Effect of Nitric Oxide on Proline Metabolism in Cucumber Seedlings under Salinity Stress

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ADDITIONAL INDEX WORDS. NaCl stress, Cucumis sativus, exogenous substance, pyrroline-5-carboxylate synthetase, proline dehydrogenase

ABSTRACT. Nitric oxide (NO), an endogenous signaling molecule in plants and animals, mediates responses to abiotic and biotic stresses. This study was conducted in a nutrient solution to investigate 1) the effects of exogenous sodium nitroprusside (SNP), an NO donor, on free proline (Pro) and protein content; and 2) the enzymes involved in Pro metabolism [pyrroline-5-carboxylate synthetase (P5CS) and proline dehydrogenase (PDH)] in cucumber (Cucumis sativus) seedling leaves and roots under NaCl stress. The results showed that the increases in free Pro and protein were significantly higher in the 50 mM NaCl solution but highly significant with the addition of 100 μM SNP to the 50 mM NaCl solution for the entire treatment period. Moreover, leaves maintained higher levels of free Pro and protein content than roots throughout the experiments. The P5CS activity increased in the saline treatment compared with the control, and this increase was greater in the 50 mM NaCl + 100 μM SNP solution than in the other treatments. On the other hand, the PDH activity was inhibited under NaCl stress but the reduction in activity was greater in the 50 mM NaCl + 100 μM SNP solution than in the others. These findings suggest that Pro metabolism was significantly altered during the exogenously applied NO under salt stress and that this alteration prompted the accumulation of higher levels of free Pro, which, in turn, maintained the turgor in the cucumber seedlings and helped protect them from salt stress. Moreover, the toxic effects generated by 50 mM NaCl were partially overcome by the application of NO, which could be used as a potential growth regulator to improve plant salinity tolerance. Therefore, it was concluded that NO could alleviate salinity damage in cucumber seedlings by regulating Pro metabolism. Overall, the adverse effects of salt stress could be lessened by the exogenous application of NO to cucumber seedlings.

Salinity is a major environmental factor limiting plant growth, productivity, and distribution. Salinity stress can trigger complex physiological and biochemical responses and affect almost all processes in plants. Under salt stress, cells experience water deficiency, or osmotic stress. Numerous studies have shown that plants under salt stress can actively accumulate small molecule metabolites and inorganic ions, which contribute to growth and development in plants. However, soluble metabolites are more important than inorganic ions for plants when they suffer environmental stresses. Proline, glycine betaine, and soluble sugars are examples of low-molecular-weight solutes that accumulate in plants. Pro is one of the most commonly accumulated osmolytes in plants under salinity conditions (Delauney and Verma, 1993). The enzyme pyrroline-5-carboxylate synthetase plays a key role in controlling the level of Pro accumulation in plants (Ashraf and Foolad, 2007). An increasing number of studies have shown that Pro accumulation in plants under stress signal (Khedr et al., 2003; Szabados and Savouré, 2009). In addition, the intermediates of Pro metabolism induce gene expression and reduce oxidation injuries from osmotic stress (Hong et al., 2000). Although Pro-inducible genes have been identified (Satoh et al., 2002), the mechanisms of Pro action are not fully understood. Pro is synthesized in plants through two pathways: the glutamate (Glu) approach and the ornithine approach by original amino acids. Pro biosynthesis from Glu appears to be the predominant pathway, especially under stress conditions (Delauney and Verma, 1993). The enzyme pyrroline-5-carboxylate synthetase plays a key role in controlling the level...
of Pro in plants and catalyzes the first two steps of Pro biosynthesis from Glu (Hu et al., 1992). P5CS has been isolated from several plants (Fujita et al., 1998; Hu et al., 1992; Yoshida et al., 1995), and a correlation between the induction of P5CS and the accumulation of Pro has been shown in Arabidopsis thaliana and rice [Oryza sativa (Yoshida et al., 1995, 1997)]. The Pro-catabolizing enzymes Pro oxidase and proline dehydrogenase also influence the level of Pro accumulation. Greater accumulation of Pro could be caused by the inhibition of these enzymes (Kandpal et al., 1981).

