



Original Article

Mechanisms of Drought Tolerance in *Cornulaca monacantha* Del.

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Article Info

Article history :

Received 23/5/2016

Received in revised form 10/6/2016

Accepted 19/6/2016

Keywords:

*Cornulaca monacantha*

Carbohydrate

Proline

Glycinebetaine

Na<sup>+</sup>

Abstract

Samples of shoot system of *Cornulaca monacantha* were collected from two sand embankments location; Rommana (North Sinai) and Khamisa (Siwa Oasis) in Egypt during wet and dry seasons. Degree of succulence, chlorophyll a, chlorophyll b, carotenoids, total carbohydrate, soluble and non-soluble carbohydrates, crude protein, total soluble protein, proline, glycinebetaine, total phenols, Na<sup>+</sup>, K<sup>+</sup>, Ca<sup>++</sup> and Mg<sup>++</sup> were determined. Degree of succulence, chlorophyll a, chlorophyll b, carotenoids, soluble carbohydrate, soluble protein, total phenols, Ca<sup>++</sup> and Mg<sup>++</sup> were decreased in dry season. On the other hand, total carbohydrate, non-soluble carbohydrate, proline, glycinebetaine, Na<sup>+</sup> and K<sup>+</sup> were increased in dry season. *Cornulaca monacantha* inhabiting Rommana was higher in degree of succulence, chlorophyll b, glycinebetaine, total phenols, soluble protein, Ca<sup>++</sup> and Mg<sup>++</sup> and lower in the other characters (except in Na<sup>+</sup> was non-significant) than that inhabiting Khamisa.

1. Introduction

Drought stress is one of the premier limitations to global agricultural production due to complexity of the water-limiting environment and changing climate. To overcome water deficit, plants have included a series of mechanisms; morphological, physiological, biochemical, cellular and molecular levels (Fang and Xiong 2015). Drought tolerance is the ability of plant to withstand with reduced water potential of the plant tissue. Drought tolerance mechanisms allow the plant to function or at least survive, at reduced water potential (Verlues *et al.*, 2014). The overproduction of different types

of compatible solutes is one of the most common stress tolerance strategies in plants (Serraj and Sinclair 2002). Compatible solutes are highly soluble, low molecular weight and non-toxic even at high concentration. They protect plants from stress by different means viz., contribution in osmoregulation, scavenging reactive oxygen species and stabilization of membranes (Farooq *et al.*, 2009). In addition to compatible solutes, inorganic ions accumulation reduce the osmotic potential and improve cell water retention in response to water stress (Morgan 1984; Rhodes and Samaras 1994).

Glycinebetaine as an organic osmolyte plays an im-

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portant role in enhancing plant tolerance under abiotic stress including drought (Quan *et al.*, 2004). Beside the direct protective roles of glycinebetaine either through positive effects on enzyme and membrane integrity or as an osmoprotectant, it may protect cells from environmental stresses indirectly by participating in signal transduction pathways (Subbarao *et al.*, 2000). Also, proline acts as an osmoprotectant in plants subjected to drought conditions (Yamada *et al.*, 2005). Relative water content and total protein were decreased by water stress, while free amino acids were increased in *Morus alba* L. (Ramanjulu and Sudhakar, 1997). Soluble carbohydrates are considered as important metabolites in plants under drought stress (Arabzadeh, 2012). Drought stress imposed a significant increase in soluble carbohydrate and decrease in soluble protein in *Cicer arietinum* cultivars (Mafakheri *et al.*, 2011).

*Cornulaca monacantha* is a desert plant, woody shrub, richly branched, usually with long internodes, that is now included in family Amaranthaceae (formerly Chenopodiaceae). This study aims to shed the light on the adaptive responses of *C. monacantha* as drought tolerance plant. In this study, succulence degree, plant pigments (chl. a, chl. b and carotenoids), total carbohydrate, soluble carbohydrate, crude protein, soluble protein, total phenols, proline, glycine betaine, Na<sup>+</sup>, K<sup>+</sup>, Ca<sup>++</sup> and Mg<sup>++</sup> were determined in shoot system of *C. monacantha* growing naturally at Rommana (North Sinai) and Khamisa (Siwa Oasis) during wet and dry season.

## 2. Materials and methods

### 2.1. Study area

The two studied locations (Rommana and Khamisa)

are sandy embankments habitat. The first one lies on the western part of North Sinai, farthest by about 35 km from Suez Canal. The second one lies on the western part of Siwa Oasis, farthest by about 60 km from the Lybian boarder. The geographical position of the collected samples and meteorological data were represented in Table 1. Meteorological data for Rommana (L1) and Khemaisa (L2) were collected from the nearest station from each one during 2013-2014 as represented in Table 2 using ETo calc Program according to Francisco Catalán (2015).

