# A Research on Heterosis in Cultivated (Triticum spp.) $\times$ Semi-Wild Wheat Hybridization 

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Direction and magnitude of mid-parent (MP) and high-parent (HP) heterosis were detected and estimated in 18 combinations, $180 \mathrm{~F}_{1} \mathrm{~s}$ and $150 \mathrm{~F}_{2}$ s for plant height ( PH ); spike length (SL); number of spike per plant (NS), number of spikelet per spike (NSP) and number of grain per spikelet (NG). For this, 9 females ( 7 commons Triticum aestivum (L.) Em. (Thell), $\mathrm{P}_{5}=$ Aköz 867, $\mathrm{P}_{7}=$ Köse 220/39, $\mathrm{P}_{9}=$ Penjamo 62, $\mathrm{P}_{10}=$ Sivas $111 / 33, \mathrm{P}_{11}=$ Sürak 1593/51, $\mathrm{P}_{12}=$ Sertak 52 and $\mathrm{P}_{13}=$ Yektay 406 and 2 durums (Triticum durum Desf., $\mathrm{P}_{1}=$ Kunduru $414 / 44$ and $\mathrm{P}_{4}=$ Kunduru 1149) and 4 semi wild males ( $\mathrm{P}_{2}=$ T. dicoccum., $2 \mathrm{n}=28 ; \mathrm{P}_{3}=T$. carthlicum; $\mathrm{P}_{6}=$ T. vavilovi and $\mathrm{P}_{8}=$ T. spelta., $2 \mathrm{n}=42$ ) were hybridized. The MP was varied between $33.1 \%\left(\mathrm{P}_{13} \times \mathrm{P}_{8}\right)$ at the $\mathrm{F}_{1}$ for the NG and $152.6 \% \mathrm{P}_{7} \times \mathrm{P}_{8}$ at the $\mathrm{F}_{2}$ for the NS; the HP was ranged between $71.2 \%\left(\mathrm{P}_{7} \times \mathrm{P}_{8}\right)$ at the F 2 for the NG and $135.3 \%\left(\mathrm{P}_{7} \times \mathrm{P}_{8}\right)$ at the $\mathrm{F}_{1}$ for the NS. Heterosis effect was appeared mostly on the SL and PH; but fixed the highest on the NS and NSP in the each generation. The semi wilds were ordered as to be $\mathrm{P}_{8}>\mathrm{P}_{6}>\mathrm{P}_{3}>\mathrm{P}_{2}$. Statistically significant correlations were fixed between (MP- $\mathrm{F}_{1}$ ) and traits; SL and PH traits were found have been impressed as to be more adventageous and stable.

Keywords: mid-parent heterosis, high-parent heterosis, genotypic diversity, parental selection

## Kültür Buğdayı (Triticum spp.) $\times$ Yarı Yabani Buğday Melezlemesinde Heterosis Üzerine Bir Araştırma

Bitki boyu (PH), başak uzunluğu (SL), bitkide başak sayısı (NS), başakta başakçık sayısı (NSP) ve başakçıkta tane sayısı (NG) bakımından anaç ortalaması (MP) ile üstün anaç (HP) heterosisinin şiddeti ve yönü 18 kombinasyon, $180 \mathrm{~F}_{1}$ ve $150 \mathrm{~F}_{2}$ bitkisinde saptanmıştır. Bunun için, 9 ana (7 ekmeklik Triticum aestivum (L.) Em. (Thell), $\mathrm{P}_{5}=$ Aköz 867, $\mathrm{P}_{7}=$ Köse 220/39, $\mathrm{P}_{9}=$ Penjamo 62, $\mathrm{P}_{10}=$ Sivas $111 / 33, \mathrm{P}_{11}=$ Sürak 1593/51, $\mathrm{P}_{12}=$ Sertak 52 and $\mathrm{P}_{13}=$ Yektay 406 ve 2 makarnalık Triticum durum Desf., $\mathrm{P}_{1}=$ Kunduru $414 / 44, \mathrm{P}_{4}=$ Kunduru 1149) ile 4 yarı yabani baba ( $\mathrm{P}_{2}=$ T. dicoccum., $2 \mathrm{n}=28 ; \mathrm{P}_{3}=$ T. carthlicum; $\mathrm{P}_{6}=T$. vavilovi and $\mathrm{P}_{8}=$ T. spelta., $2 \mathrm{n}=42$ ) anaç melezlenmiştir. MP; \% $-94.1\left[\mathrm{~F}_{2}\right.$ de NS için $\mathrm{P}_{5} \times \mathrm{P}_{3}$ ] ile \% 87.3 [ $\mathrm{F}_{1}$ 'de NG için ( $\mathrm{P}_{13} \times \mathrm{P}_{6}$ ]; HP ise \% -85.6 [ $\mathrm{F}_{2}$ 'de NG için $\mathrm{P}_{10} \times \mathrm{P}_{8}$ ] ile \% 185.6 [ $\mathrm{F}_{1}$ 'de NS için $\mathrm{P}_{7} \times \mathrm{P}_{8}$ ] arasında; çoğunlukla SL ve PH 'de ortaya çıkmış; ancak, en yüksek değerler NS ile NSP'nin $\mathrm{F}_{1}$ ve $\mathrm{F}_{2}$ 'nde saptanmış; yarı yabaniler ise $P_{8}>P_{6}>P_{3}>P_{2}$ şeklinde sıralanmışlardır. NG dışındaki diğer özelliklerle (MP- $F_{1}$ ) arasında istatistiksel olarak önemli korelasyonlar saptanmış; SL ve PH'nın daha avantajlı ve kararlı olduğu izlenimi edinilmiştir.

Anahtar kelimeler: anaç ortalaması heterosis, üstün anaç heteosisi, genotipik farklılık, anaç seçimi.

## Introduction

"Wheat heterosis" has long been recognized and it was first time used by Briggle, (1963). The utilization of heterosis depends mainly upon the direction and magnitude of heterosis (Agnus, 1977). Estimation of the MP and HP may be useful in identifying true heterotic crosses (Singh et al. 2004). These parameters may be a useful indicator in predicting the breeding value in any wheat breeding program. It also provides information about combining ability of the parents and their usefulness in breeding programs (Sharma et al. 1986). The
use of MP as a predictor mean $\mathrm{F}_{1}$ was suggested earlier owing to its simple assessment and because of reliable information about the line breeding programs. Thus, estimates of MP and HP as well as mean of the $\mathrm{F}_{1}$ relative to the control can be considered encouraging for large-scale development and global acceptance of hybrid wheats. The choice of parental combinations yielding superior hybrids is the most important aspect in hybrid wheat breeding. So, analysis of the relative importance of general combination ability
(GCA) and special combination ability (SCA) provides knowledge of the type of gene action involved in the expression of traits and it allows inferences about optimum allocation of resources in hybrid breeding. Combining abilities (GCA and SCA) need to be tested to select the best parents for the process of hybrid wheat seed production. They are connected both with extension of breeding programmes and high costs. Commercially exploitable yield advantages of hybrids will also depend on the HP and on advances in line development, where considerable progress can be expected in the future. To select potential parents for hybrid production, testcrosses in various environments are required because SCA variance is one of the main component (Oettler et al. 2003). Additionaly, the tight correlations of GCA suggest that the probability of obtaining superior hybrids is greater when crossing the highest yielding parents. Thus, hybrid wheat breeding should be respectively efficient based only on selection for parental performance and a respectively small number of testcrosses involving outstanding parental lines. However, despite useful predictions of the combinations abilities, it is not possible to find out when used the semi wilds or/and wild wheats as parents (Walton, 1971) due to their responses of unstabilities and changabilities. In spite of numerous and different heterosis findings were reported for heading date, plant height and weight, 1000-kernel weight, number of spike and spikelet and grain yield (Briggle, 1963; Knott, 1965; Fonseca and Patterson, 1968; Shamsuddin, 1985; Sharma and Gill, 1986; Cox and Murphy, 1990; Cregan and Bush, 1992; Picket, 1993; Larik et al. 1995; Khan and Khan, 1996; Picket and Galway, 1997; Morgan, 1998; Prasad et al. 1998; Rajaram, 2000; Bashir, 2002; Singh et al. 2004; Dreisigacker et al. 2005). There are no scientific information or literature findings on the cultivated $\times$ semi wild wheats. On the other hand, hybridization among the parents at the different ploidy level always gave interesting results in plant breeding. Especially, using or applicate these results can be bring new possibilites to the plant production, especially in wheat breeding. Aims of this paper are $i$ ) to provide and observe of the heterotic information in the combinations and progenies, ii) to detect and estimate of its
direction and magnitude, iii) to assessment of the heterotic effect in terms of the male parents.

