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37~42.线粒体融合蛋白-2基因增强人乳腺癌T47D细胞对小白菊内酯的敏感性[J].邱梅清,佟仲生,贾勇圣,刘晓东,陈悦.中国肿瘤生物;

线粒体融合蛋白-2基因增强人乳腺癌T47D细胞对小白菊内酯的敏感性 点此下载全文

邱梅清 佟仲生 贾勇圣 刘晓东 陈悦

天津医科大学 附属肿瘤医院 乳腺内科,教育部乳腺癌防治重点实验室,天津市肿瘤防治重点实验室,天津 300060; 天津医科大学 乳腺癌防治重点实验室,天津市肿瘤防治重点实验室,天津 300060; 天津医科大学 附属肿瘤医院 乳腺内科,教育部乳腺癌防治重。验室,天津 300060; 天津医科大学 附属肿瘤医院 乳腺内科,教育部乳腺癌防治重点实验室,天津市肿瘤防治重点实验室,天津 300071

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摘要:

目的: 探讨线粒体融合蛋白-2(mitofusin-2, Mfn-2)基因表达对人乳腺癌T47D细胞对小白菊内酯敏感性的影响。 方法:细胞系(T47D、MDA-MB-231、MCF-7、MDA-MB-435及HCC38)中 Mfn-2 mRNA的表达。LipofectamineTM 2000体外转人乳腺癌T47D细胞,real-time PCR和Western blotting检测T47D细胞中 Mfn-2 mRNA和蛋白的表达,MTT法检测T47D细胞的调亡率及线粒体膜电位。 结果: 与正常乳腺细胞相比, Mfn-2 mRNA在乳腺癌HCC38细胞系中高表达,在T47D等其他细胞系中的后,T47D细胞中 Mfn-2 mRNA和蛋白的表达均明显上调。与EGFP转染组相比, pEGFP-Mfn-2转染组T47D细胞在小白菊内酯。是降低\[(47.93±2.21)% vs(56.93±2.05)%,P<0.05\]。流式细胞术检测结果显示: 50 mmol/L小白菊内酯作用下,pEG相比,T47D细胞调亡率升高 \[(71.2±2.1)% vs(38.8±2.6)%,P<0.05\],而线粒体膜电位明显降低\[(1.6±0.1)% vs(5.C EGFP-Mfn-2转染可增强T47D 细胞对小白菊内酯的敏感性。

关键词: 乳腺癌 T47D细胞 线粒体融合蛋白-2 小白菊内酯 敏感性

Mitofusin-2 gene enhances sensitivity of human breast cancer T47D cells to parthenolide Download Full

Qiu Meiging Tong Zhongsheng Jia Yongsheng Liu Xiaodong Chen Yue

Department of Breast Oncology & Key Laboratory of Breast Cancer Prevention and Therapy of Ministry of Educa Cancer Prevention and Therapy of Tianjin, Affiliated Tumor Hospital, Tianjin Medical University, Tianjin 300060, Oncology & Key Laboratory of Breast Cancer Prevention and Therapy of Ministry of Education & Key Laboratory Therapy of Tianjin, Affiliated Tumor Hospital, Tianjin Medical University, Tianjin 300060, China; Department of Br Laboratory of Breast Cancer Prevention and Therapy of Ministry of Education & Key Laboratory of Cancer Preve Affiliated Tumor Hospital, Tianjin Medical University, Tianjin 300060, China; Department of Breast Oncology & Ke Cancer Prevention and Therapy of Ministry of Education & Key Laboratory of Cancer Prevention and Therapy of Hospital, Tianjin Medical University, Tianjin 300060, China; College of Pharmacy, Nankai University, Tianjin 3000

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Abstract:

Objective:To investigate the effect of mitofusin-2 (Mfn-2) gene expression on sensitivity of human breast c parthenolide. Methods: The expressions of Mfn-2 mRNA in various breast cancer cell lines (T47D, MDA-MB-231, HCC38) were detected by real-time PCR. Plasmids pEGFP and pEGFP-Mfn-2 were transfected into human breast LipofectamineTM 2000 in vitro. The expression levels of Mfn-2 mRNA and protein in T47D cells were detected by blotting. MTT assay was used to detect the proliferation of T47D cells. The cell apoptotic rate and mitochondrial cells were measured by flow cytometry. Results: Compared with that in normal breast cells, Mfn-2 mRNA was h cancer HCC38 cell line, and lowly expressed in the other cell lines, such as T47D etc. After pEGFP-Mfn-2 transfe levels of Mfn-2 mRNA and rotein were significantly up-regulated in T47D cells. Compared with the pEGFP transfe 2 transfection group showed a significant decrease in surivival rate of T47D cells under the treatment of parther 2.21\]% vs \[[56.93 \pm 2.05\] %, P<0.05). Flow cytometry results showed that the apoptotic rate of T47D cells L mol/L parthenolide was significantly increased in pEGFP-Mfn-2 transfection group compared with that in pEGFP 2.1\] % vs \[[38.8 \pm 2.6\] %, P<0 05). However, the mitochondrial membrane potential was significantly decreast transfection group (\[[1.6 \pm 0.1\] \] % vs \[[5.0 \pm 0.5\] %, P<0.05). Conclusion: pEGFP-Mfn-2 transfection can enhance lls to parthenolide.

Keywords: breast cancer T47D cell mitofusin-2 parthenolide sensitivity

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