



Product Data Sheet

PD-1/PD-L1-IN 3

 $\begin{tabular}{lll} \textbf{Cat. No.:} & HY-103048 \\ \begin{tabular}{lll} \textbf{CAS No.:} & 1629654-95-0 \\ \begin{tabular}{lll} \textbf{Molecular Formula:} & $C_{89}H_{126}N_{24}O_{18}S$ \\ \end{tabular}$

Molecular Weight: 1852.17

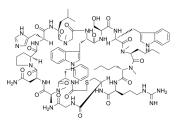
Sequence Shortening: Maa-FANPHL-Sar-WSW-Nle-Nle-RCG (Disulfide bridge: Maa1-Cys15)

Target: PD-1/PD-L1

Pathway: Immunology/Inflammation

Storage: Please store the product under the recommended conditions in the Certificate of

Analysis.



BIOLOGICAL ACTIVITY

Description	PD-1/PD-L1-IN 3, a macrocyclic peptide, is a potent and selective inhibitor of the PD-1/PD-L1 and CD80/PD-L1 interactions extracted from patent WO2014151634A1, compound No.1. PD-1/PD-L1-IN 3 interferes with PD-L1 binding to PD-1 and CD80 by binding to PD-L1, with IC $_{50}$ s of 5.60 nM and 7.04 nM, respectively. PD-1/PD-L1-IN 3 can be used for the research of various diseases, including cancer and infectious diseases ^[1] .
IC ₅₀ & Target	IC50: 5.60 nM (PD-1/PD-L1); 7.04 nM (CD80/PD-L1) ^[1]
In Vitro	PD-1/PD-L1-IN 3 (0.1 nM-10 μ M) inhibits the binding of PD-1 and CD80 to PD-L1, with IC $_{50}$ s of 5.60 nM and 7.04 nM $^{[1]}$. PD-1/PD-L1-IN 3 blocks the binding of recombinant PD-L1-Ig to Jurkat-PD-1 cells, and also block binding of recombinant PD-1-Ig to either L2987 or LK35.2-hPD-L1, with IC $_{50}$ s of 26 nM, 12 nM, and 3.5 nM, respectively $^{[1]}$. PD-1/PD-L1-IN 3 (0.001-100 μ M) promotes IFN secretion by CMV-specific T cells in a dose-dependent manner, with an EC $_{50}$ of 400 nM $^{[1]}$. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. MILLER, Michael Matthew, et al. Macrocyclic inhibitors of the pd-1/pd-l1 and cd80(b7-1)/pd-l1 protein/protein interactions. WO2014151634A1.

Caution: Product has not been fully validated for medical applications. For research use only.

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Proteins

Inhibitors