

Preparation of samples for 10x-sc/snRNA-seq

Single cell RNA-seq

- 1) Fresh cells: Provide single-cell suspension $\geq 250k$ cells (10x-scRNA + sci/10x-ATAC-seq (from same sample): $\geq 500k$ cells) with viability $>80\%$ in buffer with up to 10% serum.
- 2) Frozen cells: Provide single-cell suspension $\geq 250k$ cells (10x-scRNA + sci/10x-ATAC-seq (from same sample): $\geq 500k$ cells) on dry ice e.g. in 0.5 mL of media containing 10% DMSO.
Note: Dissociation and freezing conditions to yield high viability of single-cell preparations are cell type-dependent.

Tissues

Tissues should be flash-frozen in liquid nitrogen immediately upon resection. (Input: 10x-snRNA: 250k nuclei; 10x-snRNA + sci/10x-ATAC-seq (from same sample): $\geq 500k$ nuclei)

General Notes:

- 1) **Important:** Before preparing a large set of samples for 10x-sc/snRNA-seq, we strongly recommend submitting 1-2 test samples of the same type and using the same preparation method as the larger batch of samples to be assayed.
- 2) If you cannot fulfill the sample input requirements, please contact epigenome@ucsd.edu we will find a custom solution for you.
- 3) We prefer that samples be submitted in standard 1.5 mL microfuge tubes. If you require another format, please contact epigenome@ucsd.edu to make arrangements prior to sample submission.