**Removal of Feedback Inhibition of Δ¹-Pyrroline-5-Carboxylate Synthetase Results in Increased Proline Accumulation and Protection of Plants from Osmotic Stress**

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The Δ¹-pyrroline-5-carboxylate synthetase (P5CS; EC not assigned) is the rate-limiting enzyme in proline (Pro) biosynthesis in plants and is subject to feedback inhibition by Pro. It has been suggested that the feedback regulation of P5CS is lost in plants under stress conditions. We compared Pro levels in transgenic tobacco (Nicotiana tabacum) plants expressing a wild-type form of Vigna aconitifolia P5CS and a mutated form of the enzyme (P5CSF129A) whose feedback inhibition by Pro was removed by site-directed mutagenesis. Transgenic plants expressing P5CSF129A accumulated about 2-fold more Pro than the plants expressing V. aconitifolia wild-type P5CS. This difference was further increased in plants treated with 200 mM NaCl. These results demonstrated that the feedback regulation of P5CS plays a role in controlling the level of Pro in plants under both normal and stress conditions. The elevated Pro also reduced free radical levels in response to osmotic stress, as measured by malondialdehyde production, and significantly improved the ability of the transgenic seedlings to grow in medium containing up to 200 mM NaCl. These findings shed new light on the regulation of Pro biosynthesis in plants and the role of Pro in reducing oxidative stress induced by osmotic stress, in addition to its accepted role as an osmolyte.

Pro is known to play an important role as an osmoprotectant in plants subjected to hyperosmotic stresses such as drought and soil salinity (Delauney and Verma, 1993). Recent studies on Pro synthesis and catabolism genes have provided results that are consistent with diverse functions of Pro as a source of energy, nitrogen, and carbon, and as an osmolyte in response to dehydration (Kohl et al., 1988; Kavi Kishor et al., 1995; Peng et al., 1996; Hua et al., 1997; Zhang et al., 1997). Synthesis, accumulation, and catabolism of Pro in plants under stress may involve the loss of feedback regulation due to a conformational change in the P5CS protein (Boggess et al., 1976a, 1976b). In bacteria, Pro biosynthesis has been shown to be regulated by the end product inhibition of γ-GK activity (Smith et al., 1984). A Salmonella typhimurium mutant resistant to a toxic Pro analog (3,4-dehydro-d,l-Pro) accumulated Pro and showed enhanced tolerance to osmotic stress (Csonka, 1981). The mutation was due to a change of an Asp residue (at position 107) to Asn, rendering the γ-GK much less sensitive to inhibition by Pro (Csonka et al., 1988; Dandekar and Uratsu, 1988). We showed that the conserved Asp residue (at position 128) in the V. aconitifolia P5CS is not involved in the feedback inhibition (Zhang et al., 1995). Using site-directed mutagenesis, a replacement of Phe at position 129 by Ala was made in V. aconitifolia P5CS (P5CSF129A). This mutant enzyme was shown to retain similar kinetic characteristics as wild-type P5CS, but its feedback inhibition was virtually eliminated (Zhang et al., 1995). In this report, we demonstrate that tobacco (Nicotiana tabacum) plants carrying P5CSF129A accumulate more Pro, produce fewer free radicals, and are more tolerant to osmotic stress than plants expressing the wild-type V. aconitifolia P5CS transgene only. The P5CS transgenic seeds germinated well in a high salinity (200 mM NaCl) environment, while the wild type did not. These results demonstrated that feedback regulation of P5CS by Pro plays a role in the control of Pro biosynthesis in plants, and that Pro accumulation reduces γ-glutamyl kinase (γ-GK) and Glu-5-semialdehyde (GSA) dehydrogenase (or γ-glutamyl phosphate reductase; Hu et al., 1992; Savoure et al., 1995; Yoshiba et al., 1995). In tomato, it has been reported that there are two Pro loci in the nuclear genome: one specifies a bifunctional P5CS (tomPro2) and the other one (tomPro1) apparently encodes a poly-cistronic mRNA that directs the synthesis of γ-GK and GSA dehydrogenase as two separate peptides (Garcia-Rios et al., 1997). Two P5CS genes have also been shown to be present in both Arabidopsis and alfalfa (Strizhov et al., 1997; Ginzberg et al., 1998; Yoshiba et al., 1999). The Arabidopsis P5CS1 gene is expressed in most organs and is induced rapidly by stress (Strizhov et al., 1997; Zhang et al., 1997; Yoshiba et al., 1999). P5CS2 is expressed in dividing cell cultures and its induction by stress is dependent on protein synthesis (Ginzberg et al., 1998).

