

Field Screening of Sugarcane (*Saccharum* spp.) Mutant and Commercial Genotypes for Salt Tolerance

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Abstract

Growth and physiological attributes and sugar quality parameters are considered key criteria for screening sugarcane cultivars for salt tolerance. Maximum cane growth and yield were found in a positive check ('K88-92') as well as in cv. '(A3)AE1-18' when subjected to salt affected soil. Percent reduction in F_v/F_m , quantum efficiency of PSII (Φ_{PSII}) and water use efficiency (WUE) due to salt stress was considerably low in 'K88-92', '(A3)AE1-18' and 'KK3' which was associated with very low salt-induced reduction in net photosynthetic rate and growth characters such as shoot length, number of internodes, and internodal length as well as yield traits. In addition, brix, polarization, fiber, purity and commercial cane sugar (CCS) in '(A18)AE2-15' and '(A3)AE1-18' were well maintained under saline stress. By subjecting the data for various physiological, growth, yield and sugar quality parameters to the Ward's cluster analysis 'K88-92' (positive check), '(A3)AE1-18' and 'KK3' were identified as salt tolerant, whereas '(A11)AE1-114' and 'K97-32' as salt sensitive.

Keywords: cluster analysis, net photosynthetic rate, sugar qualities, water use efficiency, yield traits

Introduction

Sugarcane (*Saccharum* sp.) is one of the key commercial crops of the tropical and subtropical zones of the world. However, its yield is reduced to about 50% when the crop is grown on salt affected soils (Wiedenfeld, 2008). Plant physiological and biochemical responses in sugarcane grown under salt stress have been well documented. For examples, negative effects of saline stress on a number of sugarcane growth and physiological characteristics such as water potential, membrane integrity, nutritional balance, photosynthetic pigments, chlorophyll fluorescence, net photosynthetic rate, growth and yield have been reported (Cha-um and Kirdmanee, 2009; Gomathi and Thandapani, 2004, 2005). However, a great magnitude of inter-cultivar variation for salt tolerance is reported to occur in the sugarcane germplasm. For example, 'Co 85019', 'Co 94012', 'Co 94008', 'Co 86032', 'Co 92038', 'Co 85004', 'CPF-213', 'BO91', 'Co 1158', 'CoSe 92423', 'CP-4333', 'S-86-US-699', 'CP66-346' and 'H69-8235' (commercial cultivars and breeding lines) have been categorized as salt tolerant and 'CP-71-3002', 'H65-7052', 'HSF-240', 'L-116', 'Si 940-50', 'Co 85036', 'Co 97010', 'Co 95007', 'Co 97009' and 'Co 95016' as salt sensitive based on a variety of phenotypic indices (Gandonou *et al.*, 2011; Goma-

thi and Thandapani, 2004; Plaut *et al.*, 2000). However, most of the screening studies have been carried out under laboratory or greenhouse conditions. Thus, it is not sure whether degree of salt tolerance of different cultivars observed under controlled or semi-controlled conditions would be maintained under natural field conditions as the latter growth habitat is embraced with a variety of atmospheric and edaphic factors which in combination may enhance the adverse effects of a single stress. Thus, the principal aim of the present investigation was to classify some elite sugarcane genotypes in terms of their tolerance to salt stress imposed by a natural salt affected field, using some key physiological, growth, yield and quality attributes.

Materials and methods

Plant materials and salt affected treatment

Six mutant cultivars of sugarcane, '(A3)AE1-17', '(A9)AE1-103', '(A11)AE1-114', '(A13)AE1-126', '(A18)AE2-15' and 'D14' derived from γ -irradiation and ethyl methane sulfonate (EMS), and 8 commercial cultivars, i.e., 'LK92-11' (wild type), 'K95-84', 'K93-219', 'K84-200', 'K97-32', 'Mitr Phol 3', 'KK3' including a positive check ('K88-92') were propagated in MS (Murashige and Skoog, 1962) supplemented with 8.88 μ M benzyl ad-

