

论文

三七总皂苷对oxLDL诱导的人脐静脉内皮细胞CD40, VCAM-1表达的影响

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摘要:

目的 探讨三七总皂苷(TPNS)对氧化型低密度脂蛋白(oxLDL)诱导的人脐静脉内皮细胞(HUVECs)白细胞分化抗原40(CD40)、血管细胞粘附分子-1(VCAM-1)表达的影响。**方法** 原代培养HUVECs。细胞分为空白组、刺激组、小剂量(100mg/L)TPNS组、大剂量(200mg/L)TPNS组及辛伐他汀组并予相应处理。采用MTT法测定HUVECs活性,实时定量RT-PCR检测VCAM-1的基因表达,Western blot分析CD40蛋白表达量。**结果** TPNS及辛伐他汀均可以升高HUVECs的活性($P<0.05$, $P<0.001$),下调HUVECs表达免疫炎症因子CD40($P<0.05$, $P<0.01$)及VCAM-1(P 均 <0.001)的水平。且大剂量TPNS作用优于小剂量TPNS($P<0.05$)。**结论** TPNS能够减轻oxLDL对内皮细胞的损伤,降低内皮细胞免疫炎症因子的表达,在防治动脉粥样硬化中具有重要作用。

关键词: 三七总皂苷; 氧化型低密度脂蛋白; 人脐静脉内皮细胞; 白细胞分化抗原40; 血管细胞粘附分子-1

Effect of total panax notoginsenosides on expression of CD40, VCAM-1 induced by oxLDL in human umbilical vein endothelial cells

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Abstract:

Objective To explore the effect of total panax notoginsenosides (TPNS) on expression of cell differentiation antigen 40 (CD40) and vascular cell adhesion molecule-1 (VCAM-1) in human umbilical vein endothelial cells (HUVECs) induced by oxidized low density lipoprotein (oxLDL). **Methods** Primary culture HUVECs cells were divided into five groups: the control, the model, the low dose (100mg/L) the TPNS, high dose (200mg/L) TPNS and the Simvastatin group. Apart from the control group, cells in other groups were given corresponding treatment. Cytoactivity was measured by MTT. mRNA expression of VCAM-1 was detected by realtime quantitative RT-PCR. Protein expression of CD40 was measured by Western blot. **Results** TPNS and Simvastatin could up-regulate the cytoactivity of HUVECs($P<0.05$, $P<0.001$), and down-regulate expression of CD40($P<0.05$, $P<0.01$), VCAM-1 ($P<0.001$) in HUVECs induced by oxLDL. In addition, the effect of high dose TPNS was superior to the low dose TPNS ($P<0.05$). **Conclusion** TPNS can relieve injury and lower expression of immune inflammatory factors in HUVECs induced by oxLDL. TPNS has a potential effect in preventing and treating atherosclerosis.

Keywords: Total panax notoginsenosides; Oxidized low density lipoprotein; Human umbilical vein endothelial cells; Cell differentiation antigen 40; Vascular cell adhesion molecule-1

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