

Viii多糖对人肺癌A549细胞中MAGEA10表达的影响

张金禄¹, 张晓红¹, 王玉荣¹, 何超², 吴季辉², 罗成¹

1.300457 天津, 天津科技大学食品工程与生物技术学院; 2.中国科学技术大学生命科学学院

MAGEA10 Expression in Human Lung Cancer A549 Cells Influenced by Viili Polysaccharides

ZHANG Jinlu¹, ZHANG Xiaohong¹, WANG Yurong¹, HE Chao², WU Jihui², LUO Cheng¹

1. College of Food Engineering and Biotechnology, Tianjin University of Science & Technology, Tianjin 300457, China; 2. School of Life Sciences, University of Science and Technology of China

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摘要

目的

探讨Viii多糖对肿瘤特异性抗原基因MAGEA10表达量的影响及其机制。

方法

利用MTT技术检测不同浓度Viii多糖刺激A549细胞24 h、48 h、72 h后对其存活率的影响, qRT-PCR法检测Viii多糖浓度为10 mg/L、25 mg/L、50 mg/L时与正常生长A549细胞相比较MAGEA10 mRNA相对表达量的变化, 并用Western blot检测Viii多糖10 mg/L、25 mg/L、50 mg/L刺激A549细胞后MAGEA10表达抗原肽的情况。通过MAGEA10 mRNA表达量与抗原肽表达量分析探讨Viii多糖对MAGEA10表达的影响及其表达过程中的可能机制。

结果

(1) Viii多糖作用A549细胞后, 存活率下降, 在0 mg/L~50 mg/L范围内其浓度与细胞存活率呈负相关, 且作用48 h效果最显著。(2) 与未用Viii多糖刺激的空白组相比, Viii多糖浓度为50 mg/L时MAGEA10 mRNA相对表达量上升。(3) 与空白组相比, Viii多糖浓度为50 mg/L时MAGEA10蛋白表达量上升。

结论

Viii多糖对A549细胞的生长具有一定的抑制作用, 上调A549细胞中MAGEA10的转录和翻译水平, 两者机制尚需进一步研究, 但可能是相互独立的。本研究通过提高癌症的非特异性免疫来提高特异性免疫概率, 为癌症辅助治疗提供更多的科学依据。

关键词: MAGEA10 Viii多糖 A549 MAGEs

Abstract:

Objective

To discuss the effects and mechanism of MAGEA10 expression in human lung cancer A549 cells influenced by Viili polysaccharides.

Methods

MTT was used to study the survival impact of non small cell lung cancer (NSCLC) A549 with viili polysaccharides in different concentration at 24 h,48 h and 72 h. qRT-PCR was used to detect the relative expression changes of MAGEA10 mRNA with Viili polysaccharides in different concentration of 10 mg/L, 25 mg/L and 50 mg/L compared with those of the normal A549 cells. And Western blot was adopted to detect the antigen expression of MAGEA10 after stimulation of A549 with Viili polysaccharides in different concentration of 10 mg/L, 25 mg/L and 50 mg/L. Through the expression of MAGEA10 mRNA and antigen, we discussed the effects and possible mechanism of MAGEA10 expression made by Viili polysaccharides.

Results

(1) Viili polysaccharides inhibited cell proliferation of A549 and its concentration was negatively related to cell

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survival within 0 mg/L-50 mg/L and 50 mg/L was most remarkable at 48 h. (2) Compared with the blank group, the relative expression of MAGEA10 mRNA elevated at the concentration of 50 mg/L. (3) Compared with the blank group, the protein expression of MAGEA10 elevated, at the concentration of 50 mg/L.

Conclusion

Viii inhibited NSCLC A549 cell proliferation, but the increase of the gene transcription and translation levels of MAGEA10 in A549 cell needs further exploration for those two parts might be interdependent. This study aims to indicate that viii is possible to promote non specific immunity, and increase possibility of cancer cell recognition by CTLs, and possibly initiate apoptotic process of cancer cells if same epitope of MAGEA10 and corresponding TCR encounters.

Key words: MAGEA10 VIII Polysaccharides A549 Melanoma antigen-encoding genes(MAGEs)

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通讯作者: 罗成, E-mail: Luo58@yahoo.com E-mail: Luo58@yahoo.com

作者简介: 张金禄(1987-),女,硕士在读,主要从事食品营养与免疫的基础研究工作

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