enine (BA), 3% sucrose and 0.25% Phytigel[†] for 6 weeks. Single shoots were excised and roots induced on the MS medium supplemented with 2.46 μM indole butyric acid (IBA) for 2 weeks. Plantlets were cultured *in vitro* under $25\pm 2^\circ\text{C}$ ambient temperature, $60\pm 5\%$ relative humidity (RH), and $60\pm 5 \mu\text{mol m}^{-2} \text{s}^{-1}$ photosynthetic photon flux density (PPFD) provided by fluorescent lamps with a 16 h d^{-1} photoperiod. Then, the sugarcane plantlets were transferred to MS sugar-free liquid medium using vermiculite as supporting material. The number of air exchanges in the plastic chamber was adjusted to $5.13 \mu\text{mol CO}_2 \text{h}^{-1}$ by punching 32 holes on sideward of the plastic chamber (\varnothing 1 cm) and covering the holes with microporous filters (0.2 μm of pore size). For acclimatization, the plantlets were subsequently cultured in a plant growth incubator under $25\pm 2^\circ\text{C}$ ambient temperature, $60\pm 5\%$ RH, and $120\pm 5 \mu\text{mol m}^{-2} \text{s}^{-1}$ PPFD provided by fluorescent lamps with a 16 h d^{-1} photoperiod and CO_2 enrichment at $1,000\pm 100 \mu\text{mol mol}^{-1}$ (Cha-um *et al.*, 2003). Plantlets of 14 sugarcane cultivars were directly transferred to plastic bags containing clay soil ($\text{EC} = 2.687 \text{ dS m}^{-1}$; $\text{pH} = 5.5$; organic matter = 10.36%; total nitrogen = 0.17%; total phosphorus = 0.07%; total potassium = 1.19%) in 50% light intensity in a greenhouse for 1 month. Irrigation was applied as water spray. The acclimatized plants were directly transplanted to a field (30 cm plant to plant and 150 cm row to row distance) at two sites including clay soil (control) and salt affected soil (Tab. 1) at Chaiyaphum province, North-east of Thailand (Latitude $16^\circ 35' \text{N}$ and Longitude $101^\circ 55' \text{E}$) for 10 months. Chemical fertilizer (16:16:16; nitrogen: phosphorus: potassium) was applied three times, i.e., February, May and August at 0.0156 kg m^{-2} prior to stalk harvesting in November 2011. Data for plant height, stalk weight, number of stalks per plot, and total stalk weight per plot of sugarcane cultivars were recorded. Photosynthetic abilities, water use efficiency, growth characters, yield attributes, and sugar qualities were measured at the harvesting stage.

Data collection

Chlorophyll fluorescence parameters such as maximum quantum yield of PSII (F_v/F_m), photon yield of PSII (Φ_{PSII}), photochemical quenching (qP), and non-photochemical quenching (NPQ) from the youngest fully grown leaf (adaxial surface) from each plant were appraised using a fluorescence meter (FMS2, Hansatech Instrument Ltd., Norfolk, UK) following Loggini *et al.* (1999). Initial (F_0)

and maximum (F_m) fluorescence were measured on a leaf adapted to dark conditions for 30 min. The variable fluorescence yield (F_v) was calculated using an equation ($F_m - F_0$) following Maxwell and Johnson (2000).

Net photosynthetic rate (P_n ; $\mu\text{mol m}^{-2} \text{s}^{-1}$), transpiration rate (E ; $\text{mmol H}_2\text{O m}^{-2} \text{s}^{-1}$) and water use efficiency (WUE; $\mu\text{mol CO}_2 \text{mmol}^{-1} \text{H}_2\text{O}$) were measured using a Portable Photosynthesis System (Model LI 6400, LI-COR[®] Inc, Lincoln, Nebraska, USA) with an Infra-red Gas Analyser following Cha-um *et al.* (2007). Photosynthetic WUE was calculated using the equation: $\text{WUE} = P_n/E$.

Shoot height, number of internodes, intermodal length, single stalk weight, and stalk yield were measured at the harvesting stage (10 months after planting). Brix percentage was measured using a hand-held refractometer (model H-50, ATAGO, Tokyo, Japan). Polarization percentage and fiber percentage were also assayed. Purity percentage was calculated according to equation: Purity (%) = [Sucrose percentage/Brix percentage] \times 100. In addition, commercial cane sugar (CCS) was calculated following the equation: $\text{CCS} (\%) = [1.05 \times \text{Sucrose percentage}] - [0.292 \times \text{Brix percentage}]$ (Meade and Chen, 1977).

Experiment design and statistical analysis

The experiment was arranged as 14×2 factorials in a Completely Randomized Block Design (CRBD) with eight replicates ($n = 8$). Significant differences among the mean values were worked out using the Duncan's New DMRT and analyzed with the SPSS software. Data for all variables were subjected to the Ward's method of Hierarchical cluster analysis in SPSS software to classify the cultivars into salt tolerant and salt sensitive categories.

