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论著

低氧诱导因子-1 α siRNA对HaCaT细胞诱导型一氧化氮合酶表达的影响

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摘要:

目的:观察低氧诱导因子-1 α (hypoxia inducible factor -1 α , HIF-1 α)RNA干扰对低氧条件下永生化角质形成细胞株HaCaT细胞HIF-1 α 和诱导型一氧化氮合酶(inducible nitric oxide synthase, iNOS)表达的影响。方法:将HaCaT细胞分为4组:常氧对照组(无干预因素)、低氧组(低氧培养24 h)、脂质体对照组(转染空载脂质体后低氧培养24 h)、RNA干扰组(转染脂质体介导的RNA干扰序列后低氧培养24 h)。荧光实时定量PCR法检测各组HaCaT细胞的 HIF-1 α 和 iNOS mRNA表达水平,Western 印迹检测各组HaCaT细胞HIF-1 α 和 iNOS蛋白表达水平。结果:低氧组和常氧对照组 HIF-1 α mRNA的表达无明显差异($P>0.05$),而低氧组iNOS mRNA和蛋白及HIF-1 α 蛋白的表达均较常氧对照组明显增高($P<0.05$); RNA干扰组HIF-1 α 和iNOS mRNA及蛋白的表达均较脂质体对照组显著降低($P<0.05$)。结论:低氧条件下可以使HaCaT细胞HIF-1 α 和iNOS的表达增加,而抑制HIF-1 α 的表达可以使低氧条件下HaCaT细胞iNOS的表达减少。

关键词: HaCaT 低氧诱导因子-1 α siRNA 诱导型一氧化氮合酶

Effects of hypoxia inducible factor-1 α siRNA on inducible nitric oxide synthase expression in HaCaT cells

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Abstract:

Objective To observe the effect of hypoxia inducible factor -1 α (HIF-1 α) small interfering RNA (siRNA) on the expression of HIF-1 α and inducible nitric oxide synthase (iNOS) in HaCaT cells under hypoxia. Methods HaCaT cells were divided into 4 groups: the normal control group (without any treatment), the hypoxia group (under hypoxia for 24 h), the liposome control group (HaCaT cells transfected with liposome before hypoxia treatment), the RNA interference group (HaCaT cells transfected with siRNA sequences then under hypoxia for 24 h). Real-time PCR and Western blot were utilized to determine HIF-1 α and iNOS mRNA and protein expression in HaCaT cells. Results There was no significant difference of the mRNA expression of HIF-1 α between the hypoxia group and the normoxia group ($P>0.05$), but the protein expressions of HIF-1 α was increased in the hypoxic group than that in the normoxia group ($P<0.05$). Both the mRNA and protein expression of iNOS were increased in hypoxic conditions than that in the normoxia ($P<0.05$). Decreases were more significant in the mRNA and protein expression of HIF-1 α and iNOS in the RNA interference group than that in the liposome control group in HaCaT cells ($P<0.05$). Conclusion Hypoxia increased HIF-1 α and iNOS expression in HaCaT cells and inhibition of HIF-1 α expression decreased iNOS expression.

Keywords: HaCaT hypoxia-inducible factor-1 α small interfering RNA inducible nitric oxide synthase

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参考文献:

[1] Rosenberger C, Solovan C, Rosenberger A D, et al. Upregulation of hypoxia-inducible factors in normal and psoriatic skin [J]. J Invest Dermatol, 2007, 127(10): 2445-2452.

[2] Tovar-Castillo L E, Cancino-Diaz J C, Garcia-vazquez F, et al. Under-expression of VHL and over-expression of HDAC-1, HIF-1alpha, LL-37, and IAP-2 in affected skin biopsies of patients with psoriasis [J]. Int J Dermatol, 2007, 46(3): 239-246.

[3] 李勇坚,张桂英,肖嵘,等. HIF-1 α 与iNOS在银屑病皮损中的表达及其与血管生成的关系 [J]. 中南大学学报: 医学版, 2010, 35(9): 952-957.