## Material and Methods

9 females ( 7 common Triticum aestivum (L.) Em. Thell), $2 \mathrm{n}=42 ; \mathrm{P}_{5}=$ Aköz 867; $\mathrm{P}_{7}=$ Köse $220 / 39 ; \mathrm{P}_{9}=$ Penjamo 62; $\mathrm{P}_{10}=$ Sivas 111/33; $\mathrm{P}_{11}=$ Sürak 1593/51; $\mathrm{P}_{12}=$ Sertak 52; and $\mathrm{P}_{13}=$ Yektay 406 and 2 durum Triticum durum Desf., $2 \mathrm{n}=28 ; \mathrm{P}_{1}=$ Kunduru $414 / 44$ and $\mathrm{P}_{4}=$ Kunduru 1149) and 4 semi wild males ( $\mathrm{P}_{2}=T$. dicoccum, $2 \mathrm{n}=28 ; \mathrm{P}_{3}=T$. carthlicum; $\mathrm{P}_{6}=T$. vavilovii and $\mathrm{P}_{8}=T$. spelta, $2 \mathrm{n}=42$ ) were hybridized in all possible combinations without reciprocals and derived 18 combinations, 180 $\mathrm{F}_{1} \mathrm{~s}$ and (due to the non-seed setting) $150 \mathrm{~F}_{2} \mathrm{~s}$. They were grown in a randomized completely block design (RCBD) with four replications under field conditions at the University of Ankara, Faculty of Agriculture, Department of Field Crops, Experimental field during 1992/1995 in Ankara, Turkey. Males were chosen according to their differences in origin, adaptability and some dominance agromorphological characteristics such as long spike, spikelet number (Zencirci and Kün, 1996). All parents were provided from the Prof. Dr. Osman Tosun Germplasm Maintaining Bank, University of Ankara, Faculty of Agriculture, Department of Field Crops, Ankara, Turkey. Soil characteristics of the experimental site are clay structure, dark brown, $\mathrm{pH}=5.7$, lime $23.7 \%$, organic matter $1.33 \%$ and changeable potassium level is 0.028\% (Anonymous, 1999). Average temperature, rainfall and relative humidity parameters during the growing seasons are presented in Table 1. The experimental plots were designed as to be five rows of 2 m in length with the distance between rows was 20 cm and on rows was $1-2 \mathrm{~cm}$. Sowing procedure was done at the density of 250 seeds $/ \mathrm{m}^{2}$ in the beginning of October 10 competetive plants from the parents, $F_{1} s$ and $F_{2} s$ randomly selected for the observations on five agronomic characters on the basis of mean each plots as follows: PH-in cm: (measuring between the soil surface and tip of the terminal spikelet, excluding awns, on the main culm); SL-in cm (from the base of ear to the tip of terminal spikelet, excluding awns, on the main culm); NS-no (counting of the total spike number for
each plant); NSP-no (fixed by the manually counting of the total spikelet number on five randomly chosen spikes-no) and NG (mechanically harvested and balanced). The mean of the each plot was subjected to the statistical analysis, analysis of variance (ANOVA) with the MSTAT-C Statistical Software (1998) and new multiple range tests which were performed among the means as described by (Duncan, 1955). In addition, increasing ( + ) and decreasing ( - ) percentages of the MP and HP percentages were calculated as the deviation of $F_{1}$ 's value according to given formulae (Matzinger et al. 1962; Fonseca and Patterson, 1968):
[1] $\mathrm{MP}(\%)=100 \times\left(\mathrm{F}_{1}-\mathrm{MP}\right) / \mathrm{MP}$
[2] $\mathrm{HP}(\%)=100 \times\left(\mathrm{F}_{1}-\mathrm{HP}\right) / \mathrm{HP}$
Where, $\mathrm{F}_{1}=$ mean value of the trait in the hybrid, MP $=$ mean of the two parents $=\left(\mathrm{P}_{1}+\mathrm{P}_{2}\right)$ / 2 ; and $\mathrm{HP}=$ mean value of the trait in the better or high parent and calculated the heterosis values from this formulae were presented in Table 4 and 5. In addition, all obtained data were summarized as heterotic effect (+: significant or - : non significant) according to the female and male parents, and the combinations in terms of the examined traits as the MP and HP in the $\mathrm{F}_{1}$ and $\mathrm{F}_{2}$ in Table 6.

## Results

A perusal of analysis of variance indicated significant heterotic differences among the parents, combinations and the progenies for all the investigated and obtained results were presented at the Table 2 to 6 .

Heterosis for the parents: $\mathrm{P}_{7}$ was showed the significant and negative MP for the PH at the $\mathrm{F}_{1}$ and the significant and positive for the NSP at the $\mathrm{F}_{2}$. Similarly, $\mathrm{P}_{11}$ for the SL at the $\mathrm{F}_{2}, \mathrm{P}_{4}$ for the NSP at the $\mathrm{F}_{1}$, and $\mathrm{P}_{13}$ for the NG at the $\mathrm{F}_{1}$ were also expressed the significant and positive MP; however, $\mathrm{P}_{7}$ was displayed the significant HP for the NS at the $\mathrm{F}_{1}$ and $\mathrm{F}_{2}$, and the NSP at the $\mathrm{F}_{2}$ (Table 2, 3 and 6 ). $\mathrm{P}_{13}$ and $\mathrm{P}_{4}$ were also showed the statistically significant HP for the NG at the $\mathrm{F}_{1}$ and for the NS, respectively (Table $6)$. Out of nine female parents, $\mathrm{P}_{13}, \mathrm{P}_{11}, \mathrm{P}_{7}, \mathrm{P}_{4}$ were showed the statistifically significant and positive MP; and, $\mathrm{P}_{13}, \mathrm{P}_{7}, \mathrm{P}_{4}$, were exhibited HP. From them, $\mathrm{P}_{11}$ and $\mathrm{P}_{7}$ were effective on the PH and NS at the $\mathrm{F}_{1}$ and $\mathrm{F}_{2}, \mathrm{P}_{13}, \mathrm{P}_{9}$ and $\mathrm{P}_{4}$ have also been maximized of the $\mathrm{SL}, \mathrm{NG}$ and NSP (Table 6). These results are agreement
with Shamsuddin, (1990) and Sharma et al.(1984). Common wheats much more affected and contributed the heterosis and dry matter yield within the parents in this research.

Heterosis for the semi wilds: $\mathrm{P}_{8}$ was exhibited the significant and negative MP heterosis for the PH at the $\mathrm{F}_{1}$ (Table 6). In spite of $P_{3}$ and $P_{6}$ (for the NSP at the $F_{1}$ and $F_{2}$ ), they were also expressed the significant and positive MP (Table 6). But, $\mathrm{P}_{8}$ was displayed the significant and positive HP for the NS and NG at the $F_{1}$ and the $F_{2}$, respectively (Table 6). Similarly, $\mathrm{P}_{6}$ has shown the significant and positive HP for the NSP at the $\mathrm{F}_{2}$ and for the NG at $\mathrm{F}_{1}$ (Table 6). $\mathrm{P}_{3}$ had also a significant HP for the NSP at the $\mathrm{F}_{1}$ (Table 6). Semi wilds were showed three distinc groups for the contribution in terms of heterotic effect according to examined agro-morphological traits in this research such as $i$ ) $\mathrm{P}_{6}$ for the NG at the $\mathrm{F}_{1}$ and $\left.\mathrm{F}_{2}, i i\right) \mathrm{P}_{3}$ for the PH and NSP at the $\mathrm{F}_{1}$ and iii) $\mathrm{P}_{8}$ for the SL at the $\mathrm{F}_{1}$ and for the NS at the $F_{1}$ and $F_{2}$ (Table 6). In addition, it was seen that $P_{6}$ and $P_{8}$ are common participatory female parents for the all traits (Table 6).

