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Determination of Combining Ability and Genetic Diversity Using ISSR Markers to Evaluate the Genetic Variability in Wheat

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Key words:

Wheat, ISSR marker, combining ability, drought stress, GCA and SCA.

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ABSTRACT

Eight bread wheat genotypes were screened by the six ISSR primers and scored a total of 54 amplified DNA bands. The number of bands varied from 40 to 69 bands form primers ISSR 2 (6 fragments) to ISSR 5 (13 fragment). Number of polymorphic fragments varied from 1 for ISSR 4 to 8 for ISSR 1 and ISSR 5. The six ISSR primers generated 314 scorable bands across 8 studied genotypes. These primers produced a total of 54 reproducible fragments, from which 30 (55.55%) were polymorphic. The lowest genetic similarity (0.63) was obtained between P3 and P7, while, the highest genetic similarity (0.89) was scored between P7 and P8. The parents were crossed in a 8x8 half diallel scheme in 2015/2016. Parents and their 28 F1 crosses were evaluated under normal and stress conditions during 2016/2017 in two irrigation levels experiments. The mean squares were significant for all studied traits. The highest mean values were detected by parents P2, P2, P8, P2, P6 and p2 for plant height, spike length, no of spike/ plant, 1000-kernel weight, biological yield/ plant and grain yield/ plant in the combined analysis, respectively. While, the highest mean values were recorded under combined analysis with crosses P1xP6 for biological yield/ plant and the cross P2xP4 for grain yield / plant. Mean squares for combining ability estimates were highly significant for all studied traits. The ratios GCA /SCA exceeded the unity for all studied traits, revealing that additive and additive x additive types of gene action are more important than non-additive gene action in controlling these traits. The parental P6 exhibited positive and significant ĝi effects for grain yield/ plant and its attribute. The highest desirable SCA effects were obtained with the crosses P1xP6, P1xP7, P2xP4, P3xP7, P3xP8, P4xP8, P5xP8 and P6xP7 for grain yield/ plant which exhibited significant and positive ŝij effects. Positive correlation coefficients were found between genetic diversity and each of mean performance and SCA for grain yield/plant. Hence, ISSR marker can be used as a tool for determining the extent of genetic diversity among wheat genotypes and can be used to precisely predict the yield performance value for F₁ hybrids.

تحديد القدرة على التألف و التباعد الوراثي باستخدام تكنيك ISSR لتقييم التنوع الوراثي في القمح

عمار وبدان السعدون¹ وعلي عبد المقصود الحصري²وسيدهم أسعد سيدهم² ومحمود الزعبلاوى البدوى² واحمد على الحصرى² ^اقسم المحاصيل – كلية الزراعة – جامعة تكريت – العراق ²قسم المحاصيل – كلية الزراعة – جامعة بنها – مصر

الكلمات المفتاحية: استخدمت ستة بوادىء ISSR على الثمانية تراكيب وراثية تحت التجربة. أظهرت النتائج أن البادىء القدرة على الثانف، التاف، التباعد الوراثي، ISSR 2 حقق أعلى عدد من الحزم في جميع الاباء تحت الدراسة. بينما أظهر البادىء ISSR 5 أقل القدرة على التذرة على الترامية. الترامية. بينما أظهر البادىء ISSR 5 أقل القدرة على التذرة على الترامية. الترامية. الترامي، القهر البادىء ISSR 5 أقل العدرة على التذرة على المغتامية الاباء تحت الدراسة. بينما أظهرت النتائج أن البادىء ISSR 5 أقل القدرة على التألف، التباعد الوراثي، القمح. عدد من الحزم في جميع الاباء تحت الدراسة. بينما أظهر البادىء ISSR 5 أقل تكنيك ISSR 1 ألمي التذرة على المغامية المغامية الترامية. الترامية تحت الدرامية القلم البادىء ISSR 5 أول العدم الترامية الترامية القلم المغامية المغامية المنابية المعامية الترامية الترامية المغامية المغامية المعامية المغامية المعامية الترامية المعامية المغام تكنيك ISSR 1، التنوع الوراثي، القمح. عدد من الحزم. تراوح العدد الكلى لشظايا الـ ISR (ISR المختلفية (Polymorphi المرامية الترامية الترامية البادىء ISSR 1. تراوح العدد الكلى المطامية الم المختلفة (

¹ This paper is a part of MSc. Thesis for the first author

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fragments) من 1 في الباديء ISSR 4 الى 8 في الباديء ISSR 1,5. كان العدد الكلى لشظايا الـ DNA المختلفة للاباء الثمانية 30 بنسبة 55,55%.

alsadoonammar@yahoo.com و PR اوضح التحليل العنقودى Cluster analysis ان هناك علاقات وراثية للتماثل بين الاباء الثمانية و PR. اوضح التحليل العنقودى Cluster analysis ان هناك علاقات وراثية للتماثل بين الاباء الثمانية تحت الدراسة حيث تم تقسيم الاباء الى قسمين رئيسيين Two main cluster. أحتوى الاول على الاب رقم 2 اما الثانى احتوى على باقي الاباء. ولدراسة قوة الهجين والقدرة على التألف لصفات المحصول ومكوناته لثمانية آباء من القمح بالأضافة إلى 28 هجين ناتجة منها بنظام Half diallel وذلك في محطة تجارب بحوث كلية زراعة مشتهر جامعة بنها، حيث تم عمل تجربتين بمزرعة الكلية. في التجربة الأولي تم الري مرة واحدة بعد رية الزراعة مينها، حيث تم عمل تجربتين بمزرعة الكلية. في الجربة الأولي تم عشرة نباتات فردية أخذت عشوائيا من كل قطعة تجريبية وقدرت قوة الهجين لكافة الصفات المدروسة عشرة نباتات فردية أخذت عشوائيا من كل قطعة تجريبية وقدرت قوة الهجين لكافة الصفات المدروسة عشرة نباتات فردية أخذت عشوائيا من كل قطعة تجريبية وقدرت قوة الهجين لكافة الصفات المدروسة باستخدام طريقة الهجين عن قيمة متوسط الأبوين أو قيمة الأب الأفضل. وتم تحليل البيانات المدروسة هي : طول النبات (سم) – طول السنبلة – عدد سنابل النبات – وزن 1000 جبه –المحصول البيولوجى –محصول الحبوب/ نبات (سم) – طول السنبلة عدد سنابل النبات الوباء ولافت المعان الأباء والهجن معنويا لكل الصفات المدروسة تحت ظروف التحليل المشترك.