Results and discussion

Growth performances and yield traits

Almost all growth attributes measured in the sugarcane cultivars examined in the present investigation were severely affected by salt stress. There was a marked reduction shoot length in all cultivars, but the cultivars differed significantly in this attribute. Salt-induced reduction in shoot length was significantly lower in the positive check 'K88-92' (4.90% reduction) than that in the other cultivars, e.g., 14.55% in 'K95-84' and 41.41% in '(A11)AE1-114' (Tab. 2 and Fig. 1A). The cultivars also differed significantly for

Tab. 1. Soil electrical conductivity (EC_e), pH, organic matter (OM), available phosphorus (P), potassium (K), sodium (Na) and chloride (Cl) in the non-saline (control) and salt affected soils at two depths, 0-30 cm and 30-60 cm

Site	Depth (cm)	EC_e (dS m^{-1})	pH	OM (%)	P (ppm)	K (ppm)	Na (ppm)	Cl (ppm)
Control	0-30	9.37	8.07	3.05	1884	279	594	778
	30-60	6.18	8.23	1.14	504	167	530	725
Salt affected soil	0-30	37.43	6.27	0.97	460	116	1348	4134
	30-60	25.20	5.99	1.03	312	98	1061	3221

number of internodes under saline stress, e.g. number of internodes reduced to 1.19% in 'K95-84' and 24.53% in '(A18)AE2-15' under saline regime (Tab. 2). Internodal length in the salt stressed plants of all cultivars decreased significantly when compared to that in controlled plants varying from 29.41% reduction in '(A11)AE1-114' to 57.97% in 'K84-200' (Tab. 2 and Fig. 1B). Single stalk weight in each cultivar decreased significantly due to salt stress, however, there had been considerable variation for salt tolerance in the sugarcane germplasm examined in the present study in terms of single stalk weight (Tab. 2 and Fig. 1C). Yield of sugarcane per sprout in salt stressed plants of all cultivars decreased significantly varying from 22.62% reduction in 'K95-84' to 75.64% in 'K97-32' (Tab. 2 and Fig. 1D). Growth parameters are simple, rapid and sensitive method to assay when sugarcane exposed to salt stress. For example, shoot height parameter of sugarcane pot culture was reduced for 8.8% in salt stressed 'CoS 03261' (8 dS m⁻¹ EC_e) for 150 days. Leaf area reduction in 'CoS 95-255' and 'CoSe 96436' was 5.9% and 8.6% respectively. In addition, the lowest reduction percentage of dry matter in

'CoS 07250' was identified as salt tolerant variety (Saxena *et al.*, 2010). From this it is evident that only one parameter may not be suitable to classify sugarcane cultivars/lines with regard to degree of salt tolerance. For example, stalk girth, internodal length, and yield of salt tolerant sugarcane cultivars 'CP43-33' and 'CPF-213' were higher than those in salt susceptible cv. 'L116' when subjected to varying levels of saline stress for 280 days (Akhtar *et al.*, 2001). In the same study, reduction in stalk yield and brix percentage in salt tolerant cv. 'CP43-33' was reported to be significantly lower (1.88% and 20.9% reduction) than that in salt sensitive 'CP-71-3002' (75.30% and 38.75%) under 120 mM NaCl. Thus, variation in salt tolerance in sugarcane germplasm in terms of different growth parameters exists and is variable to a great extent depending on the type of cultivars.

Alteration in physiological characteristics

Maximum quantum yield of PSII (F_v/F_m) in the salt stressed plants of all sugarcane cultivars decreased significantly except in the positive check ('K88-92') and '(A3)

Tab. 2. Shoot length, number of internodes, internodal length, single stalk weight and stalk yield traits in 14 sugarcane genotypes grown under non-saline (control) and salt-affected soils for 10 months

Genotypes	Soil	Shoot length (cm)	Number of internodes	Internodal length (cm)	Single stalk weight (kg cane ⁻¹)	Stalk yield (kg sprout ⁻¹)
'LK92-11'	Control	228.3ab	23.0a	9.85b	1.75bc	24.23d
	salt-affected	186.5ghi	16.0bc	7.63c	1.30d	16.71fg
'K95-84'	control	226.8abc	21.0a	10.08b	2.15ab	33.68ab
	salt-affected	193.8efg	20.8ab	6.40d	1.85b	26.06cd
'K93-219'	control	169.3ijk	17.0b	15.69a	1.05e	13.26ghi
	salt-affected	125.0o	9.5d	9.50b	0.85f	10.60ij
'D14'	control	213.5bcd	24.8a	10.62b	2.05b	36.37a
	salt-affected	165.3jkl	14.8c	10.11b	1.75bc	24.51d
'K84-200'	control	161.8jkl	16.0bc	14.75a	0.93f	12.90fgh
	salt-affected	131.0o	10.0d	6.20d	0.55g	8.08i
'A9'	control	171.0hij	25.8a	13.18ab	1.35d	14.28fgh
	salt-affected	142.0mno	13.3cd	10.16b	0.90f	8.80i
'K97-32'	control	224.3abc	24.0a	10.56b	2.50a	36.62a
	salt-affected	138.0no	15.0bc	7.31c	0.90f	8.92i
'A18'	control	179.3ghi	13.3cd	11.53b	1.05e	11.83ghi
	salt-affected	132.3o	10.0d	7.25c	0.95f	7.96i
'A13'	control	231.5a	25.3a	8.56bc	2.10ab	26.80cd
	salt-affected	193.3efg	20.0ab	6.58d	1.70bc	18.18ef
'Mitr Phol 3'	control	206.0def	18.3b	13.13ab	1.80bc	25.04cd
	salt-affected	154.8klm	11.8cd	6.75d	1.05e	17.85ef
'A11'	control	225.3abc	17.5b	10.93b	1.90b	23.71d
	salt-affected	131.8o	9.8d	7.71c	0.70f	7.72i
'K88-92'	control	214.3bcd	20.3ab	9.13b	2.08b	29.85bc
	salt-affected	203.8def	16.8b	6.35d	1.80bc	22.54d
'KK3'	control	184.5fgh	15.5bc	13.89ab	1.90b	23.30d
	salt-affected	147.0mno	10.5d	10.29b	0.90f	16.42fg
'A3'	control	205.3def	18.7b	11.96b	1.40cd	17.67ef
	salt-affected	173.8ghi	12.8c	8.63bc	1.15e	12.94ghi