Earlier experiments suggested that Pro accumulation in plants under stress may involve the loss of feedback regulation due to a conformational change in the P5CS protein (Boggess et al., 1976a, 1976b). In bacteria, Pro biosynthesis has been shown to be regulated by the end product inhibition of γ-GK activity (Smith et al., 1984). A Salmonella typhimurium mutant resistant to a toxic Pro analog (3,4-dehydro-d,l-Pro) accumulated Pro and showed enhanced tolerance to osmotic stress (Csonka, 1981). The mutation was due to a change of an Asp residue (at position 107) to Asn, rendering the γ-GK much less sensitive to inhibition by Pro (Csonka et al., 1988; Dandekar and Uratsu, 1988). We showed that the conserved Asp residue (at position 128) in the V. aconitifolia P5CS is not involved in the feedback inhibition (Zhang et al., 1995). Using site-directed mutagenesis, a replacement of Phe at position 129 by Ala was made in V. aconitifolia P5CS (P5CSF129A). This mutant enzyme was shown to retain similar kinetic characteristics as wild-type P5CS, but its feedback inhibition was virtually eliminated (Zhang et al., 1995). In this report, we demonstrate that tobacco (Nicotiana tabacum) plants carrying P5CSF129A accumulate more Pro, produce fewer free radicals, and are more tolerant to osmotic stress than plants expressing the wild-type V. aconitifolia P5CS transgene only. The P5CS transgenic seeds germinated well in a high salinity (200 mM NaCl) environment, while the wild type did not. These results demonstrated that feedback regulation of P5CS by Pro plays a role in the control of Pro biosynthesis in plants, and that Pro accumulation reduces γ-glutamyl kinase (γ-GK) and Glu-5-semialdehyde (GSA) dehydrogenase (or γ-glutamyl phosphate reductase; Hu et al., 1992; Savoure et al., 1995; Yoshiba et al., 1995). In tomato, it has been reported that there are two Pro loci in the nuclear genome: one specifies a bifunctional P5CS (tomPro2) and the other one (tomPro1) apparently encodes a poly-cistronic mRNA that directs the synthesis of γ-GK and GSA dehydrogenase as two separate peptides (Garcia-Rios et al., 1997). Two P5CS genes have also been shown to be present in both Arabidopsis and alfalfa (Strizhov et al., 1997; Ginzberg et al., 1998; Yoshiba et al., 1999). The Arabidopsis P5CS1 gene is expressed in most organs and is induced rapidly by stress (Strizhov et al., 1997; Zhang et al., 1997; Yoshiba et al., 1999). P5CS2 is expressed in dividing cell cultures and its induction by stress is dependent on protein synthesis (Ginzberg et al., 1998).

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osmotic stress, which may be mediated by free radicals produced as a result of oxidative stress.

MATERIALS AND METHODS

Transformation of Tobacco Plants

A plasmid (pBI-P5CSF129A, Fig. 1) containing mutagenized Vigna aconitifolia P5CSF129A cDNA (Zhang et al., 1995) under the control of the cauliflower mosaic virus 35S promoter was used for tobacco (Nicotiana tabacum cv Xanthi) transformation via Agrobacterium tumefaciens. All transgenic lines tested accumulated high levels of Pro (Fig. 2); line F129A-3 was used in this study. P5CS line 136, which expressed a V. aconitifolia wild-type P5CS gene described previously (Kavi Kishor et al., 1995), was used as a control along with plants transformed with vector pBI121 only.

