# ΦΥΤΟΝ

REVISTA INTERNACIONAL DE BOTÁNICA EXPERIMENTAL INTERNATIONAL JOURNAL OF EXPERIMENTAL BOTANY

FUNDACION ROMULO RAGGIO Gaspar Campos 861, 1638 Vicente López (BA), Argentina www.revistaphyton.fund-romuloraggio.org.ar

# PGPR inoculation improves growth, nutrient uptake and physiological parameters of *Capsicum chinense* plants

La inoculación con PGPR mejora el crecimiento, la absorción de nutrientes y parámetros fisiológicos en plantas de *Capsicum chinense* 

## Castillo-Aguilar C de la C<sup>1</sup>, JJ Zúñiga-Aguilar<sup>2</sup>, AA Guzmán-Antonio<sup>2</sup>, R Garruña<sup>3</sup>

Abstract. The Habanero pepper (Capsicum chinense) is intensively cultivated in the Yucatan peninsula, México. Because of adverse environmental conditions, it required seeding in germination trays, from which six-week-old seedlings were transplanted to the soil. Adequate nursing and fertilization programmes were made to improve health and vigour before seedlings transplanting. During seed germination, we investigated the effects of inoculation with four plant growth-promoting rhizobacteria (PGPR) on growth, nutrient uptake and gas exchange of 8-week-old Capsicum chinense plants. Inoculation was made with Pseudomonas sp. -P61-, Pseudomonas sp. -A46-, Bacillus pumillus -R44-, and Paenibacillus polymyxa -BSP1.1-. The BSP1.1 strain produced the highest increase in plant height (22.52 cm), leaf area (99 cm<sup>2</sup>), fresh weight of roots (0.337 g) and shoots (2.83 g), and dry weights of roots (0.05 g) and shoots (0.43 g). Nitrogen uptake was similar (P>0.05) among treatments. However, seedlings inoculated with BSP1.1 had a higher accumulation than controls when they were exposed to P and K (50 and 40%, respectively). Likewise, seedlings inoculated with the BSP1.1 strain improved both photosynthesis (2 µmol/m²/s) and water use efficiency (7.3 µmol CO<sub>2</sub>/mmol H<sub>2</sub>O), and decreased transpiration rate (0.27 mmol/m<sup>2</sup>/s). Thus, the inoculation with the BSP1.1 strain was the best option to enhance growth and vigour of Capsicum chinense plants.

**Keywords:** Habanero pepper; Plant growth-promoting rhizobacteria; Leaf area; Biomass; Bioprotection.

Resumen. El chile habanero (Capsicum chinense) es cultivado extensivamente en la península de Yucatán México. Debido a los factores ambientales adversos, la siembra de las semillas se debió efectuar en bandejas germinadoras. Cuando las plántulas tuvieron seis semanas de edad fueron trasplantadas al suelo. Se efectuaron programas adecuados de semillero y fertilización para incrementar la salud y el vigor de las plántulas antes del trasplante. Se evaluó el efecto de la inoculación de cuatro cepas de rizobacterias promotoras del crecimiento vegetal (Pseudomonas sp. -P61-, Pseudomonas sp. -A46-, Bacillus pumillus -R44-, y Paenibacillus polymyxa -BSP1.1-) sobre el crecimiento, la absorción de nutrientes y el intercambio de gases en plantas de 8 semanas de edad de Capsicum chinense. La cepa BSP1.1 promovió el mayor incremento en altura de planta (22,52 cm), área foliar (99 cm<sup>2</sup>), peso fresco de raíz (0,337 g) y de la parte aérea (2,83 g), y peso seco de raíz (0,05 g) y de la parte aérea (0,43 g). No hubo diferencias significativas entre tratamientos en la absorción de N. Sin embargo, las plántulas inoculadas con BSP1.1 tuvieron una mayor acumulación de P y K que los controles (50 y 40%, respectivamente). Además, las plántulas inoculadas con la cepa BSP1.1 mejoraron la tasa fotosintética (2 µmol/m²/s) y el uso eficiente de agua (7,3 µmol CO<sub>2</sub>/mmol H<sub>2</sub>O), y disminuyeron la tasa de transpiración (0,27 mmol/m²/s). Así, la inoculación con la cepa BSP1.1 fue la mejor opción para aumentar el crecimiento y el vigor de las plantas de Capsicum chinense.