Heterosis for the combinations: In the research, seven combinations; namely, $\mathrm{P}_{10} \times \mathrm{P}_{6}$ for the $\mathrm{PH} ; \mathrm{P}_{9} \times \mathrm{P}_{6}, \mathrm{P}_{11} \times \mathrm{P}_{3}$ and $\mathrm{P}_{11} \times \mathrm{P}_{6}$ for the SL; $\mathrm{P}_{10} \times \mathrm{P}_{6}, \mathrm{P}_{7} \times \mathrm{P}_{6}$ for the NSP and $\mathrm{P}_{1} \times \mathrm{P}_{3}$ for the NG were showed the statistically significant and positive MP; two combinations $\mathrm{P}_{7} \times \mathrm{P}_{8}$ for the NS and $\mathrm{P}_{7} \times \mathrm{P}_{6}$ for the NSP were displayed the statistically significant and positive HP (Table 2, 3 and 6). All examined traits were exhibited the significant and positive MP but two combinations ( $\mathrm{P}_{7} \times \mathrm{P}_{8}$ and $\mathrm{P}_{7} \times \mathrm{P}_{6}$ ) were showed the significant and negative HP for the NS and NSP (Table 2, 3 and 6). Only, $\mathrm{P}_{4} \times \mathrm{P}_{3}$ showed both significant and positive MP and HP at the $F_{1}$ and $F_{2}$ for the SL (Table 6).

Heterosis for the examined traits: Mean values were found significantly for the all traits except for NS, SL and NG (Table 2, 3 and 6). The MP and HP mean values were varied for the $\mathrm{PH},-26.6$ to $16.3 \%$ for $\mathrm{F}_{1}$ and 28.0 to $11.8 \%$ for $\mathrm{F}_{2} ; 28.1$ to $-13.4 \%$ for $\mathrm{F}_{1}$ and 12.8 to $-19.3 \%$ for $\mathrm{F}_{2}$, respectively (Table 2 and 3). For the SL, 51.8 to $29.4 \%$ and 65.5 to $-66.7 \%$ for $\mathrm{F}_{1} ; 32.3$ to $-49.7 \%$ and 42.3 to $-62.0 \%$ for $\mathrm{F}_{1}$, relatively (Table 2 and 3). For the NS, 152.6 to $-21.4 \%$ for $F_{1}$ and 148.4 and $-47.6 \%$ for $F_{2}$, respectively (Table 2 and 3). For the NSP, 20.0 to $-9.2 \%$ for $F_{1} 91.6$ to $-8.4 \%$ for $F_{2}$,
respectively; 18.3 to $-23.0 \%$ for $\mathrm{F}_{1}$ and 54.3 to $73.6 \%$ for $\mathrm{F}_{2}$, respectively (Table 2 and 3). For the NG, 69.0 to $-33.1 \%$ for $\mathrm{F}_{1}$ and 86.3 to $45.9 \%$ for $\mathrm{F}_{2}$, respectively; 44.0 to $-50.3 \%$ for $F_{1}$ and 2.4 to $-58.3 \%$ for $F_{2}$, respectively (Table 2 and 3). Prasad et al. (1998) were found the maximum positively HP heterosis for the NS and NG between -25.1 to $16.9 \%$; for the SL between -33.0 to $26.7 \%$, respectively. Individually, all observed heterosis the MP and HP evolution can be summarized as follows according to examined agro-morphological traits and combinations (Table 2,3 and 6):

Plant height ( PH ): Maximum and minimum values with standardized deviations are $90.40 \pm 0.42 \mathrm{~cm} \mathrm{P}_{7} \times \mathrm{P}_{8}$ and $120.10 \pm 0.28 \mathrm{~cm}$ $\mathrm{P}_{11} \times \mathrm{P}_{3}$ at the $\mathrm{F}_{1} ; 92.80 \pm 0.26 \mathrm{~cm} \mathrm{P}_{12} \times \mathrm{P}_{8}$ and $137.84 \pm 0.78 \mathrm{~cm} \mathrm{P}_{11} \times \mathrm{P}_{6}$ at the $\mathrm{F}_{2}$ (Table 2 and 3). The MP was varied between $-26.6 \mathrm{P}_{12} \times \mathrm{P}_{8}$ at the $\mathrm{F}_{1}$ to $20.8 \% \mathrm{P}_{4} \times \mathrm{P}_{2}$ at the $\mathrm{F}_{2}$ (Table 2 and 3). For the HP it was realized between $-28.2 \mathrm{P}_{12} \times \mathrm{P}_{8}$ at the $\mathrm{F}_{1}$ to $11.9 \% \mathrm{P}_{9} \times \mathrm{P}_{6}$ at the $\mathrm{F}_{1}$ (Table 2 and 3). Totally the 33 MP ( 15 positive, 3 negative at the $F_{1}$ and 11 positive, 4 negative at the $F_{2}$ ) and the 33 HP ( 13 positive, 5 negative at the $\mathrm{F}_{1}$ and 5 positive, 10 negative at the $\mathrm{F}_{2}$ ) were recorded (Table 4). Brown et al. (1966) were informed $105.0 \%$ MP and $101.0 \%$ HP at the $\mathrm{F}_{1}$; Amaya et al. (1972) were determined the MP and HP between 6.3 to $3.5 \%$ at the $\mathrm{F}_{1}$ and 5.1 to $0.7 \%$ at the $\mathrm{F}_{2}$, respectively; Güler and Özgen, (1994) were obtained the MP and HP between -7.0 to $6.0 \%$ and -12.0 to $-2.0 \%$ at the $\mathrm{F}_{1}$, respectively; Sun et al. (1999) were reported the MP is $11.0 \%$ and the HP is $0.3 \%$ at the $\mathrm{F}_{1}$; Luo et al. (2000) were recorded them ( 13.0 to $1.3 \%$ ) for the MP at the $\mathrm{F}_{1}$ and ( 3.3 to $0.3 \%$ ) for HP at the $F_{2}$. Halloran, (1975) recorded variable heterosis of 10 to $82.0 \%$ over the MP and the HP. He was added that heterosis rarely exceed $1.0 \%$ of the MP in the PH. Larik et al. (1995) were reported that heterosis values for this trait were ranged from 0.9 to $-9.2 \%$ to the MP and -1.3 to $-14.6 \%$ for the HP. Bashir (2002) was estimated as $15.11 \%$ over the MP and $0.21 \%$ over the HP.

Spike length (SL): Minimum and maximum mean values with the standardized deviations are observed as $7.20 \pm 0.65 \mathrm{~cm} \mathrm{P}_{1} \times \mathrm{P}_{3}$ and $13.1 \pm 1.82 \mathrm{~cm} \mathrm{P}_{13} \times \mathrm{P}_{8}$ at the $\mathrm{F}_{1}$ while $5.08 \pm 0.87$ $\mathrm{cm} \mathrm{P}_{5} \times \mathrm{P}_{3}$ and $15.78 \pm 1.47 \mathrm{~cm} \mathrm{P}_{11} \times \mathrm{P}_{6}$ at the $\mathrm{F}_{2}$ (Table 2 and 3). The MP ranged between - 49.5 $\mathrm{P}_{12} \times \mathrm{P}_{8}$ at the $\mathrm{F}_{2}$ to $51.8 \% \mathrm{P}_{7} \times \mathrm{P}_{6}$ at the $\mathrm{F}_{1}$ and HP varied between -62.0 $\mathrm{P}_{12} \times \mathrm{P}_{8}$ to $150.0 \%$
$\mathrm{P}_{7} \times \mathrm{P}_{8}$ at the $\mathrm{F}_{2}$ (Table 2 and 3). Totally the 33 MP (14 positive, 4 negative in the $\mathrm{F}_{1}$ and 5 positive, 10 negative at the $F_{2}$ ) and the $33 \mathrm{HP}(8$ positive, 10 negative at the $\mathrm{F}_{1}$ and 7 positive, 8 negative at the $F_{2}$ ) were fixed (Table 4). Gyawali et al. (1968) obtained heterosis ratios for the MP and HP that lied between -1.0 to $18.0 \%$ at the $\mathrm{F}_{1}$ and between -3.0 to $17.0 \%$ at $F_{1}$, respectively. In general, obtained results are in accordance with those obtained by Singh, (1978) who reported $10 \%$ increase over the MP for the SL. It is possible to develop new wheat cultivars or hybrid combinations having long spike but dwarf PH (Li et al. 1997).

