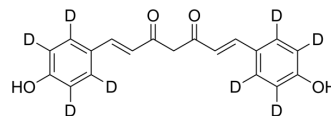


## Bisdemethoxycurcumin-d<sub>8</sub>

<b>Cat. No.:</b>	HY-N0007S		
<b>CAS No.:</b>	2470233-08-8		
<b>Molecular Formula:</b>	C <sub>19</sub> H <sub>8</sub> D <sub>8</sub> O <sub>4</sub>		
<b>Molecular Weight:</b>	316.38		
<b>Target:</b>	Apoptosis; Autophagy		
<b>Pathway:</b>	Apoptosis; Autophagy		
<b>Storage:</b>	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



### BIOLOGICAL ACTIVITY

<b>Description</b>	Bisdemethoxycurcumin-d <sub>8</sub> is the deuterium labeled Bisdemethoxycurcumin. Bisdemethoxycurcumin (Curcumin III; Didemethoxycurcumin) is a natural derivative of curcumin with anti-inflammatory and anti-cancer activities.
<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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- [2]. Lee PJ, et al. Bisdemethoxycurcumin Induces Apoptosis in Activated Hepatic Stellate Cells via Cannabinoid Receptor 2. *Molecules.* 2015 Jan 14;20(1):1277-92.
- [3]. Chen J, et al. Natural borneol enhances bisdemethoxycurcumin-induced cell cycle arrest in the G2/M phase through up-regulation of intracellular ROS in HepG2 cells. *Food Funct.* 2014 Dec 24.
- [4]. Luo C, et al. Bisdemethoxycurcumin attenuates gastric adenocarcinoma growth by inducing mitochondrial dysfunction. *Oncol Lett.* 2015 Jan;9(1):270-274.
- [5]. Li YB, et al. Bisdemethoxycurcumin Increases Sirt1 to Antagonize t-BHP-Induced Premature Senescence in WI38 Fibroblast Cells. *Evid Based Complement Alternat Med.* 2013;2013:851714.

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

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