Pre-incubation of *Sinorhizobium meliloti* with Luteolin, Methyl jasmonate and Genistein Affecting Alfalfa (*Medicago sativa* L.) Growth, Nodulation and Nitrogen Fixation under Salt Stress Conditions

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**ABSTRACT**

Salinity is among important soil stresses adversely affecting the process of nitrogen (N) fixation in leguminous plants in different parts of the world. It has been indicated that salinity can inhibit the early stages of nodulation process between bacterium and the host plant including the exchange of signal molecules (*nod* gene inducers). There has not been any research regarding the effects of *nod* gene inducers on the growth of alfalfa inoculated with *Sinorhizobium meliloti* under saline conditions. A growth chamber experiment was conducted to determine the effects of pre-incubation of *S. meliloti* with effective inducers of *nod* genes Luteolin, Methyl jasmonate and Genistein on the growth and N-fixation of two different alfalfa (*Medicago sativa* L.) cultivars (Yazdi and Hamedani) under salt stress. *Nod* gene inducers increased alfalfa growth and N fixation under normal as well under salt stressed conditions. Yazdi cultivar showed to be more tolerant to salinity than Hamedani with a higher growth rate and N fixation. Luteolin was the most effective *nod* gene inducer on plant growth and N fixation under normal and as well under salt stressed conditions. The results suggest that pre-incubation of *S. meliloti* with effective *nod* gene inducers can improve alfalfa growth and N fixation under salinity stress.

**Keywords**: Alfalfa (*Medicago sativa* L.), Genistein, Luteolin, Methyl jasmonate, *Nod* gene inducers, Salinity, *Sinorhizobium meliloti*.

**INTRODUCTION**

The process of symbiosis between legume and rhizobia is the most important source of biological nitrogen (N) fixation in agricultural systems (Kahindi *et al.*, 1997). Alfalfa is one of the important nitrogen extractors, in the form of nitrate because of its capability to produce many nodules for nitrogen fixation (Rasse and Smucker, 1999). *Sinorhizobium meliloti* is the bacteria developing symbiotic association with alfalfa (*Medicago sativa* L.) plants. The symbiotic process between *S. meliloti* and alfalfa is complex and includes a series of signal exchange between the plant and bacteria (Long, 1989; Brewin, 1991; Denarie, *et al.*, 1993; Rhijin *et al.*, 1995).

During the early stages of symbiosis and in response to signal molecules such as flavonoids and plant hormones, released through legume roots, a series of genes known as *nod* (nodulation) genes are expressed in bacteria, resulting in the formation of nodules (Rosas *et al.*, 1998). As a result of *nod* gene expression, through *nod* gene inducers, *nod* factors, lipo-chito-oligosaccharides, are produced causing root hair curling, cell division and root nodule formation. Cells of *S. meliloti* is the bacteria developing symbiotic association with alfalfa (*Medicago sativa* L.) plants. The symbiotic process between *S. meliloti* and alfalfa is complex and includes a series of signal exchange between the plant and bacteria (Long, 1989; Brewin, 1991; Denarie, *et al.*, 1993; Rhijin *et al.*, 1995).

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*meliloti* then invade the developing root nodule through infection threads, eventually forming nodule where they differentiate into bacteroids for nitrogen fixation (Long, 1989; Brewin, 1991; Denarie and Cullimore, 1993; Rhijin and Vanderleyden, 1995). Jasmonic acid and Methyl jasmonate, collectively known as jasmonates, are signaling molecules in plants, play a role in response to wounding, pathogen attack, reproduction, development, metabolic regulation and abiotic stress (Devoto and Turner, 2003; Howe, 2004; Mabood et al., 2005). Mabood et al., (2005) expressed that Jasmonates can act as signal molecules in the early stages of legume-rhizobia symbiosis, inducing nod genes of *Bradyrhizobium japonicum*. Studies have shown that Genistein is also one of the flavonoids inducing rhizobial nodulation genes in the early stages of symbiosis between soybean (*Glycine max* L.) and *B. japonicum* (Zhang et al., 1996; Pan and Smith, 2000).

Nod factors produced in the presence of Genistein can support nodule growth through regulating hormonal activities (Hirsch, 1992) and cell division (Verma, 1992). Luteolin is also one of the flavonoids inducing nodulation genes in the bacterium *S. meliloti* (Hubac et al., 1993). It is released from alfalfa imbibing seeds; it increases growth rate of *S. meliloti* (Hartwig et al., 1991) and thus plays an important role in nodule formation (Alexander, 1985). Appropriate flavonoids in root secretions are important factors in root nodule formation (Richardson et al., 1998).

