



CERTIFICATE OF ANALYSIS

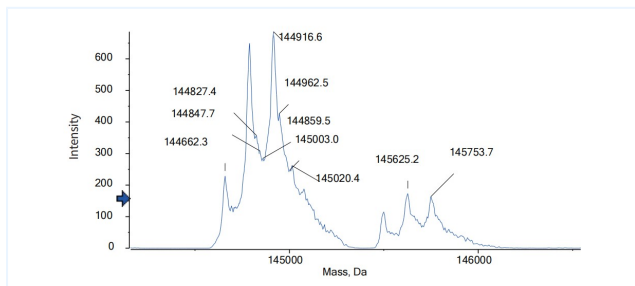


Product Details

Product name:	Anti-cMet Reference Antibody (Davutamig)	Lot.No.:	P267996
Target:	HGFR / c-Met	Catalog:	CHBA031
Target Accession:	P08581	Concentration :	1 mg/mL
Clonality:	Bispecific	Isotype:	IgG-like
Reactivity:	Human	Molecular Weight (kDa) :	144.89 kDa
Application:	Kinetics (SPR), ELISA, Bioactivity: FACS, Functional assay, Research in vivo	Endotoxin:	<1 EU/mg
Formulation:	Liquid: 100mM Pro-Ac, 20mM Arg pH 5.0 Lyophilization: 25mM histidine, 8% sucrose, 0.01% Tween80 pH6.2	Conjugation:	Unconjugated
Reconstitution:	For Powder, reconstitute with sterile, distilled water to a final concentration of 1 mg/ml. Gently shake to solubilize completely. Do not vortex.	Expression System:	CHO
Storage:	Upon receipt, store immediately at -20°C or lower for 24 months. Store aliquots at -80°C for up to 3 months. Avoid repeated freeze-thaw cycles.	Purification:	Protein A

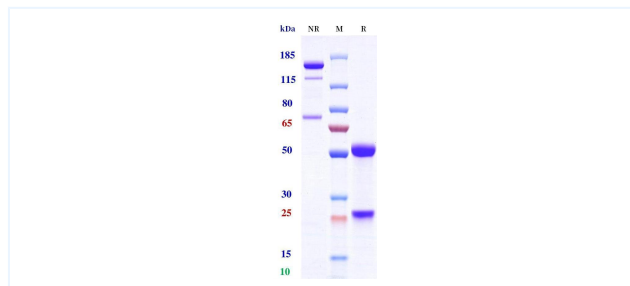
Data

MASS



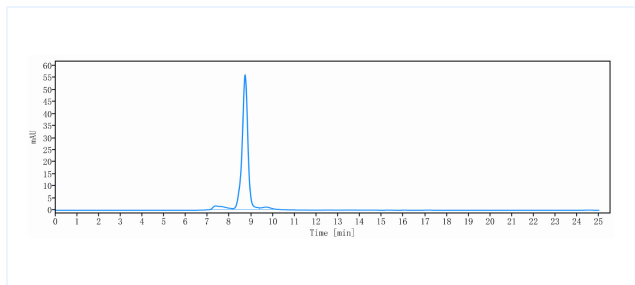
The detected molecular weight of Anti-cMet Reference Antibody (Davutamig) is 144.92 kDa.

Purity: SDS-PAGE



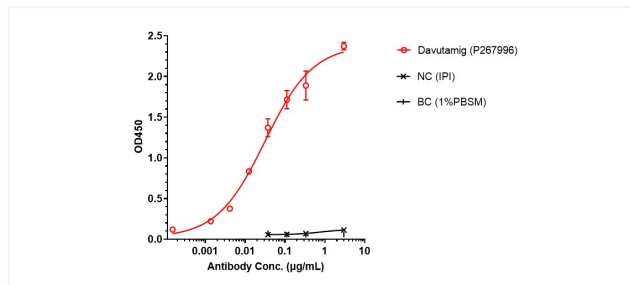
Anti-cMet Reference Antibody (Davutamig) on SDS-PAGE under reducing (R) condition. The purity of the protein is greater than 95%.

Purity: SEC-HPLC



The purity of Anti-cMet Reference Antibody (Davutamig) is 99.52%, determined by SEC-HPLC.

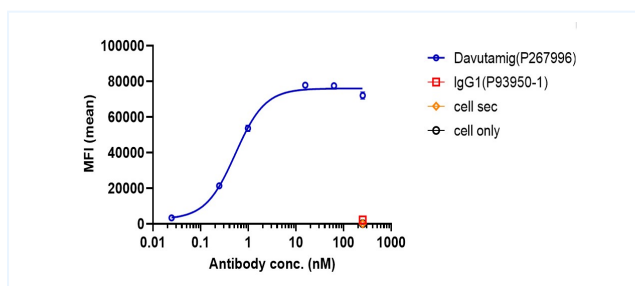
ELISA



Davutamig bound to cMet protein, and then rebounded to secondary antibodies (Anti-human-IgG-Fc-HRP), and read OD450. As shown in fig, Davutamig bound to cMet-His, and the EC₅₀ was 0.031 nM.

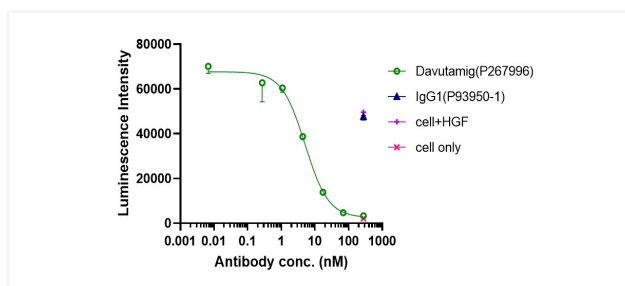
The products are for research use only. Not for use in diagnostic procedures.

Bioactivity: FACS



Davutamig bound to huMet-HEK293 cells, and then rebounded to fluorescent secondary antibodies (Anti-human IgG, Fcy PE), and test by flow cytometry. As shown in fig, Davutamig bound to huMet-HEK293 cells, and the EC50 was 0.532 nM.

Function: Luciferase



Co-incubation of Davutamig with HGF, then with the addition of Human c-MET (Luc) HEK293 Reporter cells and incubated for 6 hours. Bright-Lite was used to detect the fluorescent signal. As shown in fig, Davutamig was able to block HGF/c-Met signaling pathway, and the IC50 was 5.112 nM.

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