Product Data Sheet

BRD 4354 ditrifluoroacetate

Molecular Weight: 610.93
Target: HDAC

Pathway: Cell Cycle/DNA Damage; Epigenetics

Storage: 4°C, sealed storage, away from moisture

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

SOLVENT & SOLUBILITY

In Vitro

DMSO: 125 mg/mL (204.61 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	1.6368 mL	8.1842 mL	16.3685 mL
	5 mM	0.3274 mL	1.6368 mL	3.2737 mL
	10 mM	0.1637 mL	0.8184 mL	1.6368 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: ≥ 2.08 mg/mL (3.40 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.08 mg/mL (3.40 mM); Clear solution
- 3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (3.40 mM); Clear solution

BIOLOGICAL ACTIVITY

Description BRD 4354 (ditrifluoroacetate) is a moderately potent inhibitor of HDAC5 and HDAC9, with IC₅₀s of 0.85 and 1.88 μ M, respectively^[1].

IC₅₀ & Target HDAC5 HDAC9 $0.85 \ \mu M \ (IC_{50}) \\ 1.88 \ \mu M \ (IC_{50})$

REFERENCES

1]. Boskovic ZV, ET AL. Inhibitic	on of Zinc-Dependent Histone Deacetylas	ses with a Chemically Trigg	ered Electrophile. ACS Chem Biol. 201	6 Jul 15;11(7):1844-51.
	Caution: Product has not been fully	y validated for medical	applications. For research use on	ly.
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