### 2.2. Soil analyses

Two depths of soil (0-30 cm and 30-60 cm) supporting *C. monacantha* were collected (three replicates for each depth) to determine physical and chemical properties. Samples of plant shoot system were collected in three replicates from the two locations; Rommana and Khamisa during wet (March) and dry (August) seasons in 2014. The soil samples were air-dried, sieved through 2mm meshes and granulometric analysis was determined as described by Jackson (1967) and illustrated in Table 3. Electrical conductivity (EC), Na<sup>+</sup>, K<sup>+</sup>, Ca<sup>++</sup>, Mg<sup>++</sup>, Cl<sup>-</sup>, SO<sub>4</sub><sup>--</sup> and HCO<sub>3</sub><sup>-</sup> were estimated and determined in soil water extract (1:1). Soil water content, EC, Na<sup>+</sup>, K<sup>+</sup>, Ca<sup>++</sup> and Mg<sup>++</sup> were determined following the methods described by Rowell (1994). Chlorides were determined as described by Jackson (1967). Sulphates were determined by the turbidity method according to Rainwater and Thatcher (1960). Bicarbonate was determined by titration with sulphuric acid 0.01N according to Reitemeier (1943). All studied soil chemical analyses were represented in Table 4.

**Table 1.** The geographical position (GPS) of the collected plant samples and meteorological data.

Locations	GPS of the collected samples	GPS of the collected meteorological data
Rommana (L1)	31° 00' 583" latitude 32° 39' 352" longitude	30° 40' 156" latitude 32° 36' 287" longitude
Khamisa (L2)	29° 12' 470" latitude 25° 24' 828" longitude	29° 12' 72" latitude 25° 31' 839" longitude

**Table 2.** Meteorological data covering studied locations during 2013-2014.

Months	Precipitation mm/m		Temperature °C		Relative humidity %		Sun shine %		ETo mm/m	
	L1	L2	L1	L2	L1	L2	L1	L2	L1	L2
January	9	3	13.2	12.4	61.4	55.1	67.7	76.9	73	85
February	8	1	14.1	14.0	59.1	49.2	70.3	79.6	85	102
March	8	2	16.1	17.2	56.6	43.4	71.0	73.6	122	152
April	5	1	20.1	21.7	51.4	36.0	74.0	74.5	158	197
May	2	0	22.9	25.3	51.3	33.6	79.4	77.7	193	231
June	0	0	25.9	28.7	53.6	32.6	87.8	86.3	210	248
July	0	0	27.5	29.2	55.6	38.3	86.6	89.5	216	252
August	0	0	27.6	29.1	57.6	40.9	87.5	89.3	202	229
September	0	0	25.8	26.9	59.6	43.0	83.4	84.9	163	184
October	2	0	22.9	22.8	60.1	47.6	83.2	83.5	135	143
November	7	2	18.8	17.5	61.2	55.1	76.9	82.0	92	97
December	9	2	14.6	13.4	63.1	56.9	65.7	77.4	71	81
Total	50	11							1 720	2 001

**Table 3.** physical analysis of soil supporting *C. monacantha*.

Location	Depth cm	2-1 mm	1-0.5 mm	0.5-0.25 mm	0.25- 125 mm	0.125- 0.063 mm	< 0.063 mm	texture
Rommana	0-30	0.1	3.76	51.73	39.49	4.5	0.42	Sandy
	30-60	0	1.23	43.24	52.46	3.02	0.05	Sandy
Khamisa	0-30	0.45	4.14	29.66	47.87	17.76	0.12	Sandy
	30-60	0.2	16.72	56.75	23.82	2.45	0.06	Sandy

**Table 4.** Chemical analysis of soil supporting *C. monacantha*.

seasons	location	Depth Cm	Moisture content %	EC Mmhos	Cations meq/L				Anions meq/L		
					Na <sup>+</sup>	K <sup>+</sup>	Ca <sup>++</sup>	Mg <sup>++</sup>	Cl <sup>-</sup>	HCO <sub>3</sub> <sup>-</sup>	SO <sub>4</sub> <sup>-</sup>
Wet	Rommana	0-30	2.30	0.666	2.43	0.23	2.00	2.50	2.7	0.7	4.1
Wet	Rommana	30-60	1.22	0.825	3.35	0.26	2.25	3.25	4.3	1.0	3.5
Wet	Khamisa	0-30	0.66	0.813	6.04	0.72	1.75	1.25	4.2	1.1	3.9
Wet	Khamisa	30-60	0.53	0.495	3.74	0.31	1.00	0.50	2.5	0.9	2.3
Dry	Rommana	0-30	0.70	1.025	4.87	0.27	2.50	4.00	4.4	0.7	5.9
Dry	Rommana	30-60	0.83	0.655	2.70	0.21	2.00	2.00	2.0	0.7	4.1
Dry	Khamisa	0-30	0.07	0.435	3.04	0.28	1.00	0.25	1.0	1.0	2.6
Dry	Khamisa	30-60	0.11	0.405	2.80	0.26	1.00	0.50	1.0	1.5	2.0

### 2.3. Plant analyses

Plant samples were divided into two parts for analyses, in the first part fresh samples were used to determine degree of succulence, chlorophyll a, chlorophyll b, carotenoids and soluble protein, the second part plant samples were dried at till constant weight, then ground to fine powder to determine contents of total carbohydrate, soluble and non-soluble carbohydrate, crude protein, proline, glycinebetaine, total phenols, Na<sup>+</sup>, K<sup>+</sup>, Ca<sup>++</sup> and Mg<sup>++</sup>.