أظهرت كلا من الآباء P6,P2,P8,P2,P2,P2, بنات ومحصول النيولوجى/ نبات ومحصول حبوب النبات الفردي عدد السنابل / النبات ، وزن الـ1000 حبة ، المحصول البيولوجى/ نبات ومحصول حبوب النبات الفردي علي التوالي .كما أظهر الهجين P1xP6 علي قيم لصفة المحصول البيولوجى للنبات و الهجين P2xP4 علي التوالي .كما أظهر الهجين P1xP6 علي قيم لصفة المحصول البيولوجى للنبات و الهجين للصفات حمصول الحمية محصول البيولوجى للنبات و الهجين كربات و المعنوبي الصفات علي التوالي .كما أظهر الهجين أعاية والمالية المحصول البيولوجى للنبات و الهجين كربات و المحمول البيولوجى للنبات و الهجين P1xP4 علي الصفات علي التوالي .كما أظهر الهجين التباين الراجع للقدرة العامة والخاصة على التآلف معنوبا للصفات تحت الدراسة في كل لصفة محصول الحبوب للنبات. والممترك. وأظهرت السلالة 66 قدرة عامة علي التآلف لصفات طول السنبلة، من معاملتي الري والتحليل المشترك. وأظهرت السلالة 66 قدرة عامة علي التآلف لصفات طول السنبلة، عدد السنابل للنبات، ووزن 1000 حبة والمحصول البيولوجي للنبات و محصول حبوب النبات.أظهرت معنوبي المحمول البيولوجي للنبات و محصول حبوب النبات.أظهرت المنبلة، عدد السنابل النبات و محصول حبوب النبات.

اظهر تحليل الارتباط أن هناك معنوية بين التباعد الوراثي وكل من متوسط أداء الهجن وقوة الهجين بنوعيها أي لمتوسط الابوين والاب الافضل وتأثير القدرة الخاصة على التآلف لكل الهجن تحت الدراسة موجبة ولكن منخفضة.

من خلال هذه الدراسة يتضح ان تكنيك ISSR من المعلمات الجزيئية يمكن أن يستخدم في تحديد التباعد الوراثي بين الاباء الثمانية وتقسيمها الى مجموعات واستخدام هذا التباعد في التنبؤ بالمحصول وقوة الهجين للهجن الناتجة بين هذه الاباء.

INTRODUCTION

Wheat (*Triticum aestivum* L.) is the major cereal crop in Egypt and several other countries. World average cultivated area of wheat was 221.73* ²million hectares in 2017*; the total production was 751.36* million metric tons, with an average productivity of 3.39* metric tons

*Foreign Agricultural Service / USDA Office of Global Analysis

http://www.pecad.fas.usda.gov

-(1 metric ton per hectare = 100 grams per square meter, 1 hectare (ha) = 10,000 square meters).

hectare⁻¹. In Egypt wheat grew in 1.25* million hectares that produced 8.10* million metric tons of grains, with an average yield of 6.43* metric tons hectare⁻¹. With increasing population, it could hardly satisfy only 55% of local requirements. The increasing gap between production and consumption necessitates increasing wheat production in Egypt. To overcome this problem is to increasing the productivity of wheat through an efficient breading program.

Stresses can occur at any stage of plant growth and development, thus illustrating the dynamic nature of crop plants and their productivity is needed. Drought among abiotic stresses is the most widespread and limiting crop productivity. There are definitions of drought, which include precipitation, evapo-transpiration, potential evapotranspiration, temperature, humidity and other factors individually or in combination (**Renu and Suresh, 1998**). Also selection for genotypes with increased productivity in drought environments has been an important goal of many plant breeding programs, the biological basis for drought tolerance is still poorly understood.

The diallel cross designs are frequently used in plant breeding research to obtain information about genetic properties of parental lines or estimates of general combining ability (GCA), specific combining ability (SCA) and heritability (**Baker**, 1978; EL- Maghraby *et al.*, 2005 and Iqbal *et al.*, 2007). In addition, the diallel cross technique was reported to provide early information on the genetic behavior of these attributes in the first generation (Chowdhry *et al.*, 1992 and Topal *et al.*, 2004). Diallel analysis technique is the choice of providing such detailed genetic information for selecting breeding materials that show great promise for success (Lonnquit and Gardner, 1961).

Molecular markers that reveal polymorphism at the DNA level have been shown to be a very powerful tool for genetic diversity since they were independent of the confounding effects of environmental factors. Molecular techniques are now a valuable tool for achieving genetic variation among wheat parents is necessary to derive superior progeny from crossing and selection. Precise information on the nature and degree of genetic diversity present in wheat collections from its principal areas of cultivation would help to select parents for evolving superior varieties.

