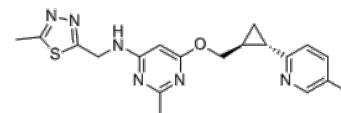


**Product Name** : MK-8189  
**Cat. No.** : PC-49494  
**CAS No.** : 1424371-93-6  
**Molecular Formula** : C<sub>19</sub>H<sub>22</sub>N<sub>6</sub>OS  
**Molecular Weight** : 382.49  
**Target** : Phosphodiesterase (PDE)  
**Solubility** : 10 mM in DMSO



## Biological Activity

MK-8189 (MK8189) is a highly potent, selective **PDE10A** inhibitor with K<sub>i</sub> of 29 pM, 500,000-fold selectivity over the other PDE enzyme families (PDE1-PDE11).

MK-8189 shows an IC<sub>50</sub> of 1.6 nM in cells recombinantly expressing full-length human PDE10A.

MK-8189 has an excellent profile against ion channels (I<sub>ks</sub>, Cav1.2, and Nav1.5 >30 μM, and functional hERG I<sub>kr</sub> IC<sub>50</sub>=33 μM). One off-target activity was identified in a broad Panlabs panel (somatostatin receptor type 2 (SSTR2) IC<sub>50</sub>=2.8 μM in a radioligand binding assay).

MK-8189 (0.25 mg/kg, oral) attenuates MK-801-induced locomotor activity in rats.

## References

Layton ME, et al. *J Med Chem.* 2023 Jan 9. doi: 10.1021/acs.jmedchem.2c01521.

**Caution: Product has not been fully validated for medical applications. Lab Use Only!**

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