Degree of succulence was calculated as fresh/dry weight according to Dehan and Tall (1978). The photosynthetic pigments (chlorophyll a, chlorophyll b and carotenoids) were determined quantitatively as described by Metzner *et al.* (1965). Soluble protein concentrations were determined by the method of Bradford (1976) using bovine serum albumin as a standard. Total and soluble carbohydrates were extracted and estimated calorimetrically by applying the general phenol-sulphoric acid method as adopted by Chaplin and Kennedy (1994). Values were expressed as g/100g dry wt. Crude protein was estimated by Kjeldahl method as described by Pregle (1945). Free proline was measured by the sulfo-salicylic acid- ninhydrin method according to Bates *et al.* (1973). Glycine-betaine content was estimated colorimetrically as described by Grieve and Grattan (1983). Total phenols were determined using Folin-Denis reagent as described by Shahidi and Nacz (1995). A known weight of plant sample was extracted by 80% ethanol, 1 ml of the extract, 0.5 ml of Folin reagent were mixed well, 1 ml of saturated Na<sub>2</sub>CO<sub>3</sub> and mixed well then 3 ml of dist. water were added. After 1 hour read the developed blue color at 725 nm by spectrophotometer using catechol as a standard. Sodium and potassium were measured by flame photometer and Calcium and magnesium by atomic absorption spectrophotometer as described by Kalra (1998).

### 2.4. Statistical analysis

Data obtained from the experiment of plant analysis were subjected to the proper statistical analysis of variance of the split plot design according to the procedure obtained by Snedecor and Cochran (1969). Mean values

of treatments were differentiated by using L.S.D at 5 % level as mentioned by Steel (1960).

## 3. Results

Degree of succulence increased significantly in *Cornulaca monacantha* during wet season, recorded the highest value (3.16) in plants inhabiting Rommana (Table 5). *C. monacantha* at Rommana was higher in degree of succulence than that at Khamisa.

Regarding plant pigments, chlorophyll a, chlorophyll b and carotenoids were decreased in dry season in *C. monacantha* as shown in Table 5. *C. monacantha* inhabiting Khamisa was higher in chlorophyll a and carotenoids, while that inhabiting Rommana was higher in chlorophyll b. The highest contents of chlorophyll a and b were recorded in *C. monacantha* growing at Khamisa in wet season 24.67 and 11.47 mg/100g fr. wt., respectively. While the highest carotenoids content was recorded in *Cornulaca* inhabiting Rommana in wet season (4.3 mg/100g fr. wt.).

In general, total carbohydrate and non-soluble carbohydrate in *C. monacantha* increased in dry season, recorded the highest values 43.73% and 39.98% in the plant of Rommana during dry season but the interaction was non-significant in total carbohydrate as illustrated in Table 5. Soluble carbohydrate generally increased in the plant during wet season (4.57%).

Proline was significantly elevated in dry season in *C. monacantha*, it increased by 35% and elevated in *C. monacantha* of Khamisa by 89% compared to that in Rommana (Table 6). The highest value of proline (112 µg/g dry wt.) was noticed in *C. monacantha* inhabiting Khamisa during dry season. As proline glycinebetaine was elevated significantly in dry season, it increased by 70% (Table 6). Glycinebetaine increased by 12% in *C. monacantha* at Rommana compared to that at Khamisa. The highest value of glycinebetaine (301.2) was noticed in the plant at Rommana in dry season.

A significant accumulation of total phenols in *C. monacantha* was noticed in wet season as shown in Table 6. *Cornulaca* at Rommana was higher in total phenols than that at Khamisa. The highest value of total phenols (224.1 mg/100g dry wt.) was recorded in the

plant at Rommana in wet season, followed by that at Khamisa in dry season (192.1 mg/100g dry wt.), Rommana in dry season (170.3 mg/100g dry wt.) and Khamisa in wet season (166.7 mg/100g dry wt.).

Crude protein had non-significant change seasonally (Table 6). It was higher accumulation in *C. monacantha* at Rommana than that at Khamisa. The highest value of crude protein (11.49%) was noticed in the plant at Rommana in wet season. Soluble protein in the plant was affected significantly by seasons and locations while the interaction between them was non-significant. Generally, it increased in wet season and was higher in *C. monacantha* at Rommana than that at Khamisa.

Sodium content was elevated significantly in dry season (Table 7). There was no significant change in the studied plant between the two locations. The highest sodium content was recorded in *Cornulaca* at Khamisa in dry season (3.62%), and the lowest content was record-

ed in *Cornulaca* at the same location in wet season (1.38%). Like sodium, potassium in *Cornulaca* was elevated significantly in dry season (Table 7). It was higher in *Cornulaca* growing at Rommana than that growing at Khamisa. Potassium content in *Cornulaca* can be arranged in descending order as those growing at Rommana in wet season (0.57%), those growing at Rommana in dry season (0.51%), those growing at Khamisa in dry season (0.39%), those growing at Khamisa in wet season (0.19%).

