PHYSIOLOGICAL RESPONSES OF DROUGHT STRESSED WHEAT PLANTS (*Triticum aestivum* L.) TREATING WITH SOME BACTERIAL ENDOPHYTES

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ABSTRACT

The main objective of this study is: Effect of two treatment of bacterial endophytes strains *Azotobacter chrocooccum* (E1) and *Pseudomonas* sp. (E2) individually whether as grains soaking and foliar application on some physiological parameters of two wheat plants (*Triticum aestivum* L.) cultivars (Sakha 93 and Gmiza 9) grown under three levels of irrigation water deficit stress 75, 50 and 25 % field capacity. The tested physiological parameters were chlorophyll pigment (chl. a, b and total) contents, relative water content (RWC), leaf water content (LWC), leaf water deficit (LWD), proline content and some major essential elements (NPK) contents. Negative impacts were obtained on the tested wheat cultivars grown under the different irrigation water deficit. Application of *Azotobacter chrocooccum* (E1) and *Pseudomonas* sp. (E2) strains individually were carried out by spray foliar and grains soaking treatments increased the tested physiological parameters for two cultivars compared with untreated plant, which could overcome the negative effects of drought stress.

Keyword: Chlorophyll, proline, endophytic bacteria strains, mineral uptake, leaf water deficit, mineral uptake and irrigation water deficit.

INTRODUCTION

Drought is one of the major environmental conditions that adversely affect plant growth, physiological, biochemical changes, including changes of the endogenous phytohormone levels and crop yield (Boyer, 1982).

Due to drought and competing water demands in Egypt have put enormous pressure on irrigation water. Conserve both the quality and quantity of water appropriate strategies will have to be developed to avoid the risk of future water supplies: Reducing irrigation water is to employ practices that improve water productivity (crop yield per unit volume of water used). Among different strategies to cope with drought issues seed priming (presowing seed treatment) is an easy, low cost and low risk technique and this approach has recently been used to overcome the drought problem in agriculture land (Iqbal and Ashraf, 2006). Although priming induced-drought tolerance has been reported in some crops, knowledge about physiological, biochemical and anatomical basis of priming induced-beneficial effects under stressful environment is still in frequent.

Plant growth-promoting bacteria include both free living and symbiotic bacteria, typically found in the soil, that facilitate the growth and development of plants (Glick *et al.*, 1999). This can occur directly to promote plant growth either by providing the plant with a compound that is synthesized

by the bacterium or by facilitating the uptake of nutrients from the soil. Thus, plant growth-promoting bacteria can directly facilitate the proliferation of plants by fixing atmospheric nitrogen; producing siderophores which can mineral solubilize and provide it to plants; synthesizing phytohormones, such as auxin, cytokinin and gibberellin, which can enhance various stages of plant growth; solubilizing minerals such as phosphorus; and synthesizing enzymes that can modulate plant growth and development (Glick, 2007).

Microbial inoculants that can promote plant growth and productivity is internationally accepted as an alternative source of N-fertilizer. It is environmental friendly and can be used to ensure a sustainable wheat production. In this bio-fertilizer technology new systems are being developed to increase the biological N₂ fixation (BNF) with cereals and other nonlegumes by establishing N₂-fixing bacteria within the roots (Cocking, 2000). Nitrogen fixation and plant growth promotion by plant growth promoting bacteria are important criteria for an effective bio-fertilizer. Inoculation of associative and free living N2-fixing bacteria have been shown to produce beneficial effects on plant growth, thus they are termed plant growth promoting rhizobacteria (PGPR) (Kloepper et al., 1980; Bashan and Holguin, 1998). Significant increases in crop yields following application of PGPR have been documented under diverse field conditions (Bashan, 1998). They have been widely reported to fix atmospheric nitrogen with grasses and cereals (Dobereiner, 1997) and enhance nutrient uptake (Lin et al., 1983; Murty and Ladha, 1988; Bashan and Holguin, 1998).

Azotobacter sp besides fixing nitrogen it is also secrete certain growth hormones such as IAA, GA and Cytokinins (Coppola, 1971) which promote vegetative growth and root development. Generally, the literature review indicates that there are possibly some positive effects of endophytic bacteria treatment on growth and reproduction of plants.

Wheat production is an essential national target to fill the gap between production and consumption. Production could be increased through cultivation of high yielding cultivars and appropriate agronomic practices (Tawfik *et al.*, 2006). It is obvious that, there found an enormous pressure on irrigation water in Egypt due to drought and competing water demands. Improvement of wheat production under irrigation water deficit (drought) has become important during recent years worldwide. Therefore, the objective of this study was to evaluate two different cultivars of wheat physiology in response to inoculation with *Azotobacter chrocooccum* (E1) and *Pseudomonas* sp. (E2) as endophytic bacterial strains under different levels of field capacity.

MATERIALS AND METHODS

The present investigation was conducted under greenhouse conditions at the Department of Agricultural Botany, Faculty of Agriculture, Kafrelsheikh University, during the two growing winter successive seasons of 2009 / 2010 and 2010 / 2011 to study the impact of certain ecophysiological and microbial studies on the effect of drought stress on some physiological features of wheat plants (*Triticum aestivum* L.).

Source of wheat cultivars:

Wheat grains (*Triticum aestivum* L.) cultivars Sakha 93 and Gmiza 9 were obtained from Wheat Research Dept. Sakha of Agricultural Research Station, Kafr El-sheaikh, Egypt.

Source of microorganisms:

Two bacterial strains [Azotobacter chrocooccum (E1) and Pseudomonas sp. (E2)] were obtained from Dr. Elsayed Belal, Associate Professor of Agricultural Microbiology, Dep. of Agric. Botany, Fac. of Agriculture, Kafrelsheikh University, where these bacterial strains were isolated in a previous study as entophytic bacteria from wheat plants (unpublished data). Cultivation of microorganisms:

Azotobacter chrocooccum (E1) and Pseudomonas sp. (E2) were cultivated in nutrient liquid medium. 200 ml nutrient liquid medium were inoculated with 2 ml of a cell suspension of (Azotobacter chrocooccum (E1) or Pseudomonas sp. (E2) (nutrient broth medium, 10⁸ cfu / ml) was incubated at 30 °C and 150 rpm for 3 days. The cultures were incubated at 30 °C and 150 rpm for 5 days. Thereafter, two bacterial strains were applied on wheat as follow:

Grain treatments

Two bacterial strains were applied at the time of sowing as seed treatment. Grains were immersed in each bacterial suspension (10⁸ cfu / ml) for 30 min.. Grains were witted with 10 % sugar syrup, and thoroughly mixed with an amount of bacterial suspension (10⁸ cfu / ml) for 30 min. enough to obtain 10⁸ cfu / per gram of grains and then air dried. Grains were then sown in each pot (10 grains / pot). On the other hand, grains wheat were immersed in a manner in 10 % sugar syrup and were thoroughly mixed with an amount of nutrient broth medium (without bacterial growth).

Wheat plant spraying

Wheat plants (20 days from sowing) were sprayed weekly intervals with bacterial suspension (10⁸ cfu / ml) from each bacterial strains.

Pots, soil preparation and wheat grains sowing:

Each pot (30 cm in diameter) contained 8 Kg of air dried clay soil. The chemical analysis was determined by conventional methods, twelve grains / pot were sown at equal distances and depth. After two weeks from sowing, the seedlings were thinned to ten seedlings / pot, three of them were kept for the morphological characters throughout the experimental period.

The soil used in this experiment was fertilized with nitrogen at rate a 360 kg/h of urea fertilizer (46% nitrogen). Super phosphate fertilizer (phosphorus 15%) was added at a rate of 240 kg/ha before sowing. Potassium was not added because the Egyptian soil is rich in this element. Chemical analysis of the soil samples were done before sowing in the two seasons, mechanical and chemical analysis of the experimental soil were determined (Metwaly, 2012) according to Page, (1982) and Klute, (1986).