Number of spike per plant (NS): For this trait, mean values with the standardized deviations were found as $3.3 \pm 1.17 \mathrm{P}_{4} \times \mathrm{P}_{3}$ and $12.0 \pm 3.19 \mathrm{P}_{7} \times \mathrm{P}_{8}$ at the $\mathrm{F}_{1} ; 2.2 \pm 0.86 \mathrm{P}_{5} \times \mathrm{P}_{3}$ and $11.8 \pm 0.11 \mathrm{P}_{7} \times \mathrm{P}_{8}$ at the $\mathrm{F}_{2}$ (Table 2 and 3). The MP and the HP were varied between -96.6 $\mathrm{P}_{5} \times \mathrm{P}_{6}$ at the $\mathrm{F}_{2}$ to $84.0 \% \mathrm{P}_{1} \times \mathrm{P}_{3}$ at the $\mathrm{F}_{1}$ and $67.3 \mathrm{P}_{5} \times \mathrm{P}_{6}$ to $185.6 \% \mathrm{P}_{7} \times \mathrm{P}_{8}$ at the $\mathrm{F}_{2}$ (Table 2 and 3). Totally, the 33 MP ( 16 positive, 2 negative in the $F_{1}$ and 7 positive, 8 negative at the $\mathrm{F}_{2}$ ) and the 33 HP ( 7 positive, 11 negative at the $F_{1}$ and 5 positive, 10 negative at the $F_{2}$ ) were fixed (Table 4). Fonseca and Patterson (1968) were reported for MP between 93.0 to $119.0 \%$ and 90.0 to $110.0 \%$ at the $F_{1}$. Winzeler et al. (1994) were determined the HP between 2.37 to $2.22 \%$ at the $\mathrm{F}_{1}$; Gyawali et al. (1968) were obtained the MP and HP estimates between -40.0 to $-21.0 \%$ and -48.0 to $-23.0 \%$ at the $F_{1}$, respectively. Similar results have been reported and confirmed by Chakraborty and Tiwari, (1995) and Larik et al. (1987) and Larik et al. (1995) for the NS.

Number of spikelets per spike (NSP): Minimum and maximum values with the standardized deviations for the number of spikelets are $17.9 \pm 0.59 \mathrm{P}_{10} \times \mathrm{P}_{8}$ and $22.2 \pm 0.93$ $\mathrm{P}_{7} \times \mathrm{P}_{6}$ at the $\mathrm{F}_{1}$ while $18.6 \pm 0.17 \mathrm{P}_{13} \times \mathrm{P}_{8}$ and $36.6 \pm 2.09 \mathrm{P}_{7} \times \mathrm{P}_{6}$ at the $\mathrm{F}_{2}$ (Table 2 and 3). The MP was ranged between -57.4 $\mathrm{P}_{5} \times \mathrm{P}_{3}$ at the $\mathrm{F}_{1}$ to $93.7 \% \quad \mathrm{P}_{5} \times \mathrm{P}_{6}$ at the $\mathrm{F}_{2}$; and the HP was varied between $-57.5 \mathrm{P}_{5} \times \mathrm{P}_{3}$ at $\mathrm{F}_{1}$ and $71.0 \%$ $\mathrm{P}_{7} \times \mathrm{P}_{6}$ at the $\mathrm{F}_{2}$ (Table 2 and 3). Totally, the 33 MP ( 15 positive, 3 negative at the $F_{1}$ and 13 positive, 2 negative at the $F_{2}$ ) and the 33 HP (11 positive, 7 negative and 12 positive, 3 negative at the $F_{2}$ ) were obtained (Table 4). Fonseca and Patterson, (1968) determined the MP and HP values between 1.1 at the $\mathrm{F}_{1}$ to
$0.8 \%$ at the $F_{2}$ and 3.7 at the $F_{1}$ and $4.3 \%$ at $F_{2}$ for this trait, respectively. Obtained findings were supported and confirmed by Singh et al. (2004); but not Prasad et al. (1972) who did not find any significant amount of the HP in the combinations. The NSP proved to be an important trait and may explain the superiority of the hybrids over the traditional varieties.

Number of grain per spikelet (NG): This trait is the most important yield determinant of wheat (Sinclair and Jamieson, 2006) and it products of plants per $\mathrm{m}^{2}$, ears per plant and grains per spike. The overall means with the standardized deviations of the crosses were varied between $27.4 \pm 0.28 \mathrm{P}_{12} \times \mathrm{P}_{8}$ to $50.8 \pm 0.66$ $\mathrm{P}_{9} \times \mathrm{P}_{6}$ at the $\mathrm{F}_{1}$ and $14.4 \pm 0.20 \quad \mathrm{P}_{11} \times \mathrm{P}_{8}$ to $59.6 \pm 0.79 \mathrm{P}_{11} \times \mathrm{P}_{6}$ at the $\mathrm{F}_{2}$ (Table 2 and 3 ). The MP was varied between $-25.0 \mathrm{P}_{13} \times \mathrm{P}_{8}$ to $87.3 \%$ $\mathrm{P}_{13} \times \mathrm{P}_{6}$ only at the $\mathrm{F}_{1}$; and the HP was ranged between -85.6 $\mathrm{P}_{10} \times \mathrm{P}_{6}$ at the $\mathrm{F}_{2}$ and $50.3 \%$ $\mathrm{P}_{12} \times \mathrm{P}_{6}$ at the $\mathrm{F}_{1}$ (Table 2 and 3). Totally, the 33 MP ( 15 positive, 3 negative at the $\mathrm{F}_{1}$ and 13 positive, 8 negative at the $\mathrm{F}_{2}$ ) and the 34 HP ( 8 positive, 10 negative at the $\mathrm{F}_{1}$ and 4 positive, 12 negative at the $F_{2}$ ) were fixed (Table 4). These results were in accordance with Garcia et al. (1991) who reported MP is $4.0 \%$ in the $\mathrm{F}_{1}$ and $4.3 \%$ in the $\mathrm{F}_{2}$ and HP is $4.3 \%$ in the $\mathrm{F}_{1}$ and $7.3 \%$ in the $\mathrm{F}_{2}$.