Calcium and magnesium in *C. monacantha* were increased significantly in wet season (Table 7). They were higher in the plant growing at Rommana than that at Khamisa. The highest calcium content was noticed in the plant growing at Rommana in wet season (3.02%), while the highest magnesium was noticed in the plant growing at Khamisa in wet season (1.82%).

**Table 5.** Succulence (fr.w/dry W.), chlorophyll a, b and carotenoids (mg/100g fr. W.), total, soluble and non-soluble carbohydrate (g/100g dry w.) in *Cornulaca monacantha*.

Season							
	Succulence	Chlorophyll a	Chlorophyll b	Carotenoids	Total carbohydrate	Soluble carbohydrate	Non-soluble carbohydrate
Wet (W)	2.44	18.31	10.53	4.12	39.64	4.57	35.16
Dry (D)	2.24	11.43	7.48	1.57	40.97	3.74	37.26
L.S.D at 5%	s	s	s	s	s	s	s
locations							
Rommana (L1)	2.97	10.41	9.56	2.67	37.62	3.41	34.24
Khamisa (L2)	1.72	19.33	8.46	3.00	42.99	4.90	38.11
L.S.D at 5%	s	s	s	s	s	s	s
Interaction between locations and seasons							
W* L1	3.16	11.95	9.60	4.30	37.03	3.12	33.91
W* L2	1.73	24.67	11.47	3.89	42.26	6.02	36.24
D* L1	2.77	8.87	9.51	1.03	38.22	3.69	34.53
D* L2	1.70	13.99	5.45	2.11	43.73	3.78	39.98
L.S.D at 5%	0.03	1.18	0.16	0.14	n.s	0.10	1.18

Where s = significant

n.s. = non-significant

**Table 6.** Proline ( $\mu\text{g/g}$  dry wt.), glycinebetaine (Mmole/g dry wt.), total phenols (mg/100g dry wt.), crude protein (g/100 g dry wt.) and soluble protein (mg/100g fr. Wt.) in *Cornulaca monacantha*.

seasons					
	Proline	Glycine betaine	Total phenols	Crude protein	Soluble protein
Wet (W)	63.5	173.6	197.7	8.23	156.09
Dry (D)	85.5	295.3	179.3	8.21	131.61
L.S.D at 5%	s	s	s	n.s	s
locations					
Rommana (L1)	51.5	247.4	195.9	8.23	126.37
Khamisa (L2)	97.5	221.5	181.2	8.21	161.57
L.S.D at 5%	s	s	s	s	s
Interaction between locations and seasons					
W* L1	44	193.7	225.1	11.49	136.64
W* L2	83	153.4	170.3	4.96	175.45
D* L1	59	301.2	166.7	9.36	116.11
D* L2	112	289.5	192.1	7.06	147.6
L.S.D at 5%	2.4	7.9	13.6	0.29	n.s

Where s = significant

n.s. = non-significant

**Table 7.** Sodium, potassium, calcium and magnesium (g/100g dry wt.) in *Cornulaca monacantha*.

Seasons				
	Sodium	Potassium	Calcium	Magnesium
Wet (W)	2.04	0.38	2.94	1.78
Dry (D)	3.04	0.45	2.20	1.68
L.S.D at 5%	s	s	s	s
Locations				
Rommana (L1)	2.58	0.54	2.78	1.76
Khamisa (L2)	2.50	0.29	2.36	1.70
L.S.D at 5%	n.s	s	s	s
Interaction between locations and seasons				
W* L1	2.70	0.57	3.02	1.74
W* L2	1.38	0.19	2.86	1.82
D* L1	2.47	0.51	2.54	1.77
D* L2	3.62	0.39	1.86	1.59
L.S.D at 5%	0.11	0.01	0.12	0.15

#### 4. Discussion

From meteorological data, it became known that Rommana location lies on arid area affected by Mediterranean weather, while Khamisa location lies on hyper-arid area (the Western Desert). Soils supporting *C. monacantha* are non-saline usually lower than 1Mmhos except at the first depth (0-30 cm) of Rommana, slightly higher than 1 Mmhos in dry season. The higher moisture content in Rommana soil due to the high precipitation and lower evapotranspiration compared with that in Khamisa especially in wet season. These climatic conditions reflected on degree of succulence to increase in *C. monacantha* growing at Rommana. A decline in succulence degree was recorded in dry season. However, *C. monacantha* growing at Khamisa showed a non-significant decrease. Gorai *et al.* (2015) noticed the decrease in relative water content in *Ephedra alata* by 17% and 30% compared with control plants after 7 days and 14 days of the drought stress treatment respectively. The decrease in chlorophyll a and b in dry season agreed with the studies of (Morsy *et al.*, 2008; Mafakheri *et al.*, 2010; Din *et al.*, 2011). The production of reactive oxygen species under drought conditions can disturb the chlorophyll biosynthesis (Herbinger *et al.*, 2002). Drought stress stimulates inhibition of photosynthetic activity in plant tissues due to imbalance between light capture and its utilization (Foyer and Noctor, 2000). Liu *et al.* (2011) found that drought stress significantly decreased chl. a+b in *Cinnamomum bodinieri*, *Broussonetia papayrifera* and *Platycarya longipes* and also decreased carotenoids in *Platycarya longipes* and *Pteroceltis tatarinowii*. Sai *et al.* (2012) reported that water deficit decreased the contents of carotenoids and anthocyanin in *Atriplex hortensis*. The accumulation of total carbohydrate and non-soluble carbohydrate in dry season agreed with Nour El-Din and Ahmed (2004). They found that carbohydrate content in *Crotalaria aegyptiaca* tend to decrease in the rainy season and to increase at the end of the dry season. Abd El-Maboud (2006) concluded that *Deverra tortuosa* as a xerophyte plant depends to large extent on the accumulation of organic intermediates; particularly carbohydrate in building osmotic potential. Soluble carbohydrate re-

