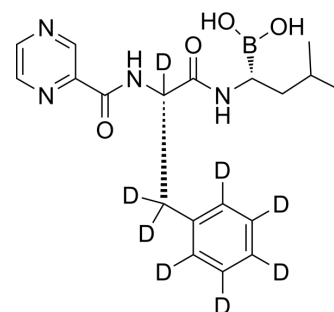


## Bortezomib-d<sub>8</sub>

<b>Cat. No.:</b>	HY-10227S
<b>Molecular Formula:</b>	C <sub>19</sub> H <sub>17</sub> D <sub>8</sub> BN <sub>4</sub> O <sub>4</sub>
<b>Molecular Weight:</b>	392.29
<b>Target:</b>	Proteasome; Apoptosis; Autophagy; NF-κB
<b>Pathway:</b>	Metabolic Enzyme/Protease; Apoptosis; Autophagy; NF-κB
<b>Storage:</b>	-20°C, stored under nitrogen * In solvent : -80°C, 6 months; -20°C, 1 month (stored under nitrogen)



### BIOLOGICAL ACTIVITY

<b>Description</b>	Bortezomib-d <sub>8</sub> is the deuterium labeled Bortezomib. Bortezomib (PS-341) is a reversible and selective proteasome inhibitor, and potently inhibits 20S proteasome (K <sub>i</sub> =0.6 nM) by targeting a threonine residue. Bortezomib disrupts the cell cycle, induces apoptosis, and inhibits NF-κB. Bortezomib is the first proteasome inhibitor anticancer agent. Anti-cancer activity[1][2].
<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]. Adams J, et al. Proteasome inhibitors: a novel class of potent and effective antitumor agents. *Cancer Res.* 1999 Jun 1;59(11):2615-22.
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- [7]. Fernández Y, et al. Chemical blockage of the proteasome inhibitory function of bortezomib: impact on tumor cell death. *J Biol Chem.* 2006 Jan 13;281(2):1107-18.

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

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