Salt Tolerance of Muskmelons as Affected by Various Salinities in Soil Culture¹

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Summary

Muskmelons (*Cucumis melo* L.) were grown in soil to determine the salt tolerance as affected by salinization of sea water, NaCl, Na₂SO₄ and MgCl₂ in Experiment I, and MgSO₄ in Experiment II, at osmotic potentials of -0.95 (only MgSO₄), -1.20, -1.70 and -2.70 bars compared with a control of -0.70 bars of base nutrient solution. Fruit fresh weight and whole plant dry weight were greatest in the control and tended to decrease in each salinity with decreasing osmotic potentials of treatment solutions. At -2.70 bars fruit fresh weight was 61.3, 55.7, 63.2, 54.0 and 35.9% compared with the control in the sea water, NaCl, Na₂SO₄, MgCl₂ and MgSO₄ series, respectively. Most of the plants at -2.70 bars died within 60 days after transplanting in the MgSO₄ series. No plants died in the other series. Growth in decreasing order was control>sea water \doteqdot NaCl \doteqdot Na₂SO₄ \doteqdot MgCl₂>MgSO₄· Na, Mg, Cl and SO₄ content in leaves and soil solution (SSo) tended to increase with decreasing osmotic potentials of treatment solutions in sodium-, magnesium-, chloride- and sulfate-salinities, respectively. EC values of SSo increased and osmotic potentials of SSo decreased as osmotic potentials of treatment solutions decreased.

Introduction

Salt tolerance of muskmelons as affected by various salinities in sand culture has been reported(14). That experiment was undertaken to explain the effect of salt source and concentration on the growth and development, using single salts added to a base nutrient solution. It was concluded that the growth in decreasing order was control>sea water $= NaCl > MgCl_2 > Na_2SO_4 = MgSO_4$ series. Also, the salt tolerance of muskmelons grown in different media (sand, soil and nutrient solution) has been studied using diluted sea water(11). It was shown that growth was more suppressed in sand than in soil culture. Thus, it is inferred that different salinities may affect plant growth suppression variously and alter the chemical properties of soil solutions in sand and soil Therefore, the present expericultures.

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This work was partly supported by Grants-in-Aid for young scientist from the Ministry of Education. ment was conducted to compare the effect of various salinities on salt tolerance of muskmelons in sand and soil cultures.

Materials and Methods

This study consisted of 2 experiments. Sea water and single salts, such as NaCl, Na₂SO₄ and MgCl₂, were added to a base nutrient solution in Experiment I and MgSO₄ was added in Experiment II.

Experiment I Uniform muskmelon seedlings, cv. Fall No.1 of Earl's Favourite in the 4 leaf stage were transplanted in wooden containers (40×40×20 cm) filled with 16 liters of paddy soil and placed in a greenhouse on Sept.7, 1977. Temperature inside the greenhouse was above 18°C. Plants were topped at the 20 th node and allowed to bear only one fruit per plant around the 10 th node. Lateral shoots and other flower buds were removed. The paddy soil used was a mixture of light clay paddy soil taken at Shizuoka and sandy clay paddy soil taken at Iwata in the ratio of 1:1 by volume. There were 13 treatments, as shown in Table

	Treatments		Treatments							
No.	Salinities	$\pi(\text{bars})^z$	Composition of base nutrient solution							
1	Base nutr. soln.	-0.70	None	2.43	1. Na ₂ HPO ₄ ·12 H ₂ O	1 mM				
2	Sea water ^y	-1.21	1.9%	3.45	2. K ₂ SO ₄	3 mM				
3		-1.70	3.8%	4.50	3. $MgSO_4 \cdot 7 H_2O$	2 mM				
4		-2.70	7.6%	6.60	4. Ca(NO ₃) ₂ ·4 H ₂ O	4 mM				
-				0.00	5. Fe 1 pp	m (Fe-EDTA)				
5	NaCl	-1.20	687 mgNaCl/liter	3.38	6. Zn 0.05 pp	m (ZnSO ₄ ·7 H ₂ O)				
6		-1.70	1,374	4.65	7. Cu 0.02 ppi	m (CuSO ₄ ·5 H ₂ O)				
7		-2.70	2,748	7.05	8. B 0.5 pp	m (H ₃ BO ₃)				
8	Na ₂ SO ₄	-1.20	1,261 mgNa ₂ SO ₄ /liter	3.66	9. Mo 0.05 pp:	m $(Na_2MoO_4 \cdot 2 H_2O)$				
9		-1.70	2,521	5. 18	10. Mn 0.5 pp	m (MnSO ₄)				
10		-2.70	5,042	8.08	pH ≑ 6. 0					
11	MgCl ₂	-1.20	1,728 mgMgCl ₂ 6H ₂ O/lis	er 3.67						
12		-1.70	3,456	5.24						
13		-2.70	6,912	8.21						

