

论著

WTK1细胞在tk位点突变和DNA损伤与修复中的应用

张建清; 张立实; 王瑞淑

深圳市疾病预防控制中心, 广东 深圳 518020

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摘要 背景与目的: 为WTK1细胞在遗传毒理学可同时应用于基因突变和DNA损伤的研究提供实验依据。材料与方法: 分别用标准诱变剂甲基磺酸甲酯(MMS)和过氧化氢(H₂O₂)处理WTK1细胞, 采用tk基因突变试验和单细胞凝胶电泳技术(Single Cell Gel Electrophoresis, SCGE)对细胞的tk位点突变和过氧化氢诱发的DNA损伤情况进行检测。结果: 甲基磺酸甲酯可诱发WTK1细胞tk位点的突变, 以诱发染色体畸变为主。过氧化氢诱发了WTK1细胞DNA的损伤, 并有剂量反应关系。随着修复孵育时间的延长, 彗星细胞尾长和彗星细胞出现率明显下降, 与对照组比较, 差异有显著性(P < 0.01)。结论: WTK1细胞可同时应用于tk基因突变和DNA损伤与修复的研究。采用该细胞株可对化合物进行基因突变和DNA损伤进行研究评价。

关键词 [WTK1细胞](#); [tk位点](#); [突变](#); [DNA损伤](#); [DNA修复](#)

Application of WTK1 Cells to tk Site Mutation and DNA Damage and Repair Capacity

ZHANG Ji an-qing; ZHANG Li -shi ; WANG Rui -shu

Center for Disease Control , Shenzhen 518020 , China

Abstract BACKGROUND & AIM: To provide a basis in tk gene mutation and DNA damage and repair capacity research in WTK1 cells in toxicology field. MATERIAL AND METHODS: The tk site mutation frequency and DNA damage as well as repair capacity were detected after WTK1 cells treated by MMS and H₂O₂ respectively. RESULTS: MMS induced tk site mutation, the mutation colonies mainly were slow growth mutants (SG—mutant) that was chromosome aberration in aberration category. H₂O₂ could induce DNA damage in WTK1 cells and showed dose-response relationship. The quantity and the tail—length of Comet cells decrease significantly with incubation time . CONCLUSION: WTK1 cells could be a useful biology material used for testing tk gene mutation, DNA damage and repair capacity induced by chemicals.

Keywords [WTK1 cells](#) [tk site](#) [mutation](#) [DNA damage](#) [DNA repair capacity](#)

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通讯作者 张建清 jianqingzh@163.net

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