NO is a small, highly diffusible, gaseous-free radical and a ubiquitous bioactive molecule that plays a key role as an intracellular intercellular messenger inducing various processes through either redox or additive chemistry in plants (Lamattina et al., 2003). These processes include senescence (Leshem et al., 1998), stress responses (Ruan et al., 2004), programmed cell death (Beligni et al., 2002), and disease resistance (Chandok et al., 2003). Along with other plant growth regulators, however, NO is a reactive nitrogen species, and studies have shown its effects on different cells are either protective or toxic, depending on its concentration and the position of action (Lamattina et al., 2003). In recent years, increasing evidence has demonstrated that NO serves as a signal in developmental, hormonal, and stress responses in plants at lower concentration (Beligni et al., 2002; He et al., 2004; Xu et al., 2010). In relation to abiotic stresses, application of the NO donor sodium nitroprusside has been shown to reduce the harmful effects of salinity on plants (Fan et al., 2010). Furthermore, NO is able to scavenge reactive oxygen species (ROS) as an antioxidant agent and alter antioxidative gene expression as a signaling molecule, thus protecting plant cells from oxidative damage (Arasimowicz and Floryszak-Wieczorek, 2007). In our previous studies, NO was shown to alleviate salt stress in cucumber seedlings by regulating free polyamines and scavenging free radicals (Fan et al., 2010). Pro levels are correlated with NO because L-arginine is a common precursor in their biosynthesis (Gao et al., 2009). The effects of exogenously applied NO on Pro accumulation and the metabolism of plants grown under salt stress are still unclear. Therefore, cucumber, an important horticultural crop, was selected as the test plant for this study because it is highly sensitive to salinity, especially during germination and the seedling stage.

In this study, we investigated the effects of exogenously applied NO on Pro content and related enzymes’ activities in cucumber seedlings under NaCl stress. The main goals of this research were to study the effects of exogenous NO application on Pro metabolism in cucumber seedlings grown under salt stress and to elucidate the physiological mechanism of the increased tolerance of cucumber plants to salt stress resulting from the exogenous application of NO.

**Materials and Methods**

The experiments were carried out in a greenhouse at Nanjing Agricultural University, Nanjing, China. Cucumber (cv. Jinchun 2) seeds were placed in sterile petri plates on filter paper moistened with distilled water. They were allowed to germinate in the dark in a thermostatically controlled chamber at 29 ± 1 °C for ≈30 h. The germinated seeds were sown in washed quartz sands. The average temperatures under natural light were 25 to 30 °C during the day and 16 to 20 °C at night. Relative humidity fluctuated between 60% and 70%. At the second fully expanded leaf stage (at 21 d after sowing), the cucumber leaves were numbered from the apical to the basal nodes. Seedlings were removed from the plastic plates, and the roots were rinsed with distilled water. Uniformly sized healthy seedlings were selected and transferred into troughs (51 × 33 × 20 cm) filled with 20 L full-strength Hoagland’s nutrient solution, which was aerated for 40 min each hour. After pre-culturing for 3 d, the seedlings were treated with one of the following methods: 1) control [CK (full-strength Hoagland’s nutrient solution)]; 2) NaCl treatment [NaCl (full-strength Hoagland’s nutrient solution containing 50 mM NaCl)]; 3) SNP treatment [NaCl + SNP (full-strength Hoagland’s nutrient solution containing 50 mM NaCl with 100 μM SNP)]; or 4) NaNO2 treatment [NaCl + NaNO2 (full-strength Hoagland’s nutrient solution containing 50 mM NaCl with 1 μM NaNO2)]. SNP was used as the NO donor. The NaNO2 treatment served as a second control because the decomposition of 100 μM SNP generates a maximum of 1 μM NO2⁻ as a byproduct. Troughs were arranged in a completely randomized block design with three replicates for each treatment (for a total of 12 troughs). All the nutrient solutions were renewed every 2 d to maintain identical concentrations.

Leaf and root samples from healthy cucumber seedlings were harvested in triplicate at 0, 2, 4, 6, and 8 d after initiation of the treatment and analyzed immediately.

Free Pro content was determined according to Bates et al. (1973). Approximately 0.5 g of leaf or root samples from each group was homogenized in 3% (w/v) sulphasalicylic acid, and the homogenate was filtered through filter paper. After adding acid ninhydrin and glacial acetic acid, the resulting mixture was heated at 100 °C for 1 h in a water bath. The reaction was then stopped with an ice bath. The mixture was extracted with toluene, and the absorbance of the fraction with the toluene aspired from the liquid phase was read at 520 nm. Pro concentration was measured with a calibration curve and expressed as micrograms of Pro per gram fresh weight.