sponded negatively by drought in *C. monacantha*. Similar results were obtained by Morsy *et al.* (2008). On the other hand, many authors found the reverse trend (Choluj *et al.*, 2008; Liu *et al.*, 2011; Mafakheri *et al.*, 2011).

Proline accumulation in *C. monacantha* in response to drought and temperature stresses is a good indicator of drought tolerance mechanism. Many researchers suggest a positive correlation between proline accumulation and drought stress in plants (Liu *et al.*, 2011; Hayat *et al.*, 2012; Cha-um *et al.*, 2013; Gorai *et al.*, 2015). In this respect, Anjum *et al.* (2012) reported that progressive stress reduced the relative water content but increased proline concentration in the two maize cultivars; dog dan-60 and nong da-95. Proline act as an osmolyte, a reactive oxygen species scavenger and a molecular chaperone stabilizes proteins structure (Szabados and Savoure 2010). Like proline, glycinebetaine accumulated in response to drought stress. Similar results were obtained by Sankar *et al.* (2007). Glycinebetaine and proline are two major organic osmolytes that accumulate in a variety of plant species in response to environmental stresses including drought. Both compounds are thought to have positive effects on enzyme and membrane integrity along with adaptive roles in mediating osmotic adjustment in plant grown under stress conditions (Ashraf and Foolad, 2007). Hoque *et al.* (2008) reported that betaine reduce oxidative damage in tobacco suspension cells by enhancing reactive oxygen species detoxification system. It has been reported that glycinebetaine in *Salsola tetrandra* is more effective as a defense against free radicals under drought than salinity stress (Abd El-Maboud and Eisa, 2016). Cha-um *et al.* (2013) concluded that exogenous application of glycinebetaine significantly alleviates water-deficit in indica rice and maintain grain yield. Total phenols are affected negatively by drought stress. It was lower in *C. monacantha* growing at Khamisa which is subjecting to more drought conditions than that growing in Rommana. A decrease of total phenols in dry season agreed with Król *et al.* (2014) who found that total phenolics were decreased in leaves and roots of grapevine when subjected to drought stress.

An increase in crude protein of *C. monacantha* inhabiting Khamisa and a decrease in that inhabiting Rommana in dry season neglected seasons effect. Regarding locations effect, *C. monacantha* inhabiting Khamisa decreased significantly in crude protein by 42% compared to that in Rommana. That decrease may be return to the reduction in soil moisture content supporting the plant at Khamisa location. Davies (1987) interpreted the decrease of protein under drought stress to might be the result of increased protease activity or due to fragmentation of proteins resulted from reactive oxygen species effect. The decrease of soluble protein in dry season agreed with those in Mafakheri *et al.* (2011).

It is noteworthy that  $\text{Na}^+$  and  $\text{K}^+$  contents of the plant increased in dry season by 1.5 and 1.2 fold. That increase may contribute to decrease the osmotic potential in plant cell in response to drought stress and help in water uptake along a soil-plant gradient. In spite of growing at poor sand dunes, non-saline soil, *C. monacantha* was able to take up large quantities of  $\text{Na}^+$  from soil and accumulate it in aboveground tissue. In addition to the reduction in  $\text{K}^+$  accumulation in the plant indicates that  $\text{K}^+$  can be substituted by  $\text{Na}^+$  as cation exchange of monovalent ions. It has been reported that  $\text{Na}^+$  could substituted for  $\text{K}^+$  in some cellular functions especially in a terrestrial life under limited  $\text{K}^+$  supply, but in others it is toxic. In the vacuole,  $\text{Na}^+$  is not toxic and can do osmotic functions, reducing the total  $\text{K}^+$  requirements and improving growth when  $\text{K}^+$  is a limiting factor (Navarro and Rubio, 2006). Epstein *et al.* (1973) concluded that  $\text{Na}^+$  competes with  $\text{K}^+$  for uptake by plant roots, supporting that  $\text{K}^+$  transporters are also the gates for  $\text{Na}^+$  entry.  $\text{Na}^+$  ions inter root cells through HKT proteins and non-selective voltage-independent cation channels, some of which (labeled CNGC) are inactivated by cyclic nucleotides (cAMP and cGMP) Pardo and Quintero (2002). Potassium is not only an essential nutrient for plant growth and development, but also contributes to the survival of plants exposed to various biotic and abiotic stresses (Wang *et al.*, 2013). The decrease of  $\text{Ca}^{++}$  and  $\text{Mg}^{++}$  levels of *C. monacantha* in dry season could be attributed to the competition between them and monovalent cations in the soil such as

