

论著

肢体缺血预处理对脂多糖诱导大鼠急性肺损伤的保护作用

宋智<sup>1</sup>, 罗万俊<sup>2</sup>, 秦岭<sup>3</sup>, 陈胜喜<sup>2</sup>

中南大学1.湘雅三医院普外三科,长沙 410013; 2.湘雅医院心胸外科,长沙 410008; 3.湘雅医院呼吸内科,长沙 410008

摘要:

目的: 研究非创伤性肢体缺血预处理(non-wounded limb ischemic preconditioning, N-LIP)对脂多糖诱导大鼠急性肺损伤(acute lung injury, ALI)的保护作用。方法: 将15只SD雌性大鼠随机分为对照组、急性肺损伤组(ALI组)、急性肺损伤+非创伤性双下肢缺血预处理组(ALI+N-LIP组), 检测N-LIP后ALI大鼠肺功能的变化、支气管肺泡灌洗液(bronchoalveolar lavage fluid, BALF)中白细胞数量和乳酸脱氢酶(lactate dehydrogenase, LDH)水平, 血清中超氧化物歧化酶(superoxide dismutase, SOD)活性和丙二醛(malondialdehyd, MDA)含量, 免疫组织化学方法检测肺组织内肺泡表面活性物质-A(pulmonary surfactant-associated protein A, SP-A)表达水平, HE染色观察肺组织病理改变。结果: 乙酰甲胆碱(methacholine, Mch)激发后ALI+N-LIP组大鼠气道阻力(airway resistance, AR)的增高程度(P<0.01)、动态顺应性(dynamic compliance, Cdyn)的减弱程度(P<0.01)均明显小于ALI组; ALI+N-LIP组BALF中的白细胞数和LDH含量均明显小于ALI组(P<0.05); ALI+N-LIP组血清中SOD活力高于ALI组(P<0.05), MDA含量明显小于ALI组(P<0.05); 免疫组织化学显示: ALI+N-LIP组肺组织中SP-A含量最高, 其次为对照组, ALI组肺组织内SP-A含量最低; 肺部HE染色显示: ALI+N-LIP组损伤程度明显轻于ALI组, 但较对照组要严重。结论: N-LIP对脂多糖诱导的ALI有保护作用, 其作用机制可能与其促进SP-A表达和抗氧化作用有关。

关键词: 急性肺损伤 脂多糖 非创伤性肢体缺血预处理 肺保护 肺泡表面活性物质相关蛋白-A

Protective effect of limb ischemic preconditioning on acute lung injury induced by lipopolysaccharide in rats

SONG Zhi<sup>1</sup>, LUO Wanjun<sup>2</sup>, QIN Ling<sup>3</sup>, CHEN Shengxi<sup>2</sup>

1.Third Department of General Surgery, Third Xiangya Hospital, Central South University, Changsha 410013;  
2.Department of Cardiothoracic Surgery, Xiangya Hospital, Central South University, Changsha 410008;  
3.Department of Respiratory, Xiangya Hospital, Central South University, Changsha 410008, China

Abstract:

Objective To explore the protective effect of noninvasive limb ischemic preconditioning (N-LIP) on acute lung injury (ALI) induced by lipopolysaccharide (LPS) in rats. Methods Fifteen female SD rats were randomly divided into a control group, an acute lung injury group (ALI group), an acute lung injury and noninvasive limb ischemic preconditioning group (ALI+N-LIP group). After ALI rats were treated with N-LIP, the changes of airway resistance (AR) and dynamic compliance (Cdyn) were tested by invasive pulmonary function system and recorded. Blood samples and bronchoalveolar lavage fluid (BALF) were collected, the amounts of white blood cell (WBC) in BALF were counted by cytometry, and the level of lactate dehydrogenase (LDH) in BALF was also examined by automatic biochemistry analyzer. The level of serum superoxide dismutase (SOD) and malondialdehyd (MDA) was examined by chromatometry. The lung tissues were acquired to observe the expression of pulmonary surfactant-associated protein-A (SP-A) and pathological changes. Results After being stimulated by methacholine (Mch), the increasing rate of AR and decreasing rate of Cdyn in the ALI+N-LIP group were less than those in the ALI group (P<0.01). The levels of WBC and LDH in BALF in the ALI+N-LIP group were much lower than those in the ALI group (P<0.05). Meanwhile, the activity of serum SOD in the ALI+N-LIP group was higher, and the level of serum MDA was lower than that in the ALI group (P<0.05). The expression of SP-A in the lung tissue in the ALI+N-LIP group was the highest in the 3 groups, while that in the ALI group was the weakest (P<0.01). Injury of the lung tissue in the ALI+N-LIP group was less than that in the ALI group, but more severe than that in the control group. Conclusion N-LIP has protective effect on acute lung injury induced by LPS in rats. The possible mechanism is related to improving the secretion of SP-A and antioxidation.

Keywords: acute lung injury; lipopolysaccharide; noninvasive limb ischemic preconditioning; pulmonary protection; pulmonary surfactant-associated protein-A

收稿日期 2009-10-29 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: 罗万俊

作者简介:

作者Email: Luo3478@yahoo.cn

参考文献:

- [1] Ng C S, Wang S, Yim A P, et al. Pulmonary dysfunction after cardiac surgery [J]. Chest, 2002, 121 (4): 1269-1277.
- [2] Sievers H H, Freund-Kaas C, Eleftheriadis S, et al. Lung protection during total cardiopulmonary bypass by isolated

扩展功能

本文信息

- Supporting info
- PDF(1637KB)
- [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
- 加入我的书架
- 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

- 急性肺损伤
- 脂多糖
- 非创伤性肢体缺血预处理
- 肺保护
- 肺泡表面活性物质相关蛋白-A

本文作者相关文章

PubMed

lung perfusion: preliminary results of a novel perfusion strategy [J]. *Ann Thorac Surg*, 2002, 74 (4): 1167-1172.