P5CS activity was assayed according to the method of Vogel and Kopac (1960) modified as follows. Approximately 0.7 mL of reaction medium containing 50 mM Tris (pH 7.5), 2 mM MgCl₂, 10 mM ATP, 1 mM NADH, 50 mM glutamic acid, and 0.1 mL of enzyme extract was incubated at 37 °C for 30 min. The reaction was then stopped with 0.3 mL of 10% (w/v) trichloroacetic acid. The color reaction was developed by incubating the solution with 0.1 mL of 0.5% (w/v) aminobenzaldehyde for 1 h. After centrifugation at 12,000 g, for 10 min, the supernatant fraction was removed and the absorbance at 440 nm was measured.

A 0.1 absorbance change in optical density at 440 nm·h⁻¹ was defined as one unit of enzyme activity.

For the extraction and assay of PDH, fresh leaves or roots (0.5 g) were homogenized in 50 mM Tris-HCl buffer containing 7 mM MgCl₂, 0.6 M KCl, 3 mM EDTA, 1 mM dithiothreitol, and 5% (w/v) polyvinylpolypyrrolidone, crosslinked, and adjusted to a pH of 7.4. The homogenate was filtered and centrifuged at 39,000 g, for 20 min at 4 °C (Roesales et al., 2007). The supernatant was used to determine PDH activity. PDH activity was assayed by reduction of NAD⁺ (or NADP⁺) at 340 nm. The reaction mixture had a pH of 10.3 and contained 0.15 mM Na₂CO₃·HCl buffer, 15 mM L-proline, and 1.5 mM NAD⁺ or NADP⁺ (Miller and Stewart, 1976).

Protein was measured following the procedure of Bradford (1976) with bovine serum albumin as the standard.

All experiments were performed in triplicate. All data were statistically analyzed with SAS (Version 8.2; SAS Institute, Cary, NC) using Duncan’s multiple range test at the 5% level of significance.
Results

Pro content. As shown in Figure 1A, the Pro content in the cucumber seedling leaves increased significantly under NaCl stress compared with the CK; Pro content was 146%, 196%, 299%, and 356% of the CK plant content at 2, 4, 6, and 8 d of treatment, respectively. Exogenous supply of SNP further increased the Pro content. The Pro content in the leaves of the cucumber seedlings grown in the SNP treatment reached the highest level at 8 d of treatment and was 12% higher than that grown under NaCl stress alone.

Similarly, NaCl stress significantly increased the Pro content of the roots (Fig. 1B). However, the leaves always maintained a higher level of Pro than the roots. A highly significant increase was observed in SNP + NaCl as compared with other treatments. Although the difference of Pro content between the salt treatment and the SNP treatment decreased after 8 d of treatment, the difference remained obvious.

Compared with NaCl stress, the application of NaNO2 to the salinized solution had no obvious effect on the Pro content in the leaves or roots.

Pyrroline-5-carboxylate synthetase activity. Salt stress caused an increase of the P5CS activity in the cucumber seedling leaves as represented in Figure 2A. SNP treatment brought out higher P5CS activity over the NaCl stress alone. The P5CS activity in the leaves treated with SNP + NaCl was significantly higher than that under NaCl stress alone at 6 and 8 d of treatment. The application of NaNO2 to the salinized solution produced an P5CS activity similar to that produced under NaCl treatment alone throughout the treatment period (Fig. 2A).

The P5CS activity of the cucumber roots increased under NaCl stress, which was significantly higher than that under CK conditions at 4 and 8 d of treatment. Application of SNP increased the P5CS activity in the roots under NaCl stress, and with SNP in solution, the activity in the roots under NaCl stress was significantly higher than that under NaCl stress alone at 4, 6, and 8 d. Combining 50 mM NaCl with 1 µM NaNO2 in solution had no obvious effects on the P5CS activity in the roots (Fig. 2B).

Proline dehydrogenase activity. Figure 3A depicted the PDH activity in leaves grown under various treatments. The PDH activity in the leaves decreased during exposure to salinity.
The difference between the NaCl treatment and the CK at 4, 6, and 8 d was significant. The addition of SNP to the salinized nutrient solution reduced the PDH activity throughout the treatment period. Applying NaNO2 to the salinized solution had no obvious effect on the PDH activity in the leaves.

The PDH activity in the roots of the cucumber seedlings showed a similar trend to that in the leaves. As compared with plants grown in CK solution, the PDH activity in the roots under salt stress decreased throughout the treatment period and reached a significant level at 4, 6, and 8 d. Application of SNP to the salinized nutrient solution decreased the PDH activity throughout the treatment period, and the PDH activity was markedly lower than that under the salt treatment alone at 2 d. The PDH activity was higher in the roots than in the leaves throughout the treatment period. Combining 50 mM NaCl with 1 μM NaNO2 in solution had no obvious effects on the PDH activity in the roots (Fig. 3B).

