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Effect of growth hormone (GH) and placental lactogen (PL) on the regulatory mechanism of maternal serum Insulin-like growth factor 1(IGF-1)concentration during pregnancy. S.Nakago, A.Kobayashi, T.Funakoshi, Y.Ueda, M.Deguchi, H.Morikawa, M.Mochizuki, Dept.Obst.and Gynec., Kobe Univ.Sch.Med., Kobe.

Aim of this study is to clarify the regulatory mechanism of IGF-1 concentrations in maternal blood during pregnancy. IGF-1 concentrations measured by IGF-1 RIA in rat maternal blood decreased from D15 to D21. Serum IGF-1 concentration in nonpregnant rat after hypophysectomy decreased significantly, and serum IGF-1 concentration was restored after GH administration to this hypophysectomized rat, but PL administration did not restore it. In the culture system using rat adult hepatocytes, medium IGF-1 increased significantly after addition of GH into the medium. But no significant change in medium IGF-1 concentrations was observed after GH addition was inhibited by the simultaneous addition of PL with GH into medium. However, no significant changes in medium IGF-1 concentrations were observed after PRL addition or after simultaneous addition of PRL with GH into the medium. These results suggest that the production of IGF-1 during pregnancy is controled by the promoting action of GH and by the suppressing action of PL.

55 Purification of gonadotropes and intracellular free calcium oscillation: Effects of gonadotropin releasing hormone and interleukin 6.

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The intracellular free calcium concentration ([Ca²⁺]i) in single gonadotropes was measured with fura-2 and a digital imaging fluorescence microscopic system to determine how interleukin-6 (IL-6) increases release of gonadotropins. IL-6 induced increase in the basal $[Ca^{2+}]i$ or the amplitude of spontaneous oscillation of $[Ca^{2+}]i$ in gonadotropes in a mixed population. Next, purified gonadotropes were prepared fluorescence-activated cell sorting (FACS) and argon laser treatment by Gonadotropes labeled with anti-LH antibody were sorted by the cells. and then cultured as monolayers for 24-48h. In this way, FACS, gonadotropes were concentrated from 5-10% to 70-85% from whole pituitary cells. After relabeling with anti-LH antibody, 100% purified gonadotropes were obtained by killing other types of cells with argon laser. Gn-RH induced almost the same responses of $[Ca^{2+}]i$ in the purified cell population as in the mixed cell population, but IL-6 did not affect $[Ca^{2+}]i$ in the purified gonadotropes. These results suggest that IL-6 affects calcium mobilization in gonadotropes indirectly via paracrine pathways.

Effects of saturated fatty acids on prostaglandin E₂ 9-ketoreductase (PGE₂-9-KR). T.Ohshige, T.Ohtsuka, M.Mibe, M.Yamaguchi, N. Mori, Dept.Obst.and Gynec., Miyazaki Medical College, Miyazaki.

NADP-dependent prostaglandin E_2 9-ketoreductase (PGE2-9-KR, EC 1.1.1. 189), which catalyzes the conversion of prostaglandin E_2 (PGE2) into prostaglandin $F_{2\alpha}$ (PGF2 $_{2\alpha}$) may regulate the intracellular PGF2 $_{2\alpha}$ /PGE2 ratio. The presence of specific inhibitors of PGE2-9-KR in human placenta has been suggested, but it is yet unclear whether regulatory mechanisms of the enzyme activity actually exist or not. On the other hand, endogenous inhibitors of 15-hydoxyprostaglandin dehydrogenase (PGDH), which catalyzes the first reaction of PGE2 inactivation, have been isolated from pregnant rabbit lung, human placenta and human milk. Recently, we observed that endogenous inhibitors of PGDH contains myristic acid 14:0, palmitic acid 16:0 and stearic acid 18:0, and that these saturated fatty acids, in that order, inhibit PGDH activity. We partially purified PGE2-9-KR from human term decidua and examined the effects of these fatty acids on the enzyme activity. Palmitic acid inhibited PGE2-9-KR activity dose-dependently, whereas the other two fatty acids had no effect. Our findings suggest that in spite of the structural similarity of these fatty acids, only palmitic acid has a specific inhibitory effect on PGE2-9-KR.