rbcL 转录物在原核体系中的成熟作用与其3′端碱基序列的关系

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摘要 对烟草(Nicotiana tobacum) rbcL 转录单位3′端前后相连的2个逆向重复序列作系统删 减并单独将此2个序列分开。在原核体系中考察这些缺失予的转录物,结果表明,成熟rbcL mRNA的积累依赖于3′端的大逆向重复序列及未翻译区的一段序列。 将带有逆向重复序列的片段插入菠菜rbcL启动子及苏氨酸终止子之间,大肠杆菌的RNA 多聚酶读过rbcL3′端的逆向重复序列而全部停止在苏氨酸终止子的终止位点,表明成熟的r bcL mRNA 不是rbcL 转录的直接产物,而是由前体 mRNA 经转录后的加工过程后形成的。

关键词 rbcL,缺失子,S1定位,转录后加工

分类号

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Abstract

There is a 15bp large reverse repeated sequence proceeded by a 7bp small one in the 3'flanking region of rbcL of Nicotiana tabacum. A 383 bp of Xbal fragme nt containing these tandemly repeats was inserted into the plasmid $p\lambda s\Delta$, at the position between the λ pand the cat gene. Then these two repeats were separated and deleted systematically to obtain various deletions. The deletion pRT65, pRT 74 and pRT83 was sequenced to determine the deleted base pairs exactly. S1 mapping analysis was adopted to investigate the transcripts of these dele tions in E. coli JM83. The results showed us that the stability of mature 306bp mRNA relied on the large repeat and a short sequence downstream. The small one w as not efficient. The regulation level of the rbcL termination was also investigated. The Xbal -EcoRI fragments from pRT65, pRT74 and pRT83 were transferred into pSP-TT* at the position between the spinach promoter and threonine terminator to construct pR T65, pRT74 and PpRT83 respectively. The results from S1 mapping analysis showed t hat the E. coli RNA threonine terminator. These results suggested that mature rb cL mRNA might be the product precisely processed from a precursor mRNA and the 3 ' flanking sequence might be the signal for precursor mRNA to be processed to the correct position.

Key words <u>rbcL</u> <u>Delection</u> <u>S1 mapping</u> <u>Posttranscriptional procession</u>

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