

[首页](#)[期刊概况](#)[编委会](#)[专家学者](#)[网上投稿](#)[过刊浏览](#)[期刊订阅](#)[广告合作](#)

中国肿瘤临床 2012, Vol. 39 Issue (21): 1635-1638 DOI: doi:10.3969/j.issn.1000-8179.2012.21.018

[临床研究](#)[最新目录](#) | [下期目录](#) | [过刊浏览](#) | [高级检索](#)

[an error occurred while processing this directive] | [an error occurred while processing this directive]

TAM 与VEGF在外周T 细胞淋巴瘤非特指型中表达及临床意义

任旭升, 王华庆, 孟祥睿, 钱正子, 周世勇, 张会来, 邱立华, 郝希山

天津医科大学附属肿瘤医院淋巴瘤科, 天津市肿瘤防治重点实验室, 中美淋巴瘤血液诊治中心(天津市300060)

Expression and Clinical Significance of Tumor-associated Macrophages and VEGF in Peripheral T-cell Lymphoma Not Otherwise Specified

Xusheng REN, Huaqing WANG, Xiangrui MENG, Zhengzi QIAN, Shiyong ZHOU, Huilai ZHANG, Lihua QIU, Xishan HAO

Department of Lymphoma, Tianjin Medical University Cancer Institute and Hospital, Key Laboratory of Cancer Prevention and Treatment of Tianjin City, Tianjin300060, China

[摘要](#)[参考文献](#)[相关文章](#)全文: [PDF \(1929 KB\)](#) [HTML \(1 KB\)](#) 输出: [BibTeX](#) | [EndNote \(RIS\)](#) [背景资料](#)

摘要 目的: 探讨肿瘤相关巨噬细胞(TAM)及血管内皮生长因子(VEGF)在外周T细胞淋巴瘤非特指型(PTCL-NOS)中的表达及临床意义。**方法:** 采用免疫组织化学方法对60例PTCL-NOS患者肿瘤组织中的CD68和VEGF进行检测, 15例正常人淋巴结组织为对照。**结果:** 肿瘤组织中CD68阳性细胞平均含量为(56.5 ± 18.6)个/高倍镜视野, 而对照组为(12.4 ± 6.2)个/高倍镜视野(P<0.01), 肿瘤组织与对照组VEGF阳性表达率分别为78.3%和26.7%(P<0.05)。TAM与骨髓侵犯、IPI评分及疗效相关(P<0.05)。TAM高表达组和低表达组的2年总生存率(overall survival, OS)分别为23.6%和55.3%(P<0.05)。VEGF的表达与肿瘤分期、骨髓侵犯和IPI评分相关(P<0.05), VEGF表达阳性组和阴性组的2年OS分别为22.9%和83.3%(P<0.01)。单变量生存分析显示VEGF表达、TAM计数、肿瘤分期、IPI评分和疗效是独立的预后影响因素(P<0.05)。多变量分析显示VEGF和疗效是独立的预后影响因素(P<0.05)。结论: TAM和VEGF在PTCL-NOS中表达明显升高, 单因素分析显示二者是PTCL-NOS的不良预后因素。多因素分析显示仅VEGF是独立的预后影响因素。

关键词: 外周T细胞淋巴瘤 肿瘤相关巨噬细胞 血管内皮生长因子 临床意义

Abstract: Objective: This work aims to detect the expression and to identify the clinical significance of tumor-associated macrophages (TAMs) and vascular endothelial growth factor (VEGF) in peripheral T-cell lymphoma not otherwise specified (PTCL-NOS). Methods: Immunohistochemistry was used to detect CD 68 and VEGF expressions in the tumor specimens of 60 cases with PTCL-NOS. A normal lymph node biopsy was used as the control sample. Results: The average content of the CD 68-positive cells was 56.5 ± 18.6 per high-power field (HPF) in the PTCL-NOS tissues and 12.4 ± 6.2 per HPF in the control sample (P<0.01). The VEGF-positive rates for the PTCL-NOS tissues and the control sample were 78.3% and 26.7% (P<0.05), respectively. TAMs were significantly correlated to bone marrow invasion, IPI score, and treatment response (P<0.05). The two-year overall survival (OS) was 23.6% and 55.3% in the groups with high-TAM- and low-TAM-expression, respectively (P<0.05). VEGF expression was closely correlated to tumor staging, bone marrow invasion, and IPI score (P<0.05). The two-year OS was 22.9% and 83.3% in the VEGF-positive and VEGF-negative groups, respectively (P<0.01). A univariate survival analysis revealed that the VEGF expression, TAMs content, tumor staging, IPI score, and treatment response were the prognostic factors of patients with PTCL-NOS (P<0.05). A multivariate Cox regression model showed that VEGF and treatment response were both independent OS predictors (P<0.05). Conclusion: TAMs and VEGF are overexpressed in PTCL-NOS. A univariate survival analysis reveals that TAMs and VEGF are both predictive factors for PTCL-NOS prognosis, and the multivariate survival analysis using Cox regression model data show that only VEGF is an independent predictive factor for OS.