In a number of studies, application of nod inducing molecules as a tool to increase nodulation and N fixation has been proved (Davis and Johnston, 1990; Bandyopadhyay et al., 1996; Pan et al., 1998). Such environmental stresses as salinity, pH and low temperature in root zone, partially or completely inhibit the early stages of symbiosis, including signal transfer between legume plants and rhizobium (Hungria et al., 1991; Ikeda, 1994; Miransari and Smith, 2007; 2008; 2009). Accordingly, the proliferation of bacteria (Brown et al., 2001) and N fixation (Bilgrami, 1989) is influenced.

In legumes, salinity by sodium chloride, in the range of 50 to 100 mM, significantly limited yield by adversely affecting plant growth, root nodules bacteria, the process of symbiosis, as well as N fixation (Zahran and Sprent, 1986; Bekki et al., 1987). Environmental stresses influence the early stages of symbiosis and nodulation, with flavonoids having their role in these stages and their effect as signal molecules in symbiosis establishment between legume-*rhizobium* having been studied. Mabood et al. (2006a) found that Jasmonates induced rhizobial nod genes and therefore enhanced the establishment of symbiosis process.

Poustini et al. (2005) studied the effects of low temperature on bean plants showing that pre-incubation of the rhizobial cells with Genistein and Methyl jasmonate increased nodulation, N fixation and growth in both optimal and suboptimal root zone temperatures. Zhang and Smith (1996) indicated that the reduction in soybean yield in low temperatures is mainly due to N limitation. Pre-incubation of *B. japonicum* with Genistein increased grain yield as well as protein. Miransari and Smith (2009) studied the effects of salinity on soybean growth and nodulation showing that pre-treating rhizobial cells with Genistein increased nodulation and growth in soybean.

Begum et al. (2001a) studied pea growth and nodulation under short growing season and found that plants, inoculated with preinduced *Rhizobium leguminosarum* bv. *viceae* with Hesperetin, Narngenin, Luteolin and Apigenin carried a higher nodule number than the control plants. With respect to the significance of salinity stress affecting legume production in different parts of the world and because salinity stress usually influences the early stages of symbiosis the following objectives were tested through this experiment. (1) The effects of salinity on alfalfa growth and nodulation, and (2) to evaluate if pre-incubating *S. meliloti* with different nod gene inducers including Luteolin, Methyl jasmonate and Genistein can alleviate the adverse effects of salinity on alfalfa growth, nodulation and nitrogenase activity under salt stress.
MATERIALS AND METHODS

To evaluate how pre-incubating *S. meliloti*, strain 1021 (provided by the Institute for Genetics, Department of Biology, University of Dresden, Germany) with different *nod* gene inducers, including Luteolin, Methyl jasmonate and Genistein, affect alfalfa growth, a growth chamber experiment was conducted. The experiment was arranged in three randomized blocks, with a 2×2×4 factorial design, two cultivars (Yazdi and Hamedani), two levels of salt (0 and 120 mM NaCl) and three inducers of nodulation (Luteolin, Methyl jasmonate and Genistein) along with control.

*S. meliloti*, strain 1021 were grown in Petri dishes at 28°C for 48 hours in Yeast Mannitol Agar (YMA) medium (mannitol 10 g, K₂HPO₄ 0.5 g, MgSO₄·7H₂O 0.2 g, NaCl 0.2 g, yeast extract 0.5 g, agar 15 g plus distilled water 1,000 ml). A single colony of *S. meliloti*, strain 1021 was cultured in 4 ml yeast mannitol broth (YMB) medium (mannitol 10 g, K₂HPO₄ 0.5 g, MgSO₄·7H₂O 0.2 g, NaCl 0.2 g, yeast extract 0.5 g and distilled water 1,000 ml). It was shaken at 28°C for 24 hours and then subcultured into 400 ml of fresh YMB medium and then shaken at 28°C for 24 hours. The Bacterial culture was distributed in four flasks containing 100 ml YMB medium. Each flask represented one treatment, namely; *S. meliloti* with no inducer addition (control); *S. meliloti* induced with Luteolin (10 µM); *S. meliloti* induced with Methyl jasmonate (50 µM); and *S. meliloti* induced with Genistein (10 µM).

Luteolin, Methyl jasmonate and Genistein stock solutions were prepared and then (except control) Luteolin, Methyl jasmonate and Genistein were added to the culture and shaken at 28°C for 1 day.

