IRAK4-IN-17

Cat. No.: HY-150594 CAS No.: 2183503-33-3 Molecular Formula: $C_{17}H_{20}F_{2}N_{8}O$ Molecular Weight: 390.39 Target: IRAK; NF-κΒ

Pathway: Immunology/Inflammation; NF-κΒ

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

Analysis.

Product Data Sheet

BIOLOGICAL ACTIVITY

Description	IRAK4-IN-17 (Compound 5) is a potent IRAK4 inhibitor with the IC $_{50}$ of 1.3 nM $^{[1]}$. IRAK4-IN-17 can be used in large B-cell
	lymphoma (DLBCL) research $^{[1]}$.

IC₅₀ & Target IRAK4 $1.3~\text{nM}~(\text{IC}_{50})$

IRAK4-IN-17 (0.7-40.1 μ M; 72 h) selectively suppresses OCI-LY10 and TMD8 cells with MYD88 L265P mutation [1]. In Vitro IRAK4-IN-17 (0.3-10 μ M; 2 h) inhibits the viability of DLBCL cells by targeting IRAK4 signaling [1].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Cell Cytotoxicity $Assay^{[1]}$

Cell Line:	OCI-LY10, TMD8, Ramos and HT cells
Concentration:	0.7-40.1 μΜ
Incubation Time:	72 hours
Result:	Suppressed OCI-LY10 and TMD8 cells with MYD88 L265P mutation (IC $_{50}$ =0.7 μ M and 1.2 μ M, respectively), but not Ramos and HT cell lines with WT MYD88 (IC $_{50}$ =11.4 μ M and 40.1 μ M, respectively).

Western Blot Analysis^[1]

Cell Line:	OCI-LY10 and TMD8 cells
Concentration:	0.3, 1, 3 and 10 μM
Incubation Time:	2 hours
Result:	Inhibited the phosphorylation of IRAK4 and the downstream molecules, IKK β and NF- κ B transcription factor (p65) in OCI-LY10 and TMD8 cells.

REFERENCES

1]. Yun Chen, et al. Design and synthesis of Imidazo[1,2-b]pyridazine IRAK4 inhibitors for the treatment of mutant MYD88 L265P diffuse large B-cell lymphoma. Eur J Med Chem. 2020 Mar 15;190:112092.						
	Caution: Product has n	ot been fully validated for me	edical applications. For research use only.			
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