Table 1. Composition of treatment solutions and base nutrient solution.

y Sca water contains 20,500 ppm Cl, 10,082 ppm Na, 2,632 ppm SO₄, 1,262 ppm Mg, 445 ppm K and 393 ppm Ca.

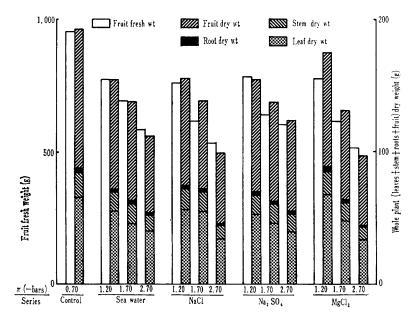


Fig.1. Effect of various salinities on fruit fresh weight and whole plant (leaves+stem+roots+fruit) dry weight. Figures below columns indicate osmotic potential (-bars) of treatment solutions.

1, consisting of control (base nutrient solution), and sea water, NaCl, Na₂SO₄ and MgCl₂ dissolved in the base nutrient solution at osmotic potentials of -0.50, -1.00 and -2.00 bars. The osmotic potential of the base nutrient solution was -0.70 bars. The sea water was taken from Shimizu (Miho).

Each treatment had 5 replications. Thus, there were 65 containers. Treatment solutions were applied to the soil medium from Sept. 11 to harvest (late Nov.). Applications (0.5 to 1 liter/container/time) were made twice on sunny days, once on cloudy days and none on rainy days. At the end of the

² Osmotic potential. The π of treament solutions includes -0.70 bars of base nutrient solution.

experiment, leaf, stem, root and fruit fresh and dry weights were measured. Extraction of soil solutions (SSo) at pF 0 to 3.8 and the other analytical methods on leaves, fruit and SSo were the same as described in earlier papers (10, 14).

Experiment II Uniform muskmelon seedlings, cv. Fall No.1 of Earl's Favourite in the 2.5 leaf stage were transplanted in wooden containers filled with 16 liters of the light clay paddy soil and placed in the greenhouse on Sept.14, 1979. MgSO₄ was dissolved in the base nutrient solution at osmotic potentials of 0 (control), -0.25, -0.50, -1.00 and -2.00 bars. Treatment solutions, as shown in Table 6, were applied to the soil medium from Sept.17 to harvest (early Dec.). The other experimental procedures and methods of analyses were the same as in Experiment I.

Results

Experiment I

Growth and fruit quality (Table 2, Fig. 1) Fruit fresh weight and whole plant, leaf, stem and root dry weight tended to be greatest in the control and decreased in each salinity as osmotic potentials of treatment

Table 2. Effect of various salinities on fruit quality of muskmelons in soil culture.

Osmotic	Salinities						
potential (bars)	Sea water	NaCl	Na ₂ SO ₄	$MgCl_2$			
So	luble solids	(%)	·· —·				
Control	14. 1e ^y						
-1.20	15.0cd	15.0cd	15. 2bcd	14.7de			
-1.70	15. 3bcd	15. 3bcd	15.4bcd	15.6bc			
-2.7 0	15.3bcd	16.0b	16.9a	15.4bcd			
	ste (degree	<u> </u>					
Control	0						
-1.20	0~0.5	0.5	0.5	0.5			
- 1.70	0.5	1	1	1			
-2.70	-2.70 0		1~1.5 2				

- * Control does not contain any additional salts and is maintained at -0.70 bars of osmotic potential.
- y Mean separation in each item by Dyncan's multiple range test, 5% level.
- * Tastes were evaluated from 0 (none) to 5: salty in sea water and NaCl; salty, bitter and astringent in Na₂SO₄; and bitter in MgCl₂.