Chlorophyll pigments measurements:

Chlorophyll content of the two wheat cultivars was determined after 40, 70 and 130 days from sowing. Chlorophyll A, B and total chlorophyll were determined in the flag leaf lamina using the spectrophotometer method described by Moran and Porath (1980).

Proline determination:

Extraction and determination of proline were performed in the flag leaf according to the method of Bates *et al.*, (1973).

Mineral nutrients (NPK) uptake:

Mineral elements [Nitrogen (N), Phosphorus (P) and Potassium (K)] were performed on the material which was dried in an electric oven at 70 $^{\circ}$ C to constant weight and following determinations were done:

- A- Total nitrogen was estimated in the digestion product, using the official Micro-Kjeldahl Method. The percentage of total nitrogen was estimated and crude protein content was calculated by the following equation:
 - Crude protein % = Total nitrogen x 5.83.
- B- Phosphorus was estimated by ascorbic acid method using the calorimetric method as described by Murphy and Riley (1962).
- C- Potassium was also estimated in the above mentioned digestion product by using flame photometer according to Jackson (1967).

Plant-water relations:

Relative water content (RWC) was measured according to Schonfeld *et al.*, (1988). Leaf water deficit (LWD) was determined and calculated using a formula according to Kalapos (1994). Leaf water content (LWC) was expressed according to Liu *et al.*, (2004).

Experimental design and statistical analysis:

The pots were arranged in a randomized complete block design with three replicates in every treatment and ten plants in each pot. Data of the physiological studies were tested by analysis of variance. Duncan's multiple range tests were used for comparisons among treatment mean (Duncan, 1955).

RESULTS AND DISCUSSION

Physiological studies:

Drought impacts include physio-biochemical responses growth, yield, membrane integrity, pigment content, osmotic adjustment water relations, and photosynthetic activity (Benjamin and Nielsen, 2006; Praba $et\ al.$, 2009). Physiological characteristics of wheat cultivars grown under different irrigation water deficit levels (75, 50 and 25 % FC) were studied. Application of two endophytes including $A.\ chrocooccum$ (E1) and $Pseudomonas\ sp.$ (E2) and the combination between them and irrigation water deficit levels were investigated. These parameters includes chlorophyll pigments (A, B and total $\mu g\ / cm^2$), relative water content, leaf water content, mineral uptake (N, P and K %), crude protein and proline content ($\mu moles\ /\ g\ dry\ weight$) were investigated.

Chlorophyll pigments:

Data presented in Tables (1, 2 and 3) cleared that, there was a negative impact on chlorophyll pigments (chl. a, b and total) content (μ g / cm²) by application of different irrigation water deficit levels (75, 50 and 25 % FC).

Chlorophyll is one of the major chloroplast components for photosynthesis, and relative chlorophyll content has a positive relationship with photosynthetic rate. The decrease in chlorophyll content under drought stress has been considered a typical symptom of oxidative stress and may be the result of pigment photo-oxidation and chlorophyll degradation. Photosynthetic pigments are important to plants mainly for harvesting light and production of reducing powers. Both the chlorophyll a and b are prone to soil dehydration (Farooq et al., 2009). Decreased or unchanged chlorophyll level during drought stress has been reported in many species, depending on the duration and severity of drought (Kyparissis et al., 1995; Zhang and Kirkham, 1996).

Environmental stresses have a direct impact on the photosynthetic apparatus, essentially by disrupting all major components of photosynthesis including the thylakoid electron transport, the carbon reduction cycle and the stomatal control of the CO2 supply, together with an increased accumulation of carbohydrates, peroxidative destruction of lipids and disturbance of water balance (Allen and Ort, 2001).

The decrease in chlorophyll under drought stress is mainly the result of damage to chloroplasts caused by active oxygen species, and related to the reduction in stomatal conductance and transpiration rate (Chartzoulakis, et al. 1999). Prolonged water stress which limited photosynthesis also led to less of sucrose phosphate synthetase activity (Vassey and Sharky, 1989 and Dubey and Singh, 1999). Furthermore, water deficit induced reduction in chlorophyll content has been ascribed to loss of chloroplast membranes, excessive swelling, distortion of the lamellae vesiculation, and the appearance of lipid droplets (Kaiser et al., 1981). Low concentrations of photosynthetic pigments can directly limit photosynthetic potential and hence primary production (Anjum, et al., 2011).

The wheat cultivars treated with the two bacterial endophytes strains (A. chrocooccum (E1) and Pseudomonas sp. (E2)) had increased significantly chlorophyll pigment content (µg / cm²) after 40 and 70 days from sowing. These results were compared with control (100 % FC) without endophytes treatment. The increasing percentage of chl. a (µg / cm²) for Gmiza 9 cultivar by application of bacterial endophytes was 13.10 % under normal irrigation water (100 % FC). On the other hand, for Sakha 93 cultivar the percentage of increasing was 15.85 % compared with control treatment. Application of A. chrocooccum (E1) and Pseudomonas sp. (E2) in combination with the different tested irrigation water deficit levels, increased significantly in chlorophyll content (µg / cm²) Gmiza 9 after 40 and 70 days from sowing. On the other hand, the results for Gmiza 9 cultivar were insignificant due to application of Pseudomonas sp. (E2) as a foliar spraying in combination with the water deficit stress percentage after 40 days from sowing during the both seasons. Auxin is recognized as a key factor, which is directly beneficial of plant. The role of microorganisms as plant growth stimulators is widespread in nature, especially in relation to a group of plant hormones that are implicated in the regulation of diverse biological processes including cell division, elongation, differentiation, root elongation and tropistic responses (Spaepen, et al., 2007). Moreover, Cytokinin stimulates the

synthesis of chlorophyll pigments (Jelić and Bogdanović, 1989). In addition, increasing in wheat cultivars treated with endophytic strains was due to the enhancement of essential elements uptake especially nitrogen element (N_2). Nitrogen is necessary for chlorophyll synthesis and as part of chlorophyll molecules, is the focal point of photosynthesis. Also, N_2 is an essential component of amino acids, which building blocks of protein.

Proline content (µmolg⁻¹):

Proline content of the leaves was significantly affected by drought stress and increased by declining the water availability in the soil. Data presented in Table (4) cleared that, all used irrigation water deficit treatments increased proline content (µmol/g) in the tested wheat cultivars leaves compared to well-irrigated plants (control 100 % FC) during the both growing seasons. Moreover, generally the increasing proline content (µmol / g) for Gmiza 9 cultivar by application of bacterial endophytes was about 25.43, 15.13 and 29.43 % under irrigation water deficit level (75, 50 and 25 % FC) respectively. On the other hand, for Sakha 93 cultivar the percentage was about 23.89, 18.02 and 53.42 % under irrigation water deficit levels (75, 50 and 25 % FC) respectively.

Plants accumulate different types of organic and inorganic solutes in the cytosol to lower osmotic potential thereby maintaining cell turgor (Rhodes and Samaras, 1994). Under drought, the maintenance of leaf turgor may also be achieved by the way of osmotic adjustment in response to the accumulation of proline, sucrose, soluble carbohydrates, glycinebetaine, and other solutes in cytoplasm improving water uptake from drying soil. The process of accumulation of such solutes under drought stress is known as osmotic adjustment which strongly depends on the rate of plant water stress. Wheat is marked by low level of these compatible solutes and the accumulation and mobilization of proline was observed to enhance tolerance to water stress (Nayyar and Walia, 2003). Of these solutes, proline is the most widely studied because of its considerable importance in the stress tolerance. Proline accumulation is the first response of plants exposed to water-deficit stress in order to reduce injury to cells. Progressive drought stress induced a considerable accumulation of proline in water stressed maize plants. The proline content increased as the drought stress progressed and reached a peak as recorded after 10 days stress, and then decreased under severe water stress as observed after 15 days of stress (Anjum et al., 2011b).

