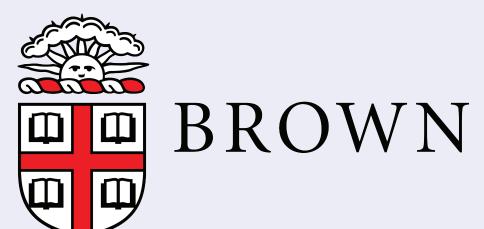
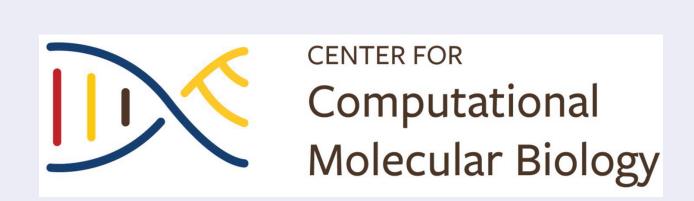
scMultiNODE: Temporal Single-Cell Data Integration across Unaligned Modalities





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Codes & Paper github.com/rsinghlab/scMultiNODE

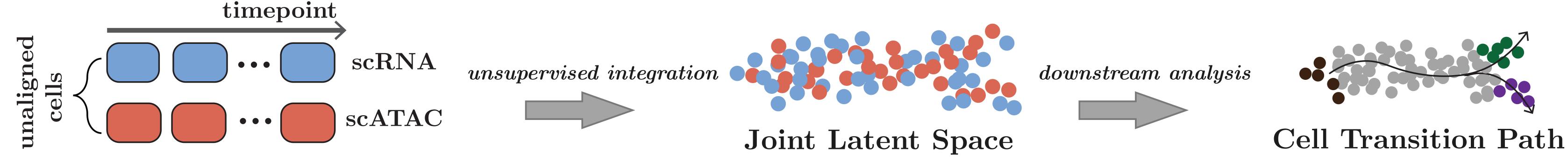
Problem

- Temporal scRNA-seq data are only profiled at discrete and sparsely spaced timepoints due to laborious and expensive lab experiments
- Obtaining different sequencing assays on the same cells across developmental stages is technically challenging



Integrate unaligned cell profiles from two modalities: gene expression and chromatin accessibility

• Retain both cell type variations & cellular dynamics during integration



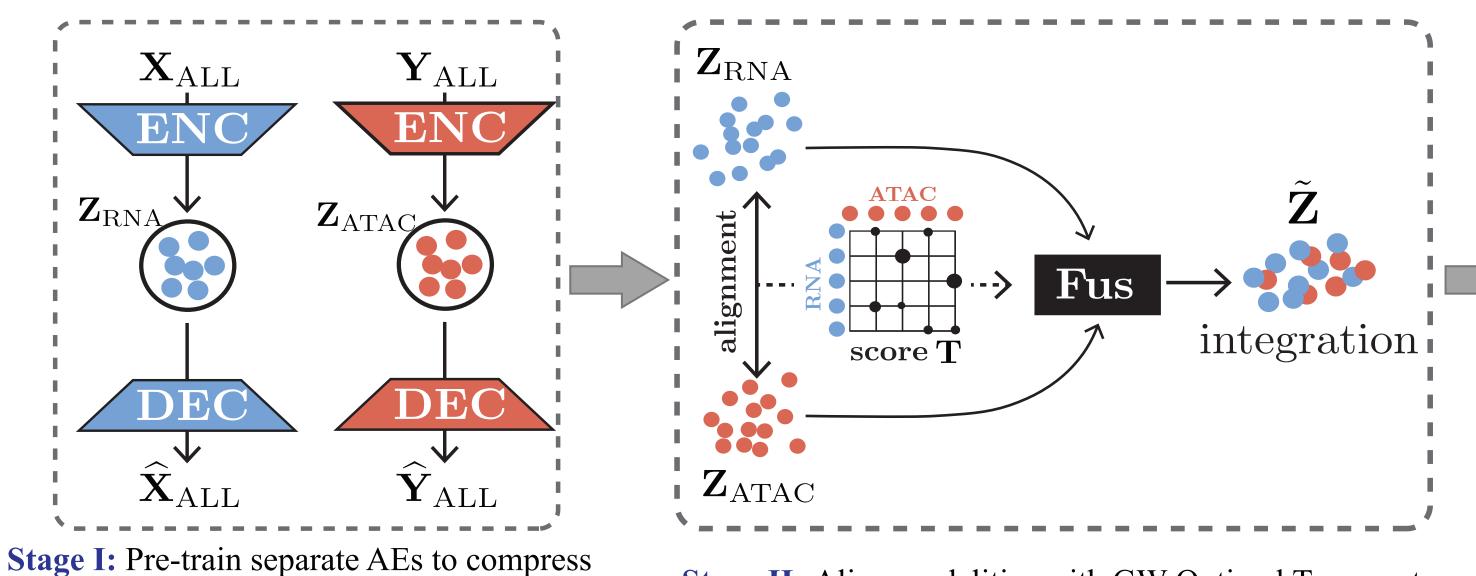
Method: single-cell Multi-Modal Neural Ordinary Differnetial Equation (scMultiNODE)