solutions decreased. At isosmotic concentrations similar suppression of growth was observed in all series. At -1.20 bars in the MgCl2 series, leaf, stem and root dry weights were especially not significantly different from the control and whole plant dry weight was greater than in the other series. Fruit fresh weight decreased to the same extent at isosmotic potentials. At -2.70 bars fruit fresh weight was 61.3, 55.7, 63.2 and 54.0% compared with the control in the sea water, NaCl, Na2SO4 and MgCl₂ series, respectively. These values were not significantly different. Fruit soluble solids seemed to be less affected by treatments, but were relatively low in the control. Fruit taste varied with the salts added to the base nutrient solution: salty in the sea water and NaCl series; salty, bitter and astringent in the Na₂SO₄ series; and bitter in the MgCl₂ series. The intensity of taste increased with decreasing osmotic potentials

Table 3-1. Effect of various salinities on Na, Mg, Cl and SO₄ content in leaves in soil culture (% of dry matter).

Osmotic	Salinities						
potential (bars)	Sea water	NaCl	Na₂SO₄	MgCl ₂			
N:	a			,			
Control	0.16efy						
-1.20	0.18ef	0.22ef	0.36de	0.14ef			
-1.70	0.35de	0.48cd	0.58c	0.10f			
-2.70	0.87a	0.65bc	0.83ab	0.08f			
M	g						
Control	1.20ef						
-1.20	1.23ef	1.09fg	1.18fg	2. 19c			
-1.70	1. 34de	1.04g	1.15fg	2.77b			
-2.70	1.40d	0.80h	0.88h	3.94a			
Ci	- ·						
Control	0. 68e			_			
-1.20	1.80d	2.45d	0.64e	2.63d			
-1.70	3.90c	3.95c	0.63e	4.31c			
-2.70	6.85a	6.20ab	0.66e	5. 7 5b			
SC	04			. —			
Control	0.86e						
-1.20	0.95e	1.87d	5.52c	0.78e			
-1.70	0.50e	0.66e	7.66b	0.53e			
-2.70	0.55e	0.40e	9.50a	0.35e			

z,y Same as Table 2.

Table 3-2. Effect of various salinities on total-N, P,
Ca and K content in leaves in soil culture
(% of dry matter).

Osmotic	Salinities						
potential (bars)	Sea water	NaCl	_ Na₂SO₄	$MgCl_2$			
To	otal-N						
Control	2.51ef*						
-1.20	2.71bcde	2.63de	3.10a	2.70bcde			
-1.70	2.89b	2.81bcd	2.61de	2.69bcde			
-2.70	2.52ef	2.86bc	2.65cde	2.33f			
P							
Control	0. 57c						
-1.20	0.74a	0.73ab	0.71ab	0.76a			
-1.70	0.71ab	0.81a	0.73ab	0.76a			
-2.70	0.68abc	0.69abc	0.75a	0.60bc			
Ca							
Control	6.92 bc						
-1.20	7.27ab	7.46a	7.05abc	6.78bc			
-1.70	7.27ab	7.54a	6.92 bc	6.66c			
-2.70	7.23ab	7.47a	5. 94 d	5.77 d			
К							
Control	1.75c						
-1.20	0.74e	1.81bc	1.98abc	1.17d			
-1.70	1.35d	2.23ab	2.00abc	1.20 d			
-2.70	2.03abc	2.30a	2.33a	1.05 de			

z,y Same as Table 2.

of treatment solutions.

Chlorosis and cupping were observed on lower leaves of the plant at -1.70 and -2.70 bars in the sea water and NaCl series. Only chlorosis appeared on lower leaves in the MgCl₂ series at -1.70 and -2.70 bars. Plants did not wither in all series. Yellowish white spots on leaves in the MgCl₂ series and interveinal chlorosis in the Na₂SO₄ series reported in the previous sand culture experiment, were not observed in the present soil culture experiment.

Major mineral elements in leaves (Table 3) Na, Mg, Cl and SO₄ content increased with decreasing osmotic potentials of treatment solutions in sodium-, magnesium-, chloride-and sulfate-salinities, respectively. Here, sea water was classified as sodium- and chloride-salinities because Na and Cl are dominant ions. Na was relatively higher in the Na₂SO₄ series than in the other sodium-

Table 4. Effect of various salinities on Na, Mg, Cl and Ca content in fruit in soil culture (% of dry matter).