PROTEIN CONTENT. The protein content in the leaves markedly increased under salt stress and continued to increase throughout the treatment period (Fig. 4A). The addition of SNP to the salinized nutrient solution increased the protein content, which was 145%, 145%, 111%, and 116% of that under NaCl stress alone at 2, 4, 6, and 8 d, respectively. Applying NaNO2 to the salinized solution had no obvious effect on the protein content in the leaves. Figure 4B showed the protein content in the roots of the cucumber seedlings in CK and in the presence of various treatments. The protein content in the roots increased under salt stress throughout the treatment period. The addition of SNP to the salinized nutrient solution clearly elevated the protein content. The protein content was higher in the leaves than in the roots throughout the treatment period. Combining 50 mM NaCl with 1 μM NaNO2 in solution had no obvious effects on the protein content in the roots.

Discussion and Conclusions

The mechanisms of salt tolerance are not yet clear, but they can be partially explained by stress-adaptation effectors that mediate ion homeostasis, osmolyte biosynthesis, ROS scavenging, water transport, and long-distance response coordination (Hasegawa et al., 2000). The typical first response of all plants to salt stress is osmotic adjustment. The accumulation of compatible solutes in the cytoplasm is considered to be a mechanism that contributes to salt tolerance (Jaleel et al., 2007). To counter salt stress, plants decrease the osmotic potential of their cells by synthesizing and accumulating compatible osmolytes such as Pro, which participate in osmotic adjustment (Ghoulam et al., 2002). The accumulation of Pro under stress in many plant species has been correlated with stress tolerance (Ashraf and Foolad,

Fig. 3. Effects of exogenous nitric oxide on proline dehydrogenase (PDH) activity in the (A) leaves and (B) roots of cucumber seedlings under 50 mM NaCl stress at 0, 2, 4, 6, and 8 d. Each histogram represents the mean value of three independent experiments, and the vertical bars indicate SE (n = 3). Significances were tested within the same day. Different letters indicate significant differences between treatments through Duncan’s multiple range test at P < 0.05; CK = control; SNP = sodium nitroprusside.

Fig. 4. Effects of exogenous nitric oxide on soluble protein content in the (A) leaves and (B) roots of cucumber seedlings under 50 mM NaCl stress at 0, 2, 4, 6, and 8 d. Each histogram represents the mean value of three independent experiments, and the vertical bars indicate SE (n = 3). Significances were tested within the same day. Different letters indicate significant differences between treatments through Duncan’s multiple range test at P < 0.05; CK = control; SNP = sodium nitroprusside.
that exogenous NO enhanced salt tolerance by increasing the oxidative damage (Fan et al., 2010). Ruan et al. (2004) reported enhanced activity of antioxidants, which protect plants from NO-induced enhancement in plants grown under salinity stress (Zhang et al., 2006). In our previous studies, we showed that NO-induced enhancement in plants grown under salinity stress might have been caused by NO-induced changes in physiological processes. NO-induced increase in growth could be related to the enhanced activity of antioxidants, which protect plants from oxidative damage (Fan et al., 2010). Ruan et al. (2004) reported that exogenous NO enhanced salt tolerance by increasing the Pro content in leaves of wheat (Triticum aestivum). Our studies showed that NO also increased the Pro content in cucumber leaves and roots under salt stress. We conclude that NO enhanced the salt tolerance of cucumber seedlings by increasing the Pro content.