The related concentrations of each inducer were selected according to the results obtained by other researchers (Capela et al., 2005; Mabood and Smith, 2005; Miransari and Smith, 2009).

Alfalfa cultivar seeds (Yazdi and Hamedani) (provided by the Institute of breeding and Providing Seeds and Seedlings of Karaj, Iran) were surface sterilized in sodium hypochlorite (2.5%) and then rinsed several times with distilled water. Seeds were germinated on plates containing water agar (1%) at 28°C for 48 hours (Howieson et al., 2000).

Five germinated seeds were planted in pots each containing sterile vermiculite, under growth chamber conditions, with light cycle of 16 (light) and 8 (dark) hours, along with day and night temperatures of 28 and 15°C, respectively. Each germinated seed was inoculated with 1 ml of fresh culture of *S. meliloti*, strain 1021, pre-incubated with the three different inducers (Beck et al., 1993).

Salt treatment was applied at sowing (Miransari and Smith, 2009). Salty Hoagland’s solution (Hoagland and Arnon, 1950), with no nitrate, containing 120 mM of NaCl (S₂) and control (S₁) were employed. Salinity concentration was selected based on salt tolerance threshold and maximum salt tolerance of alfalfa (Shannon, 1984; Fougere et al., 1991). Plants were harvested one month after sowing (Miransari and Smith, 2009) and nodule number and dry weight, shoot and root dry weight and nitrogenase enzyme activity measured by acetylene reduction method (Herdina and Silsbury, 1990) were measured. Data analysis was performed using MSTATC statistical software and comparison of means was performed with Duncan’s multiple range test.

RESULTS

Effects of Cultivar on Nodulation and Plant Growth

Analysis of variance indicated that there existed significant differences between shoot and root dry weights, nodule number (P< 0.01), and dry weight (P< 0.05) for different alfalfa cultivars. Yazdi cultivar resulted in
higher nodule number and dry weight, as well as higher shoot and root dry weights than Hamedani cultivar (Table 1).

**Effects of Inducers on Nodulation and Plant Growth**

Analysis of variance indicated that there was a significant interaction between salt and inducers on nodule number and dry weight and on shoot and root dry weights (P< 0.01). The results revealed that salinity decreased shoot and root dry weights and also nodule number as well as dry weight. Accordingly, although salinity adversely affected nodulation and plant growth, plants inoculated with rhizobium, pre-incubated with inducers, had higher nodule number and dry weight as well as shoot and root dry weights in both S1 and S2 treatments (P< 0.01) relative to control (Figures 1 and 2). The percentage of increase was higher in S2 treatment than in S1 and as well in Luteolin treatment in comparison with the other related treatments (Table 2).

**Nitrogenase Activity (Acetylene Reduction)**

Analysis of variance showed that there was a significant interaction between salt and cultivars on nitrogenase activity (P<
Figure 2: Shoot and root dry weight as affected by as well as salt and inducer treatment. Histograms with different letters are statistically different as by Duncan’s test.

Table 2: Effect of different nod gene inducers on the percentage of increase in nodulation and plant growth at 0 and 120 mM NaCl in comparison with control. Means of cultivars Yazdi and Hamedani.

<table>
<thead>
<tr>
<th>Nodule number (%)</th>
<th>Nodule dry weight (%)</th>
<th>Shoot dry weight (%)</th>
<th>Root dry weight (%)</th>
<th>Nitrogenase activity (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Inducers</td>
<td>S₁</td>
<td>S₂</td>
<td>S₁</td>
<td>S₂</td>
</tr>
<tr>
<td>Luteolin</td>
<td>57.43⁸⁺⁺⁺⁺</td>
<td>57.43⁸⁺⁺⁺⁺</td>
<td>57.43⁸⁺⁺⁺⁺</td>
<td>57.43⁸⁺⁺⁺⁺</td>
</tr>
<tr>
<td>MJ</td>
<td>102.8⁸⁺⁺⁺⁺</td>
<td>102.8⁸⁺⁺⁺⁺</td>
<td>102.8⁸⁺⁺⁺⁺</td>
<td>102.8⁸⁺⁺⁺⁺</td>
</tr>
</tbody>
</table>

S₁: Salinity treatment control; S₂: 120 mM NaCl, MJ: Methyl jasmonate.
Means followed by different letters are significantly different using Duncan’s method.
Capital letters compare levels of salt, while small ones compare different inducers within each stress level.

