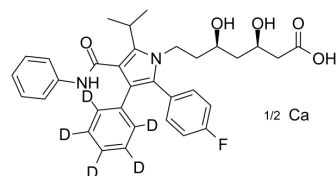


## Atorvastatin-d5 hemicalcium

<b>Cat. No.:</b>	HY-B0589S
<b>CAS No.:</b>	222412-82-0
<b>Molecular Formula:</b>	C <sub>33</sub> H <sub>30</sub> D <sub>5</sub> FN <sub>2</sub> O <sub>5</sub> ·1/2Ca
<b>Molecular Weight:</b>	583.71
<b>Target:</b>	HMG-CoA Reductase (HMGCR); Autophagy
<b>Pathway:</b>	Metabolic Enzyme/Protease; Autophagy
<b>Storage:</b>	4°C, sealed storage, away from moisture * In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture)



### BIOLOGICAL ACTIVITY

<b>Description</b>	Atorvastatin-d5 hemicalcium is the deuterium labeled Atorvastatin. Atorvastatin hemicalcium is an orally active HMG-CoA reductase inhibitor, has the ability to effectively decrease blood lipids. Atorvastatin hemicalcium inhibits human SV-SMC proliferation and invasion with IC <sub>50</sub> s of 0.39 μM and 2.39 μM, respectively <sup>[1][2][3]</sup> .
<b>In Vitro</b>	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

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- [7]. Ming-Bai Hu, et al. Atorvastatin induces autophagy in MDA-MB-231 breast cancer cells. *Ultrastruct Pathol.* Sep-Oct 2018;42(5):409-415.

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

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