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Product Data Sheet

PROTAC TYK2 degradation agent1

Cat. No.: HY-152227 CAS No.: 2921556-14-9 Molecular Formula: $C_{55}H_{69}N_{13}O_{7}S$ Molecular Weight: 1056.28 Target: JAK; PROTACs

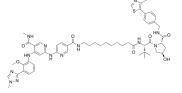
Pathway: Epigenetics; JAK/STAT Signaling; Protein Tyrosine Kinase/RTK; Stem Cell/Wnt;

PROTAC

Storage: -20°C, sealed storage, away from moisture and light

* In solvent: -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture

and light)



SOLVENT & SOLUBILITY

In Vitro

DMSO: 10 mg/mL (9.47 mM; ultrasonic and warming and heat to 60°C)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	0.9467 mL	4.7336 mL	9.4672 mL
	5 mM	0.1893 mL	0.9467 mL	1.8934 mL
	10 mM			

Please refer to the solubility information to select the appropriate solvent.

BIOLOGICAL ACTIVITY

Description PROTAC TYK2 degradation agent1 is a potent and subtype-selective TYK2 degrader. PROTAC TYK2 degradation agent1 has

autoimmune disease^[1].

DC50: 14 nM (TYK2)[1] IC₅₀ & Target

PROTAC TYK2 degradation agent1 (degrader 37) (1 µM; 6 h) selectively suppresses TYK2 protein levels^[1]. In Vitro

PROTAC TYK2 degradation agent1 has TYK2 degradation activity with DC₅₀ value of 14 nM^[1].

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

Western Blot Analysis^[1]

Cell Line:	Jurkat cells
Concentration:	1 μΜ

Incubation Time:	6 h
Result:	Suppressed TYK2 protein levels and effectively induced the degradation of TYK2 in a dose dependent manner at a low concentration range, with TYK2 degradation slightly decreasing at high concentrations.

REFERENCES

[1]. Jun-Ya Kato, et al. Discovery of a potent and subtype-selective TYK2 degrader based on an allosteric TYK2 inhibitor. Bioorg Med Chem Lett. 2023 Jan 1;79:129083.

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

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