

Effect of pentagastrin on IL-1 β induced inhibition of insulin secretion in neonatal rat islets of Langerhans

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Abstract

AIM: To observe the effect of pentagastrin (G-5) on IL-1 β induced inhibition of insulin secretion in newborn rat islet of Langerhans.

METHODS: Islets of Langerhans of 3 to 5 day old rats were isolated by collagenase digestion. The islets were maintained free floating in culture medium RPMI-1640, containing 10% (V/V) calf serum, and distributed randomly in 96-well plastic plates (6 wells in each group). There are 15 islets per well in 0.2 mL culture medium. The islets were kept at 37 °C in mixed gases of 5% CO₂ and 95% humidified air for the time required by the experimental design. Three experiments were performed in this study. (1) IL-1 β induced inhibition of insulin secretion in isolated islets of Langerhans. (2) Effect of G-5 on IL-1 β induced inhibition of insulin secretion. And (3) Effect of G-5 on the functional repair of islet B-cells inhibited by

IL-1 β . Accumulated and glucose stimulated insulin secretion was measured by radioimmunoassay in all studies. Data are presented as $\bar{x} \pm s$. Differences between groups were analyzed using the Student's *t* test. *P* < 0.05 was considered significant.

RESULTS: The function of islet B cells, which has been received IL-1 β treated for 24 h, was dose-dependently inhibited. The accumulated and glucose stimulated insulin secretion was significantly lower than that of the control group (*P* < 0.05). The inhibitory effect of IL-1 β on islet B cells can be partially reversed by G 5. Accumulated and stimulated insulin secretion of G-5 0.6 ng/mL and 0.8 ng/mL groups was significantly higher than that of IL-1 β treated alone group (*P* < 0.05). The function of islet B-cells, which received IL-1 β treatment for 24 h, could partially recover after G-5 treatment for another 24 h. But accumulated and glucose stimulated insulin secretion in groups with G-5 treatment for 10 h groups had no significant difference as compared with IL-1 β treated alone group (*P* > 0.05).

CONCLUSION: The present results indicate that G-5 may have a protective effect against the toxicity of IL-1 β on islet B-cells.

Key words: Pentagastrin; Interleukin-1; Islet, Langerhans; Insulin; Diabetes mellitus, insulin-dependent

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