010/NagalakshmiEtAl"
[4] pkgroot <- "/home/mtmorgan/SeattleIntro2010"
[5] setwd(datasrc)
[6]
[7] ## qa
[8] qaFile <- file.path(pkgroot, "data", "qa.rda")
[9] if (!file.exists(qaFile)) {
[10]     ## better: qa on the aligned reads?
[11]     fls <- list.files("SRP000227", pattern="fastq", full=TRUE)
[12]     qalst <- Map(function(fl) {
[13]         fq <- readFastq(fl)
[14]         qa(fq, lane=basename(fl))
[15]     }, fls)
[16]     qa <- do.call(rbind, qalst)
[17]     save(qa, file=qaFile)
[18] } else load(qaFile)
[19] if (interactive())
[20]     browseURL(report(qa))
[21]
[22] ## hitspergene
[23] countsFile <- file.path(pkgroot, "data", "hitspergene.rda")
[24] txdbFile <- file.path(pkgroot, "inst", "extdata", "sacCer2_sgdGene.sqlite")
[25] if (!file.exists(countsFile)) {

```

```

[26]
[27]     ## transcript ranges
[28]     library(GenomicFeatures)
[29]     if (!file.exists(txdbFile)) {
[30]         txdb <- makeTranscriptDbFromUCSC(genome="sacCer2",
[31]                                         tablename="sgdGene")
[32]         saveFeatures(txdb, txdbFile)
[33]     } else txdb <- loadFeatures(txdbFile)
[34]     exons <- exonsBy(txdb, "gene")
[35]
[36]     ## reads and counts
[37]     fls <- list.files("aln", pattern="fastq.bam$", full=TRUE)
[38]     cnt <- Map(function(fl, exons) {
[39]         ga <- readGappedAlignments(fl)
[40]         countOverlaps(exons, ga)
[41]     }, fls, MoreArgs=list(exons=exons))
[42]     hitspergene <- as(cnt, "DataFrame")
[43]     dimnames(hitspergene) <-
[44]         list(names(exons), sub(".fastq.bam", "", basename(fls)))
[45]
[46]     ## sample annotations
[47]     df <- DataFrame(Sample=rep(c("RH", "dT"), each=3),
[48]                    Replicate=rep(c("Biological", "Original", "Technical"), 2),
[49]                    SRR=c("SRR002058", "SRR002059", "SRR002061",
[50]                          "SRR002062", "SRR002051", "SRR002064"))
[51]     elementMetadata(hitspergene) <-
[52]         df[match(colnames(hitspergene), df$SRR),]
[53]     o <- with(elementMetadata(hitspergene),
[54]              order(Sample, Replicate))
[55]     hitspergene <- hitspergene[,o]
[56]
[57]     save(hitspergene, file=countsFile)
[58] } else load(countsFile)
[59]
[60] ## SRR002051.pluscvg and SRR002051.minuscvg
[61] bamFile <- file.path("aln", "SRR002051.fastq.bam")
[62] pluscvgFile <- file.path(pkgroot, "data", "SRR002051.pluscvg.rda")
[63] minuscvgFile <- file.path(pkgroot, "data", "SRR002051.minuscvg.rda")
[64] if (!file.exists(pluscvgFile) || !file.exists(minuscvgFile)) {
[65]     bam <- readGappedAlignments(bamFile)
[66]     library(BSgenome.Scerevisiae.UCSC.sacCer2)
[67]     bam@seqlengths <- seqlengths(Scerevisiae)
[68]     SRR002051.pluscvg <- coverage(grg(bam)[strand(bam) == "+"])
[69]     SRR002051.minuscvg <- coverage(grg(bam)[strand(bam) == "-"])
[70]     save(SRR002051.pluscvg, file=pluscvgFile)
[71]     save(SRR002051.minuscvg, file=minuscvgFile)

```

```
[72] } else {
[73]     load(pluscvgFile)
[74]     load(minuscvgFile)
[75] }
[76]
```

Relevant software versions are

Program: samtools (Tools for alignments in the SAM format)
Version: 0.1.11 (r851)

Program: bwa (alignment via Burrows-Wheeler transformation)
Version: 0.5.8c (r1536)
Contact: Heng Li <lh3@sanger.ac.uk>

sratoolkit.2.0b4-3-suse_linux32/fastq-dump
Version: 2.0.0

```
> sessionInfo()
```

```
R version 2.12.1 beta (2010-12-07 r53813)
Platform: i386-apple-darwin9.8.0/i386 (32-bit)
```

```
locale:
[1] C
```

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

```
other attached packages:
[1] SeattleIntro2010_0.0.33 biomaRt_2.6.0
[3] GO.db_2.4.5             hgu95av2.db_2.4.5
[5] org.Hs.eg.db_2.4.6     RSQLite_0.9-4
[7] DBI_0.2-5              AnnotationDbi_1.12.0
[9] Biobase_2.10.0
```

```
loaded via a namespace (and not attached):
[1] RCurl_1.4-3 XML_3.2-0 tools_2.12.1
```

References

- [1] U. Nagalakshmi, Z. Wang, K. Waern, C. Shou, D. Raha, M. Gerstein, and M. Snyder. The transcriptional landscape of the yeast genome defined by RNA sequencing. *Science*, 320:1344–1349, Jun 2008.