Moreover, proline promotes the production of cytokinins which improved plant growth (Shetty, et al., 1992). Proline can protect plant cells from oxidative damage by scavenging reactive oxygen species (Shao, et al., 2008). It was found in wheat that Ca appeared to reduce the devastating effects of stress by elevating the content of proline, thus improving the water status and growth of seedlings and minimizing the injury to membranes (Nayyar and Walie 2003).

Water relations:

Concerning wheat cultivars relative water content (RWC), leaf water content (LWC) and leaf water deficit (LWD), data presented in Tables $(4,\,5)$ showed that, the treatments of irrigation water deficit in the present study

decreased (RWC), leaf water content and increased leaf water deficit during the both growing season. These results were compared with control (100 % FC) treatment.

Relative water content (RWC), leaf water potential, stomatal resistance, rate of transpiration, leaf temperature and canopy temperature are important characteristics that influence plant water relations. A decrease in the relative water content (RWC) and leaf water content (LWC) in response to drought stress has been noted in wide variety of plants as reported by Nayyar and Gupta (2006). When leaves are subjected to drought, leaves exhibit large reductions in RWC and water potential. Exposure of plants to drought stress substantially decreased the leaf water potential, relative water content and transpiration rate, with a concomitant increase in leaf temperature (Siddique et al., 2001). RWC related to water uptake by the roots as well as water loss by transpiration. It is well known that leaf water status always interacts with stomatal conductance and a good correlation between leaf water potential and stomatal conductance always exists, even under drought stress. It is now clear that there is a drought-induced root-to-leaf signaling, which is promoted by soil drying through the transpiration stream, resulting in stomatal closure.

In this respect application of two bacterial endophytes *A. chrocooccum* (E1) and *Pseudomonas* sp. (E2) increased significantly wheat cultivars RWC, LWC and reduced LWD compared with untreated plants during the both seasons. The increasing percentage of RWC for Gmiza 9 cultivar by application of bacterial endophytes was about 7.16 % under normal irrigation water (100 % FC). On the other hand, for Sakha 93 cultivar the percentage was about 2.61 % compared with control treatment.

The LWD was increased by increasing levels of irrigation water deficit stress up to (25 % FC).

In this respect, application of two endophytic bacteria in combination with all irrigation water deficit levels (75, 50 and 25 % FC) increased RWC and LWC and reduced LWD during the both seasons. The obtained results were compared with each of irrigation water deficit level as alone without endophytic bacteria *A. chrocooccum* (E1) and *Pseudomonas* sp. (E2) treatments. The increasing percentage of RWC for Gmiza 9 cultivar by application of bacterial endophytes was about 14.70, 23.94 and 49.38 % under irrigation water deficit level (75, 50 and 25 % FC) respectively. On the other hand, for Sakha 93 cultivar the percentage was about 13.57, 25.79 and 54.44 % under irrigation water deficit levels (75, 50 and 25 % FC) respectively. Generally the grains soaking application of two bacterial endophytes was gave the best results, where decreased the leaf water deficit compared with other treatments.

Mineral uptakes:

It is clear from Table (6) and Fig. (1, 2, 3 and 4) that, the irrigation water deficit stress treatments (75, 50 and 25 % FC) decreased wheat cultivars uptake of nitrogen (N_2 %), phosphorus (P%) and potassium (K %) during the both seasons. These obtained results were compared to well-irrigated plants (control 100 % FC).

Generally, drought reduces both nutrient uptake by the roots and transport from the roots to the shoots, because of restricted transpiration rates and impaired active transport and membrane permeability (Viets, 1972; Alam, 1999). The decline in soil moisture also results in a decrease in the diffusion rate of nutrients in the soil to the absorbing root surface (Pinkerton and Simpson, 1986; Alam, 1999).

Nitrogen is the mineral element that plants require in the largest amounts and is a constituent of many plant cell components, including amino and nucleic acids and chlorophyll pigments. Therefore, nitrogen deficiency rapidly inhibits plant growth. Phosphorus is a constituent of nucleic acids, phospholipids, phosphoproteins, dinucleotides, and adenosine triphosphate. Hence, P is required for processes including the storage and transfer of energy, photosynthesis, the regulation of some enzymes, and the transport of carbohydrates.

Potassium is an essential factor in protein synthesis, glycolytic enzymes, and photosynthesis; an osmoticum mediating cell expansion and turgor-driven movements; and a competitor of Na⁺ under saline conditions (Marschner, 1995). Because both drought and salinity affect plant growth similarly through water deficit, K⁺ is equally important for maintaining the turgor pressure in plants under either stress. Generally, mineral fertilization not increase growth and yield parameters without sufficient water being available to the plant, and increasing soil-water availability will not increase production without adequate mineral supply. Reducing in mineral uptake under irrigation water deficit may also attributed to a decreased transpiration rate to transport mineral nutrients from root to shoot.

Grains soaking and foliar application of two different endophytic bacteria strains individually under well irrigation (100 % FC) or in combination with all irrigation water deficit stress levels (75, 50 and 25 % FC) have a positively impact on the tested mineral elements during the both growing seasons. The obtained results were compared with well-irrigated plants (control 100 % FC) treatment without endophytes treatment or each level of irrigation water deficit as alone without endophytes in the combination treatments.

However, plants treated with hormones significantly affected active absorption area, percentage active absorption area ratio, root volume and specific surface area. In general, there was no relationship between the first cropping without hormones application and second cropping with hormones application except percentage active absorption area ratio (-0.999*) which was negatively correlated.

On the other hand; cytokinins reduced the elongation of roots and the formation of lateral root (Lopez-Bucio *et al.*, 2003; Lohar *et al.*, 2004). Furthermore, Goodwin and Moris (1979) reported that cytokinins produced at the root tip of pea inhibit the lateral root formation, but support lateral stem growth. Zahir *et al.*, (2001) found that exogenous application of cytokinin at the root zone supported luxuriant growth and yield of rice. There are scanty literatures on auxin and potassium interaction, notable work was carried out by Shin *et al.* (2007).

Concerning crude protein, data in Table (6) indicate that, there is a negatively impact on crude protein content by all used irrigation water deficit treatments under the present study during the both successive seasons, where decreased as compared with non treated plants. These results are related to the increasing in N2 uptake of the tested wheat cultivars. In this respect the application of two bacterial endophytes *A. chrocooccum* (E1) and *Pseudomonas* sp. (E2) have a positive impact on N %, where significantly increased in wheat cultivars during the both seasons. The obtained results were compared with control plants without endophytes treatments. The results are in agreement with (Dobbelaere *et al.*, 2003; Cakmakc, 2005a, 2005b).

The application of two endophytic bacteria in combination with the lowest irrigation water deficit increased the N % during the both seasons. On the other side, the increasing for Sakha 93 cultivar by application of *Pseudomonas* sp. (E2) as a foliar spraying in combination with the same irrigation water deficit treatment was insignificant during the second season.

Application of two bacterial endophytes strains in combination with all other irrigation water deficit increased significantly N % during the both seasons as compared with each level of water stress treatment as alone without endophytes treatments.