Osmotic		Sali	Salinities			
potential (bars)	Sea water	NaCl	Na ₂ SO ₄	$MgCl_2$		
N.	a					
Control	0.26f*					
-1.20	0.54e	0.62 d e	0.73cd	0.26f		
-1.70	0.89bc	0.77cd	0.88 bc	0.21 f		
-2.70	1.02b	1.18a	1.19a	0.23f		
M	g					
Control	0.24 d					
-1.20	0.23 de	0.21efg	0.21efg	0.29c		
-1.70	0.22ef	0.22ef	0.21efg	0.34b		
-2.70	0.21efg	0.18h	0.19gh	0.40a		
Cl	I					
Control	0.57e					
-1.20	1.46d	1.50d	0.63e	1.54d		
-1.70	2.09c	1.99c	0.52e	2.12c		
-2.70	3.05b	3. 23a	0.52e	3.41a		
Ca	a					
Control	0. 17cd					
-1.20	0.19bcd	0.22ab	0.16 d	0.19bcd		
-1.70	0.22ab	0.25a	0.21abc	0.20cd		
-2.70	0.22ab	0.23ab	0.16 d	0. 19bcd		

z,y Same as Table 2.

salinities. Mg was much higher in the MgCl₂ series than in the other series. Cl was almost similar at isosmotic potentials in the chloride-salinity. SO₄ was much higher in the Na₂SO₄ series (5.52 to 9.50%) and somewhat higher at -1.20 bars in the NaCl series (1.87%) than in the control (0.86%). P tended to be higher in all series (0.60 to 0.81%) than in the control (0.57%). Ca was lower at -2.70 bars in the Na₂SO₄ (5.94%) and MgCl₂ (5.77%) series than in the control (6.92%). Total-N and K were not affected.

Major mineral elements in fruit (Table 4) Na, Mg and Cl content increased with decreasing osmotic potentials of treatment solutions in sodium-, magnesium- and chloridesalinities, respectively. SO₄ was present only in traces compared with other ions (data not shown). Ca was less affected by the treatments. Na and Mg, and Cl tended to

Table 5-1. Chemical properties of soil solution (pF = 0 to 3.8) at the end of the experiment.

Osmotic	Salinities					
potential (bars)	Sea water	NaCl	Na ₂ SO ₄	$MgCl_2$		
N	a (me/l)					
Controlz	24. 1h ^y					
-1.20	69.0g	93.3fg	108. 4ef	34.4h		
-1.70	131.2e	159. 0d	181.5cd	23. 5h		
-2.70	186.5c	218.0b	239.4a	29.4h		
M	g (me/l)					
Control	17. 2h					
-1.20	27.6fg	25.7fgh	15.6h	73.7c		
-1.70	42.8e	33.5efg	17.8h	143.8b		
-2.70	56. 7 d	32.1efg	19.4gh	274.7a		
Cl	(%)					
Control	0.064 f					
-1.20	0. 325e	0.430 de	0.102f	0.475d		
-1.70	0.665c	0.775c	0.081f	0.744c		
-2.70	1.055b	1.060 b	0.073f	1.453a		
SC), (%)					
Control	0.24g	••	•			
-1.20	0.25 fg	0.31defg	0.59c	0.26fg		
-1.70	0.29defg	0.34def	1.06b	0.35de		
-2.70	0.32defg	0.27ef	1.71a	0.37d		

z.y Same as Table 2.

accumulate more in the outer flesh (outer mesocarp) and rind (epicarp), respectively.

Chemical properties of SSo at the end of the experiment (Table 5) As osmotic potentials of treatment solutions decreased, Na, Mg, Cl and SO₄ concentrations increased in sodium-, magnesium-, chlorideand sulfate-salinities, respectively. Na was lower in the sea water series, higher in the NaCl series and highest in the Na2SO4 series in the sodium-salinity. Cl was relatively high in the MgCl₂ series compared with the sea water and NaCl series. P was higher in the Na₂SO₄ series and lower in the MgCl₂ series than in the control. Ca in the MgCl₂ series increased with decreasing osmotic potentials of treatment solutions and was higher than that in the control. K and EC values tended to increase and osmotic potentials tended to decrease as osmotic potentials of treatment solutions decreased. NO₃-N and pH were less affected by treatments.