Different letters in each column show significant difference at $p \leq 0.01$ by the Duncan's New Multiple Range Test (DMRT)

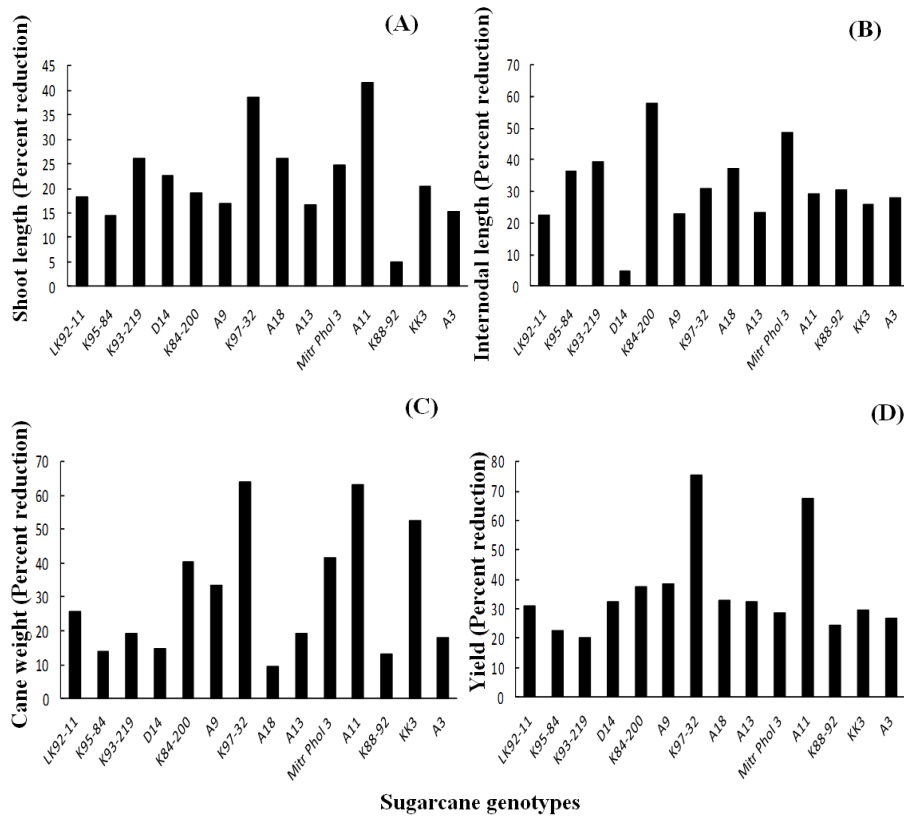


Fig. 1. Percent reduction in shoot length(A), internodal length (B), cane weight (C), and yield (D) in 14 sugarcane genotypes grown under non-saline (control) and salt-affected soils for 10 months

