#### 论著

## 威猛(VM - 26) 诱导Hela 细胞G2 期停滞及染色体损伤分析

张英辉 王美华 刘军须 郑力芬

河北医科大学基础医学研究所细胞生物室 石家庄 050017

收稿日期 修回日期 网络版发布日期:

本文采用抗癌药物VM - 26 诱导Hela 细胞周期G2 停滞,并应用染色体预凝集技术,显示其对染色体的影 响。结果表明:5μg/ mlVM - 26 作用24h 后,G2 期停滞细胞的染色体异常主要表现为染色体不规则凝缩和较严重的 染色体断裂。由此提示;VM - 26 通过抑制拓朴异构酶 II 活性,除可造成的DNA 和染色质袢的断裂外,染色体骨架 本身的断裂或也是引起细胞周期G2 期停滯和细胞死亡的原因之一。

威猛 抗癌药 染色体预凝集 染色体损伤

# INVESTIGATIONOF THE HELA CELL CHROMOSOME DAMAGES IN G2 PHASE ARRESTED BY VM - 26

Yan Yunli , Zheng Lifen , Zhang Yinghui , Wang Meihua , Liu junxu

Department of Cell Biology , Instit ute of Basic Medicine , Hebei Medical University, S hijiaz huang 050017

**Abstract** It was unclear about the changes of chromosome const ruction in G2 phase arrested by teniposide (VM - 26). To that, the premature chromosome condensation (PCC) assay was used in the present study. Hela cells were cultured for 24h with 5μg/ ml (7.75μM) VM - 26. The 本文作者相关文章 PCC were induced by mixing the Hela cells with mitotic CHL cells in 50 %polyethylene glycol. It was revealed that the chromosome aberrations in G2 phase yielded by the VM - 26 were mainly in two types, the irregular condensation of the chromosomes and the type of chromosome breaks. Such aberrations might response for the DNA st rand breaks, the chromatin breaks at the scaffold attachment s and especially the breaks across the chromosome cores, due to the inhibition of the VM - 26 on topoisomerase II. Together with the result s of the cloning efficiency by measuring the Hela cells after cultured with the VM - 26, it was thought that the chromosome damages might result in the cell dath when they lost the DNA, or chromosome repair in G2 phase.

**Keywords** teniposide antineoplastic agent s premature chromosome condensation chromosome damage.

DOI

### 本文信息

- ► <u>Supporting info</u>
- ▶ [PDF全文](169k)
- ▶[HTML全文](0k)
- ▶参考文献

服务与反馈

- ▶ 把本文推荐给朋友
- ▶加入我的书架
- ► Email Alert

相关信息

- ▶ 本刊中 包含"威猛"的 相关文章
- 阎蕴力郑力芬张英辉王美华刘军 须