

Maurocalcine

Cat. No.:	HY-P5114
CAS No.:	269745-22-4
Molecular Formula:	C ₁₅₆ H ₂₇₀ N ₅₆ O ₄₆ S ₆
Molecular Weight:	3858.55
Sequence:	Gly-Asp-Cys-Leu-Pro-His-Leu-Lys-Leu-Cys-Lys-Glu-Asn-Lys-Asp-Cys-Cys-Ser-Lys-Lys-Cys-Lys-Arg-Arg-Gly-Thr-Asn-Ile-Glu-Lys-Arg-Cys-Arg (Disulfide bridge:Cys3-Cys17;Cys10-Cys21;Cys16-Cys32)
Sequence Shortening:	GDCLPHLKLCKENKDCCKKCKRRGTNIEKRRCR (Disulfide bridge:Cys3-Cys17;Cys10-Cys21;Cys16-Cys32)
Target:	Calcium Channel
Pathway:	Membrane Transporter/Ion Channel; Neuronal Signaling
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.

BIOLOGICAL ACTIVITY

Description	Maurocalcine is an agonist of ryanodine receptor (RyR) channel types 1, 2 and 3 with cellular permeability. Maurocalcine induces [³ H]ryanodine binding on RyR1 with an EC ₅₀ value of 2558 nM. Maurocalcine exhibits a apparent affinity of 14 nM for RyR2. Maurocalcine can be applied to in vivo cell tracking or other cell imaging techniques ^{[1][2][3]} .								
In Vitro	<p>Maurocalcine (100 nM, 24 h) increases the sensitivity of RyR2 to activating [Ca²⁺]_i and decrease its sensitivity to inhibiting [Ca²⁺]_i^[1].</p> <p>Maurocalcine (5 μM, 4 h/24 h) is absolutely no sign of significant cell toxicity for HEK293^[3].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Viability Assay^[3]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>HEK293</td> </tr> <tr> <td>Concentration:</td> <td>5, 10 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>4, 24 h</td> </tr> <tr> <td>Result:</td> <td>Had absolutely no sign of significant cell toxicity for concentrations up to 5 μM whether the incubation time lasted 4 h or 24 h. Exhibited only 8.0% cell toxicity for concentration of 10 μM and a 24 h incubation time.</td> </tr> </table>	Cell Line:	HEK293	Concentration:	5, 10 μM	Incubation Time:	4, 24 h	Result:	Had absolutely no sign of significant cell toxicity for concentrations up to 5 μM whether the incubation time lasted 4 h or 24 h. Exhibited only 8.0% cell toxicity for concentration of 10 μM and a 24 h incubation time.
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REFERENCES

- [1]. De Waard S, et al. Maurocalcine and its analog MCE12A facilitate Ca²⁺ mobilization in cardiomyocytes. *Biochem J.* 2020 Oct 30;477(20):3985-3999.
- [2]. Altafaj X, et al. Maurocalcine interacts with the cardiac ryanodine receptor without inducing channel modification. *Biochem J.* 2007 Sep 1;406(2):309-15.
- [3]. Boisseau S, et al. Cell penetration properties of maurocalcine, a natural venom peptide active on the intracellular ryanodine receptor. *Biochim Biophys Acta.* 2006

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

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