Proteins

MS645

Cat. No.: HY-125232 CAS No.: 2250091-96-2 Molecular Formula: $C_{48}H_{54}Cl_2N_{10}O_2S_2$

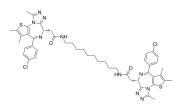
Molecular Weight: 938.04

Target: **Epigenetic Reader Domain**

Pathway: **Epigenetics**

Storage: -20°C, stored under nitrogen

* In solvent : -80°C, 6 months; -20°C, 1 month (stored under nitrogen)



Product Data Sheet

SOLVENT & SOLUBILITY

In	٧	it	ro

DMSO: 200 mg/mL (213.21 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	1.0661 mL	5.3303 mL	10.6605 mL
	5 mM	0.2132 mL	1.0661 mL	2.1321 mL
	10 mM	0.1066 mL	0.5330 mL	1.0661 mL

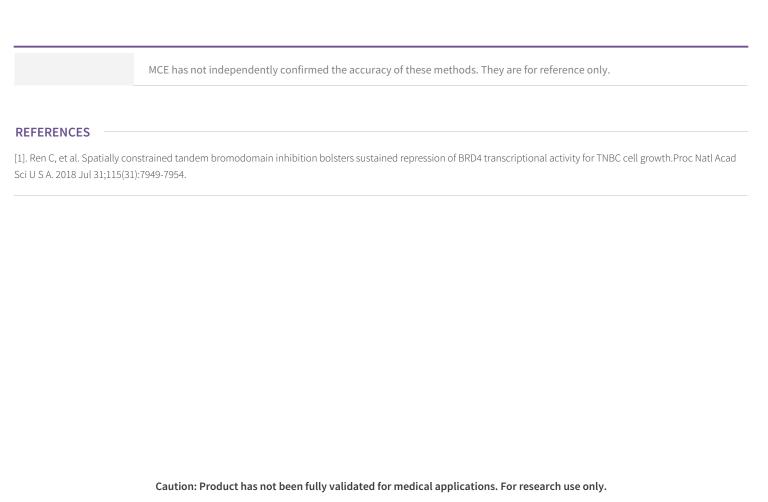
Please refer to the solubility information to select the appropriate solvent.

In Vivo

- 1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 5 mg/mL (5.33 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 5 mg/mL (5.33 mM); Clear solution

BIOLOGICAL ACTIVITY

Description	MS645 is a bivalent BET bromodomains (BrD) inhibitor with a K_i of 18.4 nM for BRD4-BD1/BD2. MS645 spatially constrains bivalent inhibition of BRD4 BrDs resulting in a sustained repression of BRD4 transcriptional activity in solid-tumor cells ^[1] .
IC ₅₀ & Target	BRD4(BD1BD2) 18.4 nM (Ki)
In Vitro	MS645 has cell growth inhibitory effects on noncancer cell lines of mouse macrophage RAW cells and nontumorigenic breast epithelial MCF10A with IC $_{50}$ s of 4.1 nM, 6.8 nM, 7.9 nM for triple-negative breast cancer (TNBC) cell lines HS5878T, BT549, and MCF $_{10A}^{[1]}$. ?MS645 (15, 30, 60 nM) results in a dramatic reduction of c-Myc expression and an increase of p21, a tumor suppressor and cell-cycle inhibitor in HCC1806 cells ^[1] .



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