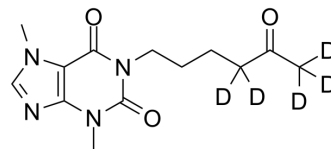


Pentoxifylline-d5

Cat. No.:	HY-B0715S2		
CAS No.:	1185995-18-9		
Molecular Formula:	C ₁₃ H ₁₃ D ₅ N ₄ O ₃		
Molecular Weight:	283.34		
Target:	Phosphodiesterase (PDE); Autophagy; HIV		
Pathway:	Metabolic Enzyme/Protease; Autophagy; Anti-infection		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



SOLVENT & SOLUBILITY

In Vitro

DMSO : ≥ 100 mg/mL (352.93 mM)
 * "≥" means soluble, but saturation unknown.

Concentration	Mass		
	1 mg	5 mg	10 mg
1 mM	3.5293 mL	17.6466 mL	35.2933 mL
5 mM	0.7059 mL	3.5293 mL	7.0587 mL
10 mM	0.3529 mL	1.7647 mL	3.5293 mL

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

BIOLOGICAL ACTIVITY

Description

Pentoxifylline-d5 (BL-191-d5) is the deuterium labeled Pentoxifylline. Pentoxifylline (BL-191), a haemorheological agent, is an orally active non-selective phosphodiesterase (PDE) inhibitor, with immune modulation, anti-inflammatory, hemorheological, anti-fibrinolytic and anti-proliferation effects. Pentoxifylline can be used for the research of peripheral vascular disease, cerebrovascular disease and a number of other conditions involving a defective regional microcirculation [1][2][3].

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

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

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