New molecular tools such as inter simple sequence repeats (ISSR) have now provided the opportunity to monitor genetic integrity at the genotype level and laboratory tests are available to determine any unintentional genetic erosion or change in genetic identity. Therefore, the investigation aimed to: use ISSR-PCR marker to detect DNA polymorphism, identify parents and estimate genetic diversity among wheat genotypes and assess the variations amongst a half diallel crosses among eight genotypes for drought avoidance and drought tolerance traits.

MATERIALS AND METHODS

Plant materials: Eight genotypes of wheat representing a wide range of diversity for several agronomic characters and drought resistance measurements were selected for the study. The names, pedigree and origin of these varieties are presented in Table (1).

NO	Entry name	Pedigree	Source
1	Yakora Rojo	Ciano 67/Sonora 6411 Klien Rendidor/3/1L815626Y-2M-1Y-0M-302M	CIMMYT
2	Gemiza 7	CMH74 A. 630/5x//Seri 82/3/Agent (Gemiza 7)	Egypt
3	Giza 168	MRI/BUG/SEPI CM933046-8M-OY-OM•2Y-O3-OGZ.	Egypt
4	Gemiza 11	BOW"S"/KVZ"S"//7C/SER182/3/GIZA 168/SAKHA61. GM7892-2GM-1GM-2GM-1GM-0GM.	Egypt
5	Sakha 93	S 92/TR 810328 S8871-1S-2S-1S-0S	Egypt
6	Sides 12	BUC//7C/ALD/5/MAYA74/ON//1160.147/3/BB/GLL /4/CHAT"S"/6/MAYA/VUL//CMH74A.630/ 4*SXSD7096-4SD-1SD-1SD 0SD	Egypt
7	Sahel 1	NS 732/PIMA//Veery'S'	ICARDA
8	13-ssd-43	Landraces	Egypt

Table (1): The name, pedigree and source of the studied parental varieties and lines.

Line No 8 was developed in the Department Of Agronomy , Faculty of Agric. at Moshtohor , Banha Univ. by Prof. Dr. M. El.Badawy.

DNA extraction: 15 seeds of parental genotypes were sowing in pots. Leaf tissue was collected from 5-7 days old germinated seedlings. Equal quantities of leaf tissue from 10 seedlings of each genotype were bulked, lyophilized and ground with a mortar. Genomic DNA was isolated and extracted using mi-plant genomic DNA Isolation Kit (Metabion).

Inter-simple sequence repeats (ISSR): After estimating the DNA samples concentration aliquots from each stock of DNA samples were diluted to a uniform concentration of 10 ng/ μ l to be used with PCR marker. Oligonucleotide sequences of the primers used in this study were selected from a set of Operon kits (Operon Technologies Inc., Alameda California, USA). A total of six primers as indicated in Table (2) were used in the detection of polymorphism among eight wheat genotypes.

Reactions were carried out in a total volume of 25 μ l containing 30 ng of genomic DNA as a template, 30 pmoles of random primer, 2mM of dNTP's mix (dATP, dCTP, dTTP and dGTP, ABgene, Surrey, UK), 10 X PCR buffer, 25 mM MgCl₂, and 2 units Taq DNA polymerase (promega, USA).

Amplifications were carried out in a thermo cycler (UNO II Biometra) programed for 94°C FOR 4 min, (one cycle); followed by 94°C for 45 sec, 38°C for 1 min, and 72°C for 1 min (35 cycle); 72°C for 10(one cycle) ,then 4°C(infinitive).

The amplification products were resolved by electrophoresis in a 2% agarose gel containing ethidium bromide (0.5 μ g/ml), visualized with ultraviolet light and photographed. DNA fragment sizes were determined by comparison with the 100bp and 1Kb DNA ladder marker (promega USA).

Field experiments: This investigation was carried out at the Experiment, Research Station of Moshtohor Faculty of Agriculture, Benha University, Kalubia Governorate, Egypt. In 2015/2016 growing season, the parents were crossed in a 8x8 diallel cross excluding reciprocals giving a total of twenty-eight crosses. In 2016/2017 two experiments using randomized complete block design with three replications were carried out. Each experiment contained the eight parents and their resulting 28 F1's. The sowing date was on 4th Dec. 2016. The first experiment was irrigated only once after planting irrigation and the second one was normally irrigated 5 irrigations. Plots of parents and F1's consisted of one row, 3 m-long, with spacing of 30 cm between rows and 20 cm between plants. The dry method of planting was used in this study. The other cultural practices of growing wheat were practiced. The amounts of total rainfall during the evaluating season were recorded in Table (2).

	Temper	rature C	БИ	Rain fall mm/month	
Months	Min.	Max.	к.н. (%)		
Dec.2016	19.7	9.2	51.3	0.5	
Jan.2017	17.7	6.1	55.9	1.6	
Feb.2017	20.4	7.8	47.2	0.8	
Mar.2017	25.8	11.4	37.3	0.4	
Apr.2017	29.1	14.4	38.9	0.3	
May.2017	34.5	19.0	32.1		

 Table 2. Monthly averages of temperature, relative humidity (R.H.) and total rain fall during 2016/2017 season at Kalubia (Moshtohor).

According to meteorological weather station Moshtohor.

Ten guarded plants from parents and the F_1 's were selected randomly from each plot for recording observations on different characters. The characters studied were, Plant height(cm), No .of spikes /plant, No .of kernels/ spike,1000- kernel weight (g), biological yield/ plant and grain yield/ plant (g).