Germination and Plant Growth

Seeds of wild-type tobacco and transgenic (pBI121, P5CS, and P5CSF129A) plants were germinated and maintained on Murashige and Skoog (MS) agar (4.3 g/L MS salts and 0.8% [w/v] Phytagar, Gibco-BRL, Cleveland) medium containing 0, 150, 200, 250, and 300 mM NaCl. The medium on which the transgenic seeds were plated contained kanamycin (200 µg/mL). Germination rate was recorded on d 14, 18, 20, and 24. Growth rate as measured by fresh weight (g) was recorded in 2-week-old seedlings. Three replicates containing about 60 seedlings each were taken for measurements. The data shown correspond to control and 200 mM NaCl-treated plants (in Fig. 5). Seeds were germinated on MS medium in a tissue culture room at 25°C ± 2°C under constant illumination (200 µmol m⁻² s⁻¹ from cool-white fluorescent tubes) and seedlings were grown in a growth chamber at 25°C ± 2°C with cycles of 16-h light and 8-h dark.

Northern- and Western-Blot Analyses

Total RNA (15 µg) isolated from the transgenic and control tobacco seedlings was electrophoresed, blotted, and hybridized with the V. aconitifolia P5CS cDNA (Hu et al., 1992) as a probe. Hybridization and washing of the filters were carried out under high-stringency conditions (Kavi Kishor et al., 1995). Western blotting was performed using polyclonal antibodies to purified V. aconitifolia P5CS protein, as described previously (Kavi Kishor et al., 1995; Zhang et al., 1995).

Growth and Salinity Treatment of Tobacco Bright Yellow 2 (BY2) Cells

Tobacco BY2 cells were maintained in MS media (4.3 g/L MS salts from Gibco-BRL, 0.1 g/L inositol, 1.0 mg/L thiamine, 0.2 mg/L 2,4-dichlorophenoxyacetic acid [2,4-D], 255 mg/L KH₂PO₄, and 30 g/L Suc). For the salinity treatment, a 5-d-old cell suspension was used and the salt concentration was adjusted with a 5.0 mM NaCl stock to final concentrations ranging from 50 to 400 mM. For time course induction of malondialdehyde (MDA) under salinity stress, the cell suspension was treated with 250 mM NaCl, and MDA contents were determined at time intervals of 5 h.

Measurement of Pro and MDA Contents

Pro concentration was determined as described previously (Kavi Kishor et al., 1995) according to the procedure of Bates et al. (1973). Values were expressed as milligrams per gram fresh weight. MDA contents were measured using a thiobarbituric acid reaction (Heath and Packer, 1968).
About 0.5 to 1.0 g of tissue was homogenized in 5 mL of 5% (w/v) trichloroacetic acid and the homogenate was centrifuged at 12,000g for 15 min at room temperature. The supernatant was mixed with an equal volume of thiobarbituric acid (0.5% in 20% [w/v] trichloroacetic acid), and the mixture was boiled for 25 min at 100°C, followed by centrifugation for 5 min at 7,500g to clarify the solution. Absorbance of the supernatant was measured at 532 nm and corrected for non-specific turbidity by subtracting the A\textsubscript{450}. MDA contents were calculated using an extinction coefficient of 155 m\textsuperscript{-1} cm\textsuperscript{-1}. Values of Pro and MDA contents were taken from measurements of three independent samples, and ses of the means were calculated.