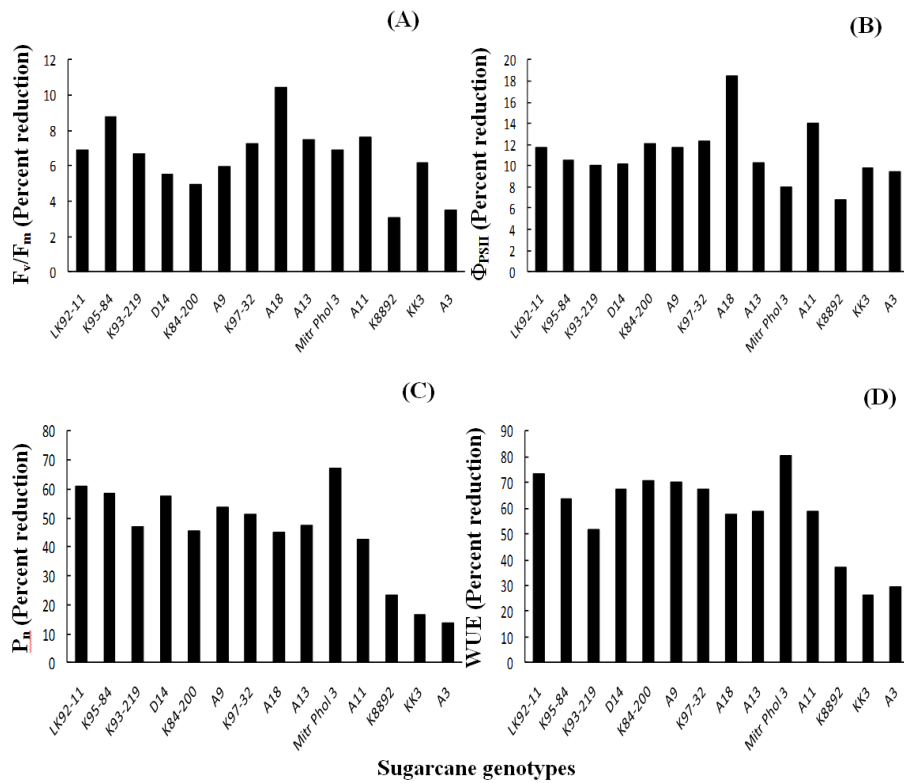


Fig. 2. Percent reduction in F_v/F_m (A), Φ_{PSII} (B), P_n (C), and WUE (D) in 14 sugarcane genotypes grown under non-saline (control) and salt-affected soils for 10 months

Tab. 3. Maximum quantum yield of PSII (F_v/F_m), photon yield of PSII (Φ_{PSII}), net photosynthetic rate (P_n) and water use efficiency (WUE) in 14 sugarcane genotypes grown under non-saline (control) and salt-affected soils for 10 months

Genotypes	Soil	F_v/F_m	Φ_{PSII}	P_n ($\mu\text{mol m}^{-2} \text{s}^{-1}$)	WUE ($\mu\text{mol CO}_2 \text{mmol}^{-1} \text{H}_2\text{O}$)
'LK92-11'	control	0.893ab	0.829a	74.1a	123.5cde
	salt-affected	0.832efg	0.732bc	29.0hi	32.6kl
'K95-84'	control	0.891ab	0.806a	74.7a	107.4fgh
	salt-affected	0.813fg	0.721bc	31.2gh	39.3jkl
'K93-219'	control	0.899ab	0.823a	74.2a	108.7fgh
	salt-affected	0.839def	0.741bc	39.5de	52.8j
'D14'	control	0.905a	0.808a	71.8a	113.0fgh
	salt-affected	0.855cde	0.726bc	30.7gh	36.6jkl
'K84-200'	control	0.882abc	0.814a	75.0a	145.8a
	salt-affected	0.839def	0.716bcd	41.0d	42.6jkl
'A9'	control	0.891ab	0.816a	74.6a	135.8abc
	salt-affected	0.838def	0.720bc	34.6fg	40.4jkl
'K97-32'	control	0.905a	0.840a	74.0a	130.6abc
	salt-affected	0.839def	0.736bc	36.0ef	42.9jkl
'A18'	control	0.896ab	0.830a	72.1a	103.0gh
	salt-affected	0.802f	0.677d	39.7de	43.6jkl
'A13'	control	0.893ab	0.804a	75.4a	116.6efg
	salt-affected	0.826efg	0.721bc	39.6de	48.1jk
'Mitr Phol 3'	control	0.897ab	0.817a	75.0a	144.9a
	salt-affected	0.835efg	0.751b	24.6i	28.4l
'A11'	control	0.897ab	0.811a	75.4a	125.9cde
	salt-affected	0.829efg	0.697cd	43.2d	51.9j
'K88-92'	control	0.893ab	0.817a	71.3a	116.9efg
	salt-affected	0.866bcd	0.761b	58.4c	73.4i
'KK3'	control	0.903ab	0.823a	74.9a	141.6ab
	salt-affected	0.847def	0.743b	62.5b	104.8gh
'A3'	control	0.886abc	0.823a	74.2a	138.2abc
	salt-affected	0.855cde	0.746b	64.0b	97.4h

