Proteins

Rifampicin-d₄

Cat. No.: HY-B0272S2 Molecular Formula: $C_{43}H_{54}D_4N_4O_{12}$

Molecular Weight: 826.96

Bacterial; Influenza Virus; Antibiotic Target:

Pathway: Anti-infection

Storage: Powder -20°C 3 years

In solvent

4°C 2 years -80°C 6 months

-20°C 1 month

Product Data Sheet

SOLVENT & SOLUBILITY

In Vitro DMF : ≥ 20 mg/mL (24.18 mM)

DMSO: $\geq 3.3 \text{ mg/mL} (3.99 \text{ mM})$

DMF:PBS(pH 7.2)(1:1) : \geq 0.5 mg/mL (0.60 mM)

Ethanol: ≥ 0.12 mg/mL (0.15 mM)

* "≥" means soluble, but saturation unknown.

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	1.2092 mL	6.0462 mL	12.0925 mL
	5 mM	0.2418 mL	1.2092 mL	2.4185 mL
	10 mM	0.1209 mL	0.6046 mL	1.2092 mL

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

BIOLOGICAL ACTIVITY

 $Rifampicin-d_4$ is the deuterium labeled Rifampicin. Rifampicin is a potent and broad spectrum antibiotic against bacterial Description pathogens. Rifampicin has anti-influenza virus activities.

In Vitro Stable heavy isotopes of hydrogen, carbon, and other elements have been incorporated into drug molecules, largely as tracers for quantitation during the drug development process. Deuteration has gained attention because of its potential to

affect the pharmacokinetic and metabolic profiles of drugs^[1].

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

REFERENCES

[1]. Russak EM, et al. Impact of Deuterium Substitution on the Pharmacokinetics of Pharmaceuticals. Ann Pharmacother. 2019;53(2):211-216.

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- [5]. Yu J, et al. Monitoring in vivo fitness of rifampicin-resistant Staphylococcus aureus mutants in a mouse biofilm infection model. J Antimicrob Chemother. 2005 Apr;55(4):528-34. Epub 2005 Mar 2.

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

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