**Palabras clave:** Chile habanero; Bacterias promotoras de crecimiento vegetal; Área foliar; Biomasa; Bioprotección.

<sup>&</sup>lt;sup>1</sup> Colegio de Postgraduados, Campus Campeche. Carretera Federal Haltunchén-Edzná Km 17.5, C.P. 24450, Sihochac, Champotón, Campeche, México.
<sup>2</sup> Unidad de Bioquímica y Biología Molecular de Plantas, Centro de Investigación Científica de Yucatán, A.C. Calle 43 No. 130, colonia Chuburná de Hidalgo, Mérida C.P. 97200, Yucatán, México.

<sup>&</sup>lt;sup>3</sup> Catedrático CONACYT-Instituto Tecnológico de Conkal. Avenida Tecnológico s/n, Conkal, C.P. 97345, Yucatán, México. renegh10@hotmail.com Address correspondence to: Ph.D. René Garruña, Tel: 52 (999) 912-41-30 ext. 126; *e-mail*: renegh10@hotmail.com ; rgarrunahe@conacyt.mx Received 11.VII.2016. Accepted 20.V.2017.

### INTRODUCTION

Plant growth-promoting rhizobacteria (PGPR) is a term applied to a group of soil bacteria associated with the rhizosphere and the root surface of many plant species. The association of PGPR with the root system can stimulate plant growth and defence responses against pathogens (Vessey, 2003). There are various mechanisms explaining how PGPR favours the growth of plants. For example, through mostly the (1) biological fixation of nitrogen (Bashan et al., 2014); (2) regulation of the synthesis of plant growth-regulators, which can promote root growth, enhancing the uptake of water and minerals from the soil; (3) decrease in ethylene content of developing or stressed plants, inducing the elongation of the root system (Ahemad & Kibret, 2014); (4) the solubilization of soilinsoluble compounds, such as calcium di-and tri-phosphates and other minerals (Marschner, 2007); (5) growth inhibition of both pathogenic and non-pathogenic microorganisms, which may compete for nutrients and antibiosis (Caballero-Mellado, 2006); and (6) biosynthesis of siderophores, which can solubilize and chelate iron from the soil (Marschner, 2007).

PGPR have been applied to commercially important crops, and their growth-promoting capacity has been evaluated (Fernández-Herrera et al., 2007). Reyes-Ramírez et al. (2014) found that the inoculation of *Capsicum chinense* seedlings with *Pseudomonas* sp. increased plant height, stem diameter and dry weight, as well as yield and size of the fruits.

Landrace genotypes of the Capsicum chinense have been traditionally cultivated in the Yucatán Peninsula under adverse environmental conditions (Garruña-Hernández et al., 2014a). It has contributed to generating one of the most pungent peppers worldwide (Bosland & Votava, 2000; Canto-Flick et al., 2008). However, adverse conditions constraint seed germination and seedling establishment, and seeds must be previously sown in pots under greenhouse conditions. The loss of seed viability and the establishment of vigorous seedlings in the field are among the main challenges in these cultivars (Garruña-Hernández et al., 2014b). The performance of plants late in their productivity period is highly dependent on the first stages of development. Thereafter, the establishment of efficient nursery and fertilization protocols is necessary to improve both the health and vigour of seedlings during the first stages of development. The use of PGPR has been considered as an alternative to fortify seedlings before transplantation (Canto-Martin et al., 2004). As a result, the aim of the current study was to evaluate the capacity of four plant growth-promoting rhizobacteria for promoting increases in both growth and vigour of Capsicum chinense seedlings, after adding them during seed germination and in the seedling nursery.