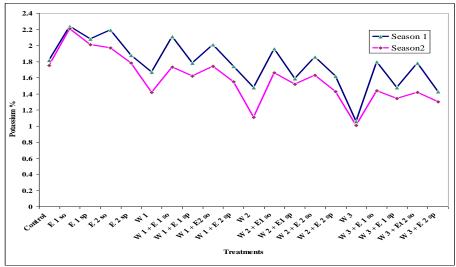


Figure 1: Potassium % of wheat cultivar Gmiza 9 as affected by different levels of irrigation water deficit and two different endophytes bacteria and their interactions during 2009 / 2010 and 2010 / 2011 seasons.

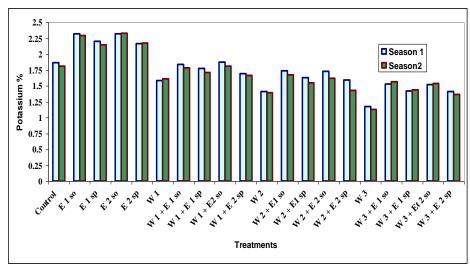


Figure 2: Potassium % of wheat cultivar Sakha 93 wheat plants as affected by different levels of irrigation water deficit and two different endophytes bacteria and their interactions during 2009 / 2010 and 2010 / 2011 seasons.

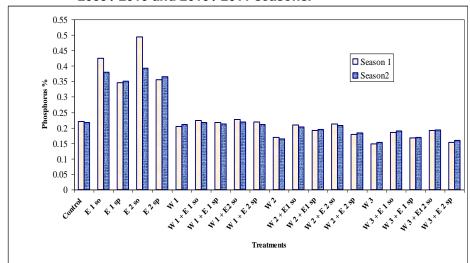


Figure 3: Phosphorus % of wheat cultivar Gmiza 9 as affected by different levels of irrigation water deficit and two different bacterial endophytes and their interactions during 2009 / 2010 and 2010 / 2011 seasons.

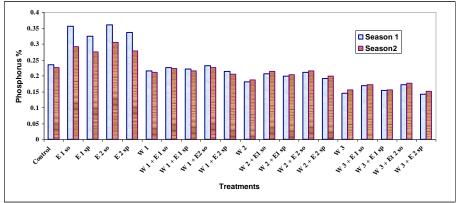


Figure 4: Phosphorus % of wheat cultivar Sakha 93 as affected by different levels of irrigation water deficit and two different bacterial endophytes and their interactions during 2009 / 2010 and 2010/2011 seasons.

Finally, the obtained results indicated that, the tested irrigation water deficit75, 50 and 25 % FC) decreased wheat cultivars uptake decreased relative water content, leaf water content and increased leaf water deficit. Moreover, chlorophyll pigments (chl. a, b and total) and mineral elements uptake (N2, P and K) were reduced. Application of endophytic bacteria strains [Azotobacter chrocooccum (E1) and Pseudomonas sp. (E2)] individually that carried out by foliar spray and grains soaking treatments increased the tested physiological parameters for the two cultivars during the both seasons. Endophuytic bacterial strains treatments play an important role in protection of wheat plants against the adverse effects of drought stress.

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الاستجابات الفسيولوجية لنباتات القمح تحت ظروف الإجهاد الجفافي باستخدام بعض البكتريا التي تنمو داخل النبات محمد مبروك العافري **، السيد بلال عبدالمنطلب بلال * و متولي محفوظ سالم متولى**
قسم النبات الزراعي- كلية الزراعة- جامعة كفرالشيخ – مصر **فرع النبات الزراعي - *فرع ميكروبيولوجي زراعية

الهدف من هذه الدراسة هو دراسة تأثير ثلاث مستويات من نقص ماء الري (75، 50 و 25% من السعة الحقلية) علي بعض الصفات الفسيولوجية لصنفين من نباتات القمح وهما سخا 93 وجميزة 9 ، اشتملت الصفات الفسيولوجية تحت الدراسة محتوي الأوراق من صبغات الكلوروفيل ومحتوي الماء النسبي ومحتوي المائي للأوراق وكذا نقص محتوي الاوراق المائي. كما اشتملت تلك الصفات علي امتصاص بعض العناصر الضرورية الكبري وهي النيتروجين والفسفور والبوتاسيوم. وأكدت النتائج المتحصل عليها ان للمستويات المختلفة من نقص ماء الري تأثير سيئا علي كل الصفات الفسيولوجية لصنفي القمح تحت الدراسة. كما أدي استخدام السلالاتين البكتيريه وهما Azotobacter chrocooccum المعاملات نقص ماء الري تحت الدراسة.

قام بتحكيم البحث

كلية الزراعة – جامعة المنصورة كلية الزراعة – جامعة المنوفية أ.د / عرفه احمد عرفه أ.د / احمد جندي اصلان Table 1: Chlorophyll a content (µg / cm²) in the flag leaf of wheat cultivars Gmiza 9 and Sakha 93 wheat plants as affected by different levels of irrigation water deficit and two bacterial endophytes and their interactions at different stages during 2009 / 2010 and 2010 / 2011 seasons.