**RESULTS**

**Expression of *V. aconitifolia* Mutated P5CS cDNA in Transgenic Tobacco Plants**

Kinetics studies of *V. aconitifolia* P5CS enzyme showed that P5CS activity is inhibited by 6 mM Pro (Hu et al., 1992; Zhang et al., 1995). To remove this allosteric inhibition, a point mutation in the P5CS cDNA was introduced so that the Phe at position 129 was replaced by Ala (Fig. 1). This mutated protein (P5CSF129A) is enzymatically active, but its feedback inhibition by Pro is virtually eliminated (Zhang et al., 1995). We placed the *V. aconitifolia* mutated P5CS cDNA under the control of cauliflower mosaic virus 35S promoter (Fig. 1) and introduced this construct into tobacco plants by *A. tumefaciens*-mediated transformation. Seeds of five independent transgenic lines were germinated and maintained on MS medium containing 0 mM NaCl or 200 mM NaCl. Pro levels in all P5CSF129A lines were almost 2-fold higher than the P5CS transgenic line (Fig. 2). The line (F129A-3) that produced the highest level of Pro both under control and salt-stressed conditions were chosen for further analyses on gene expression and oxidative damage due to osmotic stress.

The expression of P5CSF129A in transgenic plants was monitored by northern blotting (Fig. 3A). High-stringency conditions were used for RNA hybridization and washing to eliminate any possible cross-reaction with the tobacco endogenous P5CS mRNA. Expression levels of the transgene in P5CSF129A lines (Fig. 3A, lane 2) were slightly lower than that in P5CS transgenic line 136 (Fig. 3A, lane 3) which had previously been shown to express high levels of the *V. aconitifolia* P5CS gene (Kavi Kishor et al., 1995). No cross-reaction with the tobacco endogenous P5CS mRNA was detected under these conditions (Fig. 3A, lane 1). Expression levels of *V. aconitifolia* P5CSF129A and the wild-type P5CS enzyme in transgenic tobacco plants were also determined by western blotting using P5CS antibodies (Fig. 3B). A weak cross reaction of the *V. aconitifolia* P5CS antibodies with the tobacco endogenous P5CS protein at the expected size (72 kD) appeared (Fig. 3B, lane 1–2) after prolonged exposure. High levels of P5CSF129A expression were detected in the transgenic plants (Fig. 3B, lane 3–4). In agreement with mRNA levels (Fig. 3A), protein expression in P5CSF129A line (F129A-3) was also slightly lower than that in P5CS line 136 (Fig. 3B, lane 5–6). Comparison of P5CS protein levels between control and salt-treated plants indicated an increase of about 50% in P5CS protein under stress conditions. These results are consistent with our earlier report that more P5CS protein is present in plants subjected to osmotic stress (Zhang et al., 1995).

**Growth of Seedlings Expressing Mutated P5CS Devoid of Feedback Control**

Transgenic lines expressing *V. aconitifolia* wild-type P5CS and mutated enzyme (P5CSF129A) were subjected to different levels of salinity treatments. Uniform seed germination for all three groups of plants was observed in medium containing no NaCl. At concentrations of 250 and 300 mM NaCl, seed germination rate was very low for both transgenic lines, with no germination in control seeds. The
differences observed with 200 mM salt were, however, highly significant (Fig. 4). The transgenic P5CSF129A plants exhibited highest germination rate, to an extent of 60% and 68% compared with 23% and 28% in P5CS lines at d 14 (Fig. 5A) and d 18 (data not shown), respectively. On the other hand, germination of pBI121 transgenic seeds (control) was severely inhibited (i.e. only 8% and 16% germination).

Growth of seedlings in terms of fresh weight (Fig. 5B) also showed no difference in the untreated control and transgenic plants. However, the P5CSF129A plants exhibited the least inhibition of growth over P5CS and pBI121 plants under 200 mM salt stress (Figs. 4 and 5B). The inhibition of plant growth over their respective controls at d 20 was approximately 95%, 82%, and 67% in the pBI121, the P5CS, and the mutant P5CSF129A plants, respectively (data not shown). Seedling growth in terms of root proliferation with abundant root hairs was observed in the transgenic P5CSF129A plants, whereas very few control seedlings exhibited root initiation and elongation, and root hairs were conspicuously absent.