Correlation coefficients: The highest positive significant correlation coefficient was found between (MP-F2) and NS, (r=0.570) and the lowest positive significant correlation coefficient was found between ( $\mathrm{P}_{2}-\mathrm{F}_{1}$ ) and NS, ( $\mathrm{r}=0.380$ ). Four negative but non significant coefficients were found among ( $\mathrm{P}_{1}-\mathrm{F}_{1}$ ) and NG , ( $\mathrm{r}=-0.126$ ); ( $\mathrm{P}_{1}-\mathrm{F}_{2}$ ) and NG, ( $\mathrm{r}=-0.050$ ); ( $\mathrm{P}_{2}-\mathrm{F}_{2}$ ) and NS, ( $\mathrm{r}=-0.250$ ) and ( $\mathrm{HP}-\mathrm{F}_{1}$ ) and NG, ( $\mathrm{r}=-$ 0.160). Among $\left(\mathrm{P}_{1}-\mathrm{F}_{1}\right)$ and SL , $(\mathrm{r}=0.495$, $\mathrm{p}>0.01)$; $\left(\mathrm{P}_{1}-\mathrm{F}_{1}\right)$ and NSP, ( $\left.\mathrm{r}=0.500, \mathrm{p}>0.01\right)$; $\left(\mathrm{P}_{2}-\mathrm{F}_{2}\right)$ and NSP, $(\mathrm{r}=0.200, \mathrm{p}>0.01)$; $\left(\mathrm{MP}-\mathrm{F}_{1}\right)$ and PH , $(\mathrm{r}=0.450, \mathrm{p}>0.01)$; ( $\mathrm{MP}-\mathrm{F}_{1}$ ) and SL , ( $\mathrm{r}=0.570, \mathrm{p}>0.01$ ); $\left(\mathrm{MP}-\mathrm{F}_{1}\right)$ and NS, $(\mathrm{r}=0.550$, $\mathrm{p}>0.01)$; $\left(\mathrm{MP}-\mathrm{F}_{1}\right)$ and NSP, $(\mathrm{r}=0.540, \mathrm{p}>0.01)$; $\left(\mathrm{MP}_{2}\right)$ and NS, (r=0.625, p>0.01); (MP-F ${ }_{2}$ ) and $\mathrm{NG},(\mathrm{r}=0.125, \mathrm{p}>0.05)$ and $\left(\mathrm{HP}-\mathrm{F}_{1}\right)$ and NS, ( $\mathrm{r}=0.450, \mathrm{p}>0.01$ ); ( $\mathrm{HP}-\mathrm{F}_{2}$ ) and NSP, ( $\mathrm{r}=0,380, \mathrm{p}>0.01$ ); $\left(\mathrm{HP}-\mathrm{F}_{2}\right)$ and $\mathrm{NG},(\mathrm{r}=0.150$, $\mathrm{p}>0.05$ ) (Table 5). These correlations verified and supported to our research results. In the light of these findings, it can be suggested that always the MP must be preferred and for the NS and NG (for both $\mathrm{F}_{1}$ and $\mathrm{F}_{2}$ ). Bai et al.
(1992) and Zhang et al. (1985) are being supported this judgment with their research findings. Similarly, Shamsuddin, (1985) was done the identical studies and he was reported the similar significant and positive associations among the investigated various morphological, genetical and agronomic characters in his study with 20 winter and spring wheat hybrids ( $\mathrm{r}=0.460$ and $\mathrm{r}=0.450$, respectively; Kaltsikes and Larter, (1970) were also found a significant and positive correlation between the PH and yield in wheat.

## Discussion

Female and male parents can be ordered according to their heterotic effect (heterosis) from the maximum to minumum as $\mathrm{P}_{4}>\mathrm{P}_{7}>\mathrm{P}_{10}>\mathrm{P}_{5}>\mathrm{P}_{9}>\mathrm{P}_{13}>\mathrm{P}_{11}>\mathrm{P}_{12}>\mathrm{P}_{1} \quad$ and $\mathrm{P}_{3}>\mathrm{P}_{6}>\mathrm{P}_{8}>\mathrm{P}_{2}$, respectivey in this research (Table 6). $\mathrm{P}_{6}$ and $\mathrm{P}_{8}$ are the common participitory female parents for the traits. Highest significant and positive MP was taken from the $\mathrm{P}_{7} \times \mathrm{P}_{8}$ for the NS at the $\mathrm{F}_{1}$ with $152.6 \%$ and for HP was also taken from the same combintion with $135.3 \%$ for the NS at the $\mathrm{F}_{1}$ (Table 4). It may have been the result of interaction between the parental cytoplasm (genotypic diversity) and applied agronomic practices or environmental effects. It was seen that the more genetic diversity the more and violently MP. This testimony was observed and verified by Shamsuddin, (1985). Semi wilds showed less heterosis than the cultivars; but much more affected from the PH for the MP and SL for the HP; however, not at the $\mathrm{F}_{1}$ and $\mathrm{F}_{2}$. According to data, it can be suggested that only MP values must be used without paying the attention for the $\mathrm{F}_{1}$ or $\mathrm{F}_{2}$ to be able to get a higher hybrid wheat yield. This case also been reported by Briggle, 1967; Singh and Singh, 1971; Krishna and Ahmad, 1992; Picket, 1993; Sinclair and Jamieson, 2006). On the other hand, it showed that the gene pool of spelt wheat $\left(\mathrm{P}_{8}\right)$ could be potentially useful for the selection of parents for the hybrid wheat production. This result verified by Winzeler et al. 1993). But, there are several problems that to be solved and needs to be more improved so despite the very high heterosis effect no short statued varieties exist at the present, close and hard, tightness of the glumes pollination or pollination easly could be hamperring if used as a pollinator. However, a breeding effort to
select for improved parents within the gene pool of spelt appears useful for future $F_{1}$ hybrids between spelt and wheat. These findings can be applied to the hybrid wheat breeding as a cultivars x semi wild wheats. First of all, plant height traits can be shortened with the heterotic effect of this type hybridization. Obtained resits research showed that to be able to get maximum level of heterosis is possible only for MP at the level of $\mathrm{F}_{1}$. If this process continue to the out of this limits (esp. for HP at the $F_{1}$ or $F_{2}$ ) yield level is being celarly decreased ! On the other hand, with their genetical backgrounds or by genepools semiwild wheats can be use against to stress conditions such as disease and pest resistance.

Similarly, with the addition of the modern biotechnological approaches apomixis might be hope to facilitate seed production; llinkage maps and DNA based molecular markers can be considered a promising tool to predict hybrid performance, at the same time it could be a powerful tool for the identification of genetically diversified sources such as wild or semi wild wheats in the long run.

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Table 1. Meteorological data for 1992-1995 growing seasons (monthly average).

|  | Average Temperature ( ${ }^{\circ} \mathrm{C}$ ) |  |  |  |  | Rainfall (mm) |  |  |  |  | Relative Humidity (\%) |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Years |  |  |  |  | Years |  |  |  |  | Years |  |  |  |  |
| Months | 1992 | 1993 | 1994 | 1995 | $\underset{1926-92}{\text { LTA }^{\S}}$ | 1992 | 1993 | 1994 | 1995 | $\begin{array}{r} \text { LTA }^{\S} \\ 1926-92 \end{array}$ | 1992 | 1993 | 1994 | 1995 | $\begin{array}{r} \text { LTA }^{\S} \\ 1926-92 \end{array}$ |
| January | -4.0 | -4.0 | 3.8 | 3.3 | -0.1 | 4.9 | 28.8 | 30.2 | 33.6 | 40.9 | 76.5 | 79.5 | 75.5 | 33.6 | 78.0 |
| February | -3.0 | -0.7 | 1.8 | 5.2 | 1.3 | 5.7 | 33.4 | 33.6 | 10.8 | 34.9 | 69.5 | 72.8 | 74.2 | 10.8 | 74.0 |
| March | 3.9 | 5.7 | 6.8 | 6.7 | 5.4 | 50.3 | 22.4 | 18.4 | 92.6 | 35.6 | 70.1 | 61.3 | 60.1 | 92.6 | 65.0 |
| April | 11.4 | 10.4 | 14.0 | 9.9 | 11.2 | 40.2 | 28.1 | 30.7 | 61.6 | 40.3 | 59.0 | 56.5 | 55.0 | 61.6 | 59.0 |
| May | 16.2 | 15.3 | 17.0 | 17.6 | 15.9 | 1.6 | 88.1 | 39.0 | 30.8 | 51.3 | 45.4 | 65.1 | 56.5 | 30.8 | 57.0 |
| June | 19.0 | 19.7 | 20.6 | 21.8 | 19.8 | 54.9 | 13.1 | 6.6 | 60.8 | 32.6 | 57.6 | 52.5 | 47.2 | 60.8 | 51.0 |
| July | 20.5 | 22.9 | 24.2 | 23.1 | 23.1 | 29.9 | 3.5 | 5.0 | 10.7 | 13.5 | 55.4 | 45.3 | 44.4 | 10.7 | 44.0 |
| August | 23.4 | 23.1 | 23.5 | 23.4 | 23.0 | 19.9 | 11.1 | 1.1 | 3.7 | 10.3 | 46.0 | 49.7 | 46.7 | 3.7 | 42.0 |
| September | 16.7 | 19.3 | 22.8 | 19.0 | 18.4 | 2.6 | 11.1 | 6.3 | 12.7 | 17.4 | 52.0 | 49.7 | 44.0 | 12.7 | 47.0 |
| October | 15.1 | 15.0 | 16.0 | 11.6 | 12.8 | 35.1 | 1.8 | 30.0 | 27.8 | 24.4 | 60.1 | 45.8 | 60.9 | 27.8 | 58.0 |
| November | 5.4 | 4.1 | 5.6 | 3.4 | 7.3 | 47.0 | 33.6 | 67.5 | 61.6 | 36.4 | 68.4 | 65.7 | 75.0 | 61.6 | 70.0 |
| December | -0.6 | 4.0 | 0.5 | 2.4 | 2.3 | 37.9 | 33.0 | 20.6 | 52.3 | 45.6 | 77.4 | 76.6 | 78.8 | 52.3 | 78.0 |

