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摘要: Na⁺/H⁺反向转运蛋白可调控细胞质pH值、钠离子浓度和细胞体积,从而减轻盐胁迫对植物的伤害。利用发根农杆菌 (Agrobacterium rhizogenes) 介导法,向大豆根系导入由CaMV35S启动子调控的Na⁺/H⁺反向转运蛋白编码基因GmNHX1的cDNA序列,通过该基因的过量表达,提高大豆的耐盐性。通过潮霉素筛选、GUS染色及RT-PCR检测,确认获得了转GmNHX1基因的大豆发状根。对转基因发状根耐盐性分析表明:在100、150和200 mmol·L⁻¹的NaCl胁迫下,置于MS固体培养基中的转基因离体发状根的长度和重量增加值均显著大于对照。带有转基因发状根的子叶及复合体植株在盐胁迫条件下也具有较强的生存能力。试验证明,过表达GmNHX1基因能够显著提高转基因发状根的耐盐性,该结果为利用GmNHX1基因进行大豆耐盐性的改良提供了依据。

Abstract: Na⁺/H⁺ antiporter is responsible for the regulation of cytoplasmic pH, sodium concentration and cell volume for plants to cope with salt stress. To enhance the salt tolerance of soybean, the GmNHX1 cDNA that encodes a soybean Na⁺/H⁺ antiporter protein was driven by the Cauliflower Mosaic Virus 35S promoter and overexpressed in soybean roots via Agrobacterium rhizogenes mediated transformation. Hpt screening, GUS and RT-PCR analysis all confirmed that GmNHX1 was successfully integrated into the genome of soybean hairy roots. Salt tolerance analysis showed that the growth in length and weight of the transgenic hairy roots on MS agar medium were significantly better than the non-transgenic control, when supplemented with 100, 150 and 200 mmol·L⁻¹ NaCl. Both the cotyledons and composite plants with transgenic hairy roots survived better than the non-transgenic control when subjected to salinity stress. These results support the possibility of using GmNHX1 to improve salt tolerance in soybean.

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