

Gene network reconstruction using single cell transcriptomic data reveals key factors for autophagic process

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Abstract

Single-cell RNA sequencing (scRNA-seq) is a powerful tool to study heterogeneity and dynamic changes in cell populations. Data explosion fueled by technological development requires more efficient approach to process scRNA-seq data. Current algorithms for scRNA-seq data lack scalability, which can lead to exponential increase in processing time. To process large-scale scRNA-seq data we propose a new approach called SHARP by applying ensemble random projection (RP). RP reduces the dimensionality of data by trading a controlled amount of error for faster processing time. RP approximately preserves cell-to-cell distances while achieving a substantial cost reduction for computing, a desired property in handling large-scale scRNA-seq data. Ensembled average enhances modeling cell-to-cell distance by compensating errors of individual RP results. Applying SHARP to clustering using various scRNA-seq datasets including 43,745 cells from murine kidneys, we found SHARP runs at least 40 times faster than previous algorithms while maintaining a better clustering performance.

Keywords: *SHARP: Single-Cell RNA-Seq Hyper-Fast and Accurate Processing via Ensemble Random Projection*

Biography

2005 Ph.D. in Electronics and Computer Science, University of Southampton, Southampton, UK

2006 - 2006 Postdoctoral Fellow, University of Copenhagen, Denmark

2007 – 2010 Postdoctoral Fellow, University of California, San Diego, Department of Chemistry & Biochemistry

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