

论著

利用补丁甲基化技术构建特殊报告载体检测DNA甲基化对FCER1G基因启动子的调控活性

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摘要: 目的: 构建一种特殊的荧光素报告载体检测DNA甲基化对FCER1G基因启动子调控序列的调控活性。方法: 用补丁甲基化技术构建特殊的含完全甲基化和模拟甲基化的FCER1G启动子调控序列的PGL-3荧光素报告载体; 通过比较完全甲基化与模拟甲基化荧光素报告载体活性来检测DNA甲基化对FCER1G基因启动子调控序列的调控活性。结果: 成功构建特殊的完全甲基化与模拟甲基化FCER1G启动子调控序列荧光素报告载体; 并检测出完全甲基化与模拟甲基化荧光素报告载体活性比为(0.36±0.07):1 ($P<0.001$)。结论: FCER1G启动子调控序列是甲基化敏感位点, FCER1G启动子受DNA甲基化调控。

关键词: 补丁甲基化 FCER1G基因 启动子调控序列 DNA甲基化

Construction of special reporter to detect DNA methylation regulatory activity in FCER1G gene promoter through patch-methylation

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Abstract: Objective: To construct a special luciferase reporter to detect DNA methylation regulatory activity in FCER1G gene promoter regulatory element.

Methods: We constructed special full and mock methylated FCER1G gene promoter regulatory luciferase reporters by patch-methylation, and detected DNA methylation regulatory activity by comparing the luciferase activity of full-methylated luciferase reporters with mock-methylated reporters.

Results: We successfully constructed the full and mock methylated FCER1G gene promoter regulatory luciferase reporters. The ratio of luciferase activity between the full methylated and the mock methylated was (0.36±0.07):1 ($P<0.001$).

Conclusion: FCER1G promoter activity is methylation-sensitive and is regulated by DNA methylation.

Keywords: patch methylation FCER1G promoter regulatory element DNA methylation

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