Data preprocessing and creation of the data objects pasillaGenes and pasillaExons

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Abstract

This vignette describes the steps that were followed for the generation of the data objects contained in the package *pasilla*.

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1 Downloading the files

We used the RNA-Seq data from the publication by Brooks et al. [1]. The experiment investigated the effect of siRNA knock-down of pasilla, a gene that is known to bind to mRNA in the spliceosome, and which is thought to be involved in the regulation of splicing. The data set contains 3 biological replicates of the knockdown as well as 4 biological replicates for the untreated control. Data files are publicly available in the NCBI Gene Expression Omnibus under the accession GSE18508[1]. The read sequences in FASTQ format were extracted from the NCBI short read archive file (.sra files), using the sra toolkit[2].

2 Read alignment and filtering

The reads in the FASTQ files were aligned using tophat version 1.2.0 with default parameters against the reference Drosophila melanogaster genome. Table 1 summarizes the read number and alignment statistics.

[1] `http://www.ncbi.nlm.nih.gov/projects/geo/query/acc.cgi?acc=GSE18508`

[2] `http://www.ncbi.nlm.nih.gov/books/NBK47540/#SRA_Download_Guid_B.5_Converting_SRA_for`

	file	type	number of lanes	total number of reads	exon counts
1	treated1fb	single-read	5	35158667	15679615
2	treated2fb	paired-end	2	12242535 (x2)	15620018
3	treated3fb	paired-end	2	12443664 (x2)	12733865
4	untreated1fb	single-read	2	17812866	14924838
5	untreated2fb	single-read	6	34284521	20764558
6	untreated3fb	paired-end	2	10542625 (x2)	10283129
7	untreated4fb	paired-end	2	12214974 (x 2)	11653031

Table 1: Read numbers and alignment statistics. The column *exon counts* refers to the number of reads that could be uniquely aligned to an exon.

The reference genome fasta files were obtained from the Ensembl ftp server[3]. We ran `bowtie-build` to index the fasta file. For more information on this procedure see the bowtie webpage[4]. The indexed form is required by bowtie, and thus tophat.

```
wget ftp://ftp.ensembl.org/pub/release-62/fasta/drosophila_melanogaster/ \
dna/Drosophila_melanogaster.BDGP5.25.62.dna_rm.toplevel.fa.gz

gunzip Drosophila_melanogaster.BDGP5.25.62.dna_rm.toplevel.fa.gz
bowtie-build Drosophila_melanogaster.BDGP5.25.62.dna_rm.toplevel.fa \
    d_melanogaster_BDGP5.25.62
```

We generated the alignment BAM file using tophat. For the single-reads data:

```
tophat bowtie_index reads1.fastq,reads2.fastq,...,readsN.fastq
```

For the paired-end data:

```
tophat -r inner-fragment-size bowtie_index \
    reads1_1.fastq,reads2_1.fastq,...,readsN_1.fastq \
    reads1_2.fastq,reads2_2.fastq,...,readsN_2.fastq
```

More information on tophat is provided on its webpage[5]. The SAM alignment files from which *pasilla* was generated are available at `http://www-huber.embl.de/pub/DEXSeq/analysis/brooksetal/bam/`.

3 Exon count files

To generate the per-exon read counts, we first needed to define the exonic regions. To this end, we downloaded the file Drosophila_melanogaster.BDGP5.25.62.gtf.gz from Ensembl[6]. The script `dexseq_prepare_annotation.py` contained in the *DEXSeq* package was used to extract the exons of the transcripts from the file, define new non-overlapping exonic regions and reformat it to create the file Dmel.BDGP5.25.62.DEXSeq.chr.gff contained in pasilla/extdata. For example, for this file we ran:

[3] `http://www.ensembl.org/info/data/ftp/index.html`
[4] `http://bowtie-bio.sourceforge.net/tutorial.shtml`
[5] `http://tophat.cbcb.umd.edu/tutorial.html`
[6] `ftp://ftp.ensembl.org/pub/release-62/gtf/drosophila_melanogaster`

```
wget ftp://ftp.ensembl.org/pub/release-62/gtf/ \
drosophila_melanogaster/Drosophila_melanogaster.BDGP5.25.62.gtf.gz

gunzip Drosophila_melanogaster.BDGP5.25.62.gtf.gz
python dexseq_prepare_annotation.py Drosophila_melanogaster.BDGP5.25.62.gtf \
    Dmel.BDGP5.25.62.DEXSeq.chr.gff
```

To count the reads that fell into each non-overlapping exonic part, the script `dexseq_count.py`, which is also contained in the *DEXSeq* package, was used. It took the alignment results in the form of a SAM file (sorted by position in the case of a paired end data) and the `gtf` file Dmel.BDGP5.25.62.DEXSeq.chr.gff and returned one file for each biological replicate with the exon counts. For example, for the file treated1.bam, which contained single-end alignments, we ran:

```
samtools index treated1.bam
samtools view treated1.bam > treated1.sam
python dexseq_count.py Dmel.BDGP5.25.62.DEXSeq.chr.gff \
    treated1.sam treated1fb.txt
```

For the file treated2.bam, which contained paired-end alignments:

```
samtools index treated2.bam
samtools view treated2.bam > treated2.sam
sort -k1,1 -k2,2n treated2.sam > treated2_sorted.sam
python dexseq_count.py -p yes Dmel.BDGP5.25.62.DEXSeq.chr.gff \
    treated2_sorted.sam treated2fb.txt
```

The output of the two HTSeq python scripts is provided in the *pasilla* package:

```
> library("pasilla")

> inDir = system.file("extdata", package="pasilla", mustWork=TRUE)
> dir(inDir)

 [1] "Dmel.BDGP5.25.62.DEXSeq.chr.gff" "geneIDsinsubset.txt"
 [3] "pasilla_gene_counts.tsv"         "treated1fb.txt"
 [5] "treated2fb.txt"                  "treated3fb.txt"
 [7] "untreated1fb.txt"                "untreated2fb.txt"
 [9] "untreated3fb.txt"                "untreated4fb.txt"
```

The Python scripts are built upon the HTSeq library[7].