**Removal of Feedback Inhibition of P5CS Results in Higher Levels of Pro Accumulation in Transgenic Plants**

To determine if the observed phenotypic differences in germination and seedling growth were related to Pro levels in these plants, we measured Pro contents in respective seedlings grown under both normal and salt-stressed conditions. When germinated on medium containing no salt, the P5CSF129A plants were found to produce about 2-fold more Pro than the P5CS plants, which in turn synthesized 5- to 6-fold more Pro than the pBI121 seedlings (Fig. 5C). Under salt stress conditions, the P5CSF129A plants accumulated about 2 times more Pro than the P5CS plants and 3-fold more than the pBI121 seedlings. This suggests that the wild-type P5CS enzyme is subject to feedback inhibition by the end product Pro under stress conditions, because removal of this feedback regulation rendered at least a 2-fold increase in Pro content.

**Osmotic Stress Induces Free Radical Production in Plant Cells That Can Be Reduced by Pro**

To further understand how Pro accumulation helps plant cells deal with osmotic stress, we examined the relationship between osmotic stress and oxidative stress. For this, we used tobacco BY2 cells because of the uniformity of the mass of tissue, and measured free radical levels in cells treated with different concentrations of NaCl. MDA is a major cytotoxic product of lipid peroxidation and has been used extensively as an indicator of free radical production (Kunert and Ederer, 1985). MDA levels increased linearly from 24 to 62 µg/g fresh weight of cells with the increase in NaCl concentration over the range from 50 to 300 mM NaCl (Fig. 6A). In cells treated with 250 mM NaCl, MDA accumulated within 5 h and continued to be produced at a slower rate over 24 h (Fig. 6B). These results suggest that oxidative stress accounts, at least in part, for the damage caused to the plant cells by osmotic stress.

It has been proposed that Pro may act as a free radical scavenger to protect plants from damage by oxidative stress caused during osmotic stress (Alia et al., 1993; Smirnoff, 1993). To test this hypothesis, we measured the damage by free radicals to wild-type tobacco BY2 cells during salt stress with and without the addition of exogenous Pro. As shown in Figure 7A, salinity stress created by 250 mM NaCl for 8 h caused MDA accumulation from 20 to 46 µg/g fresh weight. This value was effectively reduced by 40% in the presence of 120 mM Pro in the culture medium. These data indicate that the supply of exogenous Pro significantly reduces the accumulation of free radicals in cells under osmotic stress. Taking advantage of transgenic plants that produce high levels of endogenous Pro, we carried out measurements on free radical levels in these plants with and without salt stress.
Minor differences in MDA levels were found among the three groups of plants when grown under normal conditions. Treatment with 200 mM NaCl caused about a 2-fold elevation of MDA in wild-type and pBI121- plants. A significantly lower MDA content was found in transgenic P5CSF129A plants (14.3 μg/g) than in control plants (Fig. 7B). The MDA level in P5CS plants were found to be intermediate (17.8 μg/g). These data indicate that high concentrations of Pro synthesized endogenously in transgenic plants may provide a means to reduce the levels of free radicals generated during osmotic stress. This observation demonstrated an additional role of Pro in reducing damage from oxidative stress generated by osmotic stress.

Figure 5. Effect of V. aconitifolia P5CS and P5CSF129A gene expression on seed germination, seedling growth, and Pro accumulation in transgenic tobacco plants treated with 200 mM NaCl. Values presented are the average of three independent samples each containing about 200 seeds. White bars, Control; black bars, 200 mM NaCl.

Figure 6. Free radical production as measured by MDA content in tobacco BY2 cells 5 d after subculturing. A. Effect of NaCl concentrations on MDA content measured after 8 h of salinity treatment. B, Time course of MDA accumulation in cells treated with 250 mM NaCl.