${ }^{\text {§ }}$ Long term averages.;
Source: Republic of the Environment and Forest Ministry, General Directorate of Meteorology

Table 2. Means with the standardized errors in the combinations and parents in $F_{1}$ and $F_{2}$.

| Combinations and Parents | PH (cm) |  | SL (cm) |  | NS (no) |  | NSP (no) |  | NG (no) |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | $\mathrm{F}_{1}$ | $\mathrm{F}_{2}$ | $\mathrm{F}_{1}$ | $\mathrm{F}_{2}$ | $\mathrm{F}_{1}$ | $\mathrm{F}_{2}$ | $\mathrm{F}_{1}$ | $\mathrm{F}_{2}$ | $\mathrm{F}_{1}$ | $\mathrm{F}_{2}$ |
| $\mathbf{P}_{1} \times \mathrm{P}_{2}$ | $104.21 \pm 0.06 \mathrm{~A}$ | - | $8.1 \pm 0.28$ B | - | $5.8 \pm 1.56$ B | - | $20.7 \pm 0.85 \mathrm{~A}$ | - | $39.0 \pm 4.01 \mathrm{~B}$ | - |
| $\mathrm{P}_{1}$ | $93.3 \pm 0.13 \mathrm{CD}$ |  | $7.4 \pm 0.43 \mathrm{C}$ |  | $3.4 \pm 0.11 \mathrm{C}$ |  | $21.3 \pm 0.63 \mathrm{~A}$ |  | $40.3 \pm 0.15$ B |  |
| $\mathrm{P}_{2}$ | $85.9 \pm 0.45 \mathrm{E}$ |  | $5.6 \pm 0.25 \mathrm{E}$ |  | $5.9 \pm 0.24 \mathrm{~B}$ |  | $20.1 \pm 0.51 \mathrm{~A}$ |  | $22.3 \pm 0.12 \mathrm{E}$ |  |
| $\mathbf{P}_{1} \times \mathbf{P}_{3}$ | $109.57 \pm 0.25 \mathrm{~A}$ | $119.24 \pm 0.30 \mathrm{~A}$ | $7.2 \pm 0.65 \mathrm{C}$ | $7.38 \pm 0.64$ B | $4.6 \pm 1.01 \mathrm{~B}$ | 5.6 $\pm 0.08 \mathrm{~B}$ | $19.8 \pm 0.50 \mathrm{~A}$ | $30.4 \pm 3.61 \mathrm{~A}$ | $39.6 \pm 0.88$ B | $30.4 \pm 0.21 \mathrm{D}$ |
| $\mathrm{P}_{1}$ | $100.64 \pm 0.82 \mathrm{CD}$ |  | $6.5 \pm 0.30 \mathrm{D}$ |  | $2.0 \pm 0.37 \mathrm{D}$ |  | $19.5 \pm 0.24 \mathrm{~A}$ |  | $30.1 \pm 0.63 \mathrm{D}$ |  |
| $\mathrm{P}_{3}$ | $105.63 \pm 0.25 \mathrm{CD}$ |  | $6.9 \pm 0.82 \mathrm{D}$ |  | $3.0 \pm 0.88 \mathrm{C}$ |  | $19.7 \pm 0.36 \mathrm{~A}$ |  | $33.5 \pm 0.47 \mathrm{C}$ |  |
| $\mathbf{P}_{4} \times \mathbf{P}_{3}$ | $118.87 \pm 0.40 \mathrm{AD}$ | $108.60 \pm 0.61 \mathrm{C}$ | $7.9 \pm 0.93 \mathrm{C}$ | $7.36 \pm 0.26$ B | $3.3 \pm 1.17 \mathrm{C}$ | $5.2 \pm 0.24$ B | $22.2 \pm 0.93 \mathrm{~A}$ | $21.4 \pm 2.06 \mathrm{C}$ |  | $21.8 \pm 0.69 \mathrm{DE}$ |
| $\mathrm{P}_{4}$ | $111.87 \pm 0.11 \mathrm{D}$ |  | $6.2 \pm 0.10 \mathrm{D}$ |  | $3.7 \pm 0.46 \mathrm{C}$ |  | $19.7 \pm 0.18 \mathrm{~A}$ |  | $32.4 \pm 0.26 \mathrm{C}$ |  |
| $\mathrm{P}_{3}$ | $110.78 \pm 0.36 \mathrm{D}$ |  | $7.0 \pm 0.65 \mathrm{C}$ |  | $4.7 \pm 0.52 \mathrm{C}$ |  | $19.8 \pm 0.20 \mathrm{~A}$ |  | $32.4 \pm 0.55 \mathrm{C}$ |  |
| $\mathbf{P}_{4} \times \mathrm{P}_{2}$ | $113.75 \pm 0.92 \mathrm{~A}$ | $105.16 \pm 0.85 \mathrm{C}$ | $9.6 \pm 0.57 \mathrm{~B}$ | $6.90 \pm 0.50 \mathrm{C}$ | $4.6 \pm 0.58 \mathrm{C}$ | $4.8 \pm 0.80$ B | $21.0 \pm 0.67 \mathrm{~A}$ | $19.4 \pm 1.81 \mathrm{C}$ | $33.0 \pm 0.25 \mathrm{C}$ | $30.4 \pm 0.63$ D |
| $\mathbf{P}_{4}$ | $109.30 \pm 0.15 \mathrm{D}$ |  | $8.9 \pm 0.97 \mathrm{~B}$ |  | $2.8 \pm 0.19 \mathrm{D}$ |  | $20.0 \pm 0.25 \mathrm{~A}$ |  | $33.7 \pm 0.37 \mathrm{C}$ |  |
| $\mathrm{P}_{2}$ | $94.90 \pm 0.39 \mathrm{CD}$ |  | $6.5 \pm 0.33 \mathrm{D}$ |  | $5.3 \pm 0.26$ B |  | $19.6 \pm 0.30 \mathrm{~A}$ |  | $33.4 \pm 0.28 \mathrm{C}$ |  |
| $\mathbf{P}_{5} \times \mathbf{P}_{3}$ | $117.12 \pm 0.04 \mathrm{~A}$ | $104.48 \pm 0.23 \mathrm{C}$ | $8.1 \pm 0.74$ B | $5.08 \pm 0.87 \mathrm{C}$ | $4.3 \pm 1.24 \mathrm{C}$ | $2.2 \pm 0.86 \mathrm{D}$ | $18.8 \pm 0.14$ B | $21.6 \pm 1.40 \mathrm{C}$ | $43.4 \pm 0.74 \mathrm{~A}$ | $26.0 \pm 0.19 \mathrm{DE}$ |
| $\mathrm{P}_{5}$ | $112.70 \pm 0.58$ B |  | $8.6 \pm 0.50 \mathrm{~B}$ |  | $4.1 \pm 0.13 \mathrm{C}$ |  | $20.3 \pm 0.19 \mathrm{~A}$ |  | $35.2 \pm 0.66 \mathrm{BC}$ |  |
| $\mathbf{P}_{3}$ | $109.58 \pm 0.81 \mathrm{C}$ |  | $7.5 \pm 0.29 \mathrm{C}$ |  | $4.3 \pm 0.36 \mathrm{C}$ |  | $21.1 \pm 0.17 \mathrm{~A}$ |  | $38.0 \pm 0.90$ B |  |
| $\mathbf{P}_{5} \times \mathrm{P}_{6}$ | $114.36 \pm 0.67 \mathrm{~A}$ | $109.46 \pm 0.56 \mathrm{C}$ | $11.0 \pm 1.21 \mathrm{~A}$ | $11.52 \pm 1.49 \mathrm{~A}$ | $4.1 \pm 1.40 \mathrm{~B}$ | $3.4 \pm 0.51 \mathrm{C}$ | $20.4 \pm 0.55 \mathrm{~A}$ | $33.2 \pm 2.92 \mathrm{~A}$ | $41.0 \pm 0.59 \mathrm{~A}$ | $43.0 \pm 0.60 \mathrm{C}$ |
| $\mathrm{P}_{5}$ | $111.32 \pm 0.64 \mathrm{~B}$ |  | $8.1 \pm 0.13 \mathrm{~B}$ |  | $2.0 \pm 0.18 \mathrm{D}$ |  | $20.1 \pm 0.50 \mathrm{~A}$ |  | $42.0 \pm 0.13 \mathrm{~A}$ |  |