[3] 李国虎, 陈胜喜, 罗万俊, 等. 体外循环心脏直视手术中心脏缺血预处理对肺的保护作用 [J]. *湖南医科大学学报*, 1998, 23 (1): 41-43.

LI Guohu, CHEN Shengxi, LUO Wanjun, et al. The protective effects of cardiac ischemic preconditioning on lung in cardiac operation with cardiopulmonary bypass [J]. *Bulletin of Hunan Medical University*, 1998, 23 (1): 41-43.

[4] 张春芳, 陈胜喜, 郭海周, 等. 缺血预处理对猪肺隔离药物灌注肺损伤的保护作用 [J]. *中南大学学报: 医学版*, 2005, 30 (1): 64-67.

ZHANG Chunfang, CHEN Shengxi, GUO Haizhou, et al. Protective mechanism of ischemic preconditioning to the lung ischemia-reperfusion injury [J]. *Journal of Central South University. Medical Science*, 2005, 30 (1): 64-67.

[5] Kodavanti U P, Hauser R, Christiani D C, et al. Pulmonary responses to oil fly ash particles in the rat differ by virtue of their specific soluble metals [J]. *Toxicol Sci*, 1998, 43 (2): 204-212.

[6] Wolfrum S, Nienstedt J, Heidbreder M, et al. Calcitonin gene related peptide mediates cardioprotection by remote preconditioning [J]. *Regul Pept*, 2005, 127 (1/3): 217-224.

[7] Weinbrenner C, Schulze F, Sarvary L, et al. Remote preconditioning by infrarenal aortic occlusion is operative via  $\delta 1$ -opioid receptors and free radicals in vivo in the rat heart [J]. *Cardiovasc Res*, 2004, 61 (3): 591-599.

[8] Li G, Labruto F, Sirsjo A, et al. Myocardial protection by remote preconditioning: the role of nuclear factor kappa-B p105 and inducible nitric oxide synthase [J]. *Eur J Cardiothorac Surg*, 2004, 26 (5): 968-973.

[9] Brzozowski T, Konturek P C, Konturek S J, et al. Ischemic preconditioning of remote organs attenuates gastric ischemia-reperfusion injury through involvement of prostaglandins and sensory nerves [J]. *Eur J Pharmacol*, 2004, 499 (1/2): 201-213.

[10] Konstantinov I E, Arab S, Li J, et al. The remote ischemic preconditioning stimulus modifies gene expression in mouse myocardium [J]. *J Thorac Cardiovasc Surg*, 2005, 130 (5): 1326-1332.

[11] Jones S P, Girod W G, Palazzo A J, et al. Myocardial ischemia reperfusion injury is exacerbated in absence of endothelial cell nitric oxide synthase [J]. *Am J Physiol*, 1999, 276 (5Pt2): 1567-1573.

[12] Kuroki Y, Takahashi H, Chiba H, et al. Surfactant proteins A and D: disease markers [J]. *Biochim Biophys Acta*, 1998, 1408 (2/3): 334-345.

[13] Jobe A H, Ikegami M. Biology of surfactant [J]. *Clin Perinatol*, 2001, 28 (3): 655-669.

[14] Poynter S E, LeVine A M. Surfactant biology and clinical application [J]. *Crit Care Clin*, 2003, 19 (3): 459-472.

[15] Lyra P P, Diniz E M. The importance of surfactant on the development of neonatal pulmonary diseases [J]. *Clinics (Sao Paulo)*, 2007, 62 (2): 181-190.

[16] Wright J R, Youmans D C. Degradation of surfactant lipids and surfactant protein A by alveolar macrophages in vitro [J]. *Am J Physiol*, 1995, 268 (5 Pt 1): 772-780.

[17] Bates S R, Fisher A B. Surfactant protein A is degraded by alveolar macrophages [J]. *Am J Physiol*, 1996, 271 (2 Pt 1): 258-266.

[18] Goss C H, Brower R G, Hudson L D, et al. Incidence of acute lung injury in the United States [J]. *Crit Care Med*, 2003, 31 (6): 1607-1611.

[19] Gunther A, Ruppert C, Schmidt R, et al. Surfactant alteration and replacement in acute respiratory distress syndrome [J]. *Respir Ras*, 2001, 2 (6): 353-364.

#### 本刊中的类似文章

1. 金丽艳, 徐军美, 贺志鹰, 阮文燕, 柴湘平. 乌司它丁对油酸致急性肺损伤大鼠肺组织血红素氧化酶-1的影响[J]. *中南大学学报(医学版)*, 2007, 32(04): 675-678

2. 刘敬臣, 王海棠, 王维. 丙氨酰谷氨酰胺对脓毒症大鼠肺损伤的保护作用[J]. *中南大学学报(医学版)*, 2008, 33(12): 1095-1100

3. 王瑞珂, 赵双平, 杨新平, 蔡宏伟\*. 乌司他丁在酸吸入性急性肺损伤早期治疗中的作用[J]. *中南大学学报(医学版)*, 2004, 29(3): 305-308

4. 赵双平\*, 郭曲练, 王瑞珂, 王锴. 沐舒坦对盐酸吸入后大鼠氧化抗氧化系统的影响[J]. *中南大学学报(医学版)*, 2004, 29(5): 586-588

5. 赵双平, 郭娇, 郭曲练, 张重, 叶治. 不同浓度七氟醚预处理对内毒素性急性肺损伤大鼠肺组织的影响[J]. *中南大学学报(医学版)*, 2010, 35(9): 921-