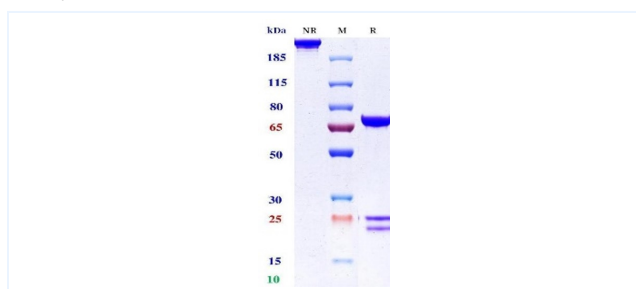


## Product Details

<b>Product name:</b>	<b>Anti-LAG-3 &amp; PD-1 (Emb-02 Biosimilar)</b>	<b>SKU:</b>	<b>BIO1013SM</b>
<b>Target Name:</b>	<b>LAG-3 &amp; PD-1</b>	<b>Size:</b>	<b>100ug/ 1mg/ 5mg</b>
<b>Target Uniprot:</b>	<b>P18627 &amp; Q15116</b>	<b>Concentration:</b>	<b>Lyophilized</b>
<b>Clone#:</b>	<b>Emb-02 (Bispecific)</b>	<b>Isotype:</b>	<b>Fab-IgG</b>
<b>Reactivity:</b>	<b>Human</b>	<b>Calculated M.W.:</b>	<b>237.5 kDa</b>
<b>Application:</b>	<b>ELISA, Bioactivity: FACS, Functional assay, Research in vivo</b>	<b>Endotoxin:</b>	<b>&lt;0.001 EU/ug</b>
<b>Formulation:</b>	<b>100 mM Pro-Ac 20mM Arg pH 5.0</b>	<b>Conjugation:</b>	<b>None</b>
<b>Storage:</b>	<b>-20°C for 2 years under sterile conditions; -20°C for 1 year under sterile conditions; Avoid repeated freeze-thaw cycles.</b>	<b>Expression System:</b>	<b>CHO</b>
<b>Reconstitution:</b>	<b>Dissolve with sterile ddH<sub>2</sub>O</b>	<b>Purification:</b>	<b>Protein A</b>

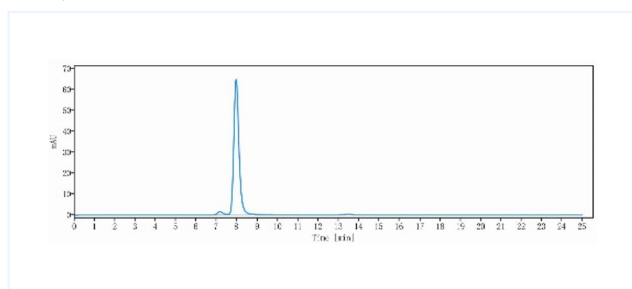
## Data

### Purity: SDS-PAGE



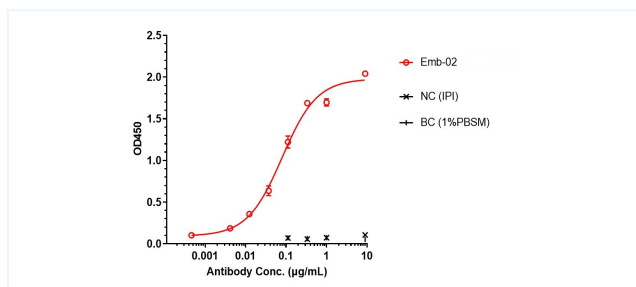
Anti-LAG-3 & PD-1 Reference Antibody (Emb-02) on SDS-PAGE under reducing (R) condition. The purity of the protein is greater than 95%.