LI Yongjian, ZHANG Guiying, XIAO Rong, et al. Expression of iNOS and HIF-1 α with angiogenesis in affected skin biopsies from patients with psoriasis [J]. Journal of Central South University. Medical Science, 2010, 35(9): 952-957.

[4] 刘重霄,刘勇,师蔚,等. 缺氧条件下谷氨酸对鼠脑星形胶质细胞血管内皮生长因子表达的影响 [J]. 南方医科大学学报, 2010, 30(3): 435-438.

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- expressions in hypoxic rat astrocytes in vitro [J]. Journal of Southern Medical University, 2010,30(3):435-438.
- [5] Brown J M, Wilson W R. Exploiting tumour hypoxia in cancer treatment [J]. Nat Rev Cancer, 2004,4(6): 437-447.
- [6] Lee J W, Bae S H, Jeong J W, et al. Hypoxia-inducible factor (HIF-1) alpha: its protein stability and biological functions [J]. Exp Mol Med. 2004 ,36(1):1-12.
- [7] Tao J, Yang J, Wang L, et al. Expression of GLUT-1 in psoriasis and the relationship between GLUT-1 upregulation induced by hypoxia and proliferation of keratinocyte growth [J]. J Dermatol Sci, 2008,51(3): 203-207.
- [8] Kwon Y W, Kwon K S, Moon H E, et al. Insulin-like growth factor-II regulates the expression of vascular endothelial growth factor by the human keratinocyte cell line HaCaT [J]. J Invest Dermatol, 2004, 123(1): 152-158.
- [9] 杨井,陶娟,李延,等.缺氧对HaCaT细胞HIF-1 α 、GLUT-1表达的影响及与细胞增殖的关系 [J]. 中国皮肤性病学杂志,2009,23 (10):621-632.

YANG Jing, TAO Juan, LI Yan, et al. Effect of hypoxia on the expression of HIF-1 α and GLUT-1 in HaCaT cell line and the Relationship with the cell proliferation [J]. The Chinese Journal of Dermatovenereology, 2009, 23(10): 621-632.

[10] Kessler J, Hahnel A, Wichmann H, et al. HIF-1 α inhibition by siRNA or chetomin in human malignant glioma cells: effects on hypoxic radioresistance and monitoring via CA9 expression [J]. BMC Cancer, 2010, 10: 605.

[11] Frank S, Stallmeyer B, Kampfer H, et al. Nitric oxide triggers enhanced induction of vascular endothelial growth factor expression in cultured keratinocytes (HaCaT) and during cutaneous wound repair [J]. FASEB J, 1999, 13(14):2002-2014.

[12] Ormerod A D, Copeland P, Shah S A. Treatment of psoriasis with topical NG-monomethyl-L-arginine, an inhibitor of nitric oxide synthesis [J]. Br J Dermatol, 2000, 142(5): 985-990."

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[J]. 中南大学学报(医学版), 2006, 31 (05) : 635-639

2. 陈明亮¹, 张桂英², 易梅¹, 陈潇³, 李吉¹, 谢红付¹, 陈翔¹.长波紫外线对人皮肤成纤维细胞增殖及NO/iNOS系统的影响[J]. 中南大学学报(医学版), 2009,34(08): 705-711

3. 刘惠宁¹, 蔡净亭¹, 林秋华², 何可人¹, 余蓉¹.Caveolin-1与绒毛膜癌侵袭力之间的关系[J]. 中南大学学报(医学版), 2008,33(04): 331-337

4. 罗红¹, 胡冬煦², 陈平¹.抑制A549细胞COX-2表达的D-siRNAs的合成[J]. 中南大学学报(医学版), 2007,32(03): 437-442

5. 吕辉, 贺智敏*.RNA干扰技术的演进及其在基因功能和基因治疗研究中的应用[J]. 中南大学学报(医学版), 0,(): 102-105

6. 吕辉, 贺智敏*.RNA干扰技术的演进及其在基因功能和基因治疗研究中的应用[J]. 中南大学学报(医学版), 2005,30(1): 102-105

7. 李勇坚^{1,2}, 张桂英², 肖嵘², 陈欢², 文海泉².HIF-1 α 与iNOS在银屑病皮损中的表达

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