Key words: Peripheral T-cell lymphoma Tumor-associated macrophage Vascular endothelial growth factor Clinical significance

服务

[把本文推荐给朋友](#)[加入我的书架](#)[加入引用管理器](#)[E-mail Alert](#)[RSS](#)[作者相关文章](#)

引用本文:

. TAM 与VEGF在外周T 细胞淋巴瘤非特指型中表达及临床意义[J]. 中国肿瘤临床, 2012, 39(21): 1635-1638.

. Expression and Clinical Significance of Tumor-associated Macrophages and VEGF in Peripher-al T-cell Lymphoma Not Otherwise Specified[J]. Chinese Journal of Clinical Oncology, 2012, 39(21): 1635-1638.

链接本文:

http://118.145.16.228:8081/Jweb_zgzllc/CN/doi:10.3969/j.issn.1000-8179.2012.21.018 或 http://118.145.16.228:8081/Jweb_zgzllc/CN/Y2012/V39/I21/1635

没有本文参考文献

- [1] 李 慧. 肿瘤干细胞对肿瘤血管生成的作用及调控机制的最新研究进展[J]. 中国肿瘤临床, 2012, 39(9): 493-496.
- [2] 刘 艳,高 健,师宜荃,赵立武,刘易欣. **CD105**和**VEGF**在水泡状胎块中的表达及意义[J]. 中国肿瘤临床, 2012, 39(9): 570-573.
- [3] 唐海林, 苏 坚, 邓 敏, 曾 希, 廖前进, 周秀田, 苏 琦. 胃癌组织中**miR-222**与**TIMP3**的表达及临床意义[J]. 中国肿瘤临床, 2012, 39(4): 194-196.
- [4] 肖炜明, 吴克艳, 龚卫娟, 卜 平. 胃癌组织中**CD1a**
+
树突状细胞密度和血管内皮生长因子表达及其相关性*[J]. 中国肿瘤临床, 2012, 39(21): 1623-1625.
- [5] 李少雷,陈晋峰,郑庆锋,吴 楠,阎 石,王 洋,张建芝,杨 跃. **Endostatin VEGF-C**和**VEGFR-3**在非小细胞肺癌及其淋巴结组织中的表达与意义[J]. 中国肿瘤临床, 2011, 38(24): 1519-1523.
- [6] 顾 涛,付占昭,付宝红,张丽娜,张绍华,曹晓艳. 非小细胞肺癌高龄患者三维适形放疗前后血清**VEGF TGF-β1**水平的测定及临床意义[J]. 中国肿瘤临床, 2011, 38(24): 1565-1567.
- [7] 史东升,周静敏,马淑萍. 紫龙金对人非小细胞肺癌**A549**细胞生长及**VEGF**表达的影响[J]. 中国肿瘤临床, 2011, 38(20): 1267-1270.
- [8] 解瑞玲, 董伯升, 庞慧, 成媛, 尚可, 李醒亚. 肿瘤相关巨噬细胞在食管癌组织浸润及其对预后的影响[J]. 中国肿瘤临床, 2011, 38(2): 83-86 .
- [9] 廖红,蔡绮纯,王潇潇,蔡清清,林旭滨,林桐榆,夏忠军,李志铭,姜文奇,黄慧强. **CD68**阳性淋巴瘤相关巨噬细胞在滤泡性淋巴瘤间质中的密度及其临床意义的研究[J]. 中国肿瘤临床, 2011, 38(18): 1081-1084.
- [10] 王立平,赵杨,申兴斌. **PTTG**、**bFGF**、**VEGF-C**在大肠正常黏膜及大肠癌中的表达及意义[J]. 中国肿瘤临床, 2011, 38(13): 788-792.

友情链接



版权所有 ©2013 《中国肿瘤临床》编辑部

地址: 天津市河西区体院北环湖西路肿瘤医院内 300060

电话/传真: (022)23527053 E-mail: cjco@cjco.cn cjcotj@sina.com 津ICP备1200315号