#### MATERIALS AND METHODS

Biological materials and culture conditions. Four bacterial strains isolated from the potato rhizosphere grown in the Valley of Toluca (P61, Pseudomonas tolaasii; R44, Bacillus pumilus) or on an andisol soil from the Valley of Allende (BSP1.1, Paenibacillus polymyxa; A46, Pseudomonas tolaasii), Mexico, were tested in this study. The research was conducted under greenhouse conditions at the Yucatan Scientific Research Center (CICY), Mexico. Seeds of the orange ("Akil") variety of Capsicum chinense were sown in polystyrene trays with sterilized sphagnum peat moss (Premier Mexico, Mexico). After sowing, the seeds were inoculated with a broth culture of the respective PGPR isolates (P61, A46, R44, or BSP1.1), containing 10<sup>7</sup> colony-forming units of bacteria (10<sup>7</sup> CFU/ mL broth). The control seeds were soaked with distilled water. Two weeks after sowing, the inoculated seedlings were dipped again for five min into broth cultures of the respective PGPR strain (107 CFU/mL broth). Control seedlings without bacterial inoculation were treated accordingly. The fertilization programme began 25 days after seed sowing, by submerging the trays twice a week in a solution with 18:30:15 (N:P:K). Measurement of growth and physiological parameters was performed eight weeks after seed sowing.

**Growth parameters.** Eight weeks after sowing, plant diameter and height were measured using a digital Vernier scale, and leaf area was determined using an area metre (LiCor LI-3100, Nebraska, USA). Fresh (FW) and dry (DW) weight were measured for individual organs: leaves and stem (shoot), and roots were separated and weighed Thereafter, they were placed in a forced-air oven at 70 °C until obtaining a constant mass (approximately 72 h).

Determination of nutrient content in plants. All glassware was washed three times with concentrated nitric acid, and rinsed with MilliQ deionized water (18.2 M $\Omega$ ) three times each. Five mL of 70% nitric acid ACS (JT Baker, Phillipsburg, USA) were added to wash Teflon-covered receptacles. Then, the receptacles were heated in a microwave oven (CEM, MARS-Xpress) for 10 min at 200 °C. After washing, the receptacles were rinsed thoroughly with MilliQ deionized water (18.2 M $\Omega$ ) as described above. The protocol reported by Bhandari & Amarasiriwardena (2000) was followed for digestion and analysis of nutrients. In brief, Capsicum chinense plants were harvested eight weeks after sowing and rinsed thoroughly with tap water. Then, whole plants were dried at 70 °C for three days and ground to fine dust in an electric mill. For nutrient analysis, 0.5 g samples of milled plants tissues were mixed with 9 mL of nitric acid (TraceUltra Sigma) and 1 mL of hydrogen peroxide (TraceUltra Sigma), and digestion proceeded for 15 min at 200 °C. After digestion, the receptacles remained inside the microwave oven until they reached

room temperature, and then each was depressurized in a gas extraction cabinet. All samples were diluted 1:25 with MilliQ deionized water (18.2 M $\Omega$ ) prior to analysis. It was conducted by using an inductively coupled plasma optical emission spectrometry (ICP-OES, IRIS Intrepid II XDL model, Thermo®). Blanks of water and nitric acid were processed in the same manner. Nitrogen concentrations in whole plants were measured in ground samples using the Kjeldahl method.

Gas exchange analyses. The plant material was naturally illuminated during 11 h/day. The maximum photosynthetic photon flux density (PPFD) was 1000 µmol/m²/s at noon. The relative humidity (RH) was ca. 75% and the average daily temperature was 27 °C. Those parameters were measured using data loggers (HOBO H08-004-02; Onset Computer Corp., Bourne, MA, USA) and Quantum sensors (LI-190SB; LI-COR, Lincoln, NE, USA) placed on the top of and inside the greenhouse. Gas exchange analyses were conducted inside a greenhouse under relevant growth conditions by a non-invasive method, using a portable infrared gas analyser system (IRGA; LICOR, LI-6400, Nebraska, USA). At 12:00 h, 30 fully expanded young leaves from each treatment were placed in the gas-exchange leaf chamber of the IRGA LI-6400 (ten plants per treatment were exclusively for gas exchange analyses). Photosynthesis  $(A_{N})$  and transpiration (E) were estimated under the greenhouse conditions described above. Water use efficiency was subsequently calculated  $(A_N/E)$ .

