

: NLX-101

**Molecular Formula :** C<sub>19</sub>H<sub>21</sub>ClF<sub>2</sub>N<sub>4</sub>O

Molecular Weight : 394.85

: PC-73308

: 635323-95-4

: 5-HT Receptor

: 10 mM in DMSO

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Product Name

Cat. No.

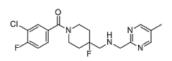
CAS No.

Target

Solubility

## **Data Sheet**

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## **Biological Activity**

NLX-101 (F15599) is a potent, highly selective post-synaptic 5-HT1A receptor agonist with Ki of 3.4 nM.

NLX-101 (F15599) is over 1000-fold selective with respect to the other 5-HT1 receptor subtypes, 5-HT2-7 receptor families, and also numerous GPCRs, transporters, ion channels, and enzymes.

NLX-101 (F15599) activates h5-HT1A receptors with an efficacy superior to that of the prototypical 5-HT1A agonist (+/-)-8-OH-DPAT in cellular model of signal transduction.

NLX-101 (F15599), unlike 5-HT, more potently and efficaciously stimulated G( $\alpha$ i) than G( $\alpha$ o) activation.

NLX-101 (F15599) potently activated ERK1/2 phosphorylation and strongly induced c-fos mRNA expression in rat prefrontal cortex.

NLX-101 (F15599) attenuates muscular hyperalgesia in reserpine-induced myalgia model.

## References

Maurel JL, et al. J Med Chem. 2007 Oct 4;50(20):5024-33.

Newman-Tancredi A, et al. Br J Pharmacol. 2009 Jan;156(2):338-53.

Assié MB, et al. Int J Neuropsychopharmacol. 2010 Nov;13(10):1285-98.

Caution: Product has not been fully validated for medical applications. Lab Use Only! E-mail: tech@probechem.com