Input (e.g., scRNA-seq + scATAC-seq)

- Gene expression $\mathbf{X}^{(t)}$ at measured timepoints $t \in \mathcal{T}_{RNA} \subset \{0, 1, \cdots\}$
- Chromatin accessibility $\mathbf{Y}^{(t)}$ at measured timepoints $t \in \mathcal{T}_{ATAC} \subset \{0, 1, \dots\}$

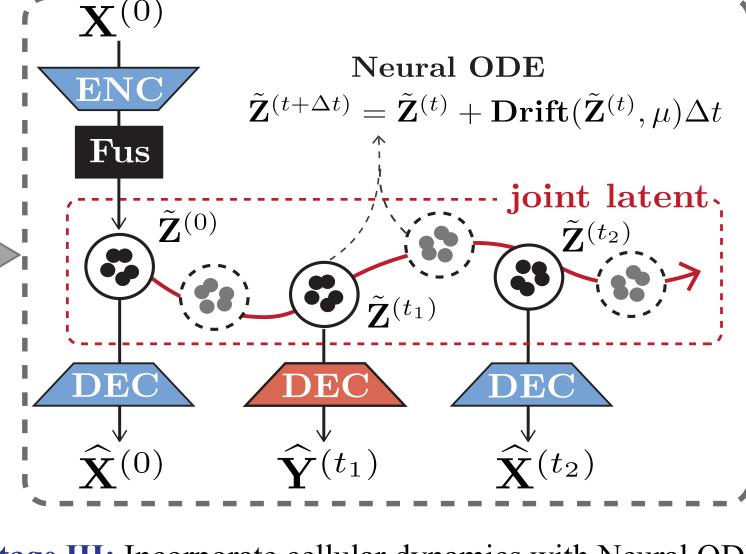
Advantage

- Quantization GW enables efficient alignment across large-scale multi-modal data
- Integrations capture both cell type variations and developmental dynamics



Goal

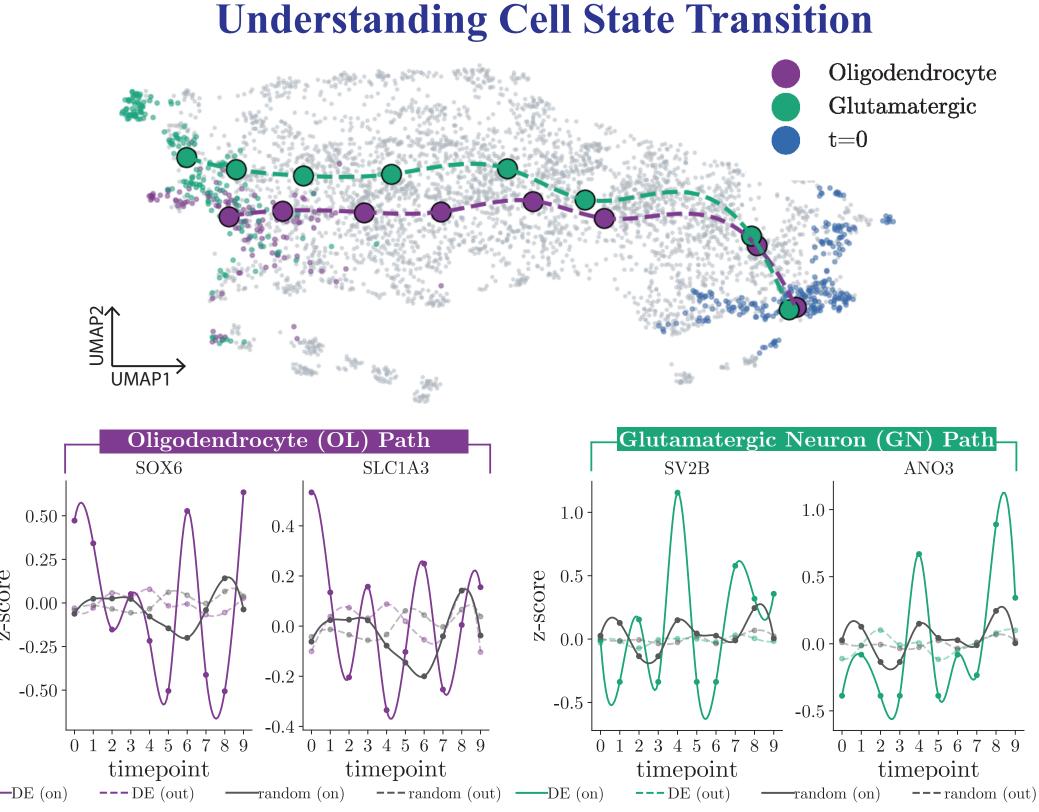
Stage II: Align modalities with GW Optimal Transport



