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探讨CK7 CK20和TTF-1在判断转移性腺癌原发病灶中的应用价值

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Value of Combined Detection of CK7, CK20, and TTF-1 Expression in Evaluating the Origin of Metastatic Carcinoma

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摘要

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摘要 了解CK7、CK20及TTF-1在腺癌中的表达情况，探讨联合应用这3种标志物对判断转移性腺癌原发部位的价值。方法：采用组织芯片和免疫组织化学的方法，检测CK7、CK20与TTF-1在229例腺癌及10例食管鳞癌组织中的表达。结果：3种抗体在不同组织来源肿瘤中的表达可以组合成8种模式：CK7+/CK20-/TTF-1+；甲状腺乳头状癌阳性率较高（50%）；CK7+/CK20-/TTF-1-；涎腺腺癌（94.44%）、乳腺小叶癌（85.71%）、卵巢浆液性癌（71.43%）、胆囊腺癌（63.64%）、宫颈腺癌（61.54%），阳性率较高；CK7+/CK20+/TTF-1+；阳性率均不高；CK7+/ CK20+/TTF-1-；阳性率均不高；CK7- /CK20-/TTF-1+；阳性率均不高；CK7- /CK20-/TTF-1-；阳性率均不高；CK7-/CK20+/TTF-1+；阳性率均不高；CK7-/CK20+/TTF-1-；阳性率较高；CK7-/CK20+/TTF-1+；阳性率均不高；CK7-/CK20+/TTF-1-；肠腺癌（66.67%）阳性率较高；结论：联合检测CK7、CK20及TTF-1抗体有助于判断部分转移性腺癌的器官来源，缩小原发肿瘤的搜寻范围。

关键词： [CK7](#) [CK20](#) [TTF-1](#) [转移性腺癌](#) [原发部位](#) [组织芯片](#)

Abstract: To investigate the expression of CK7, CK20, and TTF-1 in adenocarcinoma, and summarize a valuable regular pattern to determine their organic source. Methods: A total of 229 cases of adenocarcinoma tumor tissues and 10 cases of esophageal squamous cell carcinoma were studied by the tissue array technology and immunohistochemistry for CK7, CK20, and TTF-1. Results: Eight types of combined patterns of CK7, CK20, and TTF-1 in different cancers were observed: CK7+ / CK20- / TTF-1+; papillary thyroid carcinoma (50 %); CK7+ / CK20- / TTF-1-; salivary gland adenocarcinoma (94.44 %), lobular carcinoma (85.71 %), ovarian serous carcinoma (71.43 %), gallbladder adenocarcinoma (63.64 %), and cervical adenocarcinoma (61.54 %), high positive rates; CK7+ / CK20+ / TTF-1+; none had a high positive rate; CK7+ / CK20+ / TTF-1-; none had a high positive rate; CK7- / CK20- / TTF-1+; none had a positive rate; CK7- / CK20- / TTF-1-; prostate cancer (100 %), esophageal squamous cell carcinoma (100 %), and renal cell carcinoma (90.90 %), high positive rates; CK7+ / CK20+ / TTF-1+ : none had a positive rate; and CK7- / CK20+ / TTF-1-; intestinal adenocarcinoma (66.67 %), high positive rate. Conclusion: The combined detection of CK7, CK20, and TTF-1 expression helps determine the origins of some metastatic adenocarcinomas, minimize the range of such origins, and establish the groundwork for the origin evaluation of adenocarcinomas.

Key words: [CK7](#) [CK20](#) [TTF-1](#) [Metastatic carcinoma](#) [Primary site](#) [Tissue array](#)

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