

Product Data Sheet

Inhibitors

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Glucagon (22-29)

Cat. No.: HY-P3617

CAS No.: 32204-93-6

Molecular Formula: $C_{_{49}}H_{_{71}}N_{_{11}}O_{_{12}}S$ Molecular Weight: 1038.22

Sequence Shortening: FVQWLMNT

Target: Others
Pathway: Others

Storage: Please store the product under the recommended conditions in the Certificate of

Analysis.

BIOLOGICAL ACTIVITY

Description	Glucagon (22-29) is partial agonist of Glucagon (19–29). Glucagon specifically inhibits the Ca^{2+} pump in liver plasma membranes independently of adenylate cyclase activation ^{[1][2]} .
In Vitro	Glucagon (22-29) produces only a 5-15% maximal inhibition of (Ca ²⁺ -Mg ²⁺) ATPase, with a low potency (K _i =1 μM), similar to that of native glucagon ^[1] . Glucagon (22-29) (10 nM) evokes an early small decrease of cell contraction when added alone and an 18% significant positive inotropic effect when added in combination with 30 nM glucagon. Glucagon (22-29) thus appears to act as a partial agonist of miniglucagon ^[2] . Glucagon (22-29) consists of the COOH-terminal part of miniglucagon but lacks the three Ala1g-Gln20-Asp21 residues ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

 $[1]. \ Mallat\ A, et\ al.\ A\ glucagon\ fragment\ is\ responsible\ for\ the\ inhibition\ of\ the\ liver\ Ca2+\ pump\ by\ glucagon.\ Nature.\ 1987\ Feb\ 12-18;325(6105):620-2.$

[2]. Pavoine C, et al. Miniglucagon [glucagon-(19-29)] is a component of the positive inotropic effect of glucagon. Am J Physiol. 1991 May;260(5 Pt 1):C993-9.

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

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