Different letters in each column show significant difference at $p \leq 0.01$ by the Duncan's New Multiple Range Test (DMRT)

AE1-17' in which reduction in this trait had been only 3.02% and 3.50%, respectively. In contrast, percent reduction in F_v/F_m in the other sugarcane cultivars ranged from 4.95% in 'K84-200' to 10.46% in '(A18)AE2-15' (Fig. 2A). Photon yield of PSII (Φ_{PSII}), net photosynthetic rate (P_n) and water use efficiency ($WUE = P_n/E$) in all sugarcane cultivars declined significantly under saline stress (Tab. 3). Leaf P_n and WUE were found to be very sensitive parameters in sugarcane cultivars subjected to salt affected soil, especially in salt sensitive cultivars. The Φ_{PSII} in the salt stressed sugarcane plants decreased significantly, e.g., 8.04% reduction in 'Mitr Phol 3', 6.78% in 'K88-92' (6.78%), 9.76% in 'KK3' (9.76%) and 9.39% in '(A3)AE1-17' (Fig. 1B). Similarly, percent reduction in P_n salt stressed plants of 'K88-92', 'KK3' and '(A3)AE1-17' was 23.13%, 16.53% and 13.71%, respectively (Fig. 1C). Also, WUE in the salt stressed plants of 'K88-92' (37.20%), 'KK3' (26.01%) and '(A3)AE1-17' (29.51%) was 37.2%, 26.01%, and 29.51%, respectively (Fig. 1D). In the present study, P_n (42.69-67.23%) and WUE (51.43-80.43%) in salt sensitive cultivars were also sensitive than F_v/F_m (4.95-10.46%) and Φ_{PSII} (10.06-18.51%). *In vitro*

salt tolerant screening, F_v/F_m , Φ_{PSII} , quantum efficiency of PSII (qP) and P_n in salt stressed plantlets (200 mM NaCl) were sharply decreased. Only Φ_{PSII} and qP in salt stressed 'K88-1' or 'K88-92' cultivar were maintained at <10% diminishing (Cha-um *et al.*, 2012). Also, the salt tolerant cultivar, 'K88-92' was applied as positive check in the field trial. The photosynthetic abilities in salt stressed plants have been potentially classified the salt tolerance in sugarcane. For example, diminishing of F_v/F_m in salt tolerant genotypes, 'C 92038' (9.28%) and 'Co 85004' (6.85%) of sugarcane in mature and reproductive stage was lower than that in salt sensitive, 'Si 94050' (17.35%) and 'Co 85036' (22.46%) when subjected to 1% NaCl (EC_e 7 dS m^{-1}) (Gomathi *et al.*, 2010; Gomathi and Rakkiyapan, 2011). Vasantha *et al.* (2010) reported that the P_n in 8 sugarcane cultivars declined significantly but to a varying extent when treated with salt stress of EC_e 8 dS m^{-1} . Percent reduction in P_n cvs. 'Co 94012' and 'Co 94008' had been 4.57% and 4.56%, respectively, whereas that in other cultivars had been from 11.57% in 'Co 86032' to 46.25% in 'Co 97009' (Vasantha *et al.*, 2010). It has been reported

that the decline in P_n depends on salt tolerance abilities, level of salt stress and period of salt treatment.

Quality of sugarcane and cluster analysis

Brix percentage of five sugarcane cultivars, 'K93-219', 'K84-200', '(A9)AE1-103', 'K97-32' and 'KK3' significantly decreased when plants were subjected to salt affected soil, i.e., varying from 17.49% in 'K84-200' to 26.58% in 'KK3', whereas that of other cultivars remained almost unchanged (Tab. 4 and Fig. 3). Polarization, fiber, purity and commercial cane sugar (CCS) percentages in salt stressed plants of all sugarcane cultivars also decreased significantly, especially in '(A13)AE1-126' (34.53%, 16.26%, 11.56% and 38.0% reduction, respectively), and 'KK3' (36.79%, 35.12%, 16.42% and 41.19%, respectively) (Tab. 4 and Fig. 3). In contrast, the reduction in these quality parameters in the positive salt tolerant check 'K88-92' was relatively lower, i.e., 13.57%, 10.14%, 3.88% and 14.33% reduction, respectively. The data for physiological attributes, plant growth, yield potential and quality traits of sugarcane genotypes were subjected to the multivariate

cluster analysis, through which it was possible to identify salt tolerant cluster as '(A3)AE1-11' and 'KK3' where in the positive check 'K88-92' was also present (Fig. 4). Also, '(A11)AE1-114' was classified as salt susceptible where in the negative check 'K97-32' was presented. Lingle and Wiegand (1997) have reported that brix, polarization and purity percentage of sugarcane cultivar 'CP 70-321' declined depending on soil salinity of a salt affected field (2-10 dS m⁻¹ EC_e). Also, brix, sucrose, purity and CCS in salt tolerant cultivars, 'C 92038' and 'Co 85004', have been reported to be maintained better than those in salt susceptible cultivars, 'Si 94050' and 'Co 85036' (Gomathi and Thandapani, 2005). In a previous study, the multivariate cluster analysis (Ward's method) has been found to be very effective in screening some commercial sugarcane cultivars for salt tolerance under controlled conditions (Cha-um *et al.*, 2012). However, in the recent report, changes in physiological and growth characters, yield attributes and quality of sugarcane were evaluated in a salt affected field.

