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Genome analysis

scDAPA: detection and visualization of dynamic alternative polyadenylation from single cell RNA-seq data

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Abstract

Motivation: Alternative polyadenylation (APA) plays a key post-transcriptional regulatory role in mRNA stability and functions in eukaryotes. Single cell RNA-seq (scRNA-seq) is a powerful tool to discover cellular heterogeneity at gene expression level. Given 3′ enriched strategy in library construction, the most commonly used scRNA-seq protocol—10× Genomics enables us to improve the study resolution of APA to the single cell level. However, currently there is no computational tool available for investigating APA profiles from scRNA-seq data.

Results: Here, we present a package scDAPA for detecting and visualizing dynamic APA from scRNA-seq data. Taking bam/sam files and cell cluster labels as inputs, scDAPA detects APA dynamics using a histogram-based method and the Wilcoxon rank-sum test, and visualizes candidate genes with dynamic APA. Benchmarking results demonstrated that scDAPA can effectively identify genes with dynamic APA among different cell groups from scRNA-seq data.

Availability and implementation: The scDAPA package is implemented in Shell and R, and is freely available at https://scdapa.sourceforge.io.

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Supplementary information: Supplementary data are available at *Bioinformatics* online.

1 Introduction

Alternative polyadenylation (APA) is increasingly recognized as an important regulation mechanism for many biological processes (e.g. cell development, differentiation and proliferation) and molecular functions (e.g. mRNA stability, translation efficiency and localization) via dynamically using different polyadenylation sites during maturation of nuclear pre-mRNA (Chen *et al.*, 2017). More and more evidence shows widespread occurrences of cell type-specific APA in eukaryotes (Cao *et al.*, 2019; Hwang *et al.*, 2017; Velten *et al.*, 2015). To profile APA dynamics among different cell types, one traditional way is to dissociate tissues and purify cell types before performing Poly(A)-tag sequencing (Cao *et al.*, 2019), which may introduce unexpected cellular stress and affect the APA profile

consequently. An alternative way is using methods like cTag-PAPERCLIP (Hwang *et al.*, 2017), which could directly profile individual cell types from tissues without enzymatic digestion and fluorescent-activated cell sorting. The common limitation of these two strategies is that they cannot deal with unknown cell types or rare cell subpopulations. The ideal way to differentiate APA profiles from different cell types is to perform single cell RNA-seq (scRNA-seq), which can isolate cells of different states and/or types by dissecting their transcriptome profiles (Saliba *et al.*, 2014; Zheng *et al.*, 2017). By integrating a conventional scRNA-seq protocol and a 3′ enriched bulk population RNA-seq protocol, Velten *et al.* (2015) developed a method BATseq to investigate the APA profile at single cell resolution. However, the low sensitivity and complex steps of BATseq limit its wide application (Chen *et al.*, 2017). Promising

experimental protocols for quantifying APA at single cell level are still lacking.

One commonly used scRNA-seq protocol, i.e. 10× Genomics, applying a 3′ selection/enriched strategy in library construction, provides the potential of quantifying APA dynamics at single cell resolution (Chen *et al.*, 2017; Saliba *et al.*, 2014; Ye *et al.*, 2019). In our recent study (Ye *et al.*, 2019), we found out 3′ ends extracted from the 10× Genomics scRNA-seq data are quite adjacent to authentic poly(A) sites or 3′ UTR annotations defined by sequencing experiments, and investigated the role of APA in acute myeloid leukemia using several scRNA-seq datasets from 10× Genomics, demonstrating the high validity and value of existing scRNA-seq data in performing APA analysis. However, currently there is no easy-to-use computational tool available for investigating APA profiles from scRNA-seq data. In this work, we developed a package scDAPA for exploration and visualization of APA dynamics in different cell types and conditions from scRNA-seq data.

2 Implementation

The scDAPA consists of three major steps (Supplementary Fig. S1): (i) 3′ ends extraction and annotation; (ii) dynamic APA detection; and (iii) dynamic APA visualization. First, scDAPA takes the sequence alignment results (a bam/sam file) and cell type information (a cell barcode-cell type data in csv format) as inputs, and extracts valid mapping records (uniquely mapped in genomic region, with valid cell barcode and not PCR duplicates etc.) into different files of distinct cell types. 3′ ends of extracted reads are annotated to genes based on the given genome annotation file (a gff/gtf file). Second, the dynamic APA is detected among different cell types within the same biological samples or among different samples of the same cell type. Briefly, we use a histogram-based method to divide the dispersed 3′ ends into distinct bins with the same width (default 100 bp), and a Site Distribution Difference (SDD) index is calculated to quantify the APA difference between conditions as $SDD = \sum_{n=1}^{N} |p_A^n - p_B^n|/2$, $SDD \in [0, 1]$, where N is the number of bins, A and B denote two different cell groups, and p_A^n represents the percentage of 3′ ends located in the nth bin of a specific gene in cell group A. Then, the Wilcoxon rank-sum test is applied to measure the significance of differential APA usage. A gene with a SDD value above a given cutoff (e.g. 0.2) and a p-value below a given cutoff (e.g. 0.05) will be recognized a gene with significant differential APA usage (DE-APA gene). For multiple statistical tests, the Benjamini-Hochberg method is employed to control the false discovery rate. Last, scDAPA allows intuitive visualization of APA profiles of candidate genes under different conditions, which presents diverse isoforms of a target gene, and a smooth density plot alongside the gene to show the 3′ ends distribution. The Supplementary Material contains a user manual of the scDAPA package.