Pro accumulation in plant tissues has been attributed to 1) an increase in Pro biosynthesis; 2) a decrease in Pro degradation; 3) a decrease in protein synthesis or Pro use; and 4) the hydrolysis of proteins (Ashraf and Foolad, 2007). Hare et al. (1999) described in detail the molecular regulation mechanisms of Pro synthesis and degradation. Under salt stress, Pro is synthesized mainly through the glutamic acid pathway. Glu is the major amino acid involved in Pro synthesis, and transgenic tobacco (Nicotiana tabacum) plants with a reduced expression of cytosolic glutamine synthetase accumulated less Pro than non-transformed plants under salt stress (Kavikishore et al., 1995). P5CS is a rate-limiting enzyme for the biosynthetic pathway in higher plants and is feedback-inhibited by Pro (Zhang et al., 1995). In this pathway, Pro is synthesized from Glu through P5C by two successive reductions, which are catalyzed by P5CS and P5C reductase (Hare et al., 1999). High levels of Pro synthesis in stressed plants under field conditions could favor a better recovery. Also, it has been reported that higher Pro accumulation in P5CS-transformed tobacco plants reduces free radical levels, as measured by malondialdehyde content, in response to osmotic stress (Parvanova et al., 2004). Transgenic tobacco plants overexpressing P5CS have shown an increased concentration of Pro and resistance to both drought and salinity stresses (Kishor et al., 1995). Salt stress, dehydration stress, or abscisic acid can induce the expression of P5CS (Silva-Ortega et al., 2008; Sripiyowanich et al., 2010). Vendruscolo et al. (2007) reported the effects of water deficit on wheat plants transformed with the Vigna aconitifolia P5CS cDNA and found that the tolerance to water deficit in transgenic plants was mainly the result of mechanisms that protect plants against oxidative stress, not to osmotic adjustment. Uchida et al. (2002) proved that SNP induced the mRNA expression of P5CS by a Northern blot analysis, and this provided the basis for our research on the adjustment of Pro metabolism under salt stress. In this study, we found that salt stress improved the P5CS activity of cucumber, and NO further increased P5CS activity in leaves and roots, in accordance with Ruan et al. (2004). Hong et al. (2000) demonstrated that the feedback regulation of P5CS played a role in controlling the level of Pro in plants under both normal and stress conditions. This contradicts the continued synthesis of Pro by plants under stress. P5CS is subject to allosteric regulation by Pro (Hu et al., 1992; Zhang et al., 1995). The inhibition of feedback regulation of P5CS is lost during stress, and this increases Pro accumulation. We observed an increase in P5CS activity and Pro content only under salt stress alone or under salt stress with SNP in solution.

Conversely, Pro content also depends on its degradation (Yoshida et al., 1997). PDH, the key enzyme that catalyzes Pro degradation in mitochondria, acts from transcriptional expression to protein synthesis and finally catalyzes Pro into P5C, which is then converted into glutamic acid by P5C dehydrogenase (P5CDH). These steps constitute the principal processes of oxidative Pro degradation (Hare et al., 1999). In A. thaliana, AtPDH was the only enzyme acting as a functional PDH (Ashraf and Foolad, 2007). Our studies showed that exogenous NO reduced the PDH activity in cucumber leaves and roots, resulting in a reduction of Pro degradation in cucumber seedlings under salt stress. In response to water stress, Pro accumulation is dependent on the transcriptional activation and translation of NAD(P)H-dependent P5CS to produce P5C (Yoshida et al., 1995). Furthermore, Pro degradation is controlled by PDH and P5CDH. Parre et al. (2007) identified the signaling components involved in the regulation of Pro metabolism under water stress in A. thaliana and indicated that PLC (phosphoinositide-specific phospholipase C)-based signaling was a committed step in Pro biosynthesis under salinity stress but not under mannitol stress.

Our results showed that the Pro content in leaves was higher than that in roots based on a combination of the changing trends of P5CS and PDH activity. This could be attributed to serious metabolic problems such as deficits of substrates and intermediates involved in Pro synthesis, caused by the presence of salt on the outside of roots in the nutrient solution, a condition to which leaves are subjected indirectly. Thus, we observed that the P5CS activity in roots was higher than that in leaves and that the PDH activity in roots was lower than that in leaves. In contrast, Pro accumulation was higher in leaves than in roots.

Exogenous NO increased the protein content in leaves and roots and thereby increased salt tolerance. With its hydrophobic end proteins and hydrophilic end-binding hydrones, Pro has a strong hydrating ability. This forces the proteins to bind more water molecules, preventing dehydration and denaturation (Hoekstra et al., 2001). A small increase in Pro could be caused by the hydrolysis of Pro-rich proteins (Song et al., 2005). We observed protein and Pro accumulation, indicating that the accumulation of Pro may come from the hydrolysis of Pro-rich proteins under salt stress. Moreover, protein accumulation was not caused by the use of Pro for protein synthesis. Buhl and Stewart (1983) also found that salt treatment dramatically increased the synthesis of Pro from Glu and that its use in oxidation and protein synthesis decreased by 50% and 60%, respectively, in barley (Hordeum vulgare) under salinity stress.

Based on the results from this study, it can be concluded that exogenous SNP increased salt tolerance in cucumber seedlings by adjusting the biosynthesis of Pro; as the P5CS activity increased and the PDH activity decreased, the accumulation of Pro accelerated. SNP could be used as a potential growth regulator to improve plant growth under salinity stress. Furthermore, the application of NO2- (a byproduct from the decomposition of SNP) did not produce effects similar to those of SNP.

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