DISCUSSION

Biosynthesis of Pro in living cells is subject to several control mechanisms. In Escherichia coli and yeast, GK and GSADH are synthesized as separate peptides and form a heterodimer. GK is feedback inhibited by the end product of the pathway, Pro. In animals and plants, GK and GSADH are encoded by a bifunctional P5CS gene. Whereas human P5CS is feedback regulated by Orn (Hu et al., 1999), plant P5CS is subject to allosteric regulation by Pro (Hu et al., 1992; Zhang et al., 1995). The activity of V. aconitifolia P5CS is reduced to 50% by 6 mM Pro (Hu et al., 1992). Plant cells under osmotic stress can accumulate up to 129 mM Pro in the cytosol (Binzel et al., 1987; DeLaunay and Verma, 1993), a concentration that would almost completely turn off the P5CS enzyme. This is contradictory to the fact that plants under stress continue to synthesize Pro and build up the Pro pool. To explain this paradox, it has been assumed that the P5CS enzyme may undergo a conformational change and lose its feedback regulation property (Boggess et al., 1976a, 1976b). We reasoned that if the feedback regulation of the wild-type P5CS is completely lost during
stress, then expression of the mutant P5CS, i.e. P5CSF129A, will not result in the synthesis of more Pro than expression of the wild-type P5CS transgene. On the other hand, if P5CS retains its feedback regulation under stress conditions, transgenic plants expressing the unregulated version of the enzyme, P5CSF129A, should accumulate much more Pro than those harboring the wild-type P5CS transgene.

Our results (Figs. 2 and 5C) show that removal of feedback inhibition in P5CSF129A resulted in a 2-fold increase in Pro compared with that in the P5CS transgenic line under both normal and stressed conditions. We conclude that feedback regulation of P5CS in plants is not completely eliminated under stress. Thus, Pro synthesis in plants under stress is regulated not only by transcriptional activation of P5CS (Hu et al., 1992; Garcia-Rios et al., 1997; Zhang et al., 1997), but also by feedback regulation by the end product of the pathway. Furthermore, a reciprocal increase in P5CS and Pro dehydrogenase during stress and recovery from stress controls the levels of Pro according to the environment (Peng et al., 1996).

In our previous report (Zhang et al., 1995), we performed site-directed mutagenesis to substitute each of the six amino acid residues between positions 126 and 131 of the P5CS peptide. Two residues were found to have a different degree of effect on the allosteric property of the enzyme. Substitution of Phe at 129 with Ala (P5CSF129A) produced the most profound effect, with an increase in the 50% inhibition values from 6 mM Pro in the wild-type P5CS to 960 mM in P5CSF129A. Substitution of Asp at position 126 (P5CSD126A) resulted in a moderate change in feedback inhibition by 86 mM Pro (Zhang et al., 1995). This demonstrates that the feedback regulation property of P5CS can be changed to different degrees by modification of different amino acid residues. It remains to be determined if such a point mutation directly affects the allosteric site or if it brings about a conformational change in the protein.

Under osmotic stress conditions, the wild-type P5CS enzyme may undergo some conformational change around the Pro feedback interaction site, leading to a partial loss of its allosteric regulation property. This “partial loss” hypothesis can explain the paradox of why plants under stress continue to build up Pro levels even after Pro concentrations reach the feedback inhibition levels. The differences between the levels of P5CS protein in both control and transgenic lines under normal and stress conditions indicate a contribution of native P5CS which is known to be induced under stress and cross-react with vigna P5CS antibody.

Accumulation of Pro in plants under stress may offer multiple benefits to the cell. We showed that free radicals are formed during osmotic stress, as measured by an increase in the MDA production. These radicals can react with many cellular constituents, including DNA, proteins, and lipids, leading to radical chain processes, crosslinks, peroxidation, membrane leakage, and the production of toxic compounds (Davies, 1995). MDA, a lipid peroxidation product, has been used widely to assess the levels of free radicals in living cells (Kunert and Ederer, 1985). In this study, we found that MDA levels increased significantly with the NaCl concentration in tobacco BY2 cells (Fig. 6, A–B). Exogenously supplied Pro significantly reduced (by 40%) the levels of free radicals in the salt-treated BY2 cells (Fig. 7A). This confirms earlier observations by Alia et al. (1993) on the production of free radicals under salinity stress.