### Purity: SEC-HPLC



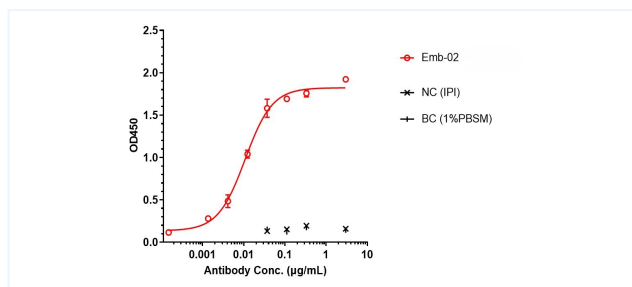
Anti-LAG-3 & PD-1 Reference Antibody (Emb-02) is 97.15%, determined by SEC-HPLC.

### ELISA



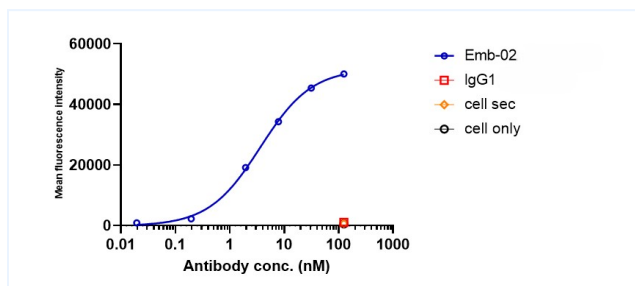
Emb-02 bound to LAG-3 protein, and then rebounded to secondary antibodies(Anti-Human-IgG-Fc-HRP) , and read OD450. As shown in fig, Emb-02 bound human LAG-3 Protein-His, and the EC50 was 0.08nM.

### ELISA



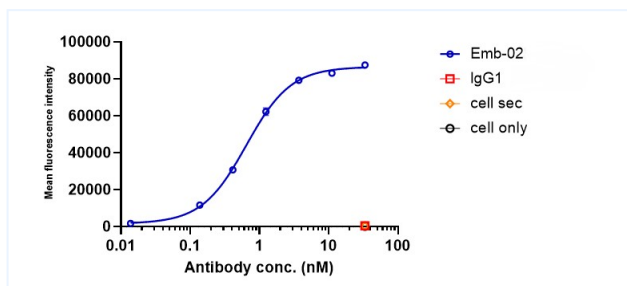
Emb-02 bound to PD-1 protein, and then rebounded to secondary antibodies(Anti-Human-IgG-Fc-HRP) , and read OD450. As shown in fig, Emb-02 bound human PD-1 Protein-His, and the EC50 was 0.01 nM.

### Bioactivity: FACS



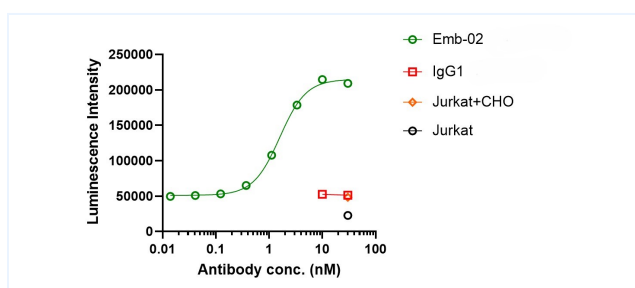
Emb-02 bound to huLAG-3-CHO-K cells, and then rebounded to fluorescent secondary antibodies(Anti-Human IgG, Fcγ PE) , and test by flow cytometry. As shown in fig, Emb-02 bound to huLAG-3-CHO-K cells, and the EC50 was 3.630 nM.

### Bioactivity: FACS



Emb-02 bound to huPD-1-Jurkat cells, and then rebounded to fluorescent secondary antibodies(Anti-Human IgG, Fcγ PE) , and test by flow cytometry. As shown in fig, Emb-02 bound to huPD-1-Jurkat cells, and the EC50 was 0.645 nM.

### Function: Luciferase



Co-incubation of Emb-02 with PD-1-NF-AT-Jurkat an CD3L-huPD-L1-CHO-K cells and incubated for 6 hours. Bright-Lite was used to detect the fluorescent signal. As shown in fig, Emb-02 was able to block the PD-1/PD-L1 signaling pathway, and the EC50 was 1.585 nM.