中国公共卫生 2014, 30(2) 206-208 DOI: 10.11847/zgggws2014-30-02-24 ISSN: 1001-

0580 CN: 21-1234/R

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论文

2,4-二羟基二苯甲酮对DMN致小鼠急性肝毒性保护作用

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摘要:

目的 探索2,4-二羟基二苯甲酮(BP-1)对二甲基亚硝胺(DMN)所致小鼠急性肝损伤的保护作用。方法 将体重18~22 g的ICR小鼠随机分为对照组、模型组、BP-1低、中、高组(200、400、800 mg/kg)灌胃给药4 d,末次给药30 min后腹腔注射给予DMN 22 mg/kg,24 h处死,测定谷丙转氨酶(ALT),谷草转氨酶(AST),乳酸脱氢酶(LDH)活性;测定肝脏还原性谷胱甘肽(GSH),氧化型谷胱甘肽(GSSG)和丙二醛(MDA)含量;观察肝脏病理组织学变化。结果 与对照组比较,模型组小鼠血清ALT、AST、LDH活性均明显升高(P<0.01),MDA含量[(0.256±0.059) μmol/g]升高(P<0.01),肝小叶出现大量坏死细胞;与模型组比较,BP-1各剂量组小鼠血清ALT、AST、LDH水平均明显下降(P<0.05),肝组织中GSH/GSSG比值升高(P<0.01),高剂量BP-1组小鼠肝脏MDA含量[(0.062±0.034) μmol/g]明显下降(P<0.01),肝小叶坏死区域减少。结论 2,4-二羟基二苯甲酮对DMN所致小鼠急性肝损伤有保护作用。

关键词: 2,4-二羟基二苯甲酮(BP-1) 肝脏毒性 二甲基亚硝胺(DMN) 保肝作用

Protective effect of 2,4-dihydroxybenzophenone on DMN induced acute hepatotoxicity in mice

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Abstract:

Objective To investigate the protective effect of 2,4-dihydroxybenzophenone (BP-1) on acute hepatotoxicity induced by dimethylnitrosamine(DMN) in mice. Methods Twenty-five male ICR mice with body weight of 18-22 g were divided into control group, model group, low, moderate, and high dose BP-1 exposure group (200,400,and 800 mg/kg).BP-1 was administrated to the mice in exposure groups for 4 days and DMN was injected into the mice 30 min after BP-1 administration on the 4th day. Twenty-four hours after the treatment, all mice were killed and activities of serum alanine aminotransferase (ALT), aspartate aminotransferase (AST), and lactate dehydrogenase (LDH) were measured. The contents of reduced glutathione (GSH), oxidized glutathione (GSSG), and malonaldehyde (MDA) were also determined. Histopathological changes of liver were observed. Results Serum ALT, AST and LDH activities of DMN group were significantly increased compared with those of the vehicle control group (P<0.01 for all). The content of MDA was increased to $0.256\pm0.059 \,\mu\text{mol/g}$ (P<0.01). Histopathological observations revealed a severe injury of liver. Serum ALT, AST and LDH activities of BP-1 group were significantly decreased compared with those of the model group (P<0.05 for all). The ratio of GSH/GSSG was increased and the content of MDA was decreased(0.062 ± 0.034 µmol/g, P<0.01). Histopathological observations also revealed an ameliorated change of liver. Conclusion The results indicate that 2,4dihydroxybenzophenone has protective effect on acute hepatotoxicity induced by DMN in mice.

Keywords: 2,4-dihydroxybenzophenone hepatic toxicity protective effect on DMN liver

收稿日期 2013-03-26 修回日期 网络版发布日期 2013-06-06

DOI: 10.11847/zgggws2014-30-02-24

基金项目:

重大新药创制科技重大专项(2009ZX09103-007)

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