### 论著

# 雌激素和高胰岛素调节胰岛素受体底物-1,2表达机制研究

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目的:研究雌激素和高浓度胰岛素对胰岛素受体底物(IRS)-1和-2表达的分子机理。方法:将IRS-1,2 基因5′调控区克隆至含荧光素酶表达载体pGL3质粒,转染HeLa细胞,加雌激素(1 nmol/L)或高浓度胰岛素 (100 nmol/L)培养,检测IRS-1,2基因5′调控区相对转录活性。结果: 高浓度胰岛素刺激细胞48 h,IRS-2 基因5'调控区相对转录活性减低,IRS-1无明显差异;雌激素处理显著增加IRS-1,2基因5'调控区的相对转录活<mark>▶加入引用管理器</mark> 性。结论: 高胰岛素可能通过作用于IRS-2基因5′调控区中胰岛素作用元件,使其转录活性降低。而雌激素则通 ▶复制索引 过作用于IRS-1,2基因5′调控序列,使其转录活性提高,增强其表达。

雌激素类; 胰岛素; 受体,胰岛素 关键词

分类号 R363

# Regulatory mechanism in expression of IRS-1 and 2 by estrogen and high concentration of insulin

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#### Abstract

<FONT face=Verdana>AIM: To study the molecular mechanism in modulation of expression of insulin receptor substrate-1 and -2 (IRS-1, -2) by estrogen and high concentration of insulin. METHODS: The 5'-regulatory regions of IRS-1 and IRS-2 gene were cloned into the pGL3 plasmid with luciferase reporter, and the clones were transfected into HeLa cells. The cells were incubated with estradiol (1 nmol/L) and high concentration of insulin (100 nmol/L). The relatively transcriptional activity of the 5'-regulatory regions of IRS-1 and IRS-2 gene was detected. RESULTS: It was found that the relatively transcriptional activity of the 5'-modulatory regions of IRS-2 reduced markedly after cells were incubated with 100 nmol/L insulin (P<0.05), but that of IRS-1 was not affected. Estradiol increased the relatively transcriptional activity of the 5'-regulatory regions of IRS-1 and -2 distinctly (P<0.05). CONCLUSIONS: High concentration of insulin decreases the expression of IRS-2 by acting on its insulin reactive element, and estradiol elevates the expression of IRS-1 and -2 by acting on their 5'-modulatory regions. </FONT>

Key words Estrogens Insulin Receptor insulin

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