Data analysis: Analysis of variance was conducted as outlined by **Steel and Torrie (1980)** for all characters. The analysis of GCA and SCA was done following the procedure given by **Griffing (1956)** using Method II Model I. The combined analysis of the two experiments was carried out whenever homogeneity of mean squares was detected (**Gomez and Gomez 1984**).

ISSR analysis: The obtained data of ISSR analysis were entered in a computer file as binary matrices where 0 stands for the absence of a band and 1 stands for the presence of a band in each individual sample. Similarity coefficients between a pair of inbred lines were calculated according to Jaccard (1908). A dendrogram tree was constructed by the UPGMA clustering algoritm from the SAHN option of NTSYS-PC version 2.1 (**Rohlf, 2000**).

RESULTS AND DISCUSSION

ISSR Polymorphism: ISSR experiments were conducted using ten specific primers. Four primers gave non-polymorphic fragments. Meanwhile six primers gave polymorphic amplification products; these are primer ISSR1, ISSR2, ISSR4, ISSR5, ISSR6 and ISSR10. Therefore, the last six primers were included in this study. Also, ISSR fingerprint for the studied genotypes were done to study the diversity and relationship between molecular marker polymorphism and mean performance. Among the 54 fragments generated, that ranged from six for ISSR2 to 12 for ISSR1. The mean of fragments per primer was 9. The size of fragments ranged from 20.00 b.p to 340.21 b.p length. Among 54 generated fragment 30 were polymorphic fragment obtained within the eight genotypes (Table 3). Other studies indicating different results **Guo-yue and Hui (2007)**, **Sharma** *et al.*(2014), **Tonk** *et al.*(2014), **Razmjoo** *et al.* (2015), **Tarinejad** *et al.*(2015), **Olgun** *et al.*(2015) and **Chaudhary** *et al.*(2016).

Primers ISSR	Sequence	TSB	TF	NPF	PPF
1	5'-AGAGAGAGAGAGAGAGYC-3'	65	12	8	66.67
2	5'-AGAGAGAGAGAGAGAGAGYG-3'	40	6	4	66.67
4	5'-ACACACACACACACACYG-3'	55	7	1	14.29
5	5'-GTGTGTGTGTGTGTGTYG-3'	69	13	8	61.54
6	5'-CGCGATAGATAGATAGATA- 3'	44	8	4	50.00
10	5'-GACAGACAGACAGACAAT-3'	41	8	5	62.50
	Total	314	54	30	-
	Mean	52.33	9.00	5.00	53.61

 Table (3) primer used in ISSR analysis of eight wheat genotypes and total number of fragments detected by each primer and number of polymorphic fragments.

Where: TSB = Total number of scorble bands, TF= Total number of fragments, NPF = Number of polymorphic fragments, PPF = Percentage of polymorphic fragments.

Genetic similarity: The genetic similarity matrix was produced from the ISSR data using **Nei and Li's formula (1979)**. Genetic similarity coefficients are presented in (Table 4). The lowest genetic similarity (0.63) was obtained between the two parental genotypes P3 and P7, while, the highest genetic similarity (0.89) was scored between the two parental genotypes P7 and P8. The over all mean for genetic similarity among all genotypes was (0.74). **Tahir (2010)** analyzed eleven wheat varieties of diverse origins with 12 selected ISSRs and found a wide range of genomic diversity among all the genotypes, indicating them as ISSR prime candidates for selective breeding for

specific traits and broadening the genetic base. Also, Abou-Deif *et al.* (2013) Analyzed 20 wheat genotypes using (ISSR) markers and found ISSR markers succeeded in distinguishing most of the 20 varieties in relation to their genetic background and geographical origin.

Cluster analysis: The dendrogram constructed from cluster analysis based on ISSR data is represented in Fig (1). The data collectively distinguished two main clusters. The first main cluster contained Gemiza 7 (P2) while, the second cluster consists of remaining

Table (4): Genetic similarity based on Nei and Li's coefficient for eight genotypes in whea
revealed by ISSR.

Rows/ cols	p1	p2	p3	p4	p5	рб	p7	p8
p1	1.000							
p2	0.750	1.000						
p3	0.740	0.640	1.000					
p4	0.765	0.700	0.884	1.000				
p5	0.700	0.702	0.727	0.837	1.000			
рб	0.760	0.694	0.681	0.782	0.833	1.000		
p7	0.647	0.681	0.630	0.733	0.825	0.767	1.000	
p8	0.686	0.761	0.674	0.777	0.875	0.814	0.897	1.000



Fig. (1): phonogram generated by UPGMA cluster analysis based on Nei and Li[,] coefficients showing clustering of eight genotypes.

seven genotypes this cluster separated into two sub-clusters: the first sub-cluster contained three genotypes Yakora (P1), Giza 168 (P3) and Gemiza 11(P4). Meanwhile, the second sub cluster contained the other four genotypes i.e. Sakha 93 (P5), Sahel 1(P7), 13-ssd-43 (P8) and Sides 12(P6). In addition, the second sub cluster divided into two sub-sub clusters the first sub-sub cluster contained P6. While, the inbred lines P5, P7 and P8 were belonging to the second sub-sub cluster as well as the two genotypes 7 and 8 being closely related.

The second experiment: Analyses of variance for yield and its components under drought and normal irrigation and combined analysis across the mention environments are presented in Table 3. Results indicated that mean squares due to irrigation treatments (Environments) were highly significant for all studied traits indicating overall differences between the two environments of study.