uiii	erent St	ayes u	Gmiza 9 Sakha 93 Gmiza 9 Sakha											
				za 9			Sakha 93							
	2009/20010		2010/2011 season				2009/20010			2010/201	1 season	mean of	93	
Treetment	sea	son		Chloro	phyll a		season			Chlorophyll a			means	mean of
Treatment	Chloro	phyll a	Means	aft	er	Means	Chloroph	yll a after	Means	after		Means		means
	after							•						
	40 days	70 days		40 days	70 days		40 days	70 days		40 days	70 days			
Control	35.09 ^E	38.81 ^{to}	36.7	32.44 ^C	35.49 ^{CD}	33.97	42.43 ^{CD}	24.79 ^c	33.61	40.01 ^B	19.99 ^C	30.00	35.34	31.81
E 1 Gs	40.42 ^{AB}	41.24 ^{AB}	40.83	41.24 ^A	40.03 ^{AB}	40.64	49.73 ^A	33.35 ^A	41.54	44.42 ^A	24.24 ^{AB}	34.33	40.74	37.94
E 1 Sp	38.66 ^{BC}	41.83 ^A	40.25	39.79 ^{AB}	38.98 ^B	39.39	46.03 ^B	32.33 ^A	39.18	43.79 ^A	23.24 ^B	33.51	39.82	36.35
E 2 Gs	41.58 ^A	41.34 ^{AB}	41.46	41.11 ^A	40.58 ^A	40.85	46.85 ^B	33.83 ^A	40.34	44.83 ^A	25.13 ^A	34.98	41.16	37.66
E 2 Sp	37.48 ^{CD}	39.94 ^{BC}	38.71	38.57 ^B	36.65 ^C	37.61	44.06 ^C	30.66 ^B	37.36	43.03 ^A	24.09 ^{AB}	33.56	38.16	35.46
0	01.10	00.01	00.71	00.07	00.00	01.01	11.00	00.00	07.00	10.00		00.00	00.10	00.10
W 1	31.17 ^{FG}	32.90 ^F	32.04	27.21 ^E	30.41 ^G	28.81	36.75 ^{GH}	18.40 ^{GHI}	27.58	35.51 ^C	16.21 ^{EFG}	25.86	30.43	26.72
W 1 + E 1 Gs	35.65 ^E	37.78 ^{DE}	36.72	32.77 ^C	35.16 ^{DE}	33.97	42.19 ^{CD}	23.76 ^{CD}	32.98	39.42 ^B	18.46 ^{CD}	28.94	35.35	30.96
W 1 + E 1 Sp	34.16 ^E	36.87 ^E	35.52	30.42 ^D	33.88 ^E	32.15	40.49 ^{DE}	22.72 ^{DE}	31.61	34.80 ^{CD}	17.27 ^{DEF}	26.04	33.84	28.83
W 1 + E2 Gs	34.49 ^E	37.20 ^{DE}	35.85	32.99 ^c	34.78 ^{DE}	33.89	42.31 ^{CD}	24.38 ^C	33.35	36.13 ^C	17.89 ^{DE}	27.01	34.87	30.18
W 1 + E 2 Sp	31.75 ^{FG}	34.11 ^F	32.93	27.46 ^E	31.92 ^F	29.69	38.99 ^{EF}	20.06 ^F	29.53	34.08 ^{CD}	17.59 ^{DE}	25.84	31.31	27.69
W 1 + L 2 3p	31.73	34.11	32.93	27.40		23.03	30.33	20.00	29.55			25.04	31.31	21.09
W 2	26.05 ^l	26.73 ^H	26.39	23.64 ^F	25.14 ^{JK}	24.39	33.46 ^l	16.02 ^K	24.74	31.58 ^{EF}	14.26 ^{HI}	22.92	25.39	23.83
W 2 + E1 Gs	31.97 ^F	32.55 ^F	32.26	29.33 ^D	29.05 ^H	29.19	38.62 ^{EFG}	19.89 ^{FG}	29.26	34.51 ^{CD}	16.54 ^{EF}	25.53	30.73	27.39
W 2 + E1 Sp	29.67 ^{GH}	27.96 ^H	28.82	26.80 ^E	27.60 ^l	27.20	37.27 ^{FGH}	21.79 ^b	29.53	33.18 ^{DE}	17.39 ^{DE}	25.29	28.01	27.41
W 2 + E 2 Gs	29.80 ^{GH}	30.86 ^G	30.33	29.89 ^D	28.75 ^{HI}	29.32	37.96 ^{FG}	22.42 ^{DE}	30.19	33.17 ^{DE}	16.78 ^{DEF}	24.98	29.83	27.59
W 2 + E 2 Sp	29.34 ^H	30.09 ^G	29.72	25.99 ^E	24.49 ^{KL}	25.24	35.50 ^H	19.11 ^{FG}	27.31	30.65 ^F	15.64 ^{FGH}	23.15	26.98	25.23
			_		_	-								
W 3	19.29 ^K	21.88 ^J	20.58	14.79 ^H	19.71 ^N	17.25	28.43 ^L	13.79 ^L	21.11	23.33 ^l	11.67 ^J	17.50	18.92	19.31
W 3 + E 1 Gs	24.97 ^{IJ}	27.27 ^H	26.12	23.95 ^F	26.19 ^J	25.07	31.61 ^J	17.87 ^{HIJ}	24.74	26.91 ^G	14.68 ^{GHI}	20.79	25.59	22.77
W 3 + E 1 Sp	23.15 ^J	26.53 ^H	24.84	23.55 ^F	23.58 ^{LM}	23.57	31.21 ^{JK}	16.91 ^{IJK}	24.06	25.39 ^H	13.85 ¹	19.62	24.21	21.84
W 3 + E 2 Gs	24.48 ^{IJ}	27.27 ^H	25.89	19.47 ^G	24.47 ^{KL}	21.97	31.12 ^{JK}	18.04 ^{HI}	24.58	26.43 ^G	14.49 ^{HI}	20.46	23.93	22.52
W 3 + E 2 Sp	20.49 ^K	24.97 ^l	22.73	16.00 ^H	22.46 ^M	19.23	29.51 ^{KL}	16.27 ^{JK}	22.89	23.92 ^{HI}	13.27 ^l	18.59	20.98	20.74

E1 = Azotobacter chrocooccum E2 = Pseudomonas sp Gs = Grains soaking Sp = foliar application control = 100 % field capacity W1 = 75% field capacity W2 = 50 % field capacity W3 = 25 % field capacity

Table 2: Chlorophyll b content (µg / Cm²) in the flag leaf of wheat cultivars Gmiza 9 and Sakha 93 wheat plants as affected by different levels of irrigation water deficit and two bacterial endophytes and their interactions at different stages during 2009 / 2010 and 2010 / 2011 seasons.

	uniere	ni siage		g 2009 /	2010 ai	14 2010	7 2011 3	seasons).					
			Gmiza 9					Sakha 93						
Treatment	2009/20010 season Chlorophyll b after		Means		0/2011 season orophyll b after Me		2009/20010 season Chlorophyll b after		Means	2010/2011 season Chlorophyll b after		Means	_	Sakha 93 mean of means
	40	70		40	70		40	70		40	70			
Control	17.72 ^E	17.56 ^{CD}	17.64	16.47 ^C	15.92 ^{CD}	16.19	19.29 ^{CD}	11.27 ^C	15.28	18.18 ^B	9.09 ^C	13.64	16.92	14.46
E 1 Gs E 1 Sp E 2 Gs E 2 Sp	20.41 ^{AB} 19.53 ^{BC} 21.00 ^A 18.93 ^{CD}	18.67 ^{AB} 18.93 ^A 18.71 ^{AB} 18.07 ^{BC}	19.54 19.23 19.86 18.50	20.93 ^A 20.19 ^{AB} 20.87 ^A 19.58 ^B	17.95 ^{AB} 17.48 ^B 18.19 ^A 16.43 ^C	19.44 18.84 19.53 18.01	22.60 ^A 20.92 ^B 21.29 ^B 20.03 ^C	15.16 ^A 14.69 ^A 15.38 ^A 13.93 ^B	18.88 17.81 18.34 16.98	20.18 ^A 19.91 ^A 20.38 ^A 19.56 ^A	11.02 ^{AB} 10.57 ^B 11.42 ^A 10.95 ^{AB}	15.60 15.24 15.90 15.26	19.49 19.04 19.69 18.26	17.24 16.53 17.12 16.12
W 1 W1+E1 Gs W1+E1 Sp W1+E2 Gs W1+E2 Sp	15.74 ^{FGH} 18.00 ^{DE} 17.25 ^E 17.42 ^E 16.03 ^{FG}	14.89 ^F 17.09 ^{DE} 16.68 ^E 16.83 ^{DE} 15.44 ^F	15.32 17.55 16.97 17.13 15.74	13.81 ^E 16.64 ^C 15.44 ^D 16.74 ^C 13.94 ^E	13.64 ^G 15.77 ^{DE} 15.19 ^E 15.59 ^{DE} 14.32 ^F	13.73 16.21 15.32 16.17 14.13	16.70 ^{GH} 19.18 ^{CD} 18.40 ^{DE} 19.23 ^{CD} 17.72 ^{EF}	8.36 ^{GHI} 10.79 ^{CD} 10.33 ^{DE} 11.08 ^C 9.12 ^F	12.53 14.99 14.37 15.16 13.42	16.14 ^C 17.92 ^B 15.82 ^{CD} 16.42 ^C 15.49 ^{CD}	7.37 ^{EFG} 8.39 ^{CD} 7.85 ^{DEF} 8.13 ^{DE} 7.99 ^{DE}	11.76 13.16 11.84 12.28 11.74	14.53 16.88 16.15 16.65 14.94	12.15 14.08 13.11 13.72 12.58
W 2 W2+E1 Gs W2+E1 Sp W2+E2 Gs W2+E2 Sp	13.15 ^I 16.14 ^F 14.98 ^{GH} 15.05 ^{GH} 14.82 ^H	12.09 ^H 14.73 ^F 12.65 ^H 13.96 ^G 13.62 ^G	12.62 15.44 13.82 14.51 14.22	12.00 ^F 14.88 ^D 13.61 ^E 15.17 ^D 13.19 ^E	11.27 ^{JK} 13.03 ^H 12.38 ^I 12.89 ^{HI} 10.98 ^{KL}	11.64 13.96 12.99 14.03 12.09	15.21 ^I 17.56 ^{EFG} 16.94 ^{FGH} 17.25 ^{FG} 16.14 ^H	7.28 ^K 9.04 ^{FG} 9.90 ^E 10.19 ^{DE} 8.68 ^{FGH}	11.25 13.30 13.42 13.72 12.41	14.36 ^{EF} 15.69 ^{CD} 15.08 ^{DE} 15.08 ^{DE} 13.93 ^F	6.48 ^{HI} 7.52 ^{EF} 7.91 ^{DE} 7.63 ^{DEF} 7.11 ^{FGH}	10.42 11.61 11.49 11.36 10.52	12.13 14.70 13.41 14.27 13.16	10.84 12.46 12.46 12.54 11.47
W 3 W3+E1 Gs W3+E1 Sp W3+E2 Gs W3+E2 Sp	9.74 ^K 12.61 ^{IJ} 11.69 ^J 12.37 ^{IJ} 10.35 ^K	9.89 ^J 12.34 ^H 12.01 ^H 12.34 ^H 11.29 ^I	9.82 12.48 11.85 12.36 10.82	7.51 ^H 12.16 ^F 11.95 ^F 9.88 ^G 8.12 ^H	8.84 ^N 11.74 ^J 10.57 ^{LM} 10.97 ^{KL} 10.07 ^M	8.18 11.95 11.26 10.43 9.09	12.92 ^L 14.37 ^J 14.19 ^{JK} 14.14 ^{JK} 13.41 ^{KL}	6.27 ^L 8.12 ^{HIJ} 7.69 ^{IJK} 8.20 ^{HI} 7.39 ^{JK}	9.59 11.25 10.94 11.17 10.40	10.60 ^I 12.23 ^G 11.54 ^{GH} 12.01 ^G 10.87 ^{HI}	5.30 ^J 6.67 ^{GHI} 6.29 ^I 6.59 ^{HI} 6.03 ^I	7.95 9.45 8.92 9.30 8.45	9.00 12.22 11.56 11.39 9.96	8.77 10.35 9.93 10.24 9.43

