

论著

Rho在二氧化硅诱导的人支气管上皮细胞
 α -平滑肌肌动蛋白表达中的作用

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摘要:

目的: 探讨Rho在二氧化硅(SiO₂)诱导人支气管上皮细胞(HBECs) α -平滑肌肌动蛋白(α -SMA)表达中的作用。方法: 以SiO₂处理的HBE细胞为研究对象, 免疫细胞化学及Western印迹检测 α -SMA表达; GST pull down分析Rho的活化情况; 干预实验中用Rho抑制剂Y27632处理细胞后, Western印迹检测SiO₂对 α -SMA表达的影响。结果: SiO₂ (0, 50, 100, 200, 300 μ g/mL)分别作用HBECs细胞72 h后, α -SMA表达增强, 其中以200 μ g/mL SiO₂组 α -SMA表达最强, 为对照组的(5.09 \pm 1.98)倍(P<0.01); 用200 μ g/mL SiO₂分别作用HBECs细胞1, 2, 6, 12, 24 h后, Rho明显活化(P<0.01); Rho抑制剂Y27632明显抑制了SiO₂诱导的 α -SMA表达, 20和30 μ mol/L的Y27632抑制率分别为68%和75%(P<0.01)。结论: Rho参与介导SiO₂诱导的HBECs中 α -SMA的表达。

关键词: 二氧化硅 Rho α -平滑肌肌动蛋白 人支气管上皮细胞

Effect of Rho signaling pathway on SiO₂ induced α -SMA expression in human bronchial epithelial cells

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Abstract:

Objective To investigate the role of Rho in SiO₂ induced α -SMA expression in human bronchial epithelial cells (HBECs). Methods HBECs were cultured and stimulated with SiO₂. Immunocytochemistry and Western blot were used to detect the expression of α -SMA. The activity of Rho was determined by GST pull down assay. In the prevention experiment, SiO₂-stimulated HBECs were incubated with Rho inhibitor Y27632, and the expression of α -SMA was examined by Western blot. Results With SiO₂ (0-300 μ g/mL) treatment, the expression of α -SMA increased gradually, and 200 μ g/mL of SiO₂ led to the highest expression of α -SMA which was (5.09 \pm 1.98) times of the expression of α -SMA in the control group (P<0.01). HBECs treated with SiO₂ (200 μ g/mL) for indicated time (1,2,6,12, and 24 h) showed an obvious increase of Rho activity (P<0.01). Y27632 inhibited SiO₂-induced α -SMA expression significantly, and the inhibition rate of 20 and 30 μ mol/L Y27632 was 68% and 75%, respectively (P<0.01). Conclusion Rho signaling pathway may mediate SiO₂ induced α -SMA expression in HBECs.

Keywords: SiO₂; Rho; α -SMA; human bronchial epithelial cell

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参考文献:

[1] Willis B C, Borok Z. TGF-beta-induced EMT: mechanisms and implications for fibrotic lung disease [J]. Am J Physiol Lung Cell Mol Physiol, 2007, 293(3): L525-L534.
[2] Coward W R, Saini G, Jenkins G. The pathogenesis of idiopathic pulmonary fibrosis [J]. Ther Adv Respir Dis, 2010, 4(6): 367-388.
[3] Camara J, Jarai G. Epithelial-mesenchymal transition in primary human bronchial epithelial cells is Smad-dependent and enhanced by fibronectin and TNF-alpha [J]. Fibrogenesis Tissue Repair, 2010, 3(1): 2.
[4] Linnala A, Kinnula V, Laitinen L A, et al. Transforming growth factor-beta regulates the expression of fibronectin and tenascin in BEAS 2B human bronchial epithelial cells [J]. Am J Respir Cell Mol Biol, 1995, 13(5): 578-585.
[5] Bodo M, Baroni T, Bellocchio S, et al. Bronchial epithelial cell matrix production in response to silica and basic fibroblast growth factor [J]. Mol Med, 2001, 34(7): 83-92.
[6] Willis B C, duBois R M, Borok Z. Epithelial origin of myofibroblasts during fibrosis in the lung [J]. Proc Am Thorac Soc, 2006, 3(4): 337-382.
[7] Yanez-mo M, Lara-Pazzi E, Selgas R, et al. Peritoneal dialysis and epithelial to mesenchymal transition of mesothelial cells [J]. N Engl J Med, 2003, 348(5): 403-413.
[8] Hanby A M, Chinery R, Poulsom R, et al. Downregulation of E-cadherin in the reparative epithelium of the human gastro intestinal tract [J]. Am J Pathol, 1996, 148(6): 723-729.

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[9] Willis B C, Liebler J M, Luby-Phelps K, et al. Induction of epithelial-mesenchymal transition in alveolar epithelial cells by transforming growth factor-beta1: potential role in idiopathic pulmonary fibrosis [J] . Am J Pathol, 2005, 166(5): 1321-1332.

[10] Xu G P, Li Q Q, Cao X X, et al. The Effect of TGF-beta1 and SMAD7 gene transfer on the phenotypic changes of rat alveolar epithelial cells [J] . Cell Mol Biol Lett, 2007,32(4): 126-131.

[11] Kim J H, Jang Y S, Eom K S, et al. Transforming growth factor beta1 induces epithelial-to- mesenchymal transition of A549 cells [J] . J Korean Med Sci, 2007, 22(5):898-904.

[12] Wu Z, Yang L, Cai L, et al. Detection of epithelial to mesenchymal transition in airways of a bleomycin induced pulmonary fibrosis model derived from an alpha-smooth muscle actin-Cre transgenic mouse [J] . Respir Res, 2007,8(1):1.

[13] Patel S, Takagi K I, Suzuki J, et al. Rho GTPase Activation is a key step in renal epithelial mesenchyma transdifferentiation [J] . J Am Soc Nephrol, 2005, 16(6):1977-1984.

[14] Cho H J, Yoo J. Rho activation is required for transforming growth factor- beta-induced epithelial-mesenchymal transition in lens epithelial cells [J] . Cell Biol Int, 2007, 31(10):1225-1230.

[15] 李蕾, 蒋炜, 王吉耀, 等. 转化生长因子 β 1对肝星状细胞迁移及细胞内Rho三磷酸鸟苷酶表达的影响 [J] . 中华消化杂志,2006, 26(8): 527-530.

LI Lei, JIANG Wei,WANG Jiyao, et al. Effects of Rho GTPase expression and migration induced by transform growth factor β 1 in hepatic stellate cells [J] . Chinese Journal of Digestion, 2006,26(8):527-530.

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