Genotypes mean squares were significant for all studied traits indicating wide diversity between all genotypes used in this work. Moreover, significant mean squares between genotypes and environment interaction were detected for No of spikes/plant, biological yield/ plant and grain yield/ plant. This result indicated that genotypes responded differently to different environments for the mention traits.

Mean squares due to parents were highly significant for all traits in drought stress, normal irrigation and combined across them indicating that these parents are differently in the aforementioned significant traits. Moreover, mean squares due to the interaction between parents and environments were significant for No of spike/ plant, Biological yield/ plant and grain yield/ plant. Such result indicated that wheat parents responded differently to stress and non-stress conditions for these traits. For the exceptional traits, insignificant mean squares between parents and environments were detected indicating that parents behaved similarly in stress and non-stress conditions.

Mean performance

Results in Table (5) showed the average of plant height, yield and its components traits at the combined across irrigation treatments. It's clear that the parental line (P_1) gave the lowest mean value for plant height. On the other hand, P2 was the tallest parent. Plant height for crosses ranged from 72.50 cm (P1xP3) to 97.75cm (P6xP7). Moreover, the crosses P2xP4, P2xP7, P3xP7, P4xP5, P4xP8 and P6xP8 did not differ significantly than the tallest hybrid P6xP7. Some farmers usually prefer higher plant due to the high price of hay.

Fable (5) Mean	square	s for yield	and its con	nponents ເ	under drou	ight stress co	ondition and
]	normal irr	igation as [•]	well as the	combined	over them.	

S.O.V.	df	plant height	spike length	No. of spikes /plant	1000 kernel weight	Biological yield/plant	Grain yield/plant	
Drought environment								
Rep	2	41.82	7.34**	17.81*	100.51**	7.96	0.3	
Genotypes(G)	35	163.82**	2.55**	102.81**	56.12**	4046.34**	141.80**	
Parent (P)	7	123.61**	2.62	95.07**	73.90**	2320.86**	119.96**	
Cross (C)	27	178.29**	2.53*	90.55**	53.37**	4247.56**	152.00**	
P vs C.	1	54.57*	2.67*	488.02**	5.76	10691.64**	19.1*	
Error	70	28.19	1.3	5.49	16.02	77.59	9.23	
Normal environ	nment							
Rep	2	7.16	1.9	3.39	5.33	2.16	6.72	
Genotypes(G)	35	176.69**	4.50**	114.25**	70.14**	4683.45**	179.64**	
Parent (P)	7	288.20**	4.69**	70.93**	114.22**	7645.45**	182.67**	
Cross (C)	27	146.43**	4.58**	125.90**	60.67**	4012.33**	180.58**	
P vs C.	1	213.00**	0.93	102.93**	17.19	2069.68**	132.89**	
Error	70	23.56	1.18	4.58	9.18	80.31	14.61	
Combined anal	ysis							
Irrigation (I)	1	5726.03**	102.78**	2681.12**	443.19**	363533.47**	3985.17**	
Rep/ I	4	24.49	4.62**	10.6	52.92**	5.06	3.51	
Genotypes(G)	35	301.97**	5.70**	155.57**	117.43**	6221.16**	230.37**	
Parent (P)	7	374.62**	6.63**	154.87**	181.57**	7009.63**	254.07**	
Cross (C)	27	285.38**	5.55**	142.26**	104.35**	5836.61**	228.07**	
P vs C.	1	241.60**	3.37*	519.60**	21.42	11084.74**	126.37**	
GxI	35	38.54	1.34	61.50**	8.83	2508.63**	91.07**	
p x I	7	37.2	0.68	11.12*	6.55	2956.68**	48.57**	
CxI	27	39.35	1.55	74.19**	9.69	2423.28**	104.51**	
P.vs.C x I	1	25.97	0.22	71.35**	1.52	1676.59**	25.62	
Error	140	25.87	1.24	5.04	12.6	78.95	11.92	

* and ** significant at 0.05 and 0.01 levels of probability, respectively.

On the other hand, this plant must be given high yield for grain and behave resistant to lodging. The highest parents mean value for spike length (12.92cm) was detected for P2. However, eight crosses P2xP3, P2xP4, P3xP4, P3xP7, P4xP6, P4xP7, P4xP8 and P7xP8 exhibited highest values for spike length. For No. of spike/ plant the parent P8 and the cross P1xP4 give the highest number of spikes/ plant. Heavier 1000-kernel weight were detected for P2, P4, P6, P1xP6, P2xP4, P2xP6, P2xP8, P4xP5, P4xP8, P5xP8, P6xP7 and P6xP8. The parental No 6 (P₆) gave the highest mean

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value for biological yield/ plant and ranked the first parents for this traits. Moreover, the cross P1xP6 exhibited the highest crosses for biological yield/ plant. Parent No 2 (P_2) and the cross P2xP4 gave the highest mean values for grain yield / plant. Therefore, these crosses could be efficient for prospective wheat breeding programs aiming at improving wheat grain yield.

