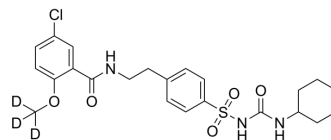


Glyburide-d₃

Cat. No.:	HY-15206S1		
CAS No.:	1219803-02-7		
Molecular Formula:	C ₂₃ H ₂₅ D ₃ ClN ₃ O ₅ S		
Molecular Weight:	497.02		
Target:	P-glycoprotein; Autophagy; Potassium Channel; CFTR; Mitochondrial Metabolism		
Pathway:	Membrane Transporter/Ion Channel; Autophagy; Metabolic Enzyme/Protease		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



BIOLOGICAL ACTIVITY

Description

Glyburide-d₃ is the deuterium labeled Glibenclamide. Glibenclamide (Glyburide) is an orally active ATP-sensitive K⁺ channel (KATP) inhibitor and can be used for the research of diabetes and obesity[1]. Glibenclamide inhibits P-glycoprotein. Glibenclamide directly binds and blocks the SUR1 subunits of KATP and inhibits the cystic fibrosis transmembrane conductance regulator protein (CFTR)[3]. Glibenclamide interferes with mitochondrial bioenergetics by inducing changes on membrane ion permeability[4]. Glibenclamide can induce autophagy[5].

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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- [2]. Fernandes MA, et al. Glibenclamide interferes with mitochondrial bioenergetics by inducing changes on membrane ion permeability. *J Biochem Mol Toxicol.* 2004;18(3):162-169.
- [3]. Heo R, et al. The anti-diabetic drug trelagliptin induces vasodilation via activation of Kv channels and SERCA pumps. *Life Sci.* 2021;283:119868.
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Caution: Product has not been fully validated for medical applications. For research use only.

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