Table 5-2. Chemical properties of soil solution (pF = to 3.8) at the end of the experiment.

Osmotic		Salinities				
potential (bars)	Sea water	NaCl	Na_2SO_4	MgCl ₂		
N	O ₃ -N (ppm)					
Control	181 bcde ^y			_		
-1.20	190bcd	160cdef	136def	136def		
-1.70	220abc	180bcde	201abc	129e f		
-2.70	221 a b	185bcde	245a	116f		
P	(ppm)					
Control	6.5c			_		
-1.20	5. 7cde	6.0cd	8.1b	4.8ef		
-1.70	5.4def	6.2cd	9.0b	4.4f		
-2.70	5.6cde	5.5cde	11.0a	5. 3de f		
Ca	(me/l)					
Control	26.1e					
-1.20	34.4d	36.3d	16.5f	48.6c		
-1.70	49.5c	48.3c	15. Of	70.1b		
-2.70	55.1 c	50.5c	14.2f	98.0a		
K	(me/l)					
Control	5.4f					
-1.20	6.8ef	7.6cdef	7.4def	8.2cde		
-1.70	11.4ab	10.0bcd	8.2cde	9.7bcd		
-2.70	13.0a	11.3ab	10.2bc	12.1ab		

z,y Same as Table 2.

Table 5-3. Chemical properties of soil solution (pF = to 3.8) at the end of the experiment.

Osmotic		Salin	nities		
potential (bars)	Sea water	NaCl	Na ₂ SO ₄	MgCl ₂	
EC	C (mS/cm)				
Control	5. 94 dy				
-1.20	11.58cd	11.90bcd	9.42cd	13.03bcd	
-1.70	17.91abcd	16.51abcd	15.86abcd	18.55abc	
-2.70	19.77abc	24.04ab	20.05abc	27.51a	
Os	motic poten	tial (bars)			
Control	- 2.73a				
-1.20	- 5.23b	- 7.20c	- 5.34b	- 6.25bc	
-1.70	-10.25ef	-11.39f	- 7.47cd	- 9.15de	
-2.70	-14.28g	$-11.74 \mathrm{f}$	-10.71ef	-15.19g	
p F	<u> </u>				
Control	6.69de				
-1.20	6.65 de f	6.90cd	7.28ab	6.72de	
-1.70	6.52ef	6.85d	7.39a	6.53ef	
-2.70	6.79d	6.70de	7.09bc	6.42f	

z,y Same as Table 2.

Osmotic	Amount of		EC	D	ry weight (g	()	Fr	uit
potential (bars)	$^{ m added}_{ m MgSO_4\cdot 7H_2O} \ ({ m mg}/l)$	(mS/cm)	Leaves	Stem	Plant top	Fresh weight (g)	Soluble ^x solids (%)	
Control ²	0	2.43	68.5ay	 15.9a	185. 2a	1,137a	14.8	
-0.95	1,922	3.20	68.1a	16. la	179.5a	1,076a	14.7	
-1.20	3,875	3.94	61.8a	14.6a	174.4a	887b	14.6	
-1.70	7,750	5. 52	53.1a	11.6b	142. 2b	777b	15.4	
-2.70	15,500	8.28	33.4b	8.1c	64.8c	408c	w	

Table 6. Effect of MgSO₄ salinity on growth of muskmelons in soil culture.

Table 7. Effect of MgSO₄ salinity on major elements in muskmelon leaves (% of dry matter).

Osmotic potential (bars)	Total-N	P	K	Ca	Mg	Na	SO ₄	Cl
Control	2.87b ^y	0.42b	1.68a	4.40a	1.85e	0.16a	3. 10b	0.79a
-0.95	2.86b	$0.46\mathbf{b}$	1.64a	3.47b	3.46d	0.08b	2.92b	0.74a
-1.20	2.76b	0.49b	1.35ab	2.80c	4.70c	0.07Ь	3.67b	0.64a
-1.70	3.05b	0.59a	0.87b	2.10d	5.47a	0.06Ъ	4.11b	0.81a
-2.70	3.98a	0.64a	1.00b	1.20e	5. 22b	0.08b	7.86a	0.78a

z,y Same as Table 2.

