



## PDE2 / PDE10-IN-1

Catalog No: tcsc0035357

Availa	able Sizes		
Size: 5mg			
Size: 10mg			
Size: 25mg			
Specif	fications		
<b>CAS No:</b> 1426833-08-0			
Formula: C <sub>15</sub> H <sub>10</sub> CIN <sub>5</sub>			
<b>Pathway:</b> Metabolic Enzy	yme/Protease		
<b>Target:</b> Phosphodieste	erase (PDE)		
Purity / Grade >98%	le:		
<b>Solubility:</b> 10 mM in DMS	5O		
Ohserved Mo	decular Weight:		

## **Product Description**

295.73

PDE2/PDE10-IN-1 is a phosphodiesterase 2 (**PDE2**) and **PDE10** inhibitor with  $IC_{50}$ s of 29 and 480 nM, respectively.

IC50 & Target: IC50: 29 nM (hPDE2A), 480 nM (rPDE10A), 5890 nM (hPDE4D), 6920 nM (hPDE11A), >10000 nM (hPDE1A, hPDE3B,



hPDE5A, hPDE6AB, hPDE7A and hPDE9A)[1]

In Vitro: PDE2/PDE10-IN-1 (Compound 6) inhibits PDE2 and PDE10, respectively, with an IC $_{50}$  value of 29 and 480 nM. PDE2/PDE10-IN-1 also inhibits PDE11A and PDE4D with IC $_{50}$ s of 6920 nM and 5890 nM, respectively. In addition PDE2/PDE10-IN-1 does not show significant inhibition of a panel of CYP450 enzymes (CYP1A2, 2C9, 2D6, 2C19, and 3A4). PDE2/PDE10-IN-1 is also inactive up to a concentration of 125  $\mu$ g/mL in a bacterial mutagenicity assay<sup>[1]</sup>.

In Vivo: The PK properties of PDE2/PDE10-IN-1 are studied in rats after 2.5 mg/kg i.v. and 10 mg/kg p.o. administration. After i.v. administration, a rapid clearance is observed ( $t_{1/2}$ =0.47 h), which is not expected based on the in vitro metabolic stability in rat liver microsomes (rLMs). Interestingly, PDE2/PDE10-IN-1 shows much slower clearance after p.o. administration ( $t_{1/2}$ =2.36 h), resulting in good bioavailability and a maximum plasma concentration ( $t_{max}$ ) of 997 ng/mL. PDE2/PDE10-IN-1 is assessed for its potential to cross the blood-brain barrier in rats after 10 mg/kg s.c. administration. PDE2/PDE10-IN-1 shows good formulatability with 10 to 20% HP $\beta$ CD at pH>3.5. The brain concentration for PDE2/PDE10-IN-1 after 1 h administration is in the range of 370-895 ng/g with high brain free fractions and brain/plasma ratios. More specifically, PDE2/PDE10-IN-1, which is orally bioavailable, occupies PDE2 with an ED<sub>50</sub> of 21 mg/kg<sup>[1]</sup>.

All products are for RESEARCH USE ONLY. Not for diagnostic & therapeutic purposes!