3 Application example

We investigated the application of scDAPA on a scRNA-seq dataset of live microglia/macrophages from pooled neuroretinas of normal and light damaged mice generated by the 10× Genomics platform (O'Koren *et al.*, 2019). We extracted the 3′ ends of valid reads from the scRNA-seq dataset and compared them to the latest mouse poly(A) site annotations from PolyA_DB 3/Ensembl (Wang *et al.*, 2018). The distribution of the distances between 3′ ends of the aligned scRNA-seq reads and their nearest poly(A) annotations is similar to that of our previous work (Fig. 1a) (Ye *et al.*, 2019), indicating a high stability of 10× Genomics scRNA-seq data in representing APA site usage. Next, we performed a cell type-to-cell type comparison to detect genes with dynamic APA across different cell groups (DE-APA genes). DE-APA genes were categorized into two groups: APA gene and non-APA gene. Referring to the annotations from PolyA_DB 3/Ensembl, a gene with only one poly(A) site is recognized as a non-APA gene, otherwise it's recognized as an APA gene. Approximately 98% of 3137 detected non-redundant DE-APA genes are annotated as APA genes (Fisher's exact test p-value
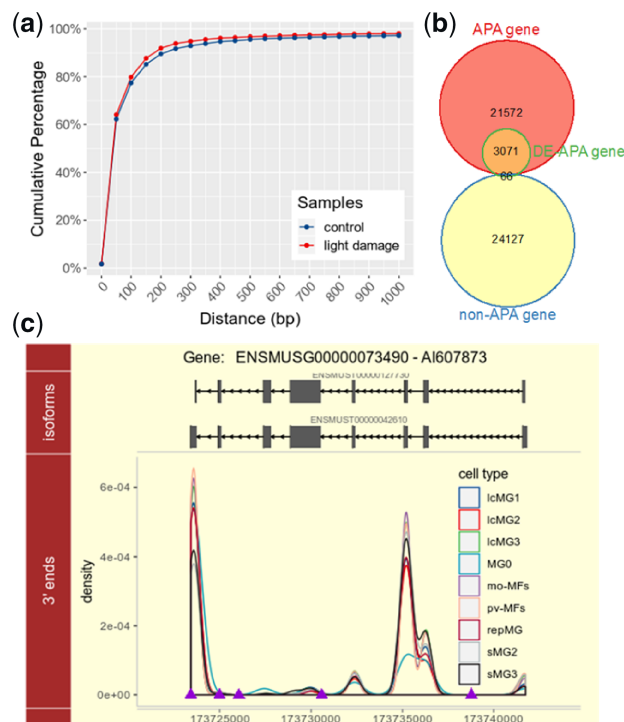

Fig. 1. An application example of scDAPA on a scRNA-seq dataset of neuroretinas from mouse. (a) Cumulative distribution of distances between 3′ ends and nearest authentic poly(A) sites; (b) categories of DE-APA genes detected from scRNA-seq data; (c) illustration of a gene *AI607873* showing dynamic APA profiles across different cell groups. Top panel, isoforms of gene *AI607873* from Ensembl; bottom panel, density distribution of 3′ ends of various cell groups; purple triangles represent the poly(A) site annotation from PolyA_DB 3. (Color version of this figure is available at *Bioinformatics* online.)

$< 2.2 \times e^{-16}$, Fig. 1b), representing a high confidence of scDAPA in dynamic APA detection using scRNA-seq data.

Additionally, in pair-wise cell type comparisons of APA usage, the cell type MG0 (microglia solely came from normal neuroretinas) always shows the highest percent of DE-APA genes compared with other nine cell types (Supplementary Fig. S2), suggesting a dramatically distinct APA preference of the nine cell types (IcMG1~3, sMG1~3, mo-MFs, pv-MFs, repMG: microglia mainly came from light damaged neuroretinas) compared to steady state MG0. Among the identified DE-APA genes, e.g. a gene *AI607873* prefers using a distal poly(A) site in MG0 compared with other nine cell types (Fig. 1c). Moreover, among the three small microglia clusters (sMG1, sMG2 and sMG3), sMG3 shows the most APA dynamics compared to MG0 (Supplementary Fig. S2). This result is consistent with the previous observation that sMG3 was particularly distinct from MG0, and was found at the final state of a trajectory analysis (O'Koren *et al.*, 2019). Considering that sMG3 is corresponding to the subretinal microglia in light damaged sample (O'Koren *et al.*, 2019), KEGG and REACTOME pathway enrichment analyses for DE-APA genes between MG0 and sMG3 were further performed. We found that these DE-APA genes were significantly over-represented in many signaling pathways, such as NF-kappa B, JAK-STAT and TNF signaling pathways (Supplementary Fig. S3a and b), which are associated with the retinal photoreceptor degeneration (Rashid *et al.*, 2018). These results demonstrate that scDAPA can effectively identify genes with dynamic APA among different cell groups from scRNA-seq data.

4 Conclusion

In summary, we developed a package scDAPA to detect and visualize dynamic APA from scRNA-seq data. We demonstrated its utilities through application to a real dataset. It is believed that scDAPA

is a useful tool in studying APA at single cell resolution, and will broadly extend the application scope of scRNA-seq data.

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