$\text{Na}^+$  which reduce  $\text{Ca}^{++}$  and  $\text{Mg}^{++}$  intake by the plant. It has been reported that  $\text{Mg}^{++}$  and  $\text{Ca}^{++}$  availability increased steadily throughout the wet season, and then decreased rapidly in dry season for three constitutive years on Barro Colorado Island (Mulkey *et al.*, 1996). In this trend, Abd El-Maboud (2006) found that  $\text{Mg}^{++}$  ions were higher in winter than summer in *Panicum turgidum* and *Deverra tortuosa*.

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## 5. Conclusion

*Cornulaca monacantha* is an example of drought tolerant plants, can grow in arid and hyper arid zones. The plant was able to overcome drought stress by the accumulation of total carbohydrate, an organic osmolytes like proline and glycinebetaine. In addition to the accumulation of  $\text{Na}^+$  as inorganic osmolyte, that may displace  $\text{K}^+$  in some metabolic processes.

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## References

- Abd El-Maboud, M.M. (2006): Ecophysiological responses of some xerophytes from Wadi El- Gafra, the Eastern Desert of Egypt. M. Sc. Thesis, Bot. Dept. Fac. Sci., Al- Azhar Univ.
- Abd El-Maboud, M.M. and Eisa, S.S. (2016): Role of internal antioxidant in the adaptation of *Salsola tetrandra* Forssk. at different habitats of the North Western Coast of Egypt. *Research Journal of Pharmaceutical, Biological and Chemical Sciences*, 7 (3): 48-55.
- Anjum, S.A., Saleem, M.F., Wang, L., Bilal, M.F. and Saeed, A. (2012): Protective role of glycinebetaine in maize against drought-induced lipid peroxidation by enhancing capacity of antioxidative system. *Australian Journal of Crop Science*, 6(4): 567-583.
- Arabzadeh, N. (2012): The effect of drought stress on soluble carbohydrates (sugars) in two species of *Haloxylon persicum* and *Haloxylon Aphyllum*. *Asian Journal of Plant Sciences*, 11(1): 44-51.
- Ashraf, M. and Foolad, M.R. (2007): Roles of glycinebetaine and proline in improving plant abiotic stress resistance. *Environmental and Experimental Botany*, 59: 206-216.
- Bates, L.; R. Waldren and I. Teare (1973): Rapid deter-



- mination of free proline for water-stress studies. *Plant and Soil*, 39, 205-207.
- Bradford, M. (1976): A rapid and sensitive method for the quantitation of microgram quantities of protein utilizing the principle of protein-dye binding. *Analytical Biochemistry*, 72, 248-254.
- Chaplin, M.F. and Kennedy, J.F. (1994): Carbohydrate analysis: A practical approach. 2<sup>nd</sup> Ed. Oxford Univ., Press Oxford, New York, Tokyo, pp. 344.
- Choluj, D., Karwowska, R., Ciszewska, A. and Jasinska, M. (2008). Influence of long-term drought stress on osmolyte accumulation in sugar beet (*Beta vulgaris L.*) plants. *Acta Physiologia Plantarum*, 30: 679-687.
- Chu-um, S., Samphumphuang, T. and Kirdmanee, C. (2013): Glycinebetaine alleviates water deficit stress in indica rice using proline accumulation, photosynthetic efficiencies, growth performances and yield attributes. *Australian Journal of Crop Science*, 7(2): 213-218.
- Davies, K.J.A. (1987): Protein damage and degradation by oxygen radicals 1. General aspects. *The Journal Biological Chemistry*, 262: 9895-9901.
- Dehan, K. and Tall, M. (1978): Salt tolerance of the wild relatives of the cultivated tomato: response of *Solanum pennellii* to high salinity. *Irrigation Science*, 1: 71-76.
- Din, J., Khan, S.U., Ali, I. and Gurmani, A.R. (2011): Physiological and agronomic response of canola varieties to drought stress. *The Journal of Animal & Plant Sciences*, 21(1): 78-82.
- Epstein, M. (1973): Mechanisms of ion transport through plant cell membranes. *International Review of Cytology*, 34: 123-167.
- Fang, Y. and Xiong, L. (2015): General mechanisms of drought response and their application in drought resistance improvement in plants. *Cellular and Molecular Life Sciences*, 72: 673-689.
- Farooq, M., Wahid, A., Kobayashi, N., Fujita, D. and Basra, S.M.A. (2009): Plant drought stress: effects mechanisms and management. *Agronomy for Sustainable Development*, 29: 185-212.
- Foyer, C.H. and Noctor, G. (2000): Oxygen processing in photosynthesis: regulation and signaling. *New Phytologist*, 146: 359-388.
- Francisco Catalán (2015): Watering crops and landscapes, smart irrigation. Calculating irrigation needs anywhere. ETo clac. <http://www.mobogenie.com/download-eto-calc-1385120.html>
- Gorai, M., Laajili, W., Santiago, L.S. and Neffati, M. (2015): Rapid recovery of photosynthesis and water relations following soil drying and re-watering is related to the adaptation of desert shrub *Ephedra alata* subsp. *alenda* (Ephedraceae) to arid environments. *Environmental and Experimental Botany*, 109: 113-121.
- Greive, C.M. and S.R. Grattan (1983): Rapid assay for determination of water-soluble quaternary amino compounds. *Plant and Soil*, 70: 303-307.
- Hayat, S., Hayat, Q., Alyemeni, M.N., Wani, A.S., Pichtel, A. and Ahmad, A. (2012): Role of proline under changing environments. *Plant Signaling & Behavior*, 7(11): 1456-1466.
- Herbinger, K., Tausz, M., Wonisch, A., Soja, G., Sorgner, A. and Grill, D. (2002): Complex interactive effects of drought and ozone stress on the antioxidant defence systems of two wheat cultivars. *Plant Physiology and Biochemistry*, 40: 691-696.
- Hoque, M.A., M.N. Banu, Y. Nakamura, Y. Shimoishi, and Y. Murata, (2008): Proline and glycinebetaine enhance antioxidant defence and methylglyoxal detoxification systems and reduce NaCl-induced damage in cultured tobacco cells. *Journal of Plant Physiology*, 165: 813-824.
- Jackson, M.L. (1967): In "Soil chemical analysis". Prentice. Hall of India Private, New Delhi. Pp.485.
- Kalra, Y.P. (1998): Handbook of reference methods for plant analysis. CRC Press, p153-165.
- Król, A., Amarowicz, R. and Weidner, S. (2014): Changes in the composition of phenolic compounds and antioxidant properties of grapevine roots and leaves (*Vitis vinifera L.*) under continuous of long-term drought stress. *Acta Physiologiae Plantarum*, 36: 1491-1499.
- Liu, C., Liu, Y., Guo, K., Fan, D., Li, G. and Zheng, Y. (2011): Effect of drought on pigments, osmotic ad-

