

论文

Bcl-2shRNA稳定转染联合 γ 射线对胃癌SGC-7901细胞凋亡的影响

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摘要:

目的 观察Bcl-2shRNA稳定转染联合 γ 线照射对胃癌细胞SGC-7901凋亡的影响。方法 构建针对Bcl-2基因的干扰质粒pGPH1/GFP/Neo, 经脂质体介导转染SGC 7901细胞, G418筛选稳定表达的细胞株, γ 线照射后形成4组细胞, 分别命名为SGC-7901(A组)、照射/SGC-7901(B组)、Bcl-2shRNA/SGC-7901(C组)、照射/ Bcl-2shRNA/SGC-7901(D组)。CCK 8检测细胞增殖, AO/PE观察细胞凋亡, 流式细胞仪检测细胞凋亡, Western blot测定Bcl 2蛋白表达量的改变。结果 Bcl-2shRNA、照射均可抑制Bcl-2蛋白表达, 且二者有协同作用, 差异有统计学意义 ($P<0.05$); D组细胞生长慢于B组、C组细胞, B、C、D组细胞增殖抑制率分别为(27.00 \pm 5.27)%、(30.10 \pm 6.49)%、(98.40 \pm 11.35)%。A、B、C、D组细胞凋亡率分别为(3.80 \pm 0.22)%、(20.80 \pm 4.15)%、(23.20 \pm 4.34)%、(92.90 \pm 25.90)%, 差异有统计学意义 ($P<0.05$)。结论 Bcl-2基因siRNA干扰联合 γ 线照射可协同抑制SGC-7901细胞中Bcl-2的表达, 诱导细胞凋亡。

关键词: 胃癌细胞; B细胞淋巴瘤/白血病 2; RNA干扰; 细胞凋亡; 照射

Influence of stable transfection of bcl-2shRNA combined with γ irradiation on apoptosis of the gastric carcinoma cell line SGC-7901

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Abstract:

Objective To observe the effect of bcl-2 gene-specific RNA interference combined with γ irradiation on apoptosis and radiosensitivity of the gastric carcinoma cell line SGC7901. Methods The recombinant eukaryotic expression vector pGPH1/GFP/Neo designed to target the bcl-2 gene was transfected into SGC7901 cells by lipofectamin. Stable positive clones were selected with G418. After γ irradiation, the cells were divided into 4 groups: SGC-7901 (group A), irradiation/SGC-7901 (group B), Bcl-2shRNA/SGC-7901 (group C) and irradiation/Bcl 2shRNA/SGC-7901 (group D). Cell proliferation was detected by CCK-8. Cell apoptosis was observed by AO/PE. The apoptosis rate was determined by flow cytometry. Expression of bcl-2 was determined by Western blot, and compared between transfection cells and non-transfection cells. Results Western blot analysis indicated that expression of bcl-2 was suppressed by shRNA and irradiation, and they had a synergetic effect. The growth of cells in group D was obviously slower than that in the other groups($P<0.05$). Cell inhibitory rates in groups B, C and D were (27.00 \pm 5.27)%, (30.10 \pm 6.49)% and (98.40 \pm 11.35)%, respectively. Apoptosis rates in groups A, B, C and D were (3.80 \pm 0.22)%, (20.80 \pm 4.15)%, (23.20 \pm 4.15)% and (92.90 \pm 25.90)%, respectively ($P<0.05$). Conclusions Bcl-2 gene suppression by RNAi and irradiation can induce cell apoptosis, co-inhibit expression of bcl-2, and obviously increase radiosensitivity in the human gastric cell line.

Keywords: Gastric carcinoma cells; B cell lymphoma/leukemia 2; RNA interference; Apoptosis; Irradiation

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