DISCUSSION

The results finally indicate that application of inducers increased nodulation and plant growth under stressed and non-stressed conditions relative to control. Salinity can
adversely affect the early stages of symbiosis (Hungria et al., 1991; Miransari and Smith, 2007; 2009). Most of these inhibitions are associated with such root hair morphology and physiologies, as root hair growth, diameter, structure as well as curling (Miransari et al., 2006). Salt stress limits the translocation of photosynthates to the roots and thus reduces nodulation and plant growth (Miransari and Smith, 2007).

Flavonoids produced by legume plants result in the induction of rhizobial nodule genes required for infection, supporting bacterial movement toward the root, and increasing the rate of growth of bacterial cells, thus affecting root nodulation. (Phillips and Tsai, 1992). Nod gene inducers have resulted in enhanced nodulation in different legume host plants including pea (Ahlawat et al., 1998; Bandyopadhyay et al., 1996; Lira et al., 2003; Novak et al., 2002) bean (Poustini et al., 2005, 2007) and soybean (Zhang and Smith, 1995; Pan et al., 1998; Belkheir et al., 2000).

There has previously been some interesting research work carried out according to the following by Professor Donald Smith and his research team on soybean regarding the use of nod gene inducers for the incubation of B. japonicum under stress. Pre-treatment of B. japonicum with nod gene inducers which resulted in an increase in plant growth and development. Genistein and Jasmonates resulted in higher dry matter than the control under different conditions including under suboptimal root zone temperature. Increased dry matter can be resulted by enhanced N fixation as the nod gene inducers can increase nodule number as well as dry weight (Zhang and Smith 1995; 1997; Mabood and Smith; 2005; 2006b). Pan and Smith (2000) found that incubation of B. japonicum with 40 µM Genistein increased root dry weight in soybean.

Begum et al. (2001a, b) reported that pre-treatment of R. leguminosarum bv. viceae cells with appropriate flavonoid molecules

Figure 3. Nitrogenase activity as affected by salt and cultivar and salt and inducer treatments. Histograms with different letters are statistically different by Duncan’s test.
including Apigenin, Daidzein, Genistein, Hesperetin, Kaempferol, Luteolin, Naringenin and Rutin increased pea and lentil dry matter, subjected to sub optimal root zone temperatures, under greenhouse as well as field conditions. Poustini et al., (2007) used pretreated Rhizobium leguminosarum bv. phaseoli with Genistein and Methyl jasmonate to inoculate bean plants and found increase in nodule number, N content and plant dry matter. According to Miransari and Smith (2009), inoculation of soybean seeds with B. japonicum, pretreated with the nod gene inducer, Genistein, increased nodule number and dry weight, N fixation and hence shoot dry weight under salinity.

The results of this research work also indicated that the nod gene inducers resulted in higher root and shoot dry weight, nodule number and dry weight as well as in nitrogenase enzyme activity, under stressed and non-stressed conditions relative to the control. Yazdi cultivar as compared with Hamadani cultivar had higher shoot and root dry weight, as well as higher nodule number and dry weight under salinity conditions. Yazdi cultivar probably uses some such mechanisms as absorption of a lower level of sodium and chloride ions, production of osmolyets and allocation of salt to its cellular vacuoles, increasing plant resistance to salinity. It may also be able to excrete higher levels of signal molecules under salinity resulting in higher root nodulation.

Luteolin was the most effective signal molecule under salinity. Plants inoculated with pre-treated rhizobial cells with Luteolin exhibited higher nodulation, nitrogenase activity, N fixation and dry matter accumulation in both normal and salt stress conditions than plants pre-treated with rhizobial cells from Methyl jasmonate and Genistein. It is likely that the biochemical structure of Luteolin is much less affected by salt ions relative to Methyl jasmonate and Genistein. In addition, it is probably a more affective nod gene inducer activating the nodulation genes in S. meliloti. It can also be due the fact that alfalfa produces Luteolin, inducing the nod genes in S. meliloti (Hartwig et al., 1990), while Genistein produced by soybean inhibits such induction (Hirsch et al., 2001). Signal molecules act completely specifically in the induction of rhizobium nod genes. This is the reason for an indication of the specificity of legume-rhizobium symbiosis (Long, 2001).

CONCLUSIONS

The results, which to the best of the authors’ knowledge have not been previously investigated, indicate the effectiveness of the signal molecule Luteolin in the alleviation of salt stress on alfalfa growth and nodulation. Also the differences between different cultivars indicated that Yazdi cultivar was a more resistant cultivar to salinity than Hamedani. Pre-incubation of S. meliloti with, especially, Luteolin can inhibit the adverse effects of salinity on alfalfa growth and nodulation.

REFERENCES


