

WDR5-IN-6

Cat. No.: HY-154949 CAS No.: 326901-92-2

Molecular Formula: $C_{13}H_8Cl_2N_2O_2S$

Molecular Weight: 327.19 Target: WDR5

Pathway: **Epigenetics**

Storage: Powder -20°C 3 years

> In solvent -80°C 6 months

> > -20°C 1 month

Product Data Sheet

SOLVENT & SOLUBILITY

In Vitro

DMSO: 100 mg/mL (305.63 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	3.0563 mL	15.2816 mL	30.5633 mL
	5 mM	0.6113 mL	3.0563 mL	6.1127 mL
	10 mM	0.3056 mL	1.5282 mL	3.0563 mL

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

In Vivo

- 1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: 2.5 mg/mL (7.64 mM); Clear solution; Need ultrasonic
- 2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: 2.5 mg/mL (7.64 mM); Clear solution; Need ultrasonic

BIOLOGICAL ACTIVITY

Description	WDR5-IN-6 is a WDR5 inhibitor, targeting to WBM site. WDR5-IN-6 inhibits cell proliferation of neuroblastoma cell lines with potent anti-tumor activity. WDR5-IN-6 shows high synergy with OICR-9429 (HY-16993), a WDR5 inhibitor targeting to WIN site. WDR5-IN-6 can be used for reasearch in neuroblastoma ^[1] .
IC ₅₀ & Target	WDR5 ^[1]
In Vitro	WDR5-IN-6 (compound 19) inhibits the proliferation of IMR32 and LAN5 cells with EC $_{50}$ values of 12.34 and 14.89 μ M, respectively. WDR5-IN-6 shows moderate inhibitory effect on SK-N-AS cells, without effect on HEK293T cells at 20 μ M $^{[1]}$. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES					
[1]. Han QL, et al. Discovery, evaluation and mechanism study of WDR5-targeted small molecular inhibitors for neuroblastoma. Acta Pharmacol Sin. 2023 Apr;44(4):877-887.					
	Caution: Product has r	not been fully validated for n	nedical applications. For research	use only.	
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