| $\mathrm{P}_{6}$ | $116.50 \pm 0.25 \mathrm{~A}$ |  | $10.5 \pm 0.24 \mathrm{~A}$ |  | $5.0 \pm 0.40 \mathrm{~B}$ |  | $16.6 \pm 0.65$ B |  | $34.0 \pm 0.66 \mathrm{BC}$ |  |
| $\mathbf{P}_{7} \times \mathbf{P}_{6}$ | $116.19 \pm 0.29 \mathrm{~A}$ | $120.06 \pm 0.15$ A | $12.9 \pm 1.72 \mathrm{~A}$ | $12.90 \pm 1.40 \mathrm{~A}$ | $7.1 \pm 3.66 \mathrm{~A}$ | 5.6 $\pm 0.40 \mathrm{~B}$ | $22.0 \pm 0.10 \mathrm{~A}$ | $36.6 \pm 2.09 \mathrm{~A}$ | $38.0 \pm 0.56 \mathrm{~B}$ | $31.6 \pm 0.57 \mathrm{D}$ |
| $\mathbf{P}_{7}$ | $99.47 \pm 0.51 \mathrm{D}$ |  | $5.0 \pm 0.45 \mathrm{E}$ |  | $8.8 \pm 0.66 \mathrm{~A}$ |  | $18.4 \pm 0.80 \mathrm{~B}$ |  | $27.0 \pm 0.41 \mathrm{D}$ |  |
| $\mathrm{P}_{6}$ | $114.41 \pm 0.14 \mathrm{~A}$ |  | $12.0 \pm 0.78 \mathrm{~A}$ |  | $4.7 \pm 0.20 \mathrm{C}$ |  | $20.0 \pm 0.54 \mathrm{~A}$ |  | $39.3 \pm 0.29 \mathrm{~B}$ |  |
| $\mathbf{P}_{7} \times \mathbf{P}_{\mathbf{8}}$ | $90.40 \pm 0.42 \mathrm{C}$ | $99.24 \pm 0.62 \mathrm{D}$ | $12.1 \pm 1.14 \mathrm{~A}$ | $7.24 \pm 1.00 \mathrm{~B}$ | $12.0 \pm 3.19 \mathrm{~A}$ | $11.8 \pm 0.11 \mathrm{~A}$ | $20.4 \pm 0.72 \mathrm{~A}$ | $29.2 \pm 1.66 \mathrm{~A}$ | $33.3 \pm 0.13 \mathrm{C}$ | $28.8 \pm 0.16 \mathrm{DE}$ |
| $\mathbf{P}_{7}$ | $97.55 \pm 0.26 \mathrm{D}$ |  | $10.5 \pm 0.66 \mathrm{~A}$ |  | $5.1 \pm 0.75$ B |  | $16.0 \pm 0.34 \mathrm{~B}$ |  | $17.1 \pm 0.36 \mathrm{~F}$ |  |
| $\mathrm{P}_{8}$ | $95.35 \pm 0.40 \mathrm{D}$ |  | $12.3 \pm 0.19 \mathrm{~A}$ |  | $4.4 \pm 0.50 \mathrm{C}$ |  | $21.0 \pm 0.10 \mathrm{~A}$ |  | $23.2 \pm 0.88$ E |  |
| $\mathrm{P}_{9} \times \mathrm{P}_{8}$ | $119.55 \pm 0.51 \mathrm{~A}$ | - | $11.7 \pm 1.33 \mathrm{~A}$ | - | $4.1 \pm 1.34 \mathrm{C}$ | - | $21.0 \pm 0.23 \mathrm{~A}$ | - | $33.2 \pm 0.40 \mathrm{C}$ | - |
| $\mathrm{P}_{9}$ | $111.15 \pm 0.36 \mathrm{~B}$ |  | $9.5 \pm 0.57 \mathrm{~B}$ |  | $3.3 \pm 0.39 \mathrm{C}$ |  | $18.8 \pm 0.61$ B |  | $31.1 \pm 0.41 \mathrm{CD}$ |  |
| $\mathbf{P}_{8}$ | $110.30 \pm 0.47 \mathrm{~B}$ |  | $13.7 \pm 0.81 \mathrm{~A}$ |  | $5.7 \pm 0.15$ B |  | $21.2 \pm 0.50 \mathrm{~A}$ |  | $31.4 \pm 0.17 \mathrm{CD}$ |  |
| $\mathrm{P}_{9} \times \mathrm{P}_{6}$ | $115.70 \pm 0.21 \mathrm{~A}$ | $116.50 \pm 0.11 \mathrm{~A}$ | $12.6 \pm 1.26 \mathrm{~A}$ | $14.38 \pm 2.02 \mathrm{~A}$ | $6.0 \pm 2.40 \mathrm{~A}$ | $10.0 \pm 0.97 \mathrm{~A}$ | $19.5 \pm 0.18 \mathrm{~A}$ | $28.4 \pm 4.43 \mathrm{~A}$ | $50.8 \pm 0.66 \mathrm{~A}$ | $48.8 \pm 0.11$ B |
| $\mathrm{P}_{9}$ | $103.27 \pm 0.78$ BC |  | $9.9 \pm 0.76 \mathrm{~B}$ |  | $5.2 \pm 0.04 \mathrm{~B}$ |  | $18.8 \pm 0.36 \mathrm{~B}$ |  | $28.8 \pm 0.53 \mathrm{D}$ |  |
| $\mathrm{P}_{6}$ | $103.43 \pm 0.53 \mathrm{BC}$ |  | $12.8 \pm 0.94 \mathrm{~A}$ |  | $5.7 \pm 0.16 \mathrm{~B}$ |  | $20.0 \pm 0.40 \mathrm{~A}$ |  | $43.6 \pm 0.25 \mathrm{~A}$ |  |
| $\mathrm{P}_{10} \times \mathrm{P}_{6}$ | $107.95 \pm 0.89 \mathrm{AB}$ | $115.38 \pm 0.61 \mathrm{~A}$ | $8.3 \pm 1.03 \mathrm{~B}$ | $9.12 \pm 0.40 \mathrm{~B}$ | $6.8 \pm 1.25 \mathrm{~A}$ | $6.0 \pm 0.10 \mathrm{~B}$ | $18.7 \pm 0.14 \mathrm{~B}$ | 26.4 $\pm 4.48$ B | $30.2 \pm 0.51 \mathrm{D}$ | $48.4 \pm 0.39$ B |
| $\mathrm{P}_{10}$ | $102.90 \pm 0.18 \mathrm{BC}$ |  | $5.1 \pm 0.65 \mathrm{E}$ |  | $9.2 \pm 0.63 \mathrm{~A}$ |  | $18.3 \pm 0.24 \mathrm{~B}$ |  | $28.4 \pm 0.64 \mathrm{D}$ |  |
| $\mathrm{P}_{6}$ | $113.00 \pm 0.77 \mathrm{~A}$ |  | $11.4 \pm 0.71 \mathrm{~A}$ |  | $4.4 \pm 0.70 \mathrm{C}$ |  | $19.0 \pm 0.68 \mathrm{~A}$ |  | $32.0 \pm 0.34 \mathrm{CD}$ |  |
| $\mathrm{P}_{10} \times \mathrm{P}_{8}$ | $92.70 \pm 0.33 \mathrm{BD}$ | $109.22 \pm 0.51 \mathrm{C}$ | $7.8 \pm 0.55 \mathrm{C}$ | $10.80 \pm 1.07 \mathrm{~B}$ | $8.5 \pm 1.86 \mathrm{~A}$ | $4.6 \pm 0.51 \mathrm{~B}$ | $17.9 \pm 0.59 \mathrm{~B}$ | $29.2 \pm 1.66 \mathrm{~A}$ | $42.3 \pm 0.88 \mathrm{~A}$ | $14.4 \pm 0.20 \mathrm{~F}$ |
| $\mathrm{P}_{10}$ | $107.00 \pm 0.41 \mathrm{AB}$ |  | $6.6 \pm 0.25 \mathrm{D}$ |  | $6.1 \pm 0.52 \mathrm{~A}$ |  | $17.9 \pm 0.53 \mathrm{~B}$ |  | $28.8 \pm 0.95 \mathrm{D}$ |  |
| $\mathrm{P}_{8}$ | $103.40 \pm 0.99 \mathrm{BC}$ |  | $15.5 \pm 0.37 \mathrm{~A}$ |  | $6.1 \pm 0.80 \mathrm{~A}$ |  | $23.2 \pm 0.32 \mathrm{~A}$ |  | $34.5 \pm 0.31 \mathrm{BC}$ |  |

