

Visualization methods for RNA-sequencing data

SUMMARY: This is the summary for this paper.

KEY WORDS: Data visualization; Exploratory data analysis; Interactive graphics; RNA-sequencing; Statistical graphics

1. Introduction

RNA-sequencing (RNA-seq) uses next-generation sequencing (NGS) to estimate the quantity of RNA in biological samples at given timepoints. In recent years, decreasing cost and increasing throughput has rendered RNA-seq an attractive alternative to transcriptome profiling. Prior to RNA-seq, gene expression studies were performed with microarray techniques, which required prior knowledge of reference sequences. RNA-seq does not have this limitation, and has enabled a new range of applications such as transcriptome de novo assembly (Grabherr et al., 2011; Robertson et al., 2010) and detection of alternative splicing processes (Anders, Reyes, and Huber, 2012; Pan et al., 2008). Coupled with its high resolution and sensitivity, RNA-seq will likely revolutionize our understanding of the intricacies of eukaryotic transcriptomes (Wang, Gerstein, and Snyder, 2009; Zhao et al., 2014).

Gene expression data is multivariate data, and its basic form is a matrix containing mapped read counts for n rows of genes and p columns of samples. These mapped read counts provide estimations of the gene expression levels across samples. Researchers typically conduct RNA-seq studies to identify differentially expressed genes (DEGs) between treatment groups. In most popular RNA-seq analysis packages, this objective is approached with models, such as the negative binomial model (Anders and Huber, 2010; Trapnell et al., 2013; Trapnell et al., 2012; Robinson et al., 2010) and linear regression models (Law et al., 2014).

Initially, it was widely claimed that RNA-seq produced unbiased data that did not require sophisticated normalization (Wang et al., 2009; Morin et al., 2008; Marioni et al., 2008). However, numerous studies have since revealed that RNA-seq data is replete with biases and that accurate detection of DEGs is not a negligible task. Problems that complicate the analysis of RNA-seq data include nucleotide-specific and read-position specific biases (Hansen et al., 2010), biases related to gene lengths and sequencing depths (Oshlack, Robinson, and Young, 2010; Robinson and Oshlack, 2010), biases introduced during library preparation

(McIntyre et al., 2011), biases pertaining to the number of replications (Schurch et al., 2016), biases derived from overlapping senseantisense transcripts and gene isoforms (Trapnell et al., 2013), and the confounding combination of technical and biological variability (Bullard et al., 2010).

In light of these complications, researchers should analyze RNA-seq data like they would any other biased multivariate data. Simply applying models to such data is problematic because models have assumptions that they alone cannot call into question. Fortunately, data visualization enables researchers to see patterns and problems they may not otherwise detect with traditional modeling. As a result, the most effective approach to analyze data is to iterate between models and visuals, and to enhance the appropriateness of applied models based on feedback from visuals (Unwin, 1992; Shneiderman, 2002).

When visualizing RNA-seq data, we primarily want to compare the variability between replicates and between treatment groups. This is best achieved by viewing the mapped read count distributions across all genes and samples. Unfortunately, the few visual tools available in popular RNA-seq packages do not allow users to effectively view their data in this manner.

Here, we use real RNA-seq data to demonstrate that our visualization tools can detect normalization problems, DEG designation problems, and common errors in the analysis pipeline. We also show that our tools can identify genes of interest that cannot otherwise be obtained by any models. In line with modern multivariate data exploration, we emphasize that interactive graphics should be an indispensable component of RNA-seq analysis. Researchers should be able to quickly flip through plots of genes that appear promising or problematic, and link between different types of plots to quickly obtain various perspectives of their data. In this paper, we are not proposing that users completely change their approach to RNA-seq analysis. Instead, we propose that users simply modify their approach to RNA-

seq analysis by assessing the sensibility of their models with multivariate graphical tools, namely parallel coordinate plots, scatterplot matrices, and replicate line plots.

2. Parallel Coordinate Plots

Parallel coordinate plots are essential to visually verify the relationships between variables in multivariate data. A parallel coordinate plot draws each row in the data table as a line. Connections between samples with positive correlations will be flat, and connections between samples with negative correlations will be crossed. An ideal dataset will have large variability between treatment groups but little variability between replicates. Researchers can quickly confirm this with a parallel coordinate plot: There should be flat connections between replicates but crossed connections between treatment groups.

There are several packages within the BioConductor software that provide graphics for RNA-seq data analysis (Huber et al., 2015). Two of the most common graphic techniques are side-by-side boxplots and Multidimensional Scaling (MDS) plots (Love, Huber, and Anders, 2014; Risso et al., 2011; Robinson et al., 2010; Su et al., 2016; Ritchie et al., 2015; Marini, 2017). Unfortunately, these plots can hide problems that still exist in the data even after normalization and that could be better detected with parallel coordinate plots.

Figure 1 exemplifies this problem for two simulated datasets, one displayed on the left half of the figure and the other displayed on the right half of the figure. Each dataset contains two treatment groups (A and B) with three replicate samples. We cannot detect any notable differences between the left and right datasets from the side-by-side boxplots at the top of the figure as they both show fairly consistent five number summaries across their six samples. Likewise, we cannot detect any notable differences between the datasets from the MDS plots in the middle of the figure as they both suggest that the datasets are clustered by the two treatment groups, although the first replicate from treatment group A appears as an outlier in the right MDS plot.

