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IL-21在肝癌细胞株H22细胞中的表达及其活性 点此下载全文

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摘要：

摘 要 目的：构建小鼠IL-21（mIL-21）真核表达载体mIL-21-pcDNA3.1，转染肝癌H22细胞，探讨其生物学活性。mIL-21真核表达载体mIL-21-pcDNA3.1，并经DNA测序证实。脂质体法介导mIL-21-pcDNA3.1转染H22细胞，RT-PCR检测mIL-21-pcDNA3.1对T细胞增殖和NK细胞杀伤活性的影响。结果：DNA序列分析证实构建的mIL-21-pcDNA3.1正确无转染的H22细胞中有mIL-21的表达。MTT法显示，转染mIL-21的H22细胞培养上清刺激T细胞增殖的刺激指数（stimulati ConA的刺激指数为4.673±0.450，均显著高于转染空质粒组的1.465±0.103和未转染组的1.447±0.245,(P<0.01）。杀伤率为(81.66±4.26)%，显著高于转染空质粒组的（34.74±5.52)%和未转染对照组的（33.61±1.42)%。结论：m达可显著增强T细胞增殖及NK细胞的杀伤功能，为其在抗肝癌治疗中的应用奠定了基础。

关键词：肝肿瘤 mIL-21基因 H22细胞株 真核表达

IL-21 expression in hepatoma cell line H22 and its biological activity Download Fulltext

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Abstract:

Abstract Objective:To construct a recombinant eukaryotic expression vector mIL-21-pcDNA3.1 and t H22, so as to assess the biological activity of mIL-21. Methods: The gene fragment encoding mouse IL-2 was then cloned into eukaryotic expression plasmid pcDNA3.1 to form recombinant plasmid mIL-21-pcDN verified by DNA sequencing. mIL-21-pcDNA3.1 was transfected into H22 cells with lipofect regent, and its PCR and Western blotting analysis. The effects of mIL-21-pcDNA3.1 on proliferation of T cells and cytotox Results: The recombinant plasmid mIL-21-pcDNA3.1 was confirmed by DNA sequencing. The expression o by RT-PCR and Western blotting analysis. MTT results showed that stimulation index (SI) of T cells stimul supernatant was 3.412±0.312, and the SI of ConA combination stimulating group was 4.673±0.450; bo those in the mock vector transfected (1.465±0.103) and untransfected groups (1.447±0.245, P<0.01). 21-H22 cell supernatant group was (81.66±4.26)%, significantly higher than those in the mock vector tr untransfected groups ([33.61±1.42]%). Conclusion: The expression of mIL-21-pcDNA3.1 plasmid in H22 of T cells and the cytotoxicity of NK cells, which lays a foundation for its role in the research of anti-hepat

Keywords:live neoplasms mIL-21 H22 cell line eukaryotic expression