

论著

pcDNA3.1 + /MAGE-3 DNA疫苗诱导特异性抗肿瘤免疫应答的实验研究

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收稿日期 2003-6-30 修回日期 2003-10-15 网络版发布日期 2009-9-13 接受日期 2003-10-15

摘要 目的: 构建pcDNA3.1+/MAGE-3 DNA疫苗, 观察其在小鼠体内诱导特异性抗肿瘤免疫应答的能力。方法: 通过RT-PCR构建重组表达质粒pcDNA3.1+/MAGE-3; 以pcDNA3.1+/MAGE-3 DNA疫苗免疫已接种肿瘤细胞的小鼠, 每10 d重复免疫1次, 共3次, 以pcDNA3.1+、PBS为对照。末次免疫后5 d检测血清中MAGE-3抗体滴度、小鼠脾淋巴细胞的细胞毒T细胞(cytotoxic T lymphocytes, CTL)杀伤活性、细胞因子IL-2和IFN- γ 的浓度, 同时计算抑瘤率。结果: 成功构建了pcDNA3.1+/MAGE-3 DNA疫苗, 用此疫苗免疫已接种B16/MAGE-3细胞的小鼠后, 能诱导小鼠脾淋巴细胞MAGE-3特异性的杀伤活性, 脾细胞培养上清中细胞因子IL-2和IFN- γ 的浓度明显增高, 血清中抗MAGE-3抗体在1:20滴度时阳性, 肿瘤生长被显著抑制, 与pcDNA3.1+组、PBS组相比, 差异显著(P<0.01)。结论: 成功构建了pcDNA3.1+/MAGE-3 DNA疫苗, 该疫苗在小鼠体内既能激活CTL杀伤活性和CD4+ T细胞活性, 又能激活体液免疫反应, 从而诱导出特异性的抗肿瘤免疫应答。

关键词 [黑色素瘤](#); [疫苗,DNA](#); [免疫应答](#); [T淋巴细胞,细胞毒性](#)

分类号 [R363](#)

Specific antitumor immune response induced by pcDNA3.1 + /MAGE-3 DNA vaccine in mice

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Abstract

AIM: To construct pcDNA3.1+/MAGE-3 DNA vaccine and investigate the antigen-specific antitumor immune responses induced by pcDNA3.1+/MAGE-3 DNA vaccine in vivo. METHODS: C57BL/6 mice challenged with B16/MAGE-3 cells were immunized by intramuscular injection of pcDNA3.1+/MAGE-3 DNA vaccine every 10 days. pcDNA3.1+ plasmid and PBS were used as controls. After three cycles of immunization, murine splenic lymphocytes, serum, and tumor were obtained for cytotoxicity assay, detections of cytokines (IL-2 and IFN- γ), measurement of MAGE-3 antibody, and tumor inhibitory rates, respectively. RESULTS: The pcDNA3.1+/MAGE-3 DNA vaccine immunized murine lymphocytes induced specific cytotoxicity against B16/MAGE-3 cells. Significantly increased secretions of IL-2 and IFN- γ were detected. The titres of antibody against MAGE-3 were 1:1 and 1:20, while controls were negative. The tumor inhibitory rate in pcDNA3.1+/MAGE-3 group was significantly different from that in controls. CONCLUSION: The pcDNA3.1+/MAGE-3 DNA vaccine was constructed successfully. pcDNA3.1+/MAGE-3 DNA vaccine activates both cellular and humoral immune responses, and induces antigen-specific antitumor immune responses in vivo.

Key words [Melanoma](#) [Vaccines](#) [DNA](#) [Immune response](#) [T-lymphocytes](#) [cytotoxic](#)

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