

An investigation of PD-L1 expression and its association with tumor infiltrating T cells in human cervical carcinomas

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Purpose: To observe the expression of Programmed death receptor-ligand 1(PD-L1) and the association between PD-L1 expression and T cell infiltration in human cervical carcinomas.

Methods : PD-L1 and PD-1 expression was respectively determined in five cases of normal cervical tissue, 7 cases of high-level cervical intraepithelial neoplasia (CIN II-III) and 67 cases of cervical carcinomas by immunohistochemistry staining; the tumor infiltrating CD4⁺T and CD8⁺T cells were determined by immunofluorescent staining, and the apoptosis of tumor infiltrating lymphocytes was examined by TUNEL assay in those cases.

Results: No PD-L1 expressed in normal cervical epithelium; PD-L1 negatively or weakly expressed in epithelia of high grade CIN, the average relative optical density was 0.82 ± 0.75 ; and PD-L1 expressed in 70% (47/67) cervical carcinomas, the average relative optical density in superficial infiltrating (<0.5 cm) and deep infiltrating cervical squamous cell carcinomas was 2.70 ± 1.68 and 2.90 ± 1.72 . PD-1 expressed in partial tumor infiltrating lymphocytes in those cases. PD-L1 expression density of cervical carcinomas was significantly higher than that of CIN ($P < 0.01$); PD-L1 expression density of superficial invasive cervical carcinomas was slightly lower than that of deep invasive cases, but there was no significant statistic difference between them; in addition, PD-L1 expression negatively associated with the number of tumor infiltrating CD8⁺T cells ($r = -0.82$, $P < 0.01$), but not with the number of CD4⁺T cells ($r = -0.05$, $P > 0.05$). Apoptosis occurred in partial tumor infiltrating lymphocytes of cervical carcinomas.

Conclusion: Human cervical carcinoma cells express PD-L1, and it negatively associates with the number of tumor infiltrating CD8⁺T cells, but not with the number of CD4⁺T cells. PD-L1 expression of tumor cells may play role on apoptosis of tumor infiltrating lymphocytes.