## **Product** Data Sheet

## Indinavir-d<sub>6</sub>

Molecular Weight:

 $\begin{array}{lll} \textbf{Cat. No.:} & \textbf{HY-B0689S} \\ \textbf{CAS No.:} & 185897-02-3 \\ \textbf{Molecular Formula:} & \textbf{C}_{36}\textbf{H}_{41}\textbf{D}_{6}\textbf{N}_{5}\textbf{O}_{4} \\ \end{array}$ 

Target: HIV Protease; HIV; Isotope-Labeled Compounds

Pathway: Anti-infection; Metabolic Enzyme/Protease; Others

Storage: Please store the product under the recommended conditions in the Certificate of

Analysis.

619.83

## **BIOLOGICAL ACTIVITY**

Description	Indinavir- $d_6$ is the deuterium labeled Indinavir. Indinavir (MK-639; L735524) is a potent and specific HIV protease inhibitor that appears to have good oral bioavailability.
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 <sup>[1]</sup> .  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]. Stein, D.S., et al., A 24-week open-label phase I/II evaluation of the HIV protease inhibitor MK-639 (indinavir). AIDS, 1996. 10(5): p. 485-92.

[3]. Liu, F., et al., Kinetic, stability, and structural changes in high-resolution crystal structures of HIV-1 protease with drug-resistant mutations L24I, I50V, and G73S. J Mol Biol, 2005. 354(4): p. 789-800.

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

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