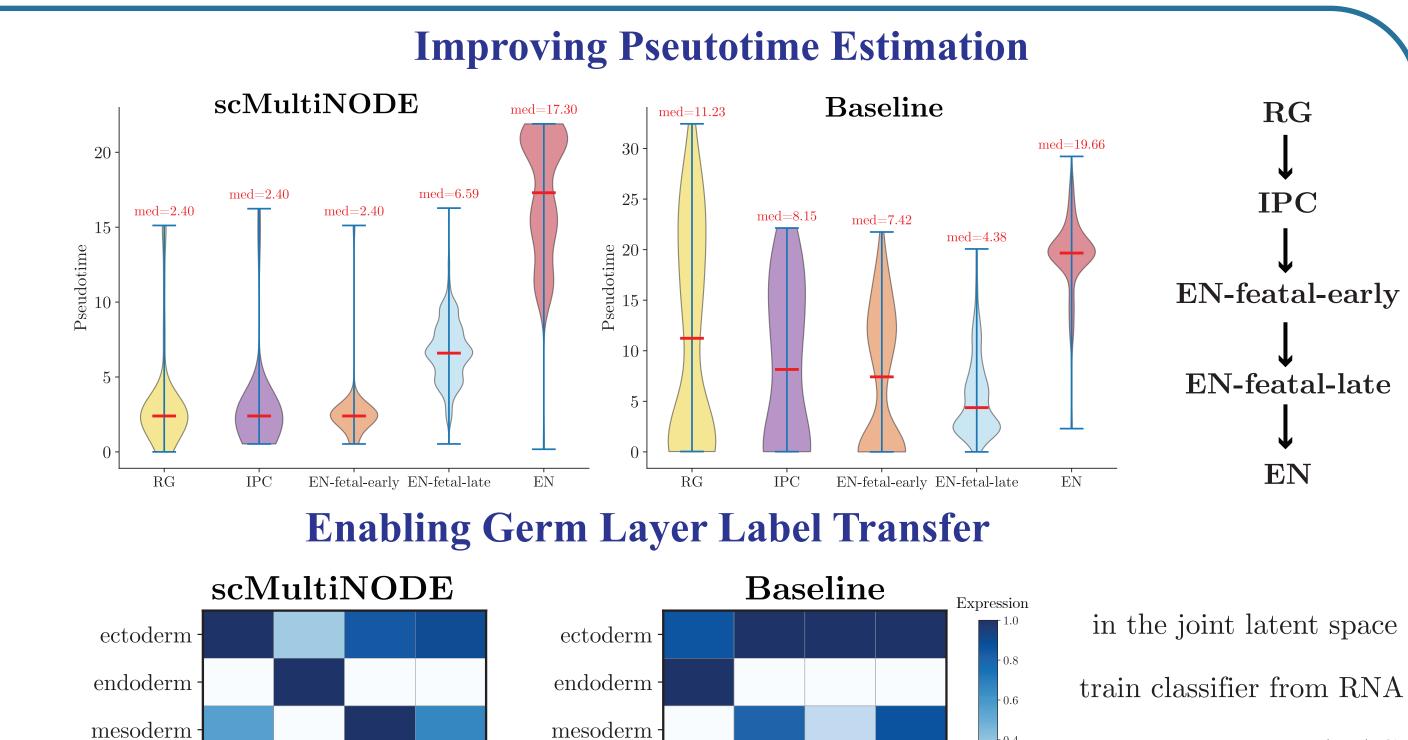
Stage III: Incorporate cellular dynamics with Neural ODE

scMultiNODE Captures Cell Type Variations & Cellular Dynamics in Integration

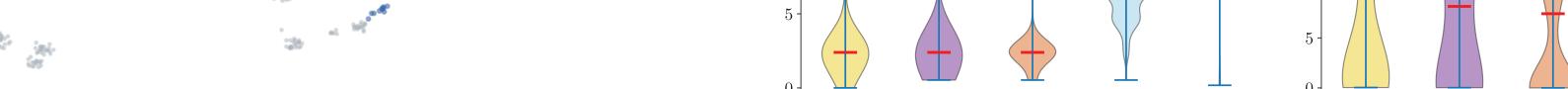
• SCOTv2 0.06UnionCom Seurat



high-dim. and sparse single-cell data



mesoderm predict labels on ATAC neuroectoderm validate with marker genes



neuroectoderm

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