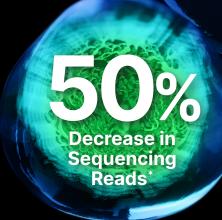


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Uninformative sequences obscure detection of low expressing transcripts and rare cell types.

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Increase in Genes Detected Per Cell*





Transcriptomic

Reads

Remove uninformative molecules from single cell libraries—doubling your transcriptomic reads

- Gain a deeper view of expression profiles of individual cells
- Deplete sequences not used for secondary analysis including: Unaligned reads, ribosomal, mitochondrial, non-variable genes
- Gain greater differentiation in cell types without perturbing cell type calls

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