	nlant	cniko	No. of	1000	Biological	Grain
Genotypes	piant beight	length	spikes	kernel	vield/ nlant	vield/ nlant
	neight	length	/plant	weight	yiciu/ plant	yiciu/ piant
P1	70.13	11.25	29.25	38.3	207.5	16.5
P2	95.92	12.92	24.5	50.58	211.17	35.25
P3	79.67	12.08	28.42	35.48	134.92	16.75
P4	80.54	12.08	32.58	46.17	188.25	24.88
P5	75.17	11.42	17.33	40.37	128.67	16.5
P6	88.75	11.92	28.83	49.23	214.33	25.42
P7	82.17	9.33	29.67	39.37	166.5	26.67
P8	83.58	11.17	33.33	39.95	199.17	23.5
1x2	73.67	11.33	16.5	38.77	140.83	18.78
1x3	72.5	9.83	30.83	35.36	167	18.58
1x4	81	12.25	32.33	40.43	192.33	25.33
1x5	86.63	11.08	28.5	37	146.17	16.67
1x6	85.75	11.92	32.17	47.65	229.17	33
1x7	76.46	10.17	28.17	37.5	149.83	27.71
1x8	75.58	10.75	23.33	38.2	123	26.33
2x3	79.5	13.33	22.33	40.53	150	21.67
2x4	92.54	12.58	14.5	50.68	115.33	37.5
2x5	78.17	11.25	25.67	42.72	187.17	22.92
2x6	89.25	12.25	17.33	49.65	209.33	30.6
2x7	91.42	11.33	26.83	41.77	174.67	19.49
2x8	87.85	12.33	21.75	46.1	174.78	25.5
3x4	78.5	13	23.17	42.13	169.17	20.58
3x5	75.92	11.83	24.83	37.93	140	12.33
3x6	82.25	11.83	29.83	41.75	202.83	20.58
3x7	90.75	12.67	22.83	44.28	181.67	27.33
3x8	79.58	11	25.08	39.48	135.5	33.83
4x5	92.33	11.92	25.67	49.2	169.67	21.83
4x6	89.67	12.58	25.58	47.92	201	27.33
4x7	86.42	12.92	28.67	44.83	174.17	26.96
4x8	95.67	12.83	19	45.8	154.67	34.83
5x6	83.42	12.08	23	44.05	159.17	25.21
5x7	82.5	10.58	20.75	42.63	129.67	20.5
5x8	81.5	10.25	27.5	47.38	155.67	27.17
6x7	97.75	11.33	25.5	46.45	204.33	31
6x8	90.42	13.17	23.33	45.82	166.33	30.79
7x8	89.96	12.58	14.25	43.27	90.83	16.25
mean of parent	81.99	11.52	27.99	42.43	181.31	23.18
mean of cross	84.53	11.82	24.26	43.19	164.08	25.02
mean of Genotype	83.97	11.75	25.09	43.02	167.91	24.61
L.S.D 5%	8.14	1.78	3.59	5.68	14.22	5.52
L.S.D 1%	10.67	2.34	4.71	7.45	18.65	7.24

Table 6. Mean performance of the genotypes for yield and its components over the studied environments.

Combining ability

The analysis of variance for combining ability for plant height, spike length, number of spikes/ plant, 1000-kernel weight, biological yield, and grain yield/ plant, under drought treatment, normal irrigation and combined analysis is presented in Table 6.

S.O.V.	df	plant height	spike length	No. of spikes /plant	1000 kernel weight	Biological yield/ plant	Grain yield/ plant
Drought envir	ronmer	nt					
GCA	7	146.79**	1.68	46.40**	58.49**	3615.79**	94.33**
SCA	28	31.56**	0.64	31.24**	8.76**	782.02**	35.50**
Error	70	9.4	0.43	1.83	5.34	25.86	3.08
GCA/SCA		4.65	2.63	1.49	6.68	4.62	2.66
Normal envir	onmen	t					
GCA	7	158.34**	3.09**	22.91**	88.19**	2226.62**	121.27**
SCA	28	34.04**	1.1	41.88**	7.18**	1394.78**	44.53**
Error	70	7.85	0.39	1.53	3.06	26.77	4.87
GCA/SCA		4.65	2.81	0.55	12.29	1.6	2.72
Combined and	alysis						
GCA	7	277.47**	4.50**	60.46**	143.88**	4199.35**	162.32**
SCA	28	56.46**	1.25**	49.71**	12.96**	1542.31**	55.40**
GCA x L	7	27.67**	0.27	8.85**	2.8	1643.06**	53.27**
SCA x L	28	9.14	0.49	23.41**	2.98	634.50**	24.63**
Error	140	8.62	0.41	1.68	4.2	26.32	3.97
GCA/SCA		4.91	3.6	1.22	11.1	2.72	2.93
GCA x							
L/GCA		0.10	0.06	0.15	0.02	0.39	0.33
SCA x							
L/SCA		0.16	0.39	0.47	0.23	0.41	0.44

 Table (7) Combining abilities mean squares for yield and its components under normal irrigation and drought stress condition as well as the combined over them.

* p< 0.05; ** p< 0.01

General (GCA) and specific (SCA) combining ability mean squares were highly significant for all studied traits in both environments as well as combined analysis except for spike length under drought and normal conditions. Such results indicated that both types of combining ability are important in the inheritance of these traits. Moreover, the ratios between GCA and SCA exceeded the unity for all studied traits, revealing that additive and additive x additive types of gene action are more important than non-additive gene action in controlling these traits. The genetic variance was previously reported to be mostly due to additive effects for plant height by Menshawy (2004) and El Hosary *et al* (2009); for spikes/ plant by El Seidy and Hamada (1997), El Borhamy (2000), Gomaa*et al* (2014); for 1000-grain weight by El Seidy and Hamada (1997), El Borhamy (2000), and for grain yield/ plant by El Seidy and Hamada (1997), El Borhamy (2000), El Borhamy (2000), Abd El-Aty and Katta (2002), El Hosary*et al* (2012), Gomaa*et al* (2014).

