



2018年11月30日 星期五

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中国肿瘤临床 > 2015, Vol. 42 > Issue (2): 82-86 DOI: doi:10.3969/j.issn.1000-8179.20141779

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112 例套细胞淋巴瘤临床病理分析*

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Clinicopathologic features of 112 patients with mantle cell lymphoma

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摘要

服务

目的: 探讨套细胞淋巴瘤(mantle cell lymphoma, MCL)的临床病理特点。方法: 收集112例MCL的临床及病理资料, 采用免疫组织化学(Envision二步法)行相关抗体标记, 荧光原位杂交技术(fluorescence in situ hybridization, FISH)对其中24例作IgH/CCND1基因断裂检测。结果: 112例(包括2例多形性和母细胞变亚型)均表达B细胞相关抗原, 94.6% (106/112)表达cyclinD1, 92.9% (104/112)表达CD5。不同免疫表型的经典型MCL的Ki-67及平均生存期无统计学差异($P>0.05$)。3例CD5-MCL未检测出IgH/CCND1基因断裂, 2例经典型MCL检测出IgH/CCND1多倍体。结论: MCL是一种具有特殊免疫表型的B细胞淋巴瘤, 多形性及母细胞变异型的预后较差, 对特殊亚型的MCL诊断有必要细分。

关键词: 套细胞淋巴瘤, 免疫组化, 荧光原位杂交, 预后

Abstract:

Objective: To explore the clinicopathologic features of 112 patients with mantle cell lymphoma (MCL). **Methods:** Data from 112 MCL cases were collected, and immunohistochemical assay was conducted. A break in the CCND1 gene was detected by fluorescence in situ hybridization (FISH). The t-test was used in the statistical analysis. **Results:** All tumor cells in the 112 cases expressed B cell-related antigen, including 1 blastoid subtype and 1 polymorphic subtype. Among all the cases, 106 expressed CD 5 and 104 expressed cyclinD1. A break in the CCND 1 gene was not found in 3 cases with CD 5-MCL. IgH/CCND 1 polyploid was found in 2 classical cases. **Conclusion:** MCL is a type of special immunophenotypic B-cell lymphoma. The prognoses of blastoid and polymorphic subtypes are poor. Special subtypes should be classified during diagnosis.

Key words: mantle cell lymphoma immunohistochemistry fluorescence in situ hybridization prognosis

收稿日期: 2014-10-23 出版日期: 2015-01-30

基金资助:

本文课题受国家临床重点专科建设项目、福建省自然基金资助项目(编号: 2012J01326)及福建省创新基金资助项目(编

号: 2012-cx-7)资助

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引用本文:

周冬梅, 陈刚, 郑雄伟, 朱伟峰, 陈宝珍. 112 例套细胞淋巴瘤临床病理分析*[J]. 中国肿瘤临床, 2015, 42(2): 82-86. Dongmei ZHOU, Gang CHEN, Xiongwei ZHENG, Weifeng ZHU, Baozhen CHEN. Clinicopathologic features of 112 patients with mantle cell lymphoma. Chinese Journal of Clinical Oncology, 2015, 42(2): 82-86.

链接本文:

<http://www.cjco.cn/CN/doi:10.3969/j.issn.1000-8179.20141779> 或 <http://www.cjco.cn/CN/Y2015/V42/I2/82>

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