E1 = Azotobacter chrocooccum E2 = Pseudomonas sp Gs = Grains soaking Sp = foliar application control = 100 % field capacity W1 = 75% field capacity W2 = 50 % field capacity W3 = 25 % field capacity

Table 3: Total chlorophyll content (μg / cm²) in the flag leaf of wheat cultivars Gmiza 9 and Sakha 93 wheat plants as affected by different levels of irrigation water deficit and two bacterial endophytes and their interactions at different stages during 2009 / 2010 and 2010 / 2011 seasons.

	Gmiza 9							<u> </u>		T				
Treatment	2009/20010 season Total chlorophyll after			2010/2011 season Total chlorophyll after		Means	2009/20010 season Total chlorophyll after		Saki Means	2010/2011 season Total chlorophyll after		Means	Gmiza 9 mean of means	Sakha 93 mean of means
	40	70		40	70		40	70		40	70			
Control	52.82 ^E	56.37 ^{CD}	54.59	48.90 ^C	51.41 ^{CD}	50.16	58.87 ^{EF}	36.07 ^C	47.47	58.19 ^B	29.08 ^C	43.59	52.38	45.53
E 1 Gs E 1 Sp E 2 Gs E 2 Sp	60.83 ^{AB} 58.19 ^{BC} 62.56 ^A 56.41 ^{CD}	59.91 ^{AB} 60.75 ^A 60.05 ^{AB} 58.01 ^{BC}	60.37 59.47 61.31 57.21	62.17 ^A 59.98 ^{AB} 61.97 ^A 58.14 ^B	57.99 ^{AB} 56.47 ^B 58.78 ^A 53.08 ^C	60.09 58.23 60.38 46.61	70.71 ^A 65.57 ^{BC} 68.14 ^{AB} 64.09 ^{CD}	48.51 ^A 47.02 ^A 49.21 ^A 44.59 ^B	59.61 56.29 58.68 54.34	64.61 ^A 63.69 ^A 65.20 ^A 62.58 ^A	35.26 ^{AB} 33.81 ^B 36.55 ^A 33.04 ^{AB}	49.94 48.75 50.88 47.81	60.23 58.85 60.85 51.91	54.78 52.52 54.78 51.08
W 1 W1+E1 Gs W1+E1 Sp W1+E2 Gs W1+E2 Sp	46.91 ^{FGH} 53.66 ^{DE} 51.42 ^E 51.92 ^E 47.78 ^{FG}	54.88 ^{DE} 53.55 ^E 54.03 ^{DE}	47.35 54.27 52.49 52.98 48.67	41.03 ^E 49.41 ^C 45.87 ^D 49.73 ^C 41.39 ^E	44.04 ^G 50.93 ^{DE} 49.07 ^E 50.38 ^{DE} 46.24 ^F	42.54 50.17 47.47 50.06 43.82	53.45 ^{HI} 61.37 ^{DE} 58.89 ^{EF} 61.54 ^{DE} 56.71 ^{FG}	26.76 ^{GHI} 34.56 ^{CD} 33.05 ^{DE} 35.46 ^C 29.17 ^F	40.11 47.97 45.97 48.50 42.94	51.65 ^C 57.34 ^B 50.62 ^{CD} 52.55 ^C 49.57 ^{CD}	23.57 ^{EFG} 26.86 ^{CD} 25.12 ^{DEF} 26.03 ^{DE} 25.59 ^{DE}	37.61 42.10 37.87 39.29 37.58	44.95 52.22 49.98 51.52 46.25	38.86 45.04 41.92 43.89 40.26
W 2 W2+E1 Gs W2+E1 Sp W2+E2 Gs W2+E2 Sp	39.20 ^I 48.11 ^F 44.56 ^{GH} 44.86 ^{GH} 44.16 ^H	38.82 ^H 47.27 ^F 40.61 ^H 44.82 ^G 43.71 ^G	38.76 47.69 42.59 44.84 43.94	35.64 ^F 44.22 ^D 40.41 ^E 45.06 ^D 39.18 ^E	36.41 ^{JK} 42.07 ^H 39.98 ^I 41.64 ^{HI} 35.48 ^{KL}	36.03 43.15 40.19 43.35 37.33	48.66 ^J 56.18 ^{FGH} 54.21 ^{GHI} 55.21 ^{GH} 51.64 ^I	23.31 ^K 28.93 ^{FG} 31.69 ^E 32.62 ^E 27.79 ^{FGH}	35.99 42.56 42.95 43.92 39.72	45.94 ^{EF} 50.19 ^{CD} 48.26 ^{DE} 48.25 ^{DE} 44.58 ^F	20.74 ^{HI} 24.06 ^{EF} 25.29 ^{DE} 24.40 ^{DEF} 22.74 ^{FGH}	33.34 37.13 36.78 36.33 33.66	37.39 45.42 41.39 44.09 40.64	34.67 39.85 39.69 40.13 36.69
W 3 W3+E1 Gs W3+E1 Sp W3+E2 Gs W3+E2 Sp	29.04 ^K 37.58 ^{IJ} 34.85 ^J 36.85 ^{IJ} 30.84 ^K	31.78 ^J 39.61 ^H 38.54 ^H 39.61 ^H 36.27 ^J	30.41 38.59 36.69 38.23 33.56	22.29 ^H 36.11 ^F 35.50 ^F 29.35 ^G 24.12 ^H	28.55 ^N 37.93 ^J 34.15 ^{LM} 35.45 ^{KL} 32.53 ^M	25.42 37.02 34.83 32.40 28.33	41.35 ^M 45.98 ^{JK} 45.39 ^{KL} 45.26 ^{KL} 42.92 ^{LM}	20.05 ^L 25.99 ^{HIJ} 24.59 ^{IJK} 26.24 ^{HI} 23.67 ^{JK}	30.70 35.99 34.99 35.75 33.29	33.93 ^I 39.15 ^G 36.94 ^{GH} 38.44 ^G 34.79 ^{HI}	16.97 ^J 21.35 ^{GHI} 20.14 ^I 21.08 ^{HI} 19.30 ^I	25.45 30.25 28.54 29.76 27.05	27.92 37.81 35.76 35.32 30.95	28.08 33.12 31.77 32.76 30.17

