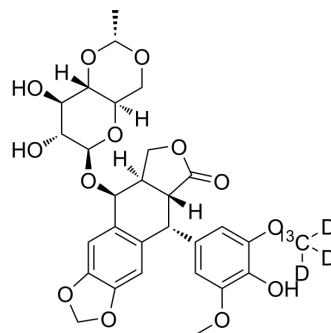


## Etoposide-<sup>13</sup>C,<sub>3</sub>D<sub>3</sub>

<b>Cat. No.:</b>	HY-13629S1
<b>Molecular Formula:</b>	C <sub>28</sub> <sup>13</sup> CH <sub>29</sub> D <sub>3</sub> O <sub>13</sub>
<b>Molecular Weight:</b>	592.57
<b>Target:</b>	Apoptosis; Topoisomerase; Bacterial; Autophagy; Mitophagy; Antibiotic; Isotope-Labeled Compounds
<b>Pathway:</b>	Apoptosis; Cell Cycle/DNA Damage; Anti-infection; Autophagy; Others
<b>Storage:</b>	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	Etoposide- <sup>13</sup> C, <sub>3</sub> D <sub>3</sub> is the <sup>13</sup> C- and deuterium labeled Etoposide. Etoposide (VP-16; VP-16-213) is an anti-cancer chemotherapy agent. Etoposide inhibits topoisomerase II, thus stopping DNA replication. Etoposide induces cell cycle arrest, apoptosis and autophagy[1].
<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>[65]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

### REFERENCES

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- [4]. Fuchs, J., et al. Comparative activity of NSC 119875, NSC 109724, NSC 123127, NSC 241240, and etoposide in heterotransplanted hepatoblastoma. *Cancer*, 1998. 83(11): p. 2400-7.
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**Caution: Product has not been fully validated for medical applications. For research use only.**

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