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论文

大鼠肾脏损伤早期HIF-1a、PPAR-y的表达及其对肾脏的保护机制

王芳1, 王东3, 蒙伶俐2, 王晓梅1, 孙少华1

- 1. 兰州大学第一医院病理研究所, 兰州 730000; 2. 宁波市第一医院病理科, 浙江 宁波 315000;
- 3. 兰州大学第一医院心血管内科, 兰州 730000

摘要:

目的 探讨经体表创伤后肾脏组织中缺氧诱导因子-1a(HIF-1a)、过氧化物酶体增殖物激活受体γ(PPAR-γ)表达的变化,及创伤后肾脏损伤与修复的作用机制。方法 采用自由落体生物撞击仪撞击大鼠脊肋区复制创伤动物模型。实验大鼠分为5组,包括非创伤对照组、创伤后1、6、12、24h组。采用免疫组织化学方法进行HIF-1a、PPAR-γ染色。结果 肾脏创伤后1、24h HIF-1a表达增强,分布于皮质远曲小管、肾盏旁小管、髓质小管;6、12h表达减弱,局限于肾盏旁小管、髓质外带小管。各组间比较差异有统计学意义(P<0.05)。PPAR-γ 1、24h呈阳性表达,分布于髓质小管上皮细胞;6、12h呈阴性表达,各组间比较差异有统计学意义(P<0.05)。结论HIF-1a、PPAR-γ可能参与了肾脏创伤后缺血、缺氧、再生、修复的过程。

关键词: 肾脏;创伤;缺氧诱导因子-1α;过氧化物酶体增殖物激活受体γ,大鼠,Wistar

Expressions of HIF-1 α and PPAR- γ in renal early trauma and their protective mechanisms on the kidney

WANG Fang1, WANG Dong3, MENG Ling-li2, WANG Xiao-mei1, SUN Shao-hua1

- 1. Institute of Pathology, The First Hospital of Lanzhou University, Lanzhou 730000, China;
- 2. Department of Pathology, The First Hospital of Ningbo, Ningbo 315000, Zhejiang, China;
- 3. Department of Cardiology, The First Hospital of Lanzhou University, Lanzhou 730000, China Abstract:

Objective To investigate expressions of hypoxia- inducible factor-1α(HIF-1α) and peroxisome proliferator-activated receptor-gamma(PPAR-γ) in traumatic kidneys, and the mechanisms of kidney trauma and repair. Methods Rats were stricken at the skin of the renal zone by a free-fall from 45cm height to establish the traumatic rat model. Then, rats were randomly divided into 5 groups: the non-traumatic control group, and 1h, 6h, 12h and 24h post-trauma groups. Expressions of HIF-1α and PPAR-γwere detected with immunohistochemical staining. Results Positive expression of HIF-1αwas increased in 1h and 24h post-trauma groups, located in cortex distal convoluted tubules, tubules near renal calyces and medullary tubules; while it was distinctly decreased in 6h and 12h post-trauma groups, located in tubules near renal calyces and tubules at the outer zone of renal medulla. There were significant differences in expression of HIF-1αamong all the groups(P<0.05). The PPAR-γ expression was positive in tubules of renal medulla in 1h and 24h post-trauma groups(P<0.05), however, negative in 6h and 12h post-trauma groups. Conclusion HIF-1αand PPAR-γ may participate in the secondary hypoxia, ischemia, regeneration and repair procedures after renal injury.

Keywords: Kidney; Trauma; Hypoxia inducible factor 1a; Peroxisome proliferator-activated receptorgamma; Rats, wistar

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通讯作者: 孙少华(1957-) 男,教授,主要从事创伤病理研究。E mail: sunshaohua2006 @sohu. com

作者简介: 王芳(1978-), 女,主治医师,硕士研究生。

作者Email:

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