E1 = Azotobacter chrocooccum E2 = Pseudomonas sp Gs = Grains soaking Sp = foliar application control = 100 % field capacity W1 = 75% field capacity W2 = 50 % field capacity W3 = 25 % field capacity

Table 4: Proline and relative water content (RWC) in the flag leaf of wheat cultivars Gmiza 9 and Sakha 93 as affected by different levels of water stress and two bacterial Endophytes and their interactions during 2009 / 2010 and 2010 / 2011 seasons.

			za 9			Sak	ha 93				Mea	ns of
	2009 /	2010 /	2009 /	2010 /	2009 /	2010 /	2009 /	2010 /	Mea	ns of		VC
Treatment	2010	2011	2010	2011	2010	2011	2010	2011				
	Proline	RWC	Proline	RWC	Proline	RWC	Proline	RWC	Gmiza 9	Sakha 93	Gmiza 9	Sakha 93
Control	0.389 ^J	80.98 ^E	0.418 ^H	87.10 ^D	0.440 ^H	98.27 ^C	0.585 ¹	93.77 ^C	0.404	0.513	84.04	96.02
E 1 Gs E 1 Sp E 2 Gs E 2 Sp	0.397 ^J 0.405 ^J 0.396 ^J 0.406 ^J	88.53 ^B 85.04 ^D 89.89 ^A 86.69 ^C	0.429 ^H 0.420 ^H 0.417 ^H 0.418 ^H	95.07 ^A 90.05 ^B 93.92 ^A 91.25 ^B	0.441 ^H 0.412 ^{HI} 0.435 ^H 0.389 ^I	100.67 ^B 98.49 ^C 103.18 ^A 96.43 ^D	0.576 ^l 0.553 ^l 0.580 ^l 0.583 ^l	99.36 ^A 96.10 ^B 99.89 ^A 94.09 ^C	0.413 0.413 0.407 0.412	0.509 0.483 0.508 0.486	91.80 87.55 91.91 88.97	100.02 97.29 101.53 95.26
W 1 W 1 + E 1 Gs W 1 + E 1 Sp W 1 + E2 Gs W 1 + E 2 Sp	0.496 ^I 0.613 ^H 0.606 ^H 0.625 ^H 0.612 ^H	76.16 ^G 85.44 ^D 80.75 ^E 86.71 ^C 81.34 ^E	0.557 ^G 0.722 ^F 0.697 ^F 0.713 ^F 0.697 ^F	71.26 ^J 86.32 ^D 84.58 ^E 88.53 ^C 82.70 ^F	0.548 ^G 0.650 ^F 0.619 ^F 0.639 ^F 0.634 ^F	82.82 ^I 96.12 ^D 94.21 ^E 96.65 ^D 92.73 ^{EF}	0.724 ^H 0.957 ^F 0.946 ^{FG} 0.951 ^F 0.906 ^G	77.95 ^l 89.73 ^D 86.85 ^E 89.41 ^D 84.68 ^F	0.527 0.668 0.652 0.669 0.655	0.636 0.804 0.783 0.795 0.770	73.71 85.88 82.67 87.62 82.02	80.39 92.93 90.53 93.03 88.71
W 2 W 2 + E1 Gs W 2 + E1 Sp W 2 + E 2 Gs W 2 + E 2 Sp	0.753 ^G 0.856 ^{EF} 0.834 ^F 0.865 ^E 0.843 ^{EF}	64.51 ^J 80.35 ^E 78.38 ^F 80.27 ^E 78.82 ^F	0.872 ^E 1.047 ^{CD} 1.003 ^D 1.035 ^{CD} 1.002 ^D	61.24 ^L 77.97 ^G 73.28 ^I 78.86 ^G 75.54 ^H	0.844 ^E 0.995 ^D 0.966 ^D 0.986 ^D 0.986 ^D	73.81 ^L 93.02 ^{EF} 89.32 ^G 91.72 ^F 89.76 ^G	1.026 ^E 1.221 ^D 1.213 ^D 1.253 ^D 1.211 ^D	65.42 ^L 87.16 ^E 83.31 ^G 86.19 ^E 80.09 ^H	0.813 0.952 0.919 0.950 0.923	0.935 1.108 1.089 1.119 1.098	62.88 79.16 75.83 79.57 77.18	69.62 90.09 86.32 88.96 84.93
W 3 W 3 + E 1 Gs W 3 + E 1 Sp W 3 + E 2 Gs W 3 + E 2 Sp	0.952 ^D 1.233 ^B 1.215 ^{BC} 1.264 ^A 1.203 ^C	49.31 ^K 74.57 ^H 71.23 ^I 75.35 ^{GH} 72.19 ^I	1.079 ^C 1.415 ^A 1.363 ^B 1.411 ^A 1.414 ^A	47.14 ^M 73.18 ^I 70.08 ^{JK} 70.82 ^J 68.90 ^K	1.070 ^C 1.436 ^A 1.347 ^B 1.420 ^A 1.363 ^B	51.96 ^M 84.68 ^H 79.78 ^J 83.12 ^I 77.24 ^K	1.327 ^C 1.567 ^A 1.548 ^{AB} 1.513 ^B 1.518 ^B	49.58 ^M 78.63 ^I 74.74 ^J 75.93 ^J 73.11 ^K	1.016 1.324 1.289 1.338 1.309	1.199 3.002 1.448 1.467 1.441	48.23 73.88 70.66 73.09 70.55	50.77 81.66 77.26 79.53 75.18

E1 = Azotobacter chrocooccum E2 = Pseudomonas sp Gs = Grains soaking Sp = foliar application control = 100 % field capacity W1 = 75% field capacity W2 = 50 % field capacity W3 = 25 % field capacity

Values within the same vertical areas with the same letter are not significantly different at 5% probability level by Duncan's Multiple Range Test

Table 5: Leaf water content (LWC) and leaf water deficit (LWD) in the flag leaf of wheat cultivars Gmiza 9 and Sakha 93 as affected by different levels of water stress and different Endophytes bacteria and their interactions during 2009 / 2010 and 2010 / 2011 seasons.