**Experimental design.** The determination of growth parameters, including height, stem diameter, leaf area, and biomass (dry and fresh weight) of root and shoot (stem and leaves) was performed using samples from 20 plants. Experimental data were collected from five independent treatments in a randomized complete block design with three replicates. One-way analysis of variance (ANOVA) was calculated for every variable with the Statistical analysis system 9.1.3 (SAS Institute Inc., Cary, NC, USA). After the F tests were significant (P<0.05), multiple comparisons of means were made using the Tukey test, with a significance level of 5%.

#### RESULTS

**Growth parameters.** Inoculation of *Capsicum chinense* seeds with the BSP1.1, R44 and P61 strains promoted a statistically significant increase in plant height (39, 35 and 25%, respectively) with respect to the control plants, while the result with the A46 strain was statistically similar to the control plants (Table 1). The leaf area of plants inoculated with the BSP1.1 and R44 strains was significantly increased by 28 and 21% with respect to the control plants, respectively (Table 1). None of the PGPR treatments promoted significant differences in the stem diameter of plants (Table 1).

 
 Table 1. Height, stem diameter and leaf area of Capsicum chinense plants inoculated with PGPR.

 Tabla 1. Altura, diámetro de tallo y área foliar de plantas de Capsicum chinense inoculadas con PGPR.

Treatments	Height (cm)	Stem diameter (mm)	Leaf area (cm²)
Control	13.80 c	2.01	71.28 c
P61	18.42 ab	2.16	78.36 bc
A46	16.10 bc	2.11	78.17 bc
R44	21.28 ab	2.20	89.96 ab
BSP1.1	22.52 a	2.20	99.00 a
LSD	6.26	1.19	15.68

Data are means. LSD = Least Significant Difference. Different letters in the same column represent statistically significant differences (Tukey  $\alpha = 0.05$ ).

**Plant biomass.** The biomass (fresh and dry weight) of 8-week-old plants, previously inoculated during seed sowing with the BSP1.1 strain was increased in both roots (75.4% in FW and 40% in DW) and shoots (43% in FW and 30% in DW) with respect to the control plants (Table 2). Inoculation with the R44 strain produced significant biomass accumulation only in the fresh weight of shoots (38%) and dry weight of roots (40%) with respect to the control plants. Inoculation with the P61 and A46 strains produced no significant differences in biomass accumulation with respect to the control treatment (Table 2).

Table 2. Biomass of Capsicum chinense plants inoculated withPGPR.

 Tabla 2. Biomasa de plantas de Capsicum chinense inoculadas con PGPR.

Treatments	Fresh weight (g)		Dry weight (g)	
	Root	Shoot	Root	Shoot
Control	0.083 b	1.62 b	0.03 b	0.30 b
P61	0.138 b	2.43 a	0.04 ab	0.33 ab
A46	0.094 b	2.16 ab	0.03 b	0.31 ab
R44	0.141 b	2.62 a	0.05 a	0.35 ab
BSP1.1	0.337 a	2.83 a	0.05 a	0.43 a
LSD	0.084	0.80	0.01	0.12

Data are means. LSD = Least Significant Difference. Different letters in the same column represent statistically significant differences (Tukey  $\alpha = 0.05$ ).

**Gas exchange analysis.** Eight-week-old plants inoculated at seed sowing with BSP1.1 had 36, 54, 32 and 66% higher photosynthesis than plants inoculated with P61, A46, and R44 and the non-inoculated (control) plants, respectively

а

(A)

(B)

(C)

(Fig. 1A). This was when photosynthetic photon flux density was at its maximum (at noon). In contrast, the non-inoculated (control) plants had a transpiration rate that was 41, 50, 14 and 58% higher than plants inoculated with P61, A46, R44 and BSP1.1, respectively (Fig. 1B). Plants inoculated with BSP1.1 had a water use efficiency 55, 62, 67 and 86% higher than that on plants inoculated with P61, A46, and R44 and the controls, respectively (Fig. 1C).

