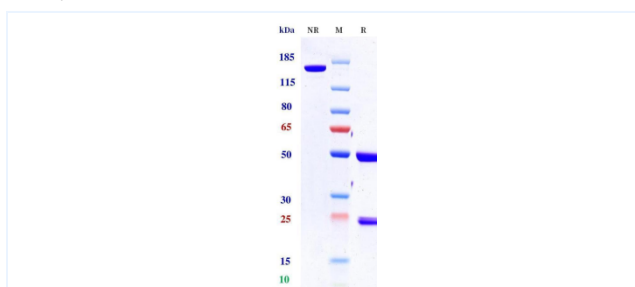


Product Details

| | | | |
|-----------------|---|--------------------|-----------------|
| Product name: | Anti-BCMA & CD3 (Elranatamab Biosimilar) | SKU: | BIO0967SM |
| Target Name: | BCMA & CD3 | Size: | 100ug/ 1mg/ 5mg |
| Target Uniprot: | Q02223 & P07766 | Concentration: | Lyophilized |
| Clone#: | Elranatamab (Bispecific) | Isotype: | IgG-like |
| Reactivity: | Human | Calculated M.W.: | 145.44 kDa |
| Application: | ELISA, Bioactivity: FACS, Functional assay, Research in vivo | Endotoxin: | <0.001 EU/ug |
| Formulation: | 100 mM Pro-Ac 20mM Arg pH 5.0 | Conjugation: | None |
| Storage: | -20°C for 2 years under sterile conditions; -20°C for 1 year under sterile conditions; Avoid repeated freeze-thaw cycles. | Expression System: | CHO |
| Reconstitution: | Dissolve with sterile ddH ₂ O | Purification: | Protein A |

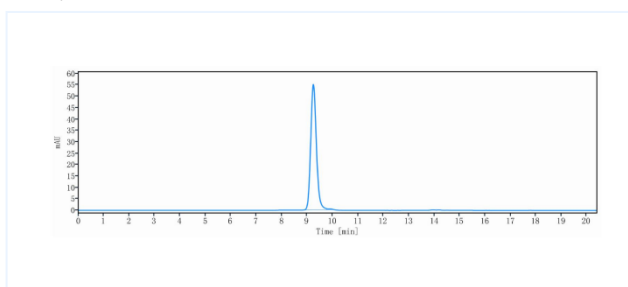
Data

Purity: SDS-PAGE



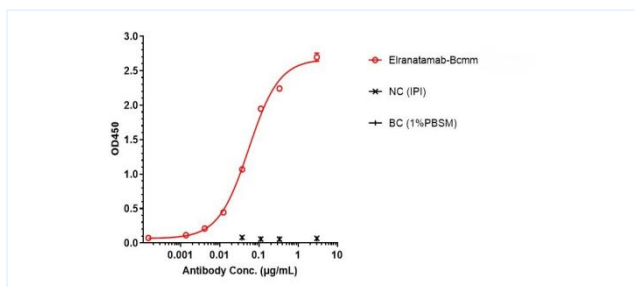
Anti-BCMA & CD3 Reference Antibody (Elranatamab) on SDS-PAGE under reducing (R) condition. The purity of the protein is greater than 95%.

Purity: SEC-HPLC



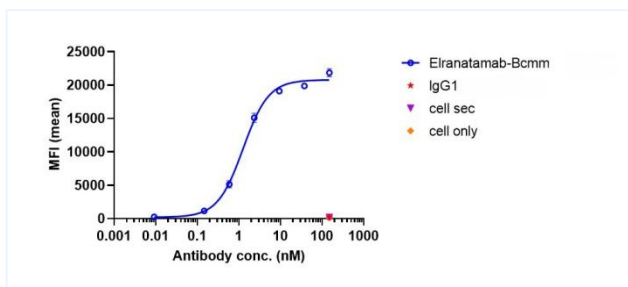
The purity of Anti-BCMA & CD3 Reference Antibody (Elranatamab) is 99.34%, determined by SEC-HPLC.

ELISA



Elranatamab-Bcmm bound to BCMA protein, and then rebounded to secondary antibodies (Anti-human-IgG-Fc-HRP), and read OD450. As shown in fig, Elranatamab-Bcmm bound to huBCMA-ECD-His, and the EC₅₀ was 0.055 nM.

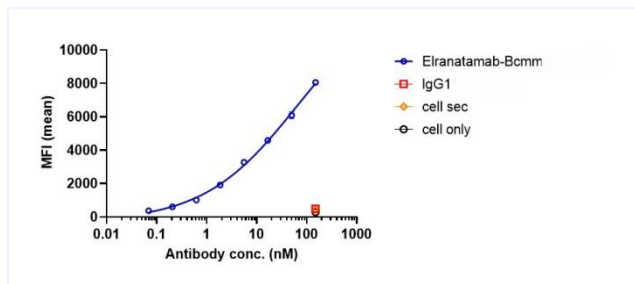
Bioactivity: FACS



Elranatamab-Bcmm bound to huBCMA-HEK293 cells, and then rebounded to fluorescent secondary antibodies (Anti-human IgG, Fc PE), and test by flow cytometry. As shown in fig, Elranatamab-Bcmm bound to huBCMA-HEK293 cells, and the EC₅₀ was 1.263 nM.

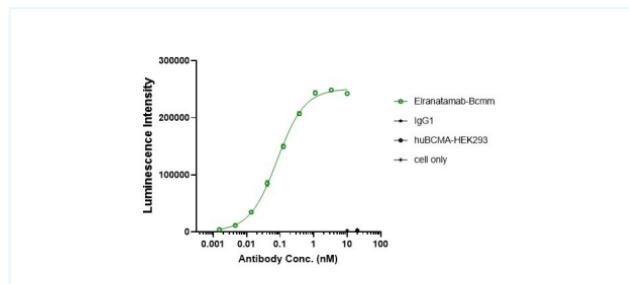
Important Note: This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications.

Bioactivity: FACS



Elranatamab-Bcmm bound to huCD3e-jurkat cells, and then rebounded fluorescent secondary antibodies (Anti-human IgG, Fcy PE) , and test by flow cytometry. As shown in fig, Elranatamab-Bcmm bound to huCD3e-jurkat cells, and the EC50 was 64.990 nM.

Function: Luciferase



Co-incubation of Elranatamab-Bcmm with Jurkat cells, then with the addition of huBCMA-HEK293 cells for 6 hours. Bright-Lite was used to detect the fluorescent signal. As shown in fig, Elranatamab-Bcmm was able to activate the NF-AT signaling pathway, and the EC50 was 0.081 nM.