In conclusion, salinity-induced decline in chlorophyll fluorescence, P_n , growth, yield and quality in salt stressed

Tab. 4. Brix percentage, polarization percentage, fiber percentage, purity and commercial cane sugar (CCS) traits in 14 sugarcane genotypes grown under non-saline (control) and salt-affected soils for 10 months

Genotypes	Soil	Brix (%)	Polarization (%)	Fiber (%)	Purity (%)	CCS (%)
'LK92-11'	Control	21.85a	13.21bc	12.45b	80.86bc	9.69bc
	salt-affected	20.30a	9.41d	10.39c	71.84c	6.12cd
'K95-84'	Control	19.20ab	16.70b	14.72a	85.89a	12.11a
	salt-affected	16.45bc	15.25b	11.85b	84.45a	11.33ab
'K93-219'	Control	18.45b	13.67bc	11.96b	80.23bc	9.87bc
	salt-affected	14.40c	10.87cd	11.26bc	74.99c	7.38c
'D14'	Control	20.75a	14.97bc	12.69b	83.07a	10.91b
	salt-affected	17.75bc	12.06c	12.52b	77.49bc	8.33c
'K84-200'	Control	19.15ab	10.56cd	12.05b	75.31c	7.33c
	salt-affected	15.80c	8.77d	10.33c	70.23c	5.61d
'A9'	Control	21.10a	15.28b	12.45b	86.30a	11.32ab
	salt-affected	16.65bc	12.82c	12.03b	84.30a	9.59bc
'K97-32'	Control	18.20b	12.76c	11.29bc	81.00b	9.29bc
	salt-affected	13.90c	9.84d	11.19bc	71.65c	6.47cd
'A18'	Control	19.00ab	13.42bc	11.55bc	81.35b	9.80bc
	salt-affected	16.75bc	12.01c	11.13bc	77.11bc	8.38c
'A13'	Control	21.10a	18.45a	14.76a	89.30a	13.71a
	salt-affected	19.40ab	12.08c	12.36b	78.98bc	8.50c
'Mitr Phol 3'	Control	19.10ab	15.55b	10.87bc	85.84a	11.94a
	salt-affected	16.55bc	13.37bc	10.06c	81.34b	9.80bc
'A11'	Control	18.55b	13.65bc	10.62bc	81.24b	10.02b
	salt-affected	16.60bc	10.92cd	10.35c	75.96c	7.64c
'K88-92'	Control	16.35bc	16.88b	13.31ab	87.74a	12.63a
	salt-affected	14.65c	14.59bc	11.99b	84.34a	10.82b
'KK3'	Control	18.25b	14.65bc	15.86a	84.95a	10.39b
	salt-affected	13.40c	9.26d	10.29c	71.00c	6.11cd
'A3'	Control	22.50a	14.51bc	13.68ab	83.67ab	10.61b
	salt-affected	19.50ab	14.01bc	12.67b	82.87ab	10.04b

Different letters in each column show significant difference at $p \leq 0.01$ by the Duncan's New Multiple Range Test (DMRT)