Nutrient uptake. Plants inoculated with BSP1.1 had a P uptake 40 and 50% higher than that on plants inoculated with A46 and the controls, respectively (Table 3). Likewise, plants inoculated with BSP1.1 had a K uptake 40% higher than that on controls (Table 3). There were no significant differences between treatments in N uptake (Table 3).

Table 3. Nutrient uptake (N:P:K) of Capsicum chinense plants inoculated with PGPR.

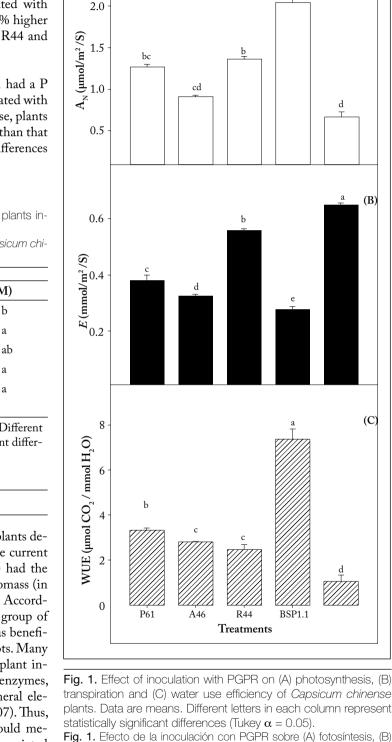
Tabla 3. Absorción de nutrientes (N:P:K) de plantas de Capsicum chinense inoculadas con PGPR.

Treatments	N (mM)	P(mM)	K (mM)
Control	907.9	2172.1 b	121.7 b
P61	1456.7	2736.5 ab	171.3 a
A46	1247.2	2611.8 b	159.8 ab
R44	1711.0	3468.8 ab	179.9 a
BSP1.1	2250.0	4352.8 a	203.3 a
LSD	2828.4	1738.7	46.2
P61 A46 R44 BSP1.1	1456.7 1247.2 1711.0 2250.0	2736.5 ab 2611.8 b 3468.8 ab 4352.8 a	171.3 a 159.8 ab 179.9 a 203.3 a

Data are means. LSD = Least Significant Difference. Different letters in the same column represent statistically significant differences (Tukey  $\alpha = 0.05$ ).

#### DISCUSSION

The efficiency of rhizobacteria to promote larger plants depends on the strain used (Khalid et al., 2004). In the current study, the strain of Paenibacillus polymyxa (BSP1.1) had the best effects, enhancing plant height, leaf area and biomass (in both roots and shoots), and promoting plant quality. According to Lee et al. (2012), Paenibacillus polymyxa is a group of PGPR, very similar to other rhizobacteria, that act as beneficial microbes improving the microbiome of plant roots. Many mechanisms are involved in efficient rhizobacteria-plant interaction. PGPR can also induce the production of enzymes, better absorption of water, and efficient use of mineral elements provided by fertilization (Egamberdiyeva, 2007). Thus, Timmusk et al. (1999) showed that P. polymyxa could metabolize cytokinins. In this way, plants and plant-associated micro-organisms have been found to contain over 30 growthpromoting compounds of the cytokinin group (Nicander et



traspiración y (C) uso eficiente de agua en plantas de Capsicum chinense. Los datos son promedios. Diferentes letras en la misma columna indican diferencias estadísticas significativas (Tukey  $\alpha = 0.05$ ).

