

论著

补充核酸内切酶III对DNA 氧化损伤影响的研究

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摘要 目的:核酸内切酶III在DNA 损伤的切补修复体系中可能发挥着重要作用,是否有效地促进DNA 损伤修复本文进行了探讨。方法:本研究采用“彗星”电泳技术分析DNA 断裂损伤,利用人体淋巴细胞和Hela 细胞并用100μmolH2O2 诱导DNA 严重损伤,其损伤分别达到19815 和320 (专用单位,AU),然后再将两种细胞各分为两组;一组为对照组,另一组为实验组补充25μl (1μg/ml) 的核酸内切酶III。在培养至30min ,60min ,180min 和240min 时观察DNA 断裂损伤情况。结果:淋巴细胞DNA断裂损伤在对照组中经过4h 培养后逐渐得到修复,DNA 损伤由初始时的19815 下降到64127 (AU);补充核酸内切酶III的实验组尽管DNA 断裂经过4h 后也得到修复,但DNA 断裂损伤率仍明显高于对照组($P < 0.05$)。Hela 细胞对H2O2 表现出较高的敏感性并有极强的自身修复能力,对照组DNA 断裂损伤从初始时的320 下降到4h 后的19 (AU),自身修复率达到9411 %,然而补充核酸内切酶III的Hela 细胞DNA 断裂损伤率在30min 、60min 、180min 、240min 时均明显高于对照组($P < 0.05$)。结论:本结果提示核酸内切酶III 在DNA 损伤修复中增加了DNA 的断裂损伤率。

关键词 [核酸内切酶III](#) [彗星电泳](#) [DNA 损伤](#) [DNA 修复](#)

THE EFFECT OF ENDONUCLEASE III SUPPLEMENTATION ON THE DAMAGE OF DNA STRAND BREAKS

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Abstract Purpose and Methods : The aim of the study was observing the effect of endonuclease IIIsupplementation on the damage of DNA strand breaks , which were measured by comet assay (SCGE) . Results : DNA strand break damage of lymphocytes and Hela cells induced by 100μmol H2O2 in controls reached 198. 5 and 320 arbitrary units respectively. The rate of DNA strand breaks of lymphocytes was gradually decreased from the beginning to 4 hour culture in control group. At the end of 4 hour culture , there was a much lower rate of DNA repair in Endonuclease IIIgroup with 33. 4 % than control group with 67. 6 % ($P < 0.05$) . The same result was found in HeLa cells. The levels of DNA strand breaks was quickly decreased from 320 to 19 arbitrary units in control group within 4 hour culture , whereas the DNA strand breaks in supplementing group with Endonuclease IIIwas kept at a high level with 172. 0 arbitrary units at 4 hour culture ($P < 0.05$) . Conclusion : The study suggested that the supplementation of endonuclease IIIcould increase the levels of DNA strand breaks.

Keywords [endonuclease III](#) [SCGE](#) [lymphocyte](#) [DNA repair](#) [DNA damage](#)

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