中南大学学报(医学版) 2010, 35(9) 964- DOI: 10.3969/j.issn.1672-

7347.2010. ISSN: 1672-7347 CN: 43-1427/R

本期目录 | 下期目录 | 过刊浏览 | 高级检索

[打印本页] [关闭]

OX40/OX40L mRNA在实验性变态反应性

神经炎中的动态变化

张宁, 李刚, 肖波, 刘运海, 蔡艳, 孙新刚, 梁静慧

中南大学湘雅医院神经内科, 长沙 410008

摘要:

目的:研究Oxford 40(OX40)和Oxford 40 ligand(OX40L)mRNA在实验性变态反应性神经炎(experimental allegic neuritis, EAN)大鼠坐骨神经、脾脏、外周血和淋巴结中的动态变化。 方法: 36只Lewis大鼠随机分为EAN 模型组和完全弗氏佐剂对照组(CFA组)。分别在第9天、第17天、第26天处死动物,采用反转录聚合酶链反应技术 检测坐骨神经根、脾脏、外周血单个核细胞和淋巴结中OX40和OX40L mRNA的表达水平。结果: EAN组大鼠在抗 ▶加入我的书架 原免疫后第17天达到发病高峰,OX40和OX40L mRNA在第9天(发病早期)和第17天时表达均较高,与第26天 (恢复期)相比,差异有统计学意义(P<0.05),各时间点与对照组相比差异有统计学意义(P<0.05);CFA 组大鼠无症状,EAN组中OX40和OX40LmRNA在坐骨神经和淋巴结中各时间点表达均升高,在外周血单个核细胞 中微量表达。结论: OX40/OX40L可能与EAN发病有关。

关键词: 实验性变态反应性神经炎 吉兰-巴雷综合征 OX40: OX40L

Dynamic change OX40/OX40L mRNA in experimental allergic neuritis

ZHANG Ning, LI Gang, XIAO Bo, LIU Yunhai, CAI Yan, SUN Xingang, LI ANG Jinghui

Department of Neurology, Xiangya Hospital, Central South University, Changsha 410008, China

Abstract:

ObjectiveTo examine the expression of mRNA of Oxford 40(OX40) and Oxford 40 ligand(OX40L) in the sciatic nerve, spleen, peripheral blood mononuclear cells and lymph nodes of experimental allegic neuritis (EAN). MethodsThirty-six Lewis rats were randomly assigned into an EAN group and a CFA group. The rats were sacrificed on 9th, 17th, and 26th day after immunization. OX40 and OX40L mRNA was detected by reverse transcription polymerase chain reaction in the spleen, sciatic nerves, peripheral blood mononuclear cells and lymphonodes. ResultsThe peak of clinical course came on 17th day after the immuuization in EAN. The mRNA expression of OX40/OX40L was higher on 8th day and 17th day than that on 26th day after the immuuization (P < 0.05). There was significant difference between the EAN group and the CFA group at the 3 time points (P<0.05); rats in the CFA group didn't have any clinical manifestations. The mRNA expression of OX40 and OX40L in the EAN group raised in the sciatic nerves and lymph nodes at the above 3 time points (P<0.05). Weak expression was seen in the peripheral blood mononuclear cells. ConclusionOX40 and OX40L may play a role in the pathogenesis of experimental allegic neuritis.

Keywords: experimental allegic neuritis; Guillain-Barre's syndrome; oxford40; oxford 40 ligand

收稿日期 2009-06-09 修回日期 网络版发布日期

DOI: 10.3969/j.issn.1672-7347.2010.

基金项目:

通讯作者: 张宁

作者简介:

作者Email: zhnxyyy63@vip.sina.com

参考文献:

- [1] Croft M, So T, Duan W, et al. The significance of OX40 and OX40L to T-cell biology and immune disease [J] .Immunol Rev, 2009,229(1):173-191.
- [2] Stoll G. Inflammatory cytokines in the nervous system: multifunctional mediators in autoimmunity and cerebral ischemia [J] . Rev Neurol, 2002,158(10Pt1): 887-891.
- [3] Maurer M, Toyka K V, Gold R. Cellular immunity in inflammatory autoimmune neuropathies [J] .

扩展功能

本文信息

- ▶ Supporting info
- PDF(1220KB)
- ▶ [HTML全文]
- ▶参考文献[PDF]
- ▶ 参考文献

服务与反馈

- ▶把本文推荐给朋友
- ▶加入引用管理器
- ▶ 引用本文
- Email Alert
- ▶ 文章反馈
- ▶浏览反馈信息

本文关键词相关文章

- >实验性变态反应性神经炎
- ▶吉兰-巴雷综合征
- ▶ OX40: OX40L

本文作者相关文章 PubMed

- Rev Neurol, 2002, 158 (12 Pt 2): S7-S15.
- [4] Croft M, So T, Duan W, et al. The significance of OX40 and OX40L to T-cell biology and immune disease [J]. Immunol Rev, 2009, 229(1):173-191.
- [5] Gramaglia I, Jember A, Pippig SD, et al. The OX40 costimulatory receptor determines the development of CD4 memory by regulating primary clonal expansion [J]. J Immunol, 2000,165 (6):3043-3050.
- [6] Weatherill A R, Maxwell J R, Takahashi C, et al. OX40 ligation enhances cell cycle turnover of Agactivated CD4+ T cells in vivo [J]. Cell Immunol, 2001, 209(1):63-75.
- [7] Rogers P R, Song J, Gramaglia I, et al. OX40 promotes Bcl-xl and Bcl-2 expression and is essential for long term survival of CD4+T cells [J]. Immunity, 2001,15(3):445-455.
- [8] Lee S W, Park Y, Song A, et al. Functional dichotomy between OX40 and 4-1BB in modulating effector CD8+ T cell responses [J]. J Immunol, 2006, 177(7): 4464-4472.
- [9] Ruby CE, Redmond W L, Haley D, et al. Anti-OX40 stimulation in vivo enhances CD8+ memory T cell survival and significantly increases recall responses [J]. Eur J Immunol, 2007,37(1): 157-166.
- [10] Bandyopadhyay S, Long M, Qui H Z, et al. Self-antigen prevents CD8 T cell effector differentiation by CD134 and CD137 dual costimulation [J] .J Immunol,2008,181(11):7728-7737.
- [11] Izmailova E, Walker R, Fitzgerald M, et al. Quantitation of peripheral blood markers of rat experimental autoimmune encephalomyelitis [J]. Autoimmunity, 2007, 40(5):355-365.
- [12] Elyaman W, Kivis kk P, Reddy J, et al. Distinct functions of autoreactive memory and effector CD4+ T cells in experimental autoimmune encephalomyelitis [J] .Am J Pathol, 2008,173(2):411-422.
- [13] Carbonia S,Aboul-Eneinb F, Waltzingera C, et al.CD134 plays a crucial role in the pathogenesis of EAE and is upregulated in the CNS of patients with multiple sclerosis [J] . J Neuroimmunol,2003,145 (1-2):1-11
- [14] Ndhlovu L C, Ishii N, Murata K, et al. Critical involvement of OX40 ligand signals in the T cell priming events during experimental autoimmune encephalomyelitis [J] . J Immunol,2001,167(5):2991-2000
- [15] Nohara C, Akiba H, Nakajima A,et al. Amelioration of experimental autoimmune encephalomyelitis with anti-OX40 ligand monoclonal antibody: a critical role for OX40 ligand in migration, but not development, of pathogenic T cells [J]. J Immunol, 2001,166(3):2108-2115.

本刊中的类似文章

Copyright by 中南大学学报(医学版)