al., 1993; Timmusk et al., 1999). In the current study, it was evident that the BSP1.1 (P. polymyxa) strain improved the quality traits of Capsicum chinense in comparison to untreated controls. In this way, the taller plants with more root biomass (BSP1.1 treatment) likely produced phytohormones that are considered to enhance root growth and surface area (i.e., bigger roots and more lateral roots and root hairs), leading to an increased plant nutrition (Richardson et al., 2009). On this last point, the BSP1.1 strain improved nutrient uptake (P and K). In this way, according to Lal & Tabacchioni (2009), P. polymyxa (formerly Bacillus polymyxa) is a non-pathogenic and endospore-forming bacillus with a wide range of properties, including nutrient fixation, plant growth promotion and soil phosphorus solubilization (Shi et al., 2009). Supporting our results, Guemouri-Athmani et al. (2000) demonstrated the nitrogen fixing ability of *P. polymyxa* (in 14 out of the 23 strains tested). However, it has not been demonstrated that plant growth promotion by *P. polymyxa* is primarily correlated with its nitrogen-fixing ability (Lal & Tabacchioni, 2009). The BSP1.1 strain tested in the current study probably had nitrogen fixing ability. In addition, the BSP1.1 strain increased the phosphorus-fixing ability. The use of phosphate-solubilizing bacteria as inoculants increases P uptake by plants (Chen et al., 2006), and P. polymyxa has been reported as a P solubilizer (De Freitas et al., 1997). It is generally accepted that the predominant mechanism of mineral phosphate solubilization is the action of organic acids synthesized by soil microorganisms (Rodríguez & Fraga, 1999). On the other hand, P. polymyxa is not reported to be a potassium-solubilizing bacteria. However, Han & Lee (2006) showed that inoculation with P. polymyxa increased the availability of P and K in the soil, the uptake of N, P and K by the roots, and the growth of peppers and cucumbers. According to Schachtman & Schroeder (1994), potassium is crucial for plant nutrition and growth. In the current study, the larger plants, suggests that plant-PGPR interaction improves P and K uptake in Capsicum chinense plants.

Based on the gas exchange results, inoculation with PGPR had a positive effect on the photosynthetic rate, leading to increased growth of Capsicum chinense plants. In a previous work with beans, inoculation with different strains of Rhizobium and Pseudomonas improved the seedling physiology (Ahmad et al., 2013). Similarly, previous studies have shown that inoculation with PGPR increases 1-aminocyclopropane-1-carboxylate (ACC) deaminase (Nadeem et al., 2009), which might have helped a greater soil water uptake by plants (Hamdia et al., 1998). Moreover, the improvement may be due to the higher dry mass of the roots. Li & Xu (2014) showed that in plants inoculated with PGPR, indole-3-acetic acid (IAA) increases. In the current study, it was observed that water use efficiency (WUE) increased due to inoculation with BSP1.1. This result indicates that the BSP1.1 strain promoted a better investment in water consumption by plants. In this way, Timmusk & Wag-

ner (1999) showed with a gnotobiotic system (with quantification of mRNA levels) that P. polymyxa isolates confer resistance to drought stress in Arabidopsis thaliana. Thus, a decrease in transpiration rate and an increase in photosynthesis (CO<sub>2</sub> assimilation rate) enhanced growth parameters (fresh mass, dry mass, leaf area and height), and subsequently improved water use efficiency of Capsicum chinense plants inoculated with P. polymyxa. The results obtained in the current study demonstrated that growth, nutrient uptake and photosynthesis activity were stimulated in Capsicum chinense plants after seeds were inoculated with PGPR during sowing. However, the effects were dependent on the used strain. Thus, of the four strains used in the current study (P61: Pseudomonas tolaasii; A46: Pseudomonas tolaasii; R44: Bacillus pumilus; BSP1.1: Paenibacillus polymyxa), the strain of *P. polymyxa* (BSP1.1) was better than the rest. Inoculation with P. polymyxa particularly improved the photosynthetic activity and water use efficiency of C. chinense plants.

These results suggest that inoculation with the BSP1.1 (*P. polymyxa*) strain could reduce the use of fertilizers and water. In addition, this research prompted the development of further experiments in the field to test productivity and fruit quality traits in seedlings inoculated with *P. polymyxa*.

#### ACKNOWLEDGEMENTS

This work was made with the support of the Plant Biochemistry and Molecular Biology Unit, of the Yucatan Scientific Research Center (CICY), Mexico. We thank the area of soil microbiology of the Colegio de Postgraduados at Campus Montecillo, who provided the rhizobacteria strains tested in this study.

