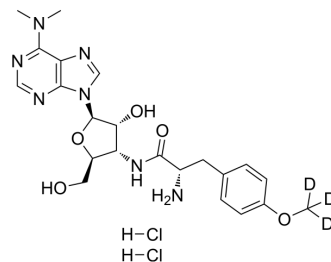


## Puromycin-d<sub>3</sub> dihydrochloride

<b>Cat. No.:</b>	HY-B1743AS
<b>Molecular Formula:</b>	C <sub>22</sub> H <sub>28</sub> D <sub>3</sub> Cl <sub>2</sub> N <sub>7</sub> O <sub>5</sub>
<b>Molecular Weight:</b>	547.45
<b>Target:</b>	Bacterial; Antibiotic; Isotope-Labeled Compounds
<b>Pathway:</b>	Anti-infection; Others
<b>Storage:</b>	-20°C, protect from light, stored under nitrogen * In solvent : -80°C, 6 months; -20°C, 1 month (protect from light, stored under nitrogen)



### BIOLOGICAL ACTIVITY

<b>Description</b>	Puromycin-d <sub>3</sub> (dihydrochloride) is the deuterium labeled Puromycin dihydrochloride. Puromycin dihydrochloride (CL13900 dihydrochloride), an aminoglycoside antibiotic, inhibits protein synthesis[1].
<b>IC<sub>50</sub> &amp; Target</b>	Aminoglycoside
<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

- [1]. Russak EM, et al. Impact of Deuterium Substitution on the Pharmacokinetics of Pharmaceuticals. *Ann Pharmacother.* 2019;53(2):211-216.
- [2]. Nathans D, et al. Puromycin inhibition of protein synthesis: incorporation of puromycin into peptide chains. *Proc Natl Acad Sci U S A.* 1964 Apr;51:585-92.
- [3]. Miyamoto-Sato E, et al. Specific bonding of puromycin to full-length protein at the C-terminus. *Nucleic Acids Res.* 2000 Mar 1;28(5):1176-82.
- [4]. Schmidt EK, et al. SUNSET, a nonradioactive method to monitor protein synthesis. *Nat Methods.* 2009 Apr;6(4):275-7.

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

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