



# TotalSeg™-C Mouse Universal Cocktail, V1.0

Catalog# / Size 199903 / 5 tests

Regulatory Status RUO

**Description** The TotalSeq™-C Mouse Universal Cocktail has been designed to react with 113 cell surface

antigens, including principal lineage antigens, and includes 9 isotype control antibodies, to aid

in the multiomic characterization of immune cells.

The lyophilized cocktail provides convenience when processing several samples at the same time or at different time points, reducing inconsistencies associated with pipetting and handling multiplex vials and samples. The antibodies in the cocktail are provided at optimized concentrations to provide a ready-to-use solution once the cocktail has been reconstituted. The TotalSeq<sup>TM</sup>-C Mouse Universal Cocktail comes packaged in convenient single-use vials.

#### **Product Details**

Verified Reactivity Mouse

Formulation Lyophilized from PBS containing stabilizers. Please reconstitute each tube with cell staining buffer

as indicated in the application notes when ready to use. Make sure the cake is completely dissolved prior to using the cocktail for staining cells. The lyophilized cocktail material can vary in

appearance and appear as either a lyophilized cake or a powder.

**Preparation** This reagent is a combination of TotalSeq<sup>™</sup>-C oligo conjugated clones at optimal concentrations

for single-cell sequencing analysis. See additional product notes for a list of clones.

Concentration Lot-specific (to obtain lot-specific concentration, please enter the lot number in our Concentration

and Expiration Lookup or Certificate of Analysis online tools.)

Storage & Handling The lyophilized antibody cocktail should be stored between 2°C and 8°C in powder form and in a

sealed container with desiccant until ready to use. Once vial is opened, it is to be reconstituted

immediately. After reconstitution, the cocktail should be used within 2 hours.

Application PG - Quality tested

Recommended Usage This panel has been optimized on mouse splenocytes and bone marrow. Using other cell types or

additional TotalSeq™ antibodies may require further optimization. Please contact our technical

service group for further information.

Buyer is solely responsible for determining whether Buyer has all intellectual property rights that are necessary for Buyer's intended uses of the BioLegend TotalSeq™ products. For example, for any technology platform Buyer uses with TotalSeq™, it is Buyer's sole responsibility to determine whether it has all necessary third party intellectual property rights to use that platform and

TotalSeq™ with that platform.

Application Notes <u>Click here for protocol</u>

Download the complete dataset for all targets

Additional Product Notes TotalSeq™ reagents are designed to profile protein expression at a single cell level following an

optimized protocol similar to the CITE-seq workflow. A compatible single cell device (e.g. <u>10x</u> <u>Genomics Chromium System and Reagents</u>) and sequencer (e.g. Illumina analyzers) are required.

Please contact <u>technical support</u> for more information, or visit <u>biolegend.com/totalseq</u>.

The barcode flanking sequences are CGGAGATGTGTATAAGAGACAGNNNNNNNNN (PCR handle), and NNNNNNNNCCCATATAAGA\*A\*A (capture sequence). N represents either randomly selected A, C, G, or T, and \* indicates a phosphorothioated bond, to prevent nuclease

degradation.

The full oligomer sequence for this product, with the specific barcode in brackets is CGGAGATGTGTATAAGAGACAGNNNNNNNNNN [Barcode] NNNNNNNNCCCATATAAGA\*A\*A.

Downlo

Download the excel file for a full list of clones and barcodes.



Download the Cell Ranger antibody reference and UMI counting csv file.

# **Antigen Details**

**Distribution** Multiple cell types and cell states

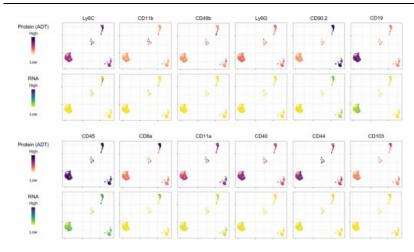
Gene ID NA

## **Related Protocols**

TotalSeg™-B or -C with 10x Feature Barcoding Technology

TotalSeq™ Universal Cocktail Reconstitution and Staining Instructions Protocol v2

### **Product Data**



Major immune cell lineages and selected cell states are identified using the TotalSeq™-C Mouse Universal Cocktail. Resting mouse splenocytes were stained with the TotalSeq™-C Mouse Universal Cocktail and processed using the 10x Genomics Single Cell 5' v1 feature barcoding kit and Illumina sequencing. Protein count data were transformed and visualized in a UMAP projection overlayed with protein expression levels for each component of the cocktail. Cell clusters were identified based on protein expression only.

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