#### REFERENCES

- Ahemad, M. & M. Kibret (2014). Mechanisms and applications of plant growth promoting rhizobacteria: Current perspective. *Jour*nal of King Saud University–Science. 26: 1-20.
- Ahmad, M., Z.A. Zahir, M. Khalid, F. Nazli & M. Arshad (2013). Efficacy of *Rhizobium* and *Pseudomonas* strains to improve physiology, ionic balance and quality of mung bean under salt-affected conditions on farmer's fields. *Plant Physiology and Biochemistry* 63:170-176.
- Bashan, Y., L.E. De-Bashan, S.R. Prabhu & J.P. Hernández (2014). Advances in plant growth-promoting bacterial inoculant technology: formulations and practical perspectives (1998–2013). *Plant and Soil* 378: 1-33.
- Bosland, P.W. & E.J. Votava (2000). Peppers: vegetable and spice capsicums. CABI publishing. Las Cruces, USA. 204 p.
- Bhandari, S.A. & D. Amarasiriwardena (2000). Closed-vessel microwave acid digestion of commercial maple syrup for the determination of lead and seven other trace elements by inductively coupled plasma-mass spectrometry. *Microchemical Journal* 64: 73-84.
- Caballero-Mellado, J. (2006). Microbiología agrícola e interacciones microbianas con plantas. *Revista Latinoamericana de Microbiología*. 48: 154-161.

- Canto-Flick, A., E. Balam-Uc, J.J. Bello-Bello, C. Lecona-Guzmán, V. Solíz-Marroquín, S. Aviléz-Viñas, E. Gómez-Uc, G. López-Puc & N. Santana-Buzzy (2008). Capsaicinoids content in Habanero pepper (*Capsicum chinense* Jacq). Hot-test cultivars. *Hort-science* 45: 1344-1349.
- Canto-Martín, J.C., S. Medina-Peralta & D. Morales-Avelino (2004). Efecto de la inoculación con Azospirrillum sp. en plantas de Chile habanero (*Capsicum chinense* Jacq). Tropical and Subtropical Agroecosystems 4: 21-27.
- Chen, Y.P., P.D. Rekha, A.B. Arun, F.T. Shen, W.A. Lai & C.C. Young (2006). Phosphate solubilizing bacteria from subtropical soil and their tricalcium phosphate solubilizing abilities. *Applied Soil Ecology* 34: 33-41.
- De Freitas, J.R., M.R. Banerjee, & J.J. Germida (1997). Phosphatesolubilizing rhizobacteria enhance the growth and yield but not phosphorus uptake of canola (*Brassica napus* L.). *Biology and Fertility of Soils* 24: 358-364.
- Egamberdiyeva, D. (2007). The effect of plant growth promoting bacteria on growth and nutrient uptake of maize in two different soils. *Applied Soil Ecology* 36: 184-189.
- Fernández-Herrera, E., M. Acosta-Ramos, F. Ponce-Gageóla, & V. Manuel-Pinto (2007). Manejo Biológico de Phytophthora capsici Leo, Fusarium oxysporum Schlechtend.:Fr y Rhizoctonia solani Kuhn en jitomate (Lycopersicon esculentum Mill). Revista Mexicana de Fitopatología. 25: 35-42.
- Garruña-Hernández, R., R. Orellana., A. Larque-Saavedra & A. Canto (2014a). Understanding the physiological responses of a tropical crop (*Capsicum chinense* Jacq.) at high temperature. *PLoS One 9: e111402*.
- Garruña-Hernández, R., L. Latournerie-Moreno, O. Ayala-Garay, J.M. Santamaría, & L. Pinzón-López (2014b). Acondicionamiento pre-siembra: una opción para incrementar la germinación de semillas de chile Habanero (*Capsicum chinense* Jacq.). Agrociencia 48: 413-423.
- Guemouri-Athmani, S., O. Berge, M. Bourrain, P. Mavingui, J.M. Thiery, T. Bhatnagar & T. Heulin (2000). Diversity of *Paenibacillus polymyxa* in the rhizosphere of wheat (*Triticum durum*) in Algerian soils. *European Journal of Soil Biology* 36: 149-159.
- Han, H.S. & K.D. Lee (2006). Effect of co-inoculation with phosphate and potassium solubilizing bacteria on mineral uptake and growth of pepper and cucumber. *Plant soil and Environment* 52: 130-136.
- Hamdia, M.A. & H.M. El-Komy (1998). Effect of salinity, gibberellic acid and *Azospirillum* inoculation on growth and nitrogen uptake of *Zea mays. Biologia Plantarum* 40: 109-120.
- Khalid, A., S. Tahir, M. Arshad & Z.A. Zahir (2004). Relative efficiency of rhizobacteria for auxin biosynthesis in rhizosphere vs. non-rhizosphere soil. *Australian Journal of Soil Research* 42: 921-926.
- Lal, S. & S. Tabacchioni (2009). Ecology and biotechnological potential of *Paenibacillus polymyxa*: a minireview. *Indian Journal of Microbiology*. 49: 2-10.
- Lee, B., M.A. Farag, H.B. Park, J.W. Kloepper, S.H. Lee & C.M. Ryu (2012). Induced resistance by a long chain bacterial volatile: elicitation of plant systemic defense by a C<sup>13</sup> volatile produced by *Paenibacillus polymyxa*. *PLoS ONE* 7: e48744.
- Li, X. & K. Xu (2014). Effects of exogenous hormones on leaf photosynthesis of *Panax ginseng*. *Photosynthetica*. 52: 152-156.

