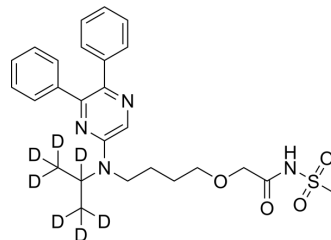


## Selexipag-d<sub>7</sub>

<b>Cat. No.:</b>	HY-14870S1		
<b>CAS No.:</b>	1265295-21-3		
<b>Molecular Formula:</b>	C <sub>26</sub> H <sub>25</sub> D <sub>7</sub> N <sub>4</sub> O <sub>4</sub> S		
<b>Molecular Weight:</b>	503.66		
<b>Target:</b>	Prostaglandin Receptor; Isotope-Labeled Compounds		
<b>Pathway:</b>	GPCR/G Protein; Others		
<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>	Selexipag-d <sub>7</sub> is the deuterium labeled Selexipag. Selexipag (NS-304) is an orally available and potent agonist for the Prostacyclin (PGI <sub>2</sub> ) receptor (IP receptor)[1][2].
<b>IC<sub>50</sub> &amp; Target</b>	IP
<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]. Kuwano K, et al. 2-[4-[(5,6-diphenylpyrazin-2-yl)(isopropyl)amino]butoxy]-N-(methylsulfonyl)acetamide (NS-304), an orally available and long-acting prostacyclin receptor agonist prodrug. *J Pharmacol Exp Ther*. 2007 Sep;322(3):1181-8.
- [3]. Mous DS, et al. Treatment of rat congenital diaphragmatic hernia with sildenafil and NS-304, selexipag's active compound, at the pseudoglandular stage improves lung vasculature. *Am J Physiol Lung Cell Mol Physiol*. 2018 May 10.

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

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