- justment and antioxidant enzymes in six woody plant species in karst habitats of southwestern China. *Environmental and Experimental Botany*, 71: 174-183.
- Mafakheri, A., Siosemardeh, A., Bahramnejad, B., Struic, P.C. and Sohrabi, Y. (2010): Effect of drought stress on yield, proline and chlorophyll contents in three chickpea cultivars. *Australian Journal of Crop Science*, 4(8): 580-585.
- Mafakheri, A., Siosemardeh, A., Bahramnejad, B., Struic, P.C. and Sohrabi, Y. (2011): Effect of drought stress and subsequent recovery on protein, carbohydrate contents, catalase and peroxidase activities in three chickpea (*Cicer arietinum*) cultivars. *Australian Journal of Crop Science*, 5(10): 1255-1260.
- Metzner, H.; H. Rau and H. Senger (1965): Untersuchungen Zur Synchronisierbarkeit einzelner Pigment-Mangel Mutanten Von *Chlorella*. *Planta*, 65 (2), 186- 194.
- Morgan, J.M. (1984): Osmoregulation and water stress in higher plants, *Annual Review of Plant Physiology*, 35: 299-319.
- Morsy, A.A., Yossef, A.M., Mosallam, H.A.M. and Hashem, A.M. (2008): Assessment of selected species along Al-Alamein-Alexandria international desert road, Egypt. *Journal of Applied Sciences Research*, 4(10): 1276-1284.
- Mulkey, S.S., Chazdon, R.L. and Smith, A.P. (1996): In "Tropical forest plant ecophysiology", 1<sup>st</sup> ed. Chapman & Hall. P478.
- Navarro, A. and Rubio, F. (2006): High-affinity potassium and sodium transport systems in plants. *Journal of Experimental Botany*, 57(5): 1149-1160.
- Nour El-Din, N.M. and Ahmed, F.A. (2004): Effect of seasonal variation on secondary metabolites and nutritive value of *Crotalaria aegyptiaca* Benth. *Egyptian Journal of Desert Researches*, 54(1): 129-147.
- Pardo, J.M. and Quintero, F.J. (2002): Plants and sodium ions: keeping company with the enemy. *Genome Biology*, 3(6): 1017.1-1017.4.
- Pregl, F. (1945): In "Quantitative organic microanalysis", 4<sup>th</sup> . Edit., J. & A. Churchill, Ltd., London.
- Quan, R.D., Shang, M., Zhang, H. and Zhang, J. (2004): Improved chilling tolerance by transformation with betA gene for the enhancement of glycinebetaine synthesis in maize. *Plant Science*, 166: 141-149.
- Rainwater, F.H. and Thatcher, L.L. (1960): In "*Methods for collection and analysis of water samples*". U.S. Geol. Survey. Water Supply, pp 301.
- Ramanjulu, S. and Sudhakar, C. (1997): Drought tolerance is partly related to amino acid accumulation and ammonia assimilation: A comparative study in two mulberry genotypes differing in drought sensitivity. *Journal of Plant Physiology*, 150: 345-350.
- Reitemeier, R.F. (1943): Semimicro analysis of saline soil solutions. *Indus and Engin. Chem., Analyt. Ed.* 15: 393-402.
- Rhodes, D. and Samaras, Y. (1994): Genetic control of osmoregulation in plants. *Cellular and Molecular Physiology of Cell Volume Regulation*, pp 347-361.
- Rowell D.L. (1994): *Soil Science: Methods and Applications*. Dept of Soil Science, Univ. of Reading. Copublished in the US with John Willey and Sons Inc.; New York, pp: 350.
- Sai, K.S., Bouraoui, Karray, N. B.N., Jaffel, K., Rejeb, M.N., Leclerc, J.C. and Ouerghi, Z. (2012): Water deficit-induced oxidative stress in leaves of Garden Orach (*Atriplex hortensis*). *Research Journal of Biotechnology*, 7(4), 46-52.
- Sankar, B., Jaleel, C.A., Manivannan, P., Kishorekumar, A., Somasundaram, R.S. and Panneerselvam, R. (2007): Drought-induced biochemical modifications and proline metabolism in *Abelmoschus esculentus* (L.) Moench. *Acta Botanica Croatica*, 66(1): 43-56.
- Serraj, R. and Sinclair, T.R. (2002): Osmolyte accumulation: can it really help increase crop yield under drought conditions" *Plant Cell Environment*, 25: 333-341.
- Shahidi, F. and Naczki, M. (1995): *Food phenolics: Sources, Chemistry, Effects, Applications*, Technomic Publishing Company Inc., Lancaster PA., pp: 231- 245.

