研究报告

水稻C2H2型锌指蛋白基因*RZF71*的克隆与表达分析

郭书巧,黄骥,江燕,张红生

南京农业大学作物遗传与种质创新国家重点实验室,南京 210095

收稿日期 2006-7-18 修回日期 2006-11-23 网络版发布日期 2007-4-12 接受日期

利用生物信息学和RT-PCR方法从水稻幼苗组织中分离了1个新的C2H2型锌指蛋白基因*RZF71*,该基因编码一条250个 氨基酸残基的多肽,含有两个典型的C2H2型锌指结构。半定量RT-PCR分析表明:RZF71在根、茎、叶和幼穗中呈组▶加入引用管理器 成性表达,在根中的表达丰度略高;在高盐和PEG6000胁迫的水稻幼苗组织中,RZF71的表达显著增强,但低温和 ABA处理对该基因的表达量影响不大。农杆菌介导的洋葱表皮细胞GFP瞬时表达实验表明: RZF71定位于细胞核内。 讨论了RZF71可能作为一个转录调控因子在水稻耐高盐和渗透胁迫中的作用。

关键词 水稻 锌指蛋白 基因克隆 非生物胁迫 表达分析 分类号

Cloning and characterization of RZF71 encoding a C2H2-type zinc finger protein from rice

GUO Shu-Qiao, HUANG Ji, JIANG Yan, ZHANG Hong-Sheng

National Key Laboratory of Crop Genetics and Germplasm Enhancement, Nanjing Agricultural University, Nanjing 210095, China

Abstract

<P>A rice zinc-finger protein gene, RZF71, encoding the C2H2-type zincfinger transcription factor was isolated from rice (Oryza sativa L. subs. Japonica) by RT-PCR approach. Gene RZF71 encodes a 25 kDa protein with 250 amino acids, which contains two typical C2H2 zinc finger domains. The expression profiling showed that RZF71 was constitutively expressed in roots, culms, leaves, and flowering spikes. The semi-quantitative RT-PCR assay showed RZF71 was strongly induced by high-salinity and 20% PEG6000 treatments, but not regulated by low temperature and ABA (abscisic acid) treatments. Tran-sient expression of the RZF71-GFP protein in onion epidermal cell showed that RZF71 was localized in cell nuclei. These results indicated that the RZF71 may play an important role in rice responses to salt and osmotic stresses as a transcription factor. </P>

Key words rice zinc finger protein gene cloning abiotic stress expression analysis

DOI: 10.1360/yc-007-0607

扩展功能

本文信息

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

服务与反馈

- ▶把本文推荐给朋友
- ▶加入我的书架
- ▶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"水稻"的 相关文章

▶本文作者相关文章

- 郭书巧
- 黄骥
- 江燕
- 张红生