

宫颈腺癌中HPV16/18感染与COX-2蛋白表达的关系

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The Relationship between HPV16/18 Infection and COX-2 Protein Expression in Cervical Adenocarcinoma

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摘要 目的探讨16、18型人乳头瘤病毒(HPV16/18)在宫颈腺癌发生发展中的作用,分析HPV16/18感染与环氧化酶-2(COX-2)蛋白表达的关系。方法采用组织微阵列技术结合原位杂交和免疫组化(SP法)检测24例慢性宫颈炎和86例宫颈腺癌HPV16/18DNA和COX-2蛋白表达情况。结果HPV16/18DNA和COX-2蛋白在宫颈腺癌中的阳性表达率分别为65.1%与86.0%,均显著高于慢性宫颈炎组织8.3%与33.3%(P<0.01)。HPV16/18感染与宫颈腺癌的病理分级和组织学类型无关,但与COX-2表达呈正相关(P<0.05)。COX-2表达与宫颈腺癌的病理分级有关,G3组COX-2蛋白阳性表达率明显高于G1组(P<0.05)。COX-2表达与宫颈腺癌组织学类型无明显相关性(P>0.05)。结论宫颈腺癌的发生与HPV16/18感染及COX-2蛋白表达异常相关,宫颈腺癌中COX-2高表达与HPV16/18感染有一定相关性,二者可能协同作用导致宫颈腺癌恶性发展。

关键词: 宫颈腺癌 HPV16/18 COX-2 原位杂交 免疫组化

Abstract: Objective To study the role of 16,18 human papilloma virus (HPV16/18) infection in the carcinogenesis of cervical adenocarcinoma, and to analyze the relationship between HPV16/18 infection and cyclooxygenase-2(COX-2) protein expression. Methods The expressions of HPV16/18DNA and COX-2 protein were determined by tissue microarray combined with *in situ* hybridization and immunohistochemistry in 86 cases of cervical adenocarcinoma and 24 cases of cervical chronic inflammation. Results The positive rates of HPV16/18DNA and COX-2 protein in cervical adenocarcinoma were 86.0% and 65.1% respectively, both of which were significantly higher than those in chronic cervical inflammation (8.3% and 33.3%)(P<0.01). There was no correlation between HPV16/18 infection and pathological grade and histological type of cervical adenocarcinoma, but there was a positive correlation between COX-2 expression and COX-2 expression (P<0.05). COX-2 expression was related to pathological grade of cervical adenocarcinoma, COX-2 expression rate in G3 group was significantly higher than that in G1 group (P<0.05). There was no significant correlation between COX-2 expression and histological type of cervical adenocarcinoma (P>0.05). Conclusion The occurrence of cervical adenocarcinoma is associated with HPV16/18 infection and COX-2 protein expression, and COX-2 high expression in cervical adenocarcinoma is associated with HPV16/18 infection, and they may have synergistic effect in causing malignant development of cervical adenocarcinoma.

Key words: Cervical adenocarcinoma HPV16/18 COX-2 *In situ* hybridization Immunohistochemistry

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