



[首页](#)

[期刊概况](#)

[编委会](#)

[期刊内容](#)

[特邀审稿](#)

[投稿指南](#)

[出版发行](#)

311-314. 胃癌患者外周血和肿瘤组织中高表达CD14⁺DR^{low/-}髓源抑制性细胞[J]. 戴夫, 李海文, 彭琼, 林沪, 李元元. 中国肿瘤生物治疗杂志, 2011, 18(3)

[胃癌患者外周血和肿瘤组织中高表达CD14⁺DR^{low/-}髓源抑制性细胞](#) [点此下载全文](#)

[戴夫](#) [李海文](#) [彭琼](#) [林沪](#) [李元元](#)

安徽医科大学 第三附属医院 消化内科, 安徽 合肥 230061; 安徽医科大学 第三附属医院 消化内科, 安徽 合肥 230061; 安徽医科大学 第三附属医院 消化内科, 安徽 合肥 230061; 解放军第302医院 生物治疗中心, 北京 100039; 解放军第302医院 生物治疗中心, 北京 100039

基金项目: 国家“十一五”科技重大专项 (No. 2009ZX10004-309)

DOI:

摘要:

目的: 探讨胃癌患者外周血和肿瘤组织中CD14⁺DR^{low/-}髓源抑制性细胞 (myeloid-derived suppressor cells, MDSCs) 的表达及其与肿瘤病理特征的关系。方法: 选取2009年3月至2010年10月安徽医科大学第三附属医院胃癌患者43例 (I期9例、II期13例、III期14例、IV期7例), 另采集26例正常健康者作为对照组。流式细胞术检测外周血、瘤组织中CD14⁺DR^{low/-}MDSCs的表达水平, 分析MDSCs表达水平与肿瘤病理特征的相关性。结果: 胃癌患者肿瘤组织中CD14⁺DR^{low/-}MDSCs的表达显著高于自身外周和健康对照者外周血的表达水平 $[2.87 \pm 1.93]%$ vs $[2.37 \pm 1.7]%$, $[0.89 \pm 0.47]%$, $P < 0.05$ 和 $P < 0.01$], 后者间差异也有统计学意义 ($P < 0.01$)。CD14⁺DR^{low/-}MDSCs的表达与胃癌的恶性程度呈正相关, 晚期胃癌组织内CD14⁺DR^{low/-}MDSCs表达明显增加 (I: II: III: IV = $(1.15 \pm 0.78)%$: $(1.71 \pm 0.92)%$: $(2.25 \pm 1.24)%$: $(4.85 \pm 2.37)%$, $P < 0.05$)。同时, 肿瘤浸润组织与非浸润组织的CD14⁺DR^{low/-}MDSCs表达也有明显差异 $[3.90 \pm 1.67]%$ vs $[2.62 \pm 1.53]%$, $P < 0.05$]。结论: CD14⁺DR^{low/-}MDSCs在胃癌患者外周血和肿瘤组织中均高表达, 与胃癌的发生、发展密切相关。

关键词: [胃癌](#) [髓源的抑制性细胞](#) [免疫逃逸](#) [免疫监视](#)

Myeloid-derived suppressor cells highly expressing CD14⁺DR^{low/-} in peripheral blood and tumor tissues of stomach carcinoma [Download Fulltext](#)

[DAI Fu](#) [LI Hai-wen](#) [PENG Qiong](#) [LIN Hu](#) [LI Yuan-yuan](#)

Department of Gastroenterology, Third Affiliated Hospital of Anhui Medical University, Hefei 230061, Anhui, China; Department of Gastroenterology, Third Affiliated Hospital of Anhui Medical University, Hefei 230061, Anhui, China; Department of Gastroenterology, Third Affiliated Hospital of Anhui Medical University, Hefei 230061, Anhui, China; Research Center for Biological Therapy, No. 302 Hospital of PLA, Beijing 100039, China; Research Center for Biological Therapy, No. 302 Hospital of PLA, Beijing 100039, China

Fund Project: Project supported by the National "the Eleventh Five-Year" Major Science and Technology Program of China (No. 2009ZX10004-309)

Abstract:

Objective: To investigate the expression of CD14⁺DR^{low/-} myeloid-derived suppressor cells (MDSCs) in peripheral blood and tumor tissues of gastric carcinoma (GC) patients and its relationship with clinicopathological of GC. Methods: Forty-three stomach carcinoma patients (9 stage I, 13 stage II, 14 stage III, 7 stage IV) were selected from Third Affiliated Hospital of Anhui Medical University (Mar. 2009 to Oct. 2010), and 26 healthy volunteers were used as control. CD14⁺DR^{low/-} MDSCs expression in peripheral blood and gastric carcinoma tissues was detected by flow cytometry, and its relationship with clinicopathological of GC was analyzed. Results: CD14⁺DR^{low/-} MDSCs expression in GC tissues was significantly higher than those in peripheral blood of GC patients and healthy controls $[2.87 \pm 1.93]%$ vs $[2.37 \pm 1.7]%$, $[0.89 \pm 0.47]%$, $P < 0.05$ and $P < 0.01$, and CD14⁺DR^{low/-} MDSCs expression in peripheral blood of GC patients was also higher than that in healthy controls ($P < 0.01$). CD14⁺DR^{low/-} MDSCs expression in GC tissues was positively correlated with their malignancy stage and significantly increased in advanced GC (stage I: II: III: IV = $(1.15 \pm 0.78)%$: $(1.71 \pm 0.92)%$: $(2.25 \pm 1.24)%$: $(4.85 \pm 2.37)%$, $P < 0.05$). Meanwhile, CD14⁺DR^{low/-} MDSCs expression in tumor infiltration tissues was significantly higher than that in tumor un-infiltration tissues $[3.90 \pm 1.67]%$ vs $[2.62 \pm 1.53]%$, $P < 0.05$). Conclusion: CD14⁺DR^{low/-} MDSCs are highly expressed in peripheral blood and GC tissues, and relates to the development and progression of GC.

Keywords: [gastric carcinoma](#) [myeloid-derived suppressor cells](#) [immune escape](#) [immune surveillance](#)

[查看全文](#) [查看/发表评论](#) [下载PDF阅读器](#)

Copyright © Biother.Org™ All Rights Reserved

主管单位: 中国科学技术协会 主办单位: 中国免疫学会、中国抗癌学会
地址: 上海市杨浦区翔殷路800号 邮政编码: 200433 京ICP备06011393号-2
本系统由北京勤云科技发展有限公司设计