The mean squares of the interaction between GCA, SCA and irrigation treatments were significant for all studied traits except both types of combining abilities x E for spike length and 1000-kernel weight. Such result indicated that the additive and non-additive types of gene action differed significantly from one environment to another for these traits. Similar results were reported by **El-Seidy and Hamada (1997), El-Seidy and Hamada (2000)**.

The ratio SCA x environment/ SCA was much higher that of GCA x irrigation/ GCA treatments for all traits indicating that non-additive effects were much more influenced by environments than additive genetic one. Such results are in harmony with those obtained by **El Hosary and Nour El Deen (2015).**

General combing ability (GCA) effects: Test of homogeneity revealed the validity of the combined analysis for the data of the two irrigation treatments. The general combining ability

effects \hat{g}_i of each parent for all studied measurements at the combined analysis are presented in Table (7).

Parent	plant height	Spike length	No of spike/ plant	1000- kernel weight	biological yield/ plant	grain yield / plant
P1	-6.39**	-0.60**	2.45**	-3.57**	5.21**	-2.21**
P2	2.85**	0.45**	-3.19**	2.42**	6.33**	2.54**
P3	-3.74**	0.19*	1.00**	-3.47**	-9.52**	-3.31**
P4	2.15**	0.65**	0.83**	2.62**	4.16**	2.26**
P5	-2.49**	-0.40**	-1.52**	-0.55*	-16.64**	-4.19**
P6	4.03**	0.32**	0.86**	3.46**	28.96**	2.78**
P7	2.39**	-0.55**	0.05	-0.77**	-7.30**	0.11
P8	1.20**	-0.05	-0.49**	-0.12	-11.21**	2.02**
L.S.D(0.05) gi	0.68	0.15	0.3	0.47	1.18	0.46
L.S.D(0.01) gi	0.89	0.19	0.39	0.62	1.55	0.6
L.S.D(0.05) gi-gj	1.29	0.28	0.57	0.9	2.25	0.87
L.S.D(0.01) gi-gj	1.69	0.37	0.74	1.18	2.95	1.15

Fable 8. Estimates of general combining ability effects for yield and its components at the
combined analysis.

* p< 0.05; ** p< 0.01

Such results are being used to compare the average performance of each parent with other genotype and facilitate selection of parents for further improvement to drought resistance. Results indicate that the parental P_1 gave desirable significant $\hat{g}i$ effects for plant height, no of spike/ plant and biological yield/ plant. P_2 exhibited significant and positive $\hat{g}i$ effects for plant height, spike length, 1000-kernel weight, biological yield/ plant and grain yield/ plant. P3 gave useful significant $\hat{g}i$ effects for plant height, spike length and no of spikes/ plant. P_4 expressed significant and positive $\hat{g}i$ effects for plant height, spike length, no of spikes/ plant, 1000-kernel weight, biological yield/ plant and grain yield/ plant. P_5 seemed good general combiner for plant height and grain. P6 exhibited positive and significant $\hat{g}i$ effects spike length, no of spikes/ plant, 1000-grain, biological yield/ plant and grain yield/ plant. Also, it is considered the best combiner for grain yield/ plant and most of its components. P7 and P8 gave positive and significant combiner for plant height.

Specific combining ability (SCA) effects

Specific combining ability effects \hat{s}_{σ} for the F₁ crosses for the studied traits in the combined analysis are presented in (Table 8).

For plant height, six crosses expressed significant and positive ŝij effects. Moreover, the cross $P_1 \times P_5$ gave the most desirable ŝij effects for plant height. However, three cross combinations i.e. P1xP2, P2xP5 and P3xP4 gave significant and negative ŝij effects for the mention trait. For spike length, five crosses in the combined analysis expressed significant and positive ŝij effects. Moreover, the cross $P_7 \times P_8$ gave the most desirable ŝij effects for this trait. For number of spikes/ plant, nine crosses expressed significant and positive ŝij effects. However, the best ŝij effects (5.29**) were detected for the cross $P_2 \times P_5$. Regarding 1000-kernel weight, five cross combinations expressed significant and positive ŝij effects. The cross P3xP7 being the highest one in this traits and recorded 5.51**. twelve crosses combinations exhibited significant and positive ŝij effects for biological yield/ plant. The best positive ŝij effects were the crosses $P_2 \times P_5$ and $P_3 \times P_7$ in the combined analysis (Table 8). Regarding to grain yield/ plant eight crosses i. e. P1xP6, P1xP7, P2xP4, P3xP7, P3xP8, P4xP8, P5xP8 and P6xP7 exhibited significant and positive ŝij effects.

