

Boost usable single cell data with CRISPRclean®

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

It's time to get the most out of your sequencing dollars.

1.5x
Increase in
Genes Detected
Per Cell*

50%
Decrease in
Sequencing
Reads*

2x
Transcriptomic
Reads

3

Simple
Steps

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

Get Started Now