Despite this, we immediately see from the parallel coordinate plots at the bottom of the figure that the left and right datasets have an important difference. The left dataset has consistent (level) lines between replicates and inconsistent (crossed) lines between treatment groups. This suggests that some of the genes (lines) have consistently low values for treatment group A and consistently high values for treatment group B, while some genes have the opposite phenomenon. As a result, these plotted genes are likely candidates for differential expression. In contrast, the right dataset does not possess this ideal structure and suggests that the genes may not be candidates for differential expression. We could not see this important distinction from the side-by-side boxplots and MDS plots because they simply provide summaries about the data on the sample level, while the parallel coordinate plot shows the sample connections for each of the 50 genes.

[Figure 1 about here.]

[Table 1 about here.]

You can experiment with fancier tables than Table 1.

We can get bold symbols using `\bm{math}`, for example, α_i .

3. Discussion

Put your final comments here.

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SUPPLEMENTARY MATERIALS

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APPENDIX

Title of appendix

Put your short appendix here. Remember, longer appendices are possible when presented as Supplementary Web Material. Please review and follow the journal policy for this material, available under Instructions for Authors at <http://www.biometrics.tibs.org>.

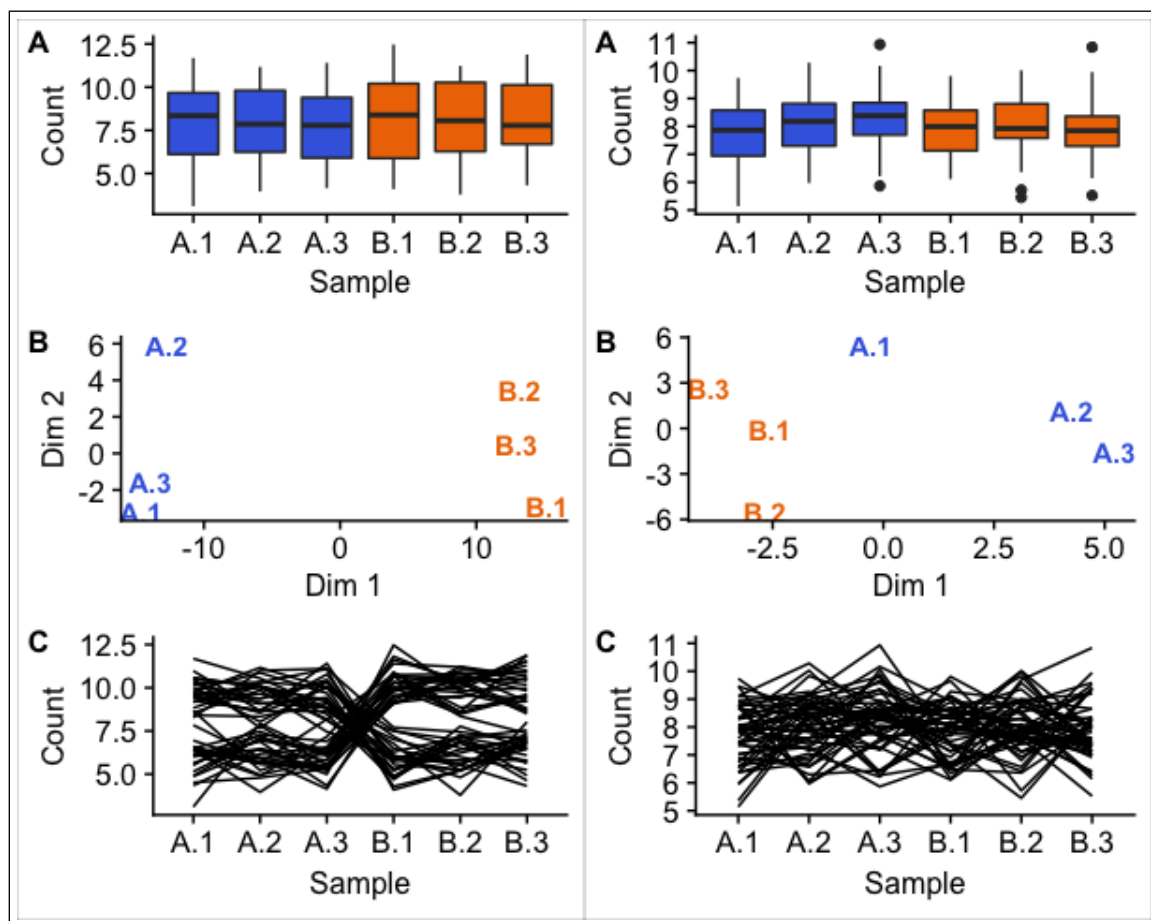


Figure 1. Caption.

Table 1*This is a simple table.*

Culture/Library prep.	Library prep. protocol	Growth condition	Flow-cell
Y1	Protocol 1	YPD	428R1
Y1	Protocol 1	YPD	4328B
Y2	Protocol 1	YPD	428R1
Y2	Protocol 1	YPD	4328B
Y7	Protocol 1	YPD	428R1
Y7	Protocol 1	YPD	4328B
Y4	Protocol 2	YPD	61MKN
Y4	Protocol 2	YPD	61MKN