cross combinations	plant height	Spike length	No of	1000-	biological	grain
			spike/	kernel	yield/	yield /
	noight	lungun	plant	weight	plant	plant
P1xP2	-6.77**	-0.27	-7.85**	-3.11*	-38.62**	-6.16**
P1xP3	-1.34	-1.51**	2.30**	-0.62	3.4	-0.5
P1xP4	1.27	0.45	3.96**	-1.63	15.05**	0.67
P1xP5	11.54**	0.32	2.48**	-1.9	-10.32**	-1.54
P1xP6	4.14*	0.44	3.76**	4.74**	27.08**	7.81**
P1xP7	-3.51	-0.44	0.57	-1.18	-15.99**	5.20**
P1xP8	-3.2	-0.35	-3.72**	-1.13	-38.92**	1.91
P2xP3	-3.58	0.95*	-0.56	-1.44	-14.72**	-2.17
P2xP4	3.57	-0.26	-8.23**	2.63*	-63.07**	8.08**
P2xP5	-6.16**	-0.55	5.29**	-2.17	29.57**	-0.05
P2xP6	-1.6	-0.27	-5.43**	0.75	6.13	0.66
P2xP7	2.21	-0.31	4.88**	-2.90*	7.73*	-7.77**
P2xP8	-0.17	0.19	0.34	0.78	11.75**	-3.6**
P3xP4	-3.88*	0.41	-3.75**	-0.03	6.61*	-2.98*
P3xP5	-1.82	0.29	0.27	-1.06	-1.75	-4.78**
P3xP6	-2.01	-0.43	2.89**	-1.25	15.48**	-3.50**
P3xP7	8.13**	1.28**	-3.30**	5.51**	30.58**	5.92**
P3xP8	-1.85	-0.89*	-0.51	0.06	-11.68**	10.51**
P4xP5	8.71**	-0.09	1.27	4.12**	14.23**	-0.85
P4xP6	-0.48	-0.14	-1.2	-1.18	-0.04	-2.32
P4xP7	-2.09	1.07**	2.70**	-0.03	9.40**	-0.02
P4xP8	8.35**	0.49	-6.43**	0.29	-6.2	5.94**
P5xP6	-2.09	0.4	-1.43	-1.87	-21.07**	2.01
P5xP7	-1.36	-0.22	-2.87**	0.94	-14.30**	-0.02
P5xP8	-1.18	-1.05*	4.42**	5.04**	15.60**	4.72**
P6xP7	7.37**	-0.19	-0.5	0.74	14.76**	3.49**
P6xP8	1.22	1.15**	-2.13*	-0.54	-19.33**	1.37
P7xP8	2.4	1.44**	-10.40**	1.14	-58.57**	-10.48**
LSD5%(sij)	3.69	0.81	1.63	2.58	6.45	2.5
LSD1%(sij)	4.84	1.06	2.14	3.38	8.45	3.28
LSD5%(sij-sik)	5.46	1.2	2.41	3.81	9.54	3.7
LSD1%(sij-sik)	7.16	1.57	3.16	5	12.51	4.85
LSD5%(sij-skL)	1.82	0.4	0.8	1.27	3.18	1.23
LSD1%(sij-skL)	2.39	0.52	1.05	1.67	4.17	1.61

Table 9. Estimates of specific combining ability effects for yield and its components 'at the combined analysis.

* p< 0.05; ** p< 0.01

It could be concluded that the previous cross combinations might be of interest in breeding programs towards the development of pure lines varieties for high biological, and grain yields/ plant under drought conditions.

The correlation between genetic distance and each of mean performance, heterosis and SCA for grain yield/plant: The correlation of GD and each of SCA and heterosis for grain yield for 28 hybrids combination are estimated. The estimated values of correlation coefficient between GD, and each of mean performance and heterosis relative to Mp as well as Bp and SCA for grain yield/plant found positive (r = 0.134, 0.067, 0.035 and 0.238, respectively) Table (10). Therefore, specified

tendency could be predicted about the relationship of GD and heterosis for grain yield/plant in this study. **El-Maghraby et al. (2005)** reported that ISSRs are useful in detecting a high level of polymorphism among wheat cultivars. **Also, Prasad** *et al.* (2000) illustrated that the utility of ISSR markers for detecting polymorphism leading to genotype identification and for estimating genetic diversity.

Crosses	genetic diversity	grain yield					
		mean	Hete	SCA			
		performance	Мр	BP	SCA		
1x2	0.25	18.78	-27.43	-46.73	-6.16		
1x3	0.26	18.58	11.78	10.95	-0.5		
1x4	0.235	25.33	22.46	1.84	0.67		
1x5	0.3	16.67	1.01	1.01	-1.54		
1x6	0.24	33	57.46	29.84	7.81		
1x7	0.353	27.71	28.38	3.91	5.2		
1x8	0.314	26.33	31.67	12.06	1.91		
2x3	0.36	21.67	-16.67	-38.53	-2.17		
2x4	0.3	37.5	24.74	6.38	8.08		
2x5	0.298	22.92	-11.43	-34.99	-0.05		
2x6	0.306	30.6	0.89	-13.18	0.66		
2x7	0.319	19.49	-37.06	-44.72	-7.77		
2x8	0.239	25.5	-13.19	-27.66	-3.6		
3x4	0.116	20.58	-1.1	-17.25	-2.98		
3x5	0.273	12.33	-25.81	-26.37	-4.78		
3x6	0.319	20.58	-2.37	-19.02	-3.5		
3x7	0.37	27.33	25.91	2.5	5.92		
3x8	0.326	33.83	68.12	43.97	10.51		
4x5	0.163	21.83	5.54	-12.23	-0.85		
4x6	0.218	27.33	8.7	7.54	-2.32		
4x7	0.267	26.96	4.61	1.09	-0.02		
4x8	0.223	34.83	44.01	40.03	5.94		
5x6	0.167	25.21	20.28	-0.82	2.01		
5x7	0.175	20.5	-5.02	-23.13	-0.02		
5x8	0.125	27.17	35.83	15.6	4.72		
6x7	0.233	31	19.04	16.25	3.49		
6x8	0.186	30.79	25.89	21.15	1.37		
7x8	0.103	16.25	-35.22	-39.06	-10.48		

 Table 10. Genetic diversity, mean performance, heterosis, heterobiltosis and specific combining ability for grain yield for all studied crosses.

The results indicated that ISSR marker can be used as a tool for determining the extent of genetic diversity among wheat genotypes and for genotypes into different groups. This study showed that GD can be used to precisely predict the yield performance and heterosis value for F_1 hybrids. For more accuracy the more primers must be used to detection the difference between parents or can use more markers.

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