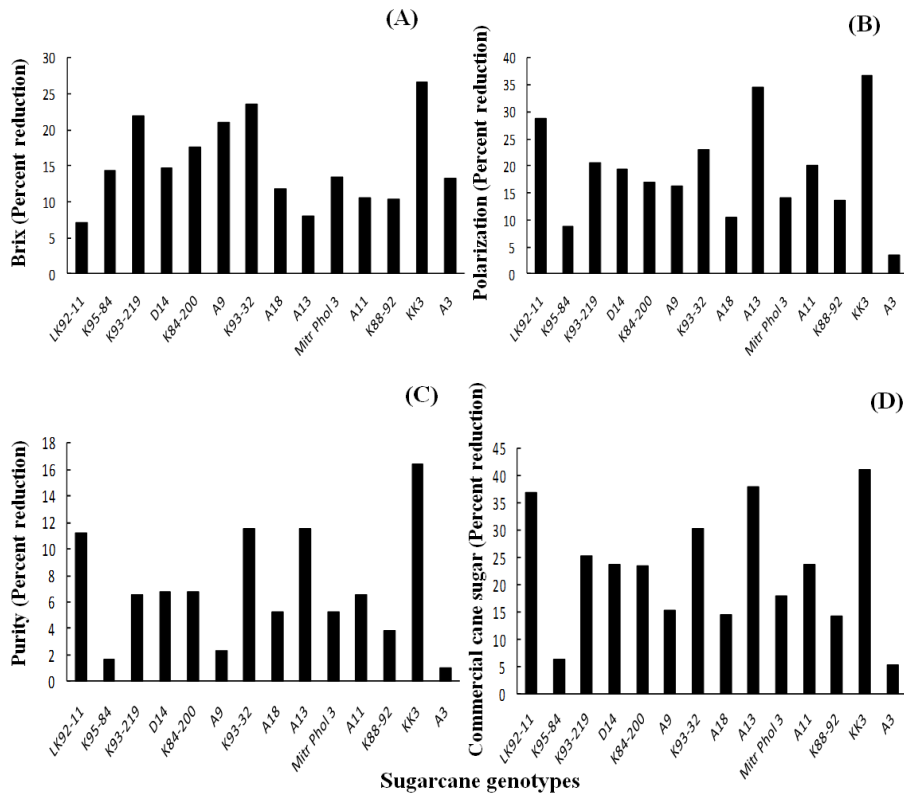


Fig. 3. Percent reduction in brix (A), polarization (B), purity (C), and commercial cane sugar (D) in 14 sugarcane genotypes grown under non-saline (control) and salt-affected soils for 10 months

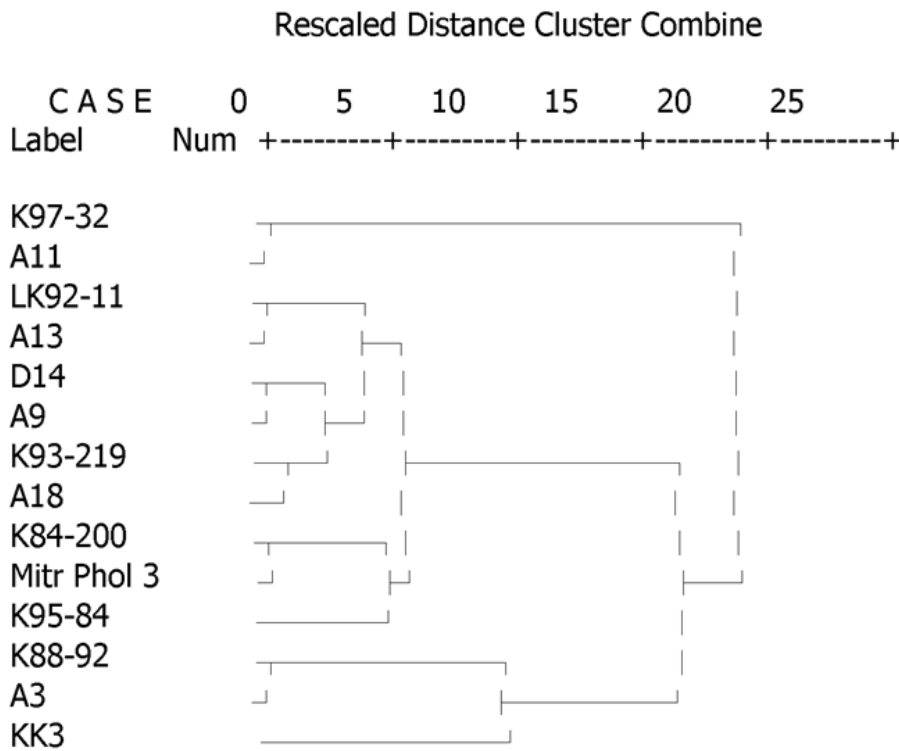


Fig. 4. Ward's dendrogram showing the classification of mutant sugarcane genotypes based on data for photosynthetic abilities, growth performances, yield and sugar quality traits (salt sensitive, 'A11' and 'K97-32', and salt tolerant, 'A3', 'KK3' and 'K88-92')

plants of sugarcane genotypes was evident. Multivariate analysis of data for all parameters allowed discriminating among the sugarcane cultivars for salt tolerance. Sugarcane cultivars '(A3)AE1-11' and 'KK3' were identified as salt tolerant using the Ward's cluster analysis as was the positive control 'K88-92'.

Acknowledgments

The authors wish to thank the Mitr Phol Co. Ltd. as a funding source and Dr. Upsorn Pliansinchai, Mitr Phol Co. Ltd. for providing a basic facility of salt affected field.

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