	auring 20	JU9 / ZU I (and 201	0/2011	seasons.							
		Gm	iza 9			Sakh	na 93					
	2009 /	2010 /	2009 /	2010 /	2009 /	2010 /	2009 /	2010 /	Gmiza	9 mean	Sakha 9	3 mean
Freatment	2010	2011	2010	2011	2010	2011	2010	2011				
	LWC	LWC	LWD	LWD	LWD	LWD	LWC	LWC	LWC	LWD	LWC	LWD
Control	2.92F	3.15FG	19.02G	12.90JK	1.37K	6.23K	2.54FG	2.83G	3.04	15.96	2.69	3.80
1 Gs	3.65A	3.82A	11.471	4.93N	00.00	0.69M	2.98A	3.26A	3.74	8.20	3.12	0.35
1 Sp	3.21CD	3.61C	14.96H	9.95LM	1.51K	3.90L	2.90A 2.70D	3.13C	3.41	12.46	2.92	2.71
2 Gs												
2 Sp	3.46B	3.69B	10.111	6.08N	00.00	0.14M	2.85B	3.20B	3.58	8.09	3.03	0.07
	3.29C	3.43D	13.31H	8.75M	3.57J	5.91K	2.61E	3.07D	3.36	11.03	2.84	4.74
/ 1 / 1+ E 1 Gs	0.441	0.041	00.045	00.075	47.40	00.055	0.401	4.000	0.00	00.40	4.04	40.00
1+ E 1 GS	2.14J	2.61L	23.84D	29.07D	17.18E	22.05E	2.19J	1.690	2.38	26.46	1.94	19.62
p	3.47B	3.40D	14.56H	13.68J	3.88J	10.27J	2.78C	2.94F	3.44	14.12	2.86	7.08
1 + E2 Gs	3.15DE	3.24E	19.25G	15.421	5.791	13.151	2.59E	2.74H	3.19	17.34	2.67	9.47
/1+E2	3.41B	3.37D	13.29H	11.47KL	3.35J	10.59J	2.73CD	3.02E	3.39	12.38	2.88	6.97
р	3.08E	3.06H	18.66G	17.30H	7.27HI	15.32H	2.53G	2.79G	3.07	17.98	2.66	11.29
12	1.99K	2.110	35.49B	38.76B	26.19B	34.58B	1.99L	1.45P	2.05	37.13	1.72	30.39
2 + E1 Gs	2.97F	3.19EF	19.65FG	22.03G	6.98HI	12.841	2.59EF	2.641	3.08	20.84	2.62	9.91
/ 2 + E1 Sp	2.79G	3.08GH	21.62E	26.72E	10.68G	16.69G	2.43H	2.38K	2.94	24.17	2.41	13.69
2 + E2 Gs	2.98F	3.04H	19.73FG	21.14G	8.28H	13.811	2.61E	2.53J	3.01	20.44	2.57	11.0
/ 2 + E 2 p	2.75G	2.85J	21.18EF	24.46F	10.24G	19.91F	2.46H	2.28L	2.80	22.82	2.37	15.08
3	1.52L	1.79P	50.69A	52.86A	48.04A	50.42A	1.49M	1.12Q	1.66	51.78	1.31	49.2
3+ E 1 Gs	2.70GH	2.77K	25.43D	26.82E	15.32F	21.37E	2.381	2.28L	2.74	26.13	2.33	18.3
3 + E1 Sp	2.501	2.44M	28.77C	29.92CD	20.22D	25.26D	2.15JK	2.12N	2.47	29.35	2.14	22.7
3 + E2 Gs	2.65H	2.951	24.65D	29.18D	16.88E	24.07D	2.351	2.36K	2.80	26.92	2.36	20.4
/ 3 + E2 Sp	2.511	2.24N	27.81C	31.10C	22.76C	26.89C	2.13K	2.18M	2.38	29.46	1.09	24.8
	2.011	L.L-714	27.010	31.100	22.700	20.000	2.101	Z. 101VI	2.00	20.70	1.00	∠-т.ч

E1 = Azotobacter chrocooccum E2 = Pseudomonas sp Gs = Grains soaking Sp = foliar application control = 100 % field capacity W1 = 75% field capacity W2 = 50 % field capacity W3 = 25 % field capacity

Table 6: Nitrogen % and crude protein in the flag leaf of wheat cultivars Gmiza 9 and Sakha 93 wheat plants as affected by different levels of water stress and two different bacterial endophytes and their interactions during 2009 / 2010 and 2010 / 2011 seasons.

during	2003 / 2010 6	and 2010 / 20									
		Gm	iza 9		Sakha 93						
Tractment	2009	/ 2010	2010	/ 2011	2009	/ 2010	2010 / 2011				
Treatment	N%	Crude Protein	N%	Crude Protein	N%	Crude Protein	N%	Crude Protein			
Control	2.67 ^C	16.69 ^C	2.33 ^C	14.58 ^C	1.84 ^D	11.52 ^D	2.21 ^{CD}	13.78 ^{CD}			
E 1 Gs	3.19 ^A	19.98 ^A	3.06 ^A	19.10 ^A	2.49 ^A	15.60 ^A	2.98 ^A	18.59 ^A			
E 1 Sp	2.99 ^B	18.67 ^B	2.79 ^B	17.43 ^B	2.11 ^C	13.19 ^C	2.72 ^B	16.99 ^B			
E 2 Gs	3.14 ^A	19.61 ^A	2.99 ^A	18.67 ^A	2.29 ^B	14.36 ^B	3.03 ^A	18.96 ^A			
E 2 Sp	3.02 ^B	18.89 ^B	2.79 ^B	17.43 ^B	2.14 ^C	13.34 ^c	2.67 ^B	16.698 ^B			
W 1	2.32 ^E	14.51 ^E	2.10 ^E	13.13 ^E	1.66 ^{EF}	10.35 ^{EF}	1.83 ^G	11.45 ^G			
W 1 + E 1 Gs	2.63 ^C	16.41 ^C	2.32 ^C	14.51 ⁶	1.86 ^D	11.59 ^D	2.22 ^{CD}	13.85 ^{CD}			
W 1 + E 1 Sp	2.64 ^C	16.48 ^C	2.26 ^{CD}	14.15 ^{CD}	1.79 ^D	11.16 ^D	2.14 ^{DE}	13.34 ^{DE}			
W 1 + E2 Gs	2.49 ^D	15.53 ^D	2.33 ^C	14.58 ^C	1.83 ^D	11.45 ^D	2.28 ^C	14.22 ^C			
W 1 + E 2 Sp	2.65 ^C	16.55 ^C	2.19 ^D	13.71 ^D	1.75 ^{DE}	10.94 ^{DE}	2.08 ^{EF}	12.97 ^{EF}			
W 2	1.77 ^H	11.08 ^H	1.84 ^ℍ	11.52 ^{HI}	1.33 ^H	8.31 ^H	1.53 ^H	9.55 ^H			
W 2 + E1 Gs	2.17 ^F	13.56 ^F	2.10 ^E	13.13 ^E	1.65 ^上	10.28 ^{EF}	1.98 ^F	12.39 ^F			
W 2 + E1 Sp	2.17 ^F	13.56 ^F	2.05 ^E	12.83 ^E	1.56 ^{FG}	9.77 ^{FG}	1.76 ^G	11.01 ^G			
W 2 + E 2 Ġs	2.10 ^F	13.13 ^F	1.97 ^F	12.32 ^F	1.63 ^{EF}	10.21 ^{EF}	1.98 ^F	12.39 ^F			
W 2 + E 2 Sp	2.15 ^F	13.42 ^F	1.97 ^F	12.32 ^F	1.48 ^G	9.26 ^G	1.74 ^G	10.86 ^G			
W 3	1.45 ^J	9.04 ^J	1.59 ^J	9.99 ^J	0.83 ^K	5.17 ^K	1.06 ^J	6.64 ^J			
W 3 + E 1 Gs	1.99 ^G	12.47 ^G	1.93 ^{FG}	12.03 ^{FG}	1.26 ^{HI}	7.88 ^{HI}	1.44 ^{HI}	8.97 ^{HI}			
W 3 + E 1 Sp	1.94 ^G	12.10 ^G	1 79 ^l	11 16'	1.18 ^{IJ}	7.36 ^{IJ}	1.40 ^l	8.75 ¹			
W 3 + E 2 Gs	1.67 ^l	10.43 ^l	1.91 ^{FGH}	11.96 ^{FGH}	1.23 ^{HIJ}	7.66 ^{HIJ}	1.48 ^{HI}	9.26 ^{HI}			
W 3 + E 2 Sp	1.79 ^H	11.16 ^H	1.87 ^{GH}	11.67 ^{GH}	1.12 ^J	7.00 ^J	1.39 ¹	8.68 ^l			

E1 = Azotobacter chrocooccum E2 = Pseudomonas sp Gs = Grains soaking Sp = foliar application control = 100 % field capacity W1 = 75% field capacity W2 = 50 % field capacity W3 = 25 % field capacity