4 Creation of the *ExonCountSet* pasillaExons

To create an *ExonCountSet* object, we started with a data frame `samples` that contained the sample annotations, as in Table 1.

```
> head(samples)
```

[7] http://www-huber.embl.de/users/anders/HTSeq/doc/overview.html

```
             condition        type
treated1fb     treated single-read
treated2fb     treated  paired-end
treated3fb     treated  paired-end
untreated1fb untreated single-read
untreated2fb untreated single-read
untreated3fb untreated  paired-end
```

We also needed the annotation file with the per exon annotation.

```
> annotationfile = file.path(inDir, "Dmel.BDGP5.25.62.DEXSeq.chr.gff")
```

With these, we could call the function `read.HTSeqCounts` to construct the object `ecs`.

```
> library("DEXSeq")
> ecs = read.HTSeqCounts(countfiles = file.path(inDir, paste(rownames(samples), "txt", sep=".")),
+            design = samples,
+            flattenedfile = annotationfile)
> sampleNames(ecs) = rownames(samples)
```

We only wanted to work with data from a subset of genes, which was defined in the following file.

```
> genesforsubset = readLines(file.path(inDir, "geneIDsinsubset.txt"))
> pasillaExons = subsetByGenes(ecs, genes=genesforsubset)
```

We added the experiment data:

```
> expdata = new("MIAME",
+    name="pasilla knockdown",
+    lab="Genetics and Developmental Biology, University of Connecticut Health Center",
+    contact="Dr. Brenton Graveley",
+    title="modENCODE Drosophila pasilla RNA Binding Protein RNAi knockdown RNA-Seq Studies",
+    url="http://www.ncbi.nlm.nih.gov/projects/geo/query/acc.cgi?acc=GSE18508",
+    abstract="RNA-seq of 3 biological replicates of from the Drosophila melanogaster
+       S2-DRSC cells that have been RNAi depleted of mRNAs encoding pasilla, a mRNA binding
+       protein and 4 biological replicates of the the untreated cell line.")
>    pubMedIds(expdata) <- "20921232"
> experimentData(pasillaExons) <- expdata
```

5 Creation of the *CountDataSet* `pasillaGenes`

The *CountDataSet* class is analogous to the *ExonCountSet* class; the latter is specifically designed to store exon level counts, while the *CountDataSet* class is useful more generally for whatever one wishes to count (e. g. ChIP peaks, gene levels counts). We made use of the function `geneCountTable` from the package *DEXSeq* to get a data frame containing the number of reads falling on each of the genes. We used the function `newCountDataSet` to create the object `pasillaGenes`.

```
> library("DESeq")
> genetable = geneCountTable(ecs)
> pasillaGenes = newCountDataSet(genetable,
+   conditions = samples)
> experimentData(pasillaGenes) = expdata
```

```
> toLatex(sessionInfo())
```

- R version 2.15.2 (2012-10-26), `i386-w64-mingw32`
- Locale: `LC_COLLATE=C`, `LC_CTYPE=English_United States.1252`, `LC_MONETARY=English_United States.1252`, `LC_NUMERIC=C`, `LC_TIME=English_United States.1252`
- Base packages: base, datasets, grDevices, graphics, methods, stats, utils
- Other packages: Biobase 2.18.0, BiocGenerics 0.4.0, DESeq 1.10.1, DEXSeq 1.4.0, lattice 0.20-13, locfit 1.5-8, pasilla 0.2.15, xtable 1.7-0
- Loaded via a namespace (and not attached): AnnotationDbi 1.20.3, DBI 0.2-5, IRanges 1.16.4, RColorBrewer 1.0-5, RCurl 1.95-3, RSQLite 0.11.2, XML 3.95-0.1, annotate 1.36.0, biomaRt 2.14.0, genefilter 1.40.0, geneplotter 1.36.0, grid 2.15.2, hwriter 1.3, parallel 2.15.2, splines 2.15.2, statmod 1.4.16, stats4 2.15.2, stringr 0.6.2, survival 2.37-2, tools 2.15.2

Table 2: The output of `sessionInfo` on the build system after running this vignette.

We saved the objects in the data directory of the package:

```
> save(pasillaExons, file=file.path("..", "..", "data", "pasillaExons.RData"))
> save(pasillaGenes, file=file.path("..", "..", "data", "pasillaGenes.RData"))
```

References

[1] A. N. Brooks, L. Yang, M. O. Duff, K. D. Hansen, J. W. Park, S. Dudoit, S. E. Brenner, and B. R. Graveley. Conservation of an RNA regulatory map between Drosophila and mammals. *Genome Research*, pages 193–202, October 2010.