PKG inhibitor peptide

| Cat. No.: | HY-P1292 | | | | | |
|----------------------|-----------------------------------------------------------------|-------|---------|---------|--|--|
| CAS No.: | 82801-73-8 | | | | | |
| Molecular Formula: | C ₃₈ H ₇₄ N ₁₈ O ₁₀ | | | | | |
| Molecular Weight: | 943.11 | | | | | |
| Sequence Shortening: | RKRARKE | | | | | |
| Target: | Others | | | H₂N∕́NH | | |
| Pathway: | Others | | | | | |
| Storage: | Sealed storage, away from moisture | | | | | |
| | Powder | -80°C | 2 years | | | |
| | | -20°C | 1 year | | | |
| | * In solvent : | | | | | |

SOLVENT & SOLUBILITY

| In Vitro | H ₂ O : 100 mg/mL (106.03 mM; Need ultrasonic) | | | | | | | |
|----------|-------------------------------------------------------------------------------|--------------------------------------------------------|---------------|-----------|------------|--|--|--|
| | Preparing Stock Solutions | Mass Solvent Concentration | 1 mg | 5 mg | 10 mg | | | |
| | | 1 mM | 1.0603 mL | 5.3016 mL | 10.6032 mL | | | |
| | | 5 mM | 0.2121 mL | 1.0603 mL | 2.1206 mL | | | |
| | 10 mM | 0.1060 mL | 0.5302 mL | 1.0603 mL | | | | |
| | Please refer to the solubility information to select the appropriate solvent. | | | | | | | |
| In Vivo | 1. Add each solvent Solubility: 100 mg | one by one: PBS /mL (106.03 mM); Clear solution; Ne | ed ultrasonic | | | | | |

| Description | PKG inhibitor peptide is an ATP-competitive inhibitor of cGMP-dependent protein kinase (PKG), with a K _i of 86 μ M ^[1] . | | | | |
|---------------------------|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|--|--|--|--|
| IC ₅₀ & Target | Ki: 86 μM (PKG) ^[1] | | | | |
| In Vitro | Summary of experiments showing that intracellular dialysis of postsynaptic cells with PKG inhibitor PKG inhibitor peptide (1 mM) failed to alter the induction of long-term depression (CCh-LTD) ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. | | | | |

REFERENCES

Product Data Sheet



[1]. Bhatnagar D, et al. Synthetic peptide analogues differentially alter the binding affinities of cyclic nucleotide dependent protein kinases for nucleotide substrates. Biochemistry. 1988 Mar 22;27(6):1988-94.

[2]. Chiung-Chun Huang, et al. Activation of muscarinic acetylcholine receptors induces a nitric oxide-dependent long-term depression in rat medial prefrontal cortex. Cereb Cortex. 2010 Apr;20(4):982-96.

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

 Tel: 609-228-6898
 Fax: 609-228-5909
 E-mail: tech@MedChemExpress.com

 Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA