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[1]陈琰琰,朱慧芬,杨道锋,等.HBx及其突变体X17-3对Hippo信号途径表达的影响[J].第三军医大学学报,2013,35(16):1704-1707. Chen Yanyan,Zhu Huifen,Yang Daofeng,et al.HBx and its mutant X17-3 regulates YAP through Hippo signaling pathway in human hepatic L02 cells[J].J Third Mil Med Univ,2013,35(16):1704-1707.

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HBx及其突变体X17-3对Hippo信号途径表达到:

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Title: HBx and its mutant X17-3 regulates YAP through Hippo

signaling pathway in human hepatic L02 cells

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摘要: 目的 初步探寻HBx及其不同突变体表达与Hippo信号途径的关系,

以及对细胞凋亡的影响。 方法 HBx及其突变体载体转染人正常 FH细胞系L02,转染48 h后,提取总蛋白,Western blot检测细胞中 Hippo信号途径MST1、YAP(yes-associated protein)的表达,磷酸化和

去磷酸化的情况。采用Annexin V/PI标记流式细胞术检测细胞凋

亡。 结果 HBx及其突变体载体转染L02细胞48 h后, Western

导航/NAVIGATE

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blot结果显示细胞MST1表达下调,p-MST1/2表达上升(P<0.05); YAP表达上升,p-YAP表达下调(P<0.05)。Annexin V/PI标记法流式 细胞术结果显示HBx及其突变体能促进L02细胞发生凋亡(P<0.05),其 介导的细胞早期凋亡尤为明显。 结论 HBx通过Hippo信号通路 途径调控下游致癌基因YAP的表达。结合HBx介导的L02细胞凋亡这一结 果说明HBx可能通过多种途径调节细胞周期。

Abstract:

Objective To investigate the relationship of HBx and its mutant X17-3 with Hippo signaling pathway, and the effects of HBx on cell cycle and apoptosis. Methods L02 cells were transfected with HBx and its mutant, respectively. After 48 h of incubation, the expression levels as well as the phosphorylation status of MST1/2 and yes-associated protein (YAP) were detected by