Experiment II

Growth (Table 6) Leaf dry weights at -0.95 to -1.70 bars were not significantly different from the control, but at -2.70 bars was much less. Stem and plant top dry weights were significantly reduced at -1.70 bars compared with the control. Fruit fresh weight decreased from -1.20 to -2.70 bars with decreasing osmotic potentials of treatment solutions. Visible symptoms of excess MgSO₄, such as interveinal necrosis and necrotic spots on leaves, were very slight at -1.20 bars, moderate at -1.70 bars and very severe at -2.70 bars. Most of the plants at -2.70 bars died by Nov. 12, about 60 days after transplanting.

Major mineral elements in leaves (Table 7) Mg increased, and Ca and K decreased as osmotic potentials of treatment solutions decreased. SO₄, total-N and P were high at lower (-1.70 and -2.70 bars) osmotic potentials of treatment solutions.

Chemical properties of SSo at the end of the experiment (Table 8) Mg and SO₄ concentrations and EC values increased and Na tended to decrease with decreasing osmotic potentials of treatment solutions. Ca and pH values were not affected by treatments.

Discussion

Fruit fresh weights at -2.70 bars were 61.3, 55.7, 63.2, 54.0 and 35.9% compared with the control in the sea water, NaCl, Na₂SO₄, MgCl₂ and MgSO₄ series, respectively. Whole plant and leaf dry weights showed similar tendencies. The estimated osmotic potential of treatment solutions which caused a 50% loss in fruit fresh weight was -3.5to -4.0 bars in the sea water, NaCl, Na₂SO₄ and MgCl₂ series, but it was -1.56 bars in the MgSO₄ series. The degree of tolerrance to salinity in sand culture was, on an average, almost half as compared with that in soil culture. Similar results were reported in green soybeans(12, 13). Hayward and Bernstein (5) pointed out that, in general, sand culture was effective in specification and other physiological studies, while soil culture was widely used for salt tolerance studies. In the previous sand(14) and present soil culture experiments, SSa (solution of sand medium) and SSo (solution of soil

z,y Same as Table 2.

^{*} Not subjected to statistical analysis.

w Plants died before harvest.

Osmotic potential (bars)	K (me/l)	Ca (me/l)	$\frac{\mathrm{Mg}}{(\mathrm{me}/l)}$	$Na \pmod{l}$	SO. (%)	EC (mS/cm)	pIH
Control ²	9.2b ^y	20.7a	22.4e	20.0b	0.26c	6.70d	5. 80a
-0.95	11.8a	21.0a	90.2d	25. 3a	0.64bc	10.38c	5. 51 a
-1.20	12.5a	20.9a	174.3c	23.4ab	0.91b	14.26b	5.51a
-1.70	12.6a	20.7a	262.0b	19.9b	1.97a	17.96a	5. 52a
-2.70	9.8b	19.8a	322.0a	13.8c	2.01a	19.52a	5.75a

Table 8. Chemical properties of soil solution (pF=0 to 3.8) at the end of the experiment.

medium) were centrifugally extracted at pF 0 to 3.8 to explain the difference in growth suppression between the two cultures. ions, anions and EC were lower in SSa, although growth suppression was more severe in sand culture. Therefore, as a preliminary experiment, SSa and SSo at various pF values were extracted from sand and soil used in the MgSO4 experiment in the previous sand and present soil cultures. Water content was 21.9, 7.1 and 1.1% in sand and 8.4, 11.7 and 23.3% in soil at pF 0 to 1. 8, 1. 8 to 3. 8 and more than 3. 8, respectively. EC values at pF 1.8 to 3.8 were 8.1, 10.8 and 13.8 mS/cm in sand culture and 8. 9, 16. 6 and 25. 7 mS/cm in soil culture at -0.70 (control), -1.20 and -2.70 bars, respectively, in the MgSO4 series. Though these values were much higher in soil than in sand culture, practically EC, cations and anions must be higher in sand than in soil culture. This is supported by the fact that Na, Mg and SO4 content in leaves was higher in sand than in soil culture.