- Snedecor, G.W. and Cochran, W.G. (1969): In "Statistical Methods?". 6<sup>th</sup> ed. Iowa State Univ., Press, Ames., Iowa, U.S.A.
- Steel, G.D.R. (1960): In "Principal and procedures of statistics?". New York McGraw-Hill Book Co., pp. 481.
- Subbarao, G.V., Nam, N.H., Chauhan, Y.S. and Johansen, C. (2000): Osmotic adjustment, water relations and carbohydrate remobilization in pigeonpea under water deficits. *Journal of Plant Physiology*, 157: 651-659.
- Szabados, L. and Savoure, A. (2010): Proline: a multifunctional amino acid. *Trends in plant Science*. 15 (2): 89-97.
- Verslues, P.E., Lasky, J.R., Juenger, T.E., Liu, T.W. and Kumar, M.N. (2014): Genome-wide association mapping combined with reverse genetics identifies new effectors of low water potential-induced proline accumulation in *Arabidopsis*. *Plant Physiology*, 164: 144-159.
- Wang, M., Zheng, Q., Shen, Q. and Guo, S. (2013): The critical role of potassium in plant stress response. *International Journal of Molecular Sciences*, 14 (4): 7370-7390.
- Yamada, M.; Morishita, H.; Urano, K.; Shiozaki, N.; Kazuko, Y.; Shinozaki, K. and Yoshida, Y. (2005): Effects of free proline accumulation in petunias under drought stress. *Journal of Experimental Botany*, 56 (417): 1975-1981.

## المخلص العربى

### أليات تحمل الجفاف فى نبات الحاد

محمد محمد عبد المعبود

قسم البيئة النباتية والمراعى - مركز بحوث الصحراء

تم تجميع عينات من المجموع الخضرى لنبات الحاد من موقعين للتكوينات الرملية: رمانة بشمال سيناء وخميسة بواحة سيوة بمصر. تم تقدير درجة العصارية ، كلوروفيل أ، كلوروفيل ب ، الكاروتينويد ، الكربوهيدرات الكلية ، الكربوهيدرات الذائبة والغير ذائبة، البروتين الخام، البروتين الكلى الذائب، البرولين، الجلايسين بيتاين، الفينولات الكلية، الصوديوم، البوتاسيوم، الكالسيوم والماغنسيوم. أظهرت النتائج انخفاض كل من درجة العصارية ، كلوروفيل أ، كلوروفيل ب، الكاروتينويد ، السكريات الذائبة، البروتين الذائب، الفينولات الكلية، الكالسيوم والماغنسيوم فى موسم الجفاف وعلى الجانب الأخر ازداد كل من الكربوهيدرات الكلية والغير ذائبة ، البرولين، الجلايسين بيتاين، الصوديوم والبوتاسيوم فى موسم الجفاف. تفوق نبات الحاد النامى بمنطقة رمانة على نظيره بمنطقة خميسة فى كل من درجة العصارية، كلوروفيل ب ، الجلايسين بيتاين، الفينولات الكلية، البروتين الذائب ، الكالسيوم والماغنسيوم بينما حدث العكس فى باقى الصفات فيما عدا الصوديوم حيث كان الاختلاف بين الموقعين غير معنوى.



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**JOESE 5**



## **Mechanisms of Drought Tolerance in *Cornulaca monacantha* Del.**

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***Reprint***

**Volume 45, Number 2 : 175-186**

**(2016)**