- Marschner, P. (2007). Plant-microbe interactions in the rhizosphere and nutrient cycling in terrestrial ecosystems. In: Marschner, P. & Rengel, Z. (eds), pp. 82-159. Nutrient cycling in terrestrial ecosystems. Springer. Berlin, Germany.
- Nadeem, S.M., Z.A. Zahir, M. Naveed & M. Arshad (2009). Rhizobacteria containing ACC-deaminase confer salt tolerance in maize grown on salt-affected fields. *Canadian Journal of Microbiology* 55: 1302-1309.
- Nicander, B., U. Stahl, P.O. Bjorkman & E. Tillberg (1993). Immunoaffinity co-purification of cytokinins and analysis by highperformance liquid chromatography with ultraviolet-spectrum detection. *Planta* 189: 312-320.
- Reyes-Ramírez, A., M. López-Arcos, E. Ruíz-Sánchez, L. Latournerie-Moreno, A. Pérez-Gutiérrez, M.G. Lozano-Contreras & M.J. Zavala-León (2014). Efectividad de inoculantes microbianos en el crecimiento y productividad de chile Habanero (*Capsicum chinense* Jacq). *Agrociencia* 48: 285-294.
- Richardson, A.E., J.M. Barea, A.M. Mcneill & C. Prigent-Combaret (2009). Acquisition of phosphorus and nitrogen in the rhizosphere and plant growth promotion by microorganisms. *Plant Soil* 321: 305-339.
- Rodríguez, H. & R. Fraga (1999). Phosphate solubilizing bacteria and their role in plant growth promotion. *Biotechnology Advances* 17: 319-339.
- Schachtman, D. P. & J.I. Schroeder (1994). Structure and transport mechanisms of a high-affinity potassium uptake transporter from higher plants. *Nature* 370: 655-658.
- Shi, Y.W., K. Lou, C. Li, L. Yang, X.Q. Wang & W.Y. Liu (2009). Effects of endophytic *Paenibacillus polymyxa* S-7 on photosynthesis, yield, and quality of sugar beet. *Chinese Journal of Applied Ecology* 20: 597-602.
- Timmusk, S. & G.H. Wagner (1999). The plant-growth-promoting rhizobacterium *Paenibacillus polymyxa* induces changes in *Arabidopsis thaliana* gene expression: a possible connection between biotic and abiotic stress responses. *Molecular Plant-Microbe Interactions* 12: 951-959.
- Timmusk, S., B. Nicander, U. Granhall & E. Tillberg (1999). Cytokinin production by *Paenibacillus polymyxa*. Soil Biology and Biochemistry 31: 1847-1852.
- Vessey, K.J. (2003). Plant growth promoting rhizobacteria as biofertilizers. *Plant and Soil* 255: 571-586.