

## CyPPA

## Chemical Properties

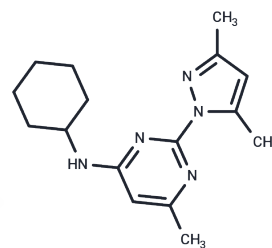
CAS No. : 73029-73-9

Formula: C<sub>16</sub>H<sub>23</sub>N<sub>5</sub>

Molecular Weight: 285.39

Storage: Powder: -20°C for 3 years | In solvent: -80°C for 1 year

Actual storage temperature shall be subject to the COA.



## Biological Description

Description	CyPPA, a subtype-selective positive modulator of SK channels, decreased spontaneous firing rate, increased the duration of the apamin-sensitive afterhyperpolarization.
Targets(IC50)	Dopamine Receptor,Potassium Channel
In vitro	CyPPA, a positive modulator of SK2 and SK3, reduces the activity of dopaminergic neurons and inhibits dopamine release with EC50 values of 14 $\mu$ M and 5.6 $\mu$ M, respectively. CyPPA counteracts hyperdopaminergic behaviors induced by methylphenidate.
In vivo	In vivo studies revealed that systemic administration of CyPPA attenuated methylphenidate-induced hyperactivity and stereotypic behaviors in mice. CyPPA induces a concentration-dependent increase in the apparent Ca <sup>2+</sup> sensitivity of channel activation, changing the EC50 (Ca <sup>2+</sup> ) from 429 nM to 59 nM

## Solubility Information

Solubility	DMSO: 99 mg/mL (346.89 mM),Sonication is recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble)
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### Preparing Stock Solutions

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	1mg	5mg	10mg
1 mM	3.504 mL	17.5199 mL	35.0398 mL
5 mM	0.7008 mL	3.504 mL	7.008 mL
10 mM	0.3504 mL	1.752 mL	3.504 mL
50 mM	0.0701 mL	0.3504 mL	0.7008 mL

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Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. Please use it as soon as possible.

Note: The dilution table applies only to solid products. For liquid products, please calculate the stock solution based on the stated concentration and/or density.

### Reference

Herrick KF, Redrobe JP, Holst D, Hougaard C, Sandager-Nielsen K, Nielsen AN, Ji H, Holst NM, Rasmussen HB, Nielsen EØ, Strøbæk D, Shepard PD, Christophersen P. CyPPA, a Positive SK3/SK2 Modulator, Reduces Activity of Dopaminergic Neurons, Inhibits Dopamine Release, and Counteracts Hyperdopaminergic Behaviors Induced by Methylphenidate. *Front Pharmacol.* 2012 Feb 13;3:11. doi: 10.3389/fphar.2012.00011. eCollection 2012. PubMed PMID: 22347859; PubMed Central PMCID: PMC3278045.

Dolga AM, Letsche T, Gold M, Doti N, Bacher M, Chiamvimonvat N, Dodel R, Culmsee C. Activation of KCNN3/SK3/K(Ca)2.3 channels attenuates enhanced calcium influx and inflammatory cytokine production in activated microglia. *Glia.* 2012 Dec;60(12):2050-64. doi: 10.1002/glia.22419. Epub 2012 Sep 21. PubMed PMID: 23002008; PubMed Central PMCID: PMC3799773.

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