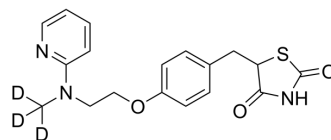


Rosiglitazone-d₃

Cat. No.:	HY-17386S
CAS No.:	1132641-22-5
Molecular Formula:	C ₁₈ H ₁₆ D ₃ N ₃ O ₃ S
Molecular Weight:	360.45
Target:	PPAR; TRP Channel; Autophagy; Ferroptosis; Isotope-Labeled Compounds
Pathway:	Cell Cycle/DNA Damage; Vitamin D Related/Nuclear Receptor; Membrane Transporter/Ion Channel; Neuronal Signaling; Autophagy; Apoptosis; Others
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Rosiglitazone-d ₃ is the deuterium labeled Rosiglitazone. Rosiglitazone (BRL 49653) is a selective, orally active PPAR γ agonist with EC50s of 30 nM, 100 nM and 60 nM for PPAR γ 1, PPAR γ 2, and PPAR γ , respectively. Rosiglitazone binds to PPAR γ with a Kd of approximately 40 nM. Rosiglitazone is also an activator of TRPC5 (EC50 \approx 30 μ M) and an inhibitor of TRPM3[1][2][3][4].
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.

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA