RIA: A NOVEL REGRESSION-BASED IMPUTATION APPROACH FOR SINGLE-CELL RNA SEQUENCING

Bang Tran, Duc Tran, Hung Nguyen, Nam Sy Vo and Tin Nguyen*
Department of Computer Science and Engineering, University of Nevada, Reno
Contact: tinn@unr.edu , Website: https://bioinformatics.cse.unr.edu/

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UNIVERSITY OF NEVADA, RENO

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Background

Advances in single-cell technologies have shifted genomics research from the analysis of bulk tissues toward a comprehensive characterization of individual cells. This holds enormous opportunities for both basic biology and clinical research. However, low amount of mRNA available within individual cells leads to the excess amount of zero counts caused by dropout events.

Objectives

Develop an imputation method, RIA, that can reliably impute missing values from single-cell data. RIA consists of two modules. The first module performs a hypothesis testing to identify the values that are likely to be impacted by the dropout events. The second module estimates the missing value using a robust regression approach.

Results

Data: 5 datasets with a total of 3,535 cells.

Metric: Adjusted Rand Index (ARI) [8], Jaccard Index [9] and Purity Index [10].

Methods: scImpute [15], MAGIC [16], t-SNE [17].

Results: RIA produces the best ARI values, preserve the transcriptomics landscape and significantly elucidates the cell lineage identification.

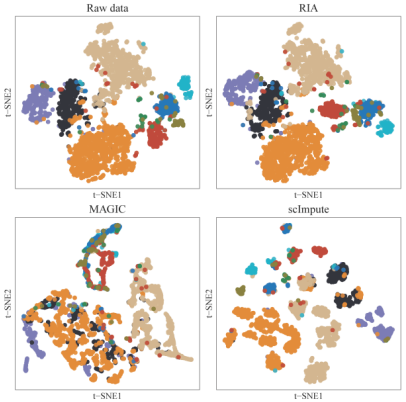

Fig. 2. RIA preserves the transcriptomics landscape for Zeisel [14] dataset.

Methodology

Hypothesis Testing and Identification of Dropout : to determine genes that are likely to be impacted by dropouts. Genes that are not impacted by dropouts, the log-transformed expression values are normally distributed. We use z-test to determine whether a zero is impacted by the dropout events. Original data is divided into two sets of genes: a set *G* that include genes affected by dropout (imputable set), and a set *M* that have high confidence of not being affected by dropout. (training set)

Regression-based Imputation:

- We select genes from the training set that are highly correlated with the gene we need to impute.
- We train the linear model using these highly-correlated genes and then estimate the missing values

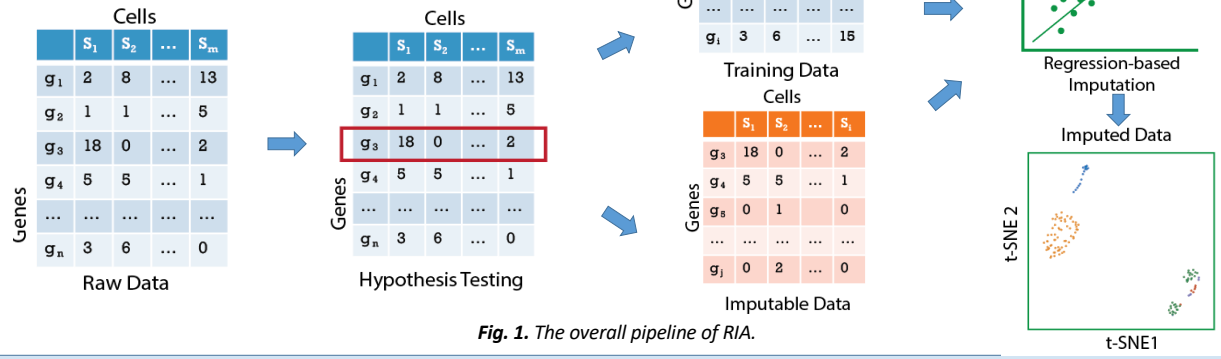

Fig. 1. The overall pipeline of RIA.

Conclusion

- Outperforms existing state-of-the-art approaches in cell group identification.
- Recover temporal trajectories in embryonic development stages
- RIA is fast and is able to impute thousands of cells with tens of thousands of genes in minutes

Future work

We plan to utilize the perturbation clustering [3],[4],[6].

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