### 论著

# 微囊藻毒素Microcystin-LR体外遗传毒性

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摘要 背景与目的: 应用人类淋巴母细胞TK6研究微囊藻毒素 (Microcystin-LR,MCLR)的体外遗传毒性。 材料与方法: MCLR体外染毒TK6细胞4 h或24 h后检测细胞毒性、微核及tk位点突变频率。 结果: 4 h染毒未引发明显细胞毒性,24 h MCLR染毒导致TK6细胞相对存活率下降,细胞微核率及TK基因突变频率明显上升,并有剂量一反应关系。最高浓度组(80 μg/ml)的细胞微核率及TK基因突变频率分别是对照组的4.8及5.1倍。 MCLR诱发tk位点两种不同表型的突变细胞集落,即正常生长集落及缓慢生长集落,并以后者为主。 结论: 24 h染毒MCLR可以诱发TK6细胞微核及基因突变,揭示MCLR可能是一种断裂剂。

关键词 微囊藻毒素 微核; 突变; TK6细胞

## Genetic Toxicity Induced by Microcystin-LR in Vitro

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Abstract BACKGROUND & AIM: Human lymphoblastoid cell line TK6 was used to investigate the in vitro genotoxicity of Microcystin-LR. MATERIAL AND METHODS: Cytotoxicity response, micronucleus(MN) and mutation frequency at tk locus induced by MCLR after 4 h or 24 h treatment were detected. RERULTS: Treatment with MCLR for 4 h did not induce a significant cytotoxic response at less than 80 μg/ml. Exposure to MCLR for 24h decreased relative survival(RS), induced both MN and TK mutation in a concentration-dependent manner. The maximum induction of MN and TK mutation were 4.8 and 5.1 times those of the control,respectively. Two distinct phenotypic colonies of TK mutants were generated , namely tk-NG and tk-SG mutant colonies but the latter dominated. CONCLUSION: MCLR was clastogenic in TK6 human lymphoblastoid cells .

### **Keywords** <u>Microcystin-LR</u> <u>micronucleus</u> <u>mutation</u> <u>TK6 cell</u>

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