人癌细胞线粒体DNA控制区序列特征分析 Sequence Feature Analysis of Mitochondrial DNA Control Region in Human Carcinoma Cells

胡义德,钱桂生,李淑平,陈维中,毛宝龄 HU Yide,QIAN Gui-sheng,LI Shu-ping,CHEN Weizhong, MAO Bao-ling

第三军医大学新桥医院全军呼吸内科研究所,重庆400037 Institute of Respiratory Diseases of Peoples ▶把本文推荐给朋友 Liberation Army, Xinqiao Hospital, Third Military Medical University, Chongqing 400037, China 收稿日期 修回日期 网络版发布日期 接受日期

为了探讨癌细胞mtDNA控制区序列的变化特征, 采用PCR产物限制性片段长度多态性(PCR-RFLP)分析与直 接测序相结合的方法,对比分析6株人癌细胞系、 6例癌患者及4例健康成人白细胞mtDNA控制区序列。发现第 16519位T→C、16 534位A→G、46位T→G和49位A→C突变,在癌细胞系和癌患者白细胞mtDNA中分别占50%(3/6)和 33.3%(2/6), 健康成人白细胞mtDNA中未见此类型突变; 第16 278位C→T突变, 在癌细胞系mtDNA中占50%(3/6), 显著高于正常人群mtDNA中此位点的多态性变异。表明癌细胞和癌患者白细胞mtDNA重链复制起点及其 相邻D环区 的特征性突变可能与细胞癌变/或癌的易感性有关。

To explore the sequence feature of mitochondrial DNA(mtDNA) control region in human carcinoma cells, polymerase chain reaction-restriction fragment length polymorphism(PCR-RFLP) and direct sequence techniques were used to analyze the sequence of mtDNA control region of 6 human carcinoma cell lines versus white blood cells which from 6 tumor patients and 4 normal adults. The T to C mutation at np 16 519, A to G mutation at np 16 534, T to G mutation at np 46, and A to C mutation at np 49 was found in 50% (3/6 cases) of carcinoma cell lines and in 33.3%(2/6 cases) of tumor patients, but it was not found in normal adults. The C to T mutation at np 16 278 was found in 50%(3/6 cases) of carcinoma cell lines, it was significantly higher than that of the polymorphism of normal population. These findings suggest that the typical mutation in the starting. area of heavy-strand replication and the first half of D-loop region might probably be associated with carcinogenesis or susceptibility of carcinoma.

关键词 人癌细胞系 线粒体DNA 控制区 核苷酸序列 突变 Key words Human carcinoma cell line mtDNA Control region Nucleotide sequence Mutation 分类号

扩展功能

本文信息

- ▶ Supporting info
- ▶ PDF(2235KB)
- ▶[HTML全文](0KB)
- ▶参考文献

服务与反馈

- ▶加入我的书架
- ▶加入引用管理器
- ▶复制索引
- ▶ Email Alert
- ▶文章反馈
- ▶浏览反馈信息

相关信息

▶ 本刊中 包含"人癌细胞系"的 相关文章

▶本文作者相关文章

- 胡义德
- 钱桂生
- 李淑平
- 陈维中
- 毛宝龄HU Yide
- QIAN Gui-sheng
- LI Shu-ping
- CHEN Wei-zhong
- MAO Bao-ling

Abstract

Key words

DOI: