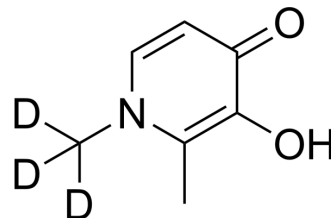


Deferiprone-d3

Cat. No.:	HY-B0568S
CAS No.:	1346601-82-8
Molecular Formula:	C ₇ H ₆ D ₃ NO ₂
Molecular Weight:	142.17
Target:	HCV; Ferroptosis; Isotope-Labeled Compounds
Pathway:	Anti-infection; Apoptosis; Others
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Deferiprone-d ₃ is the deuterium labeled Deferiprone. Deferiprone is the only orally active iron-chelating agent to be used therapeutically in conditions of transfusional iron overload[1][2].
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.

[2]. Kontoghiorghes GJ, et al. Benefits and risks of deferiprone in iron overload in Thalassemia and other conditions: comparison of epidemiological and therapeutic aspects with deferoxamine. *Drug Saf*. 2003;26(8):553-584.

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

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