Another pronounced difference between sand and soil cultures was in the Na2SO4 series where the growth in sand culture was suppressed to the same extent as in the MgSO4 series. In soil culture the growth suppression in the Na2SO4 series was not so severe as in the MgSO, series and was the same as in the sea water, NaCl and MgCl₂ series. Fruit fresh weights at -1.20, -1.70and -2.70 bars in the Na2SO4 series were 66. 4, 44. 8 and 11. 8%, and 82. 3, 69. 7 and 63.2% in sand and soil cultures, respectively, compared with the control. Na and SO4 in leaves were markedly higher in the Na₂SO₄ series than in the sea water and NaCl series. Also, Na in leaves was markedly higher in sand than in soil culture. Therefore, accumulated Na and SO₄ could be one of the causes for reduced growth in the Na₂SO₄ series in sand culture. In the nutrient solution culture experiment(15), the growth was suppressed in the Na₂SO₄ series to a lesser extent than in the MgCl₂ and MgSO₄ series.

According to some reports(9, 17, 18) the mechanism of salinity damage is different in chloride and sulfate types of salinity. The percentage of the pentose-phosphate pathway increased with increasing NaCl salinity while Na₂SO₄ salinity did not affect it(18). Higher concentrations of NaCl inhibited respiration in pea root tips(17) compared with Na₂SO₄. Therefore, in muskmelons, physiological responses may vary with salinities in spite of similar plant growth.

It has been observed by many investigators (1, 2, 3, 4, 6, 7, 8, 16) that different plant species exhibit different responses to isosmotic potentials of Cl and SO4, or Na and Mg. Some of them reported that sulfate-salinity was more toxic in sugarcane (7) and corn seedlings (8) than chloride-salinity, and that sulfate-salinity depressed growth in tomatoes more than chloride-salinity(12). The specific ion effect of Cl has been observed in green soybeans(12, 13) and honeylocust (1). Joolka et al. (6) found that grape cultivars differed in their responses to Cl and SO4 salts. The specific toxicity of Mg was stressed in red kidney beans(3), spinach, turnip, celery, Welsh onion, kidney beans and chard(16). Joshi and Naik(7) reported toxicity of Na to be greater than that of Mg. The result of this experiment revealed MgSO4 salinity to be more toxic

z.y Same as Table 2.

to muskmelons than the other salinities. This confirms the specific ion effect of Mg and SO₄ in muskmelons. Reduced growth in the MgSO₄ series compared with the Na₂SO₄ and MgCl₂ series seems to be due to synergistic effect of Mg and SO₄ ions, higher ion concentration in the medium and higher ion uptake by plants.

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各種塩類が土耕におけるメロンの耐塩性に及ぼす影響

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摘 要

メロンの耐塩性と塩の種類との関係を明らかにするため、土耕により本実験を行った。塩類源として実験 I では、海水、NaCl、Na₂SO₄、MgCl₂を、実験 II では、MgSO₄を用いて、浸透ボテンシャルをそれぞれ -0.95(MgSO₄のみ)、-1.20、-1.70、-2.70 bar とし、基本培養液で育てた対照区(-0.70 bar)と生育を比較した。果実新鮮重と全植物体乾物重は、対照区で最大となり、処理培養液の浸透ボテンシャルが低下するにつれて、それぞれの塩類源で減少した。対照区を 100% とした場合、-2.70 bar 区の果実新鮮重は、海水で 61.3%、NaCl で 55.7%、Na₂SO₄ で 63.2%,MgCl₂ で 54.0

%、 $MgSO_4$ で 35.9% となった。 $MgSO_4$ では、定植後 60 日までに大部分の株が枯死した。その他の塩類では枯死した株はなかった。本実験における生育は、対照区で最大となり、次に海水、NaCl, Na_2SO_4 , $MgCl_2$ で同程度に抑制され、 $MgSO_4$ で最も抑制された。葉及び土壌溶液中の Na, Mg, Cl, SO_4 含量は,それぞれの処理区で処理培養液の浸透ボテンシャルが低下するにつれて増加する傾向を示した。処理培養液の浸透ボテンシャルが低下すにつれ、土壌溶液の EC 値は増加し、浸透ボテンシャルは減少した。