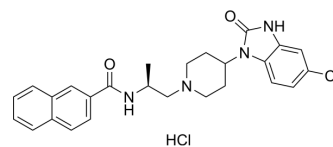


## VU0155069 hydrochloride

<b>Cat. No.:</b>	HY-108612A
<b>CAS No.:</b>	1781834-89-6
<b>Molecular Formula:</b>	C <sub>26</sub> H <sub>28</sub> Cl <sub>2</sub> N <sub>4</sub> O <sub>2</sub>
<b>Molecular Weight:</b>	499.43
<b>Target:</b>	Phospholipase
<b>Pathway:</b>	Metabolic Enzyme/Protease
<b>Storage:</b>	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	VU0155069 hydrochloride (CAY10593 hydrochloride) is a potent selective phospholipase D (PLD) inhibitor. The IC <sub>50</sub> values for PLD1 and PLD2 are 46 and 933 nM, respectively. VU0155069 hydrochloride inhibits migration of human and mouse breast cancer cell lines <sup>[1][2]</sup> .
<b>IC<sub>50</sub> &amp; Target</b>	PLD1 46 nM (IC <sub>50</sub> )
<b>In Vitro</b>	VU0155069 (0.5 μM, 1 h) significantly inhibits (R)-DOI (3 μM)-induced [ <sup>3</sup> H]PtdBut production in MCF-7 cells <sup>[3]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

### REFERENCES

- [1]. Scott SA, et al. Design of isoform-selective phospholipase D inhibitors that modulate cancer cell invasiveness. *Nat Chem Biol.* 2009 Feb;5(2):108-17.
- [2]. Lewis JA, et al. Design and synthesis of isoform-selective phospholipase D (PLD) inhibitors. Part I: Impact of alternative halogenated privileged structures for PLD1 specificity. *Bioorg Med Chem Lett.* 2009 Apr 1;19(7):1916-20.
- [3]. Barclay Z, et al. Attenuated PLD1 association and signalling at the H452Y polymorphic form of the 5-HT(2A) receptor. *Cell Signal.* 2013 Apr;25(4):814-21.

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

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