研究快报

不同启动子驱动下马铃薯蛋白酶抑制剂Pin II 转基因水稻的遗传、表达 和对粘虫抗性分析

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以不同启动子驱动的马铃薯蛋白酶抑制剂II基因(PinII-2x,PinII-4x)转基因水稻为材料,经潮霉素抗性、PCR和 Southern blot等检测对转基因水稻后代进行了遗传分析。结果显示:外源基因在68.4%的转基因植株中符合孟德 尔遗传模式,单拷贝植株率为63.6%。转基因植株后代的PinⅡ蛋白活性测定结果表明:ActI和Ubi驱动的PinⅡ-2x ≥ 复制索引 转基因水稻植株中,每克鲜叶片的Pin II蛋白含量为 $160 \mu g$ 和 $176 \mu g$,而由PIN5'驱动的Pin II - 4x为 $104 \mu g$,对照水 稻仅为20μg。ActI和Ubi驱动的*PinⅡ*表达产物对胰蛋白酶活性抑制程度分别达到37. 7%和43. 1%,明显高于*PinⅡ*自 身启动子PIN5′(29.2%)。叶片饲养粘虫幼虫的实验表明:转基因植株叶片对粘虫有抗性,但抗性达不到显著水 平,且启动子效率、Pin II表达量与抗粘虫

马铃薯蛋白酶抑制剂Ⅱ 遗传 活性 粘虫 关键词 分类号 0943

Inheritance, Expression and Armyworm resistance of Protease Inhibitor II Gene(Pin II) Driven by Different Promoters in Transgenic Rice

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Abstract

<I>The inheritance of rice lines transformed by protease inhibitor II gene under control of different promoters was investigated by analysis of hygromycin resistance, PCR and Southern blot. For segregation patterns of foreign gene, 68.4% of the transgenic rice plants were conformed to a Mendelian ratio and in which the rate of transgenic plants with single copy was 63.6%. Quantitative analysis of Pin II protein expressed in transgenic rice plants showed that Pin II protein in fresh leaves was $160\mu g/g$ for Act-<|> </|>Pin</|>|| <|>|| <|>-2x</|>, $176\mu g/g$ for Ubi-<I> Pin</I><I>II</I>-2x</I>, and $104\mu g/g$ for PIN5'-<I> Pin</l> | </l> | </l> | </l> separately while in control rice plants was only $20\mu g/g$. The inhibitory activity against tryspin of <I>Pin</I><<math>I>II</I></I> gene driven by <I>Act</I>I and Ubi promoter reached 37.7% and 43.1%, much higher than that driven by PIN5'(29.2%). Bioassay for insect resistance to armyworm (<I>Pseudaletia separata Walker</I>) revealed that transgenic plants had increased their resistance to the pest but there was not significantly different from controls, and also there was no correlation between insect resistance to armyworm and quantity of Pin II < I > </I > protein as well as promoters in transgenic rice.

Key words potato protease inhibitor II inheritance activity *Pseudaletia separata* walker

DOI:

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