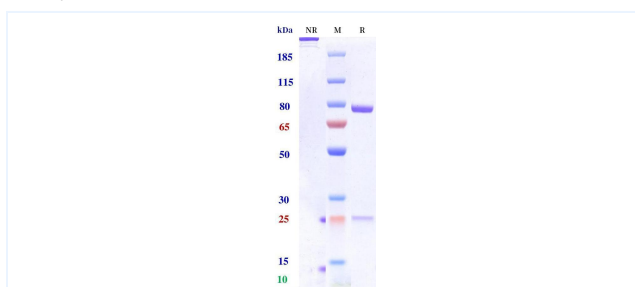


Product Details

Product name:	Anti-PD-1 & VEGF (Ivonescimab Biosimilar)	SKU:	BIO1019SM
Target Name:	PD-1 & VEGF	Size:	100ug/ 1mg/ 5mg
Target Uniprot:	Q15116 & P15692	Concentration:	Lyophilized
Clone#:	Ivonescimab (Bispecific)	Isotype:	IgG-ScFv
Reactivity:	Human	Calculated M.W.:	201.12 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

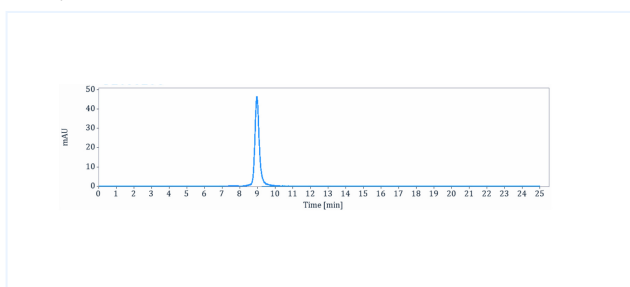
Data

Purity: SDS-PAGE



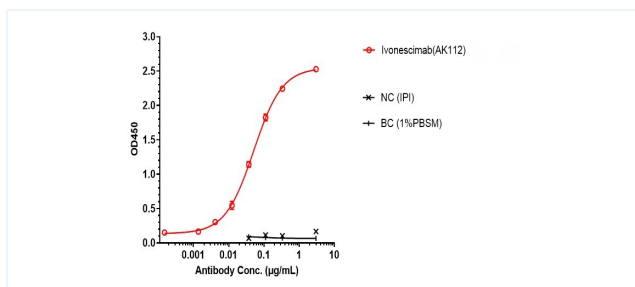
Anti-PD-1 & VEGF Reference Antibody (Ivonescimab) on SDS-PAGE under reducing (R) condition. The purity of the protein is greater than 95%.

Purity: SEC-HPLC



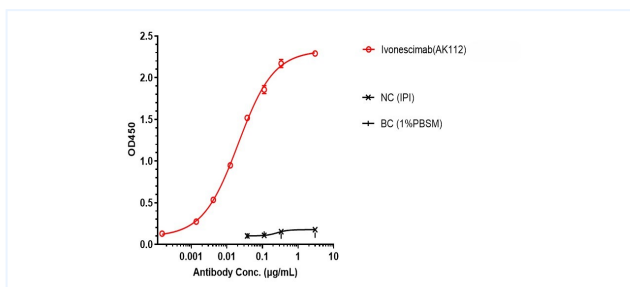
The purity of Anti-PD-1 & VEGF Reference Antibody (Ivonescimab) is 98.33%, determined by SEC-HPLC.

ELISA



Ivonescimab bound to PD-1 protein, and then rebounded to secondary antibodies(Anti-human-IgG-Fc-HRP), and read OD450. As shown in fig, Ivonescimab bound to huPD-1-His, and the EC₅₀ was 0.051 nM.

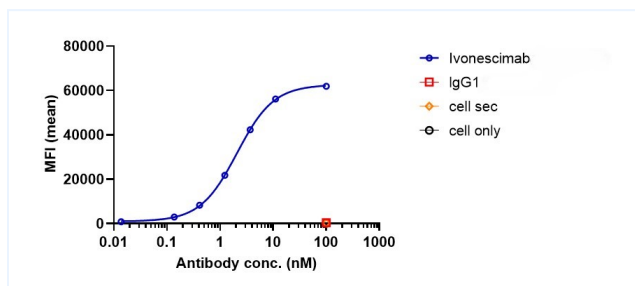
ELISA



Ivonescimab bound to VEGFA protein, and then rebounded to secondary antibodies(Anti-human-IgG-Fc-HRP), and read OD450. As shown in fig, Ivonescimab bound to huVEGFA-His, and the EC₅₀ was 0.021 nM.

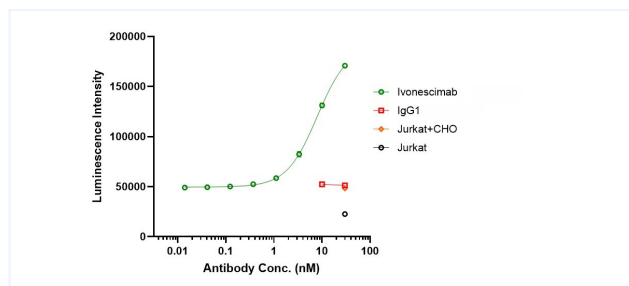
Important Note: This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications.

Bioactivity: FACS



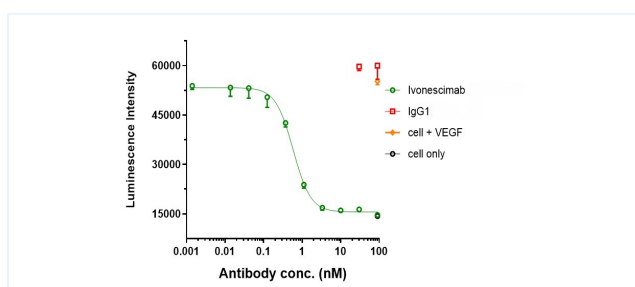
Ivonescimab bound to huPD-1-Jurkat cells, and then rebounded to fluorescent secondary antibodies (Anti-human IgG, Fcy PE), and test by flow cytometry. As shown in fig, Ivonescimab bound to huPD-1-Jurkat cells, and the EC50 was 6.274 nM.

Function: Luciferase



Co-incubation of Ivonescimab with PD-1-NF-AT-Jurkat and CD3L-huPD-L1-CHO-K cells and incubated for 6 hours. Bright-Lite was used to detect the fluorescent signal. As shown in fig, Ivonescimab was able to block the PD-1/PD-L1 signaling pathway, and the EC50 was 7.957 nM.

Function: Luciferase



Co-incubation of Ivonescimab with VEGF protein, then with the addition of VEGF-NF-AT-HEK293 cells and incubated for 6 hours. Bright-Lite was used to detect the fluorescent signal. As shown in fig, Ivonescimab can neutralize VEGF-165, and the IC50 was 7.957 nM.