Measurements of MDA contents in transgenic plants producing high levels of endogenous Pro during salinity stress showed that P5CSF129A plants produced more Pro and accumulated less MDA than P5CS transgenic or wild-type plants (Fig. 7B). That Pro levels are increased as a result of free radical generation is indicated by treating BY2 cells with plumbagin, a known free radical generator (Z. Peng and D.P.S. Verma, unpublished data). These results clearly show a role of Pro in scavenging free radicals in cells exposed to salinity. Resistance to oxidative stress can also be increased by enhanced mannitol biosynthesis in transgenic plants (Shen et al., 1997). It is possible that the increased resistance to oxidative stress is due to some indirect metabolic or physiological consequence of the accumulation of Pro and other metabolites. Overproduction of Gly betaine results in the induction of two enzymes, ascorbate peroxidase and catalase, which are known to be involved in oxidative stress resistance in Arabidopsis (Alia et al., 1999). Intermediates in Pro biosynthesis and catabo-

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**Figure 7.** Effect of exogenous and endogenous Pro on free radical production in tobacco cells and transgenic plants. A, Effect of exogenous Pro (120 mM) on MDA content in tobacco BY2 cells treated with 250 mM NaCl for 8 h. B, Effect of endogenous Pro accumulation on MDA content in 14-d-old wild-type, pBI121, P5CS, and P5CSF129A transgenic seedlings treated with 0 (control; white bars) or 200 mM NaCl (black bars).
lism, such as Gln and P5C, have also been found to induce the expression of several osmotically regulated genes in rice (Iyer and Caplan, 1998).

Accumulation of Pro in plants under stress is a result of the reciprocal regulation of two pathways: increased expression of Pro synthetic enzymes (P5CS and P5CR) and repressed activity of Pro degradation (Delauney and Verma, 1993; Peng et al., 1996). This leads to a “proline cycle,” the homeostasis of which depends on the physiological state of the tissue (Verma, 1999). Pro catabolism is catalyzed by Pro dehydrogenase and P5C dehydrogenase (Hu et al., 1996; Peng et al., 1996). Suppression of Pro degradation has been demonstrated both in radiolabeling studies (Stewart and Boggess, 1978) and gene expression experiments (Kiyosue et al., 1996; Peng et al., 1996; Verbruggen et al., 1996). Recent studies have demonstrated that high Pro concentrations are present in the phloem sap of drought-stressed alfalfa (Girousse et al., 1996) and that the expression of a Pro-specific amino acid transporter is induced in response to water deficit and salt stress (Rentch et al., 1996). Evidence for the transport of Pro to the root tip, where it accumulates during stress, has been reported (Verslues and Sharp, 1999). The data indicate that plants may have evolved a mechanism to coordinate synthesis, catabolism, and transport activities for the accumulation of Pro.

Plants well adapted to drought and saline environments manifest a variety of changes for sustained growth. The accumulation of Pro is one of the factors that facilitates this adjustment. The relative contribution of each step remains to be established. Our present results indicate that Pro synthesis in plants can be manipulated by eliminating feedback regulation of the key regulatory enzyme of the pathway, P5CS. The ability of plants to tolerate oxidative stresses imposed by osmotic stress can be significantly improved by expressing a mutant form of the enzyme in transgenic plants. Recent data have shown that expression of antisense P5CS inhibits Pro production and makes plants hypersensitive to osmotic stress (Najo et al., 1999). This is also consistent with a study on antisense Gln synthetase that reduces the Pro level and renders transgenic plants more sensitive to salt treatment (Brugiere et al., 1999). The present study also suggests that the role of Pro as a free radical scavenger may be more important in overcoming stress than in acting as a simple osmolyte. This opens a new avenue of research for metabolic engineering and stress tolerance in agriculturally important crops.

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