

## PKI(5-22)amide

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|----------------------|---|
| Cat. No.:            | HY-P3785  |
| CAS No.:             | 100853-58-5   |
| Molecular Formula:   | C <sub>84</sub> H <sub>137</sub> N <sub>29</sub> O <sub>26</sub>                          |
| Molecular Weight:    | 1969.17   |
| Sequence Shortening: | TTYADFIASGRTGRRNAI-NH2  |
| Target:              | PKA   |
| Pathway:             | Stem Cell/Wnt   |
| Storage:             | Please store the product under the recommended conditions in the Certificate of Analysis. |

### BIOLOGICAL ACTIVITY

|             |  |
|-------------|--|
| Description | PKI(5-22)amide is the active inhibitory fragment of the inhibitor of the cyclic AMP-dependent protein kinase (PKA). PKI(5-22)amide inhibits PKA activation, but fails to attenuate homologous desensitization of CRF1 receptors <sup>[1][2]</sup> .  |
| In Vitro    | PKI(5-22)amide (1 μM; 1 min) fails to block CRF1 receptor (100 nM; 30 min) desensitization in Y-79 cells permeabilized with <a href="#">Streptolysin O</a> (HY-135416) <sup>[1]</sup> .<br>MCE has not independently confirmed the accuracy of these methods. They are for reference only. |

### REFERENCES

[1]. Dautzenberg FM, et al. GRK3 mediates desensitization of CRF1 receptors: a potential mechanism regulating stress adaptation. *Am J Physiol Regul Integr Comp Physiol*. 2001 Apr;280(4):R935-46.

[2]. Reed J, et al. Conformational analysis of PKI(5-22)amide, the active inhibitory fragment of the inhibitor protein of the cyclic AMP-dependent protein kinase. *Biochem J*. 1989 Dec 1;264(2):371-80.

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

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