Table 3. Means with the standardized errors in the combinations and parents in $F_{1}$ and $F_{2}$ (cont.).

| Combinations and Parents | PH (cm) |  | SL (cm) |  | NS (no) |  | NSP (no) |  | NG (no) |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | $\mathrm{F}_{1}$ | $\mathrm{F}_{2}$ | $\mathrm{F}_{1}$ | $\mathrm{F}_{2}$ | $\mathrm{F}_{1}$ | $\mathrm{F}_{2}$ | $\mathrm{F}_{1}$ | $\mathrm{F}_{2}$ | $\mathrm{F}_{1}$ | $\mathrm{F}_{2}$ |
| $\mathbf{P}_{11} \times \mathbf{P}_{3}$ | $120.10 \pm 0.28 \mathrm{~A}$ | $117.98 \pm 0.19 \mathrm{~A}$ | $12.5 \pm 1.48 \mathrm{~A}$ | $12.2 \pm 0.62 \mathrm{~A}$ | $8.4 \pm 1.30 \mathrm{~A}$ | $5.8 \pm 0.74 \mathrm{~B}$ | $20.0 \pm 0.90 \mathrm{~A}$ | 26.8 $\pm 2.22$ B | $30.0 \pm 0.70 \mathrm{D}$ | $17.6 \pm 0.83 \mathrm{E}$ |
| $\mathbf{P}_{11}$ | $115.41 \pm 0.42 \mathrm{~A}$ |  | $13.0 \pm 0.48 \mathrm{~A}$ |  | $6.3 \pm 0.96$ A |  | $22.0 \pm 0.45 \mathrm{~A}$ |  | $32.0 \pm 0.46 \mathrm{CD}$ |  |
| $\mathrm{P}_{3}$ | $104.92 \pm 0.86 \mathrm{BC}$ |  | $9.6 \pm 0.52 \mathrm{~B}$ |  | $7.6 \pm 0.62 \mathrm{~A}$ |  | $19.0 \pm 0.96 \mathrm{~B}$ |  | $33.0 \pm 0.51 \mathrm{C}$ |  |
| $\mathbf{P}_{11} \times \mathbf{P}_{6}$ | $111.36 \pm 0.45 \mathrm{~A}$ | $137.84 \pm 0.78 \mathrm{~A}$ | $11.0 \pm 1.23 \mathrm{~A}$ | $15.8 \pm 1.47 \mathrm{~A}$ | $8.7 \pm 2.13 \mathrm{~A}$ | $6.6 \pm 0.98$ B | $21.0 \pm 0.36 \mathrm{~A}$ | $23.0 \pm 2.53 \mathrm{C}$ | $32.0 \pm 0.31 \mathrm{CD}$ | $59.6 \pm 0.79 \mathrm{~A}$ |
| $\mathrm{P}_{11}$ | $106.20 \pm 0.88 \mathrm{AB}$ |  | $11.1 \pm 0.55 \mathrm{~A}$ |  | $8.0 \pm 0.46 \mathrm{~A}$ |  | $18.8 \pm 0.55$ B |  | $31.0 \pm 0.42 \mathrm{CD}$ |  |
| $\mathrm{P}_{6}$ | $109.10 \pm 0.54 \mathrm{AB}$ |  | $8.0 \pm 0.80 \mathrm{~B}$ |  | $4.8 \pm 0.17 \mathrm{C}$ |  | $20.0 \pm 0.91 \mathrm{~A}$ |  | $33.0 \pm 0.40 \mathrm{C}$ |  |
| $\mathrm{P}_{12} \times \mathrm{P}_{6}$ | $113.60 \pm 0.29 \mathrm{~A}$ | - | $9.0 \pm 0.45$ B | - | $5.3 \pm 0.29$ B | - | $21.3 \pm 0.15 \mathrm{~A}$ | - | $45.1 \pm 0.46 \mathrm{~A}$ | - |
| $\mathrm{P}_{12}$ | $98.43 \pm 0.26 \mathrm{C}$ |  | $8.7 \pm 0.14$ B |  | $6.0 \pm 0.59 \mathrm{~A}$ |  | $17.5 \pm 0.21 \mathrm{~B}$ |  | $23.4 \pm 0.23 \mathrm{E}$ |  |
| $\mathrm{P}_{6}$ | $115.76 \pm 0.13$ B |  | $10.5 \pm 0.65 \mathrm{~A}$ |  | $4.4 \pm 0.66 \mathrm{C}$ |  | $18.0 \pm 0.13 \mathrm{~B}$ |  | $30.0 \pm 0.37 \mathrm{D}$ |  |
| $\mathrm{P}_{12} \times \mathrm{P}_{8}$ | $105.20 \pm 0.06 \mathrm{~B}$ | $92.80 \pm 0.26 \mathrm{E}$ | $10.6 \pm 0.36$ A | $6.1 \pm 0.26$ C | $6.5 \pm 0.84 \mathrm{~B}$ | $9.2 \pm 0.92 \mathrm{~A}$ | $20.3 \pm 0.50 \mathrm{~A}$ | $35.4 \pm 0.51 \mathrm{~A}$ | $27.4 \pm 0.28$ D | $31.2 \pm 0.28 \mathrm{D}$ |
| $\mathrm{P}_{12}$ | $102.95 \pm 0.83 \mathrm{BC}$ |  | $6.5 \pm 0.38 \mathrm{D}$ |  | $7.5 \pm 0.78 \mathrm{~A}$ |  | $17.3 \pm 0.96 \mathrm{~B}$ |  | $22.0 \pm 0.54 \mathrm{E}$ |  |
| $\mathbf{P}_{8}$ | $107.45 \pm 0.52 \mathrm{AB}$ |  | $14.8 \pm 0.47 \mathrm{~A}$ |  | $5.5 \pm 0.61$ B |  | $23.3 \pm 0.05 \mathrm{~A}$ |  |  |  |
| $\mathbf{P}_{13} \times \mathbf{P}_{8}$ | $115.55 \pm 0.74 \mathrm{~A}$ | $113.76 \pm 0.29 \mathrm{~B}$ | $13.1 \pm 0.82 \mathrm{~A}$ | $8.5 \pm 0.76$ B | $5.6 \pm 0.15$ B | $5.4 \pm 0.03 \mathrm{~B}$ | $21.4 \pm 0.89 \mathrm{~A}$ | $18.6 \pm 0.17 \mathrm{D}$ | $29.3 \pm 0.19 \mathrm{D}$ | $31.6 \pm 0.61 \mathrm{D}$ |
| $\mathrm{P}_{13}$ | $111.70 \pm 0.73 \mathrm{~B}$ |  | $11.5 \pm 0.52 \mathrm{~A}$ |  | $6.2 \pm 0.60$ B |  | $19.5 \pm 0.42 \mathrm{~B}$ |  | $45.3 \pm 0.80 \mathrm{~A}$ |  |
| $\mathrm{P}_{8}$ | $108.50 \pm 0.45 \mathrm{AB}$ |  | $22.2 \pm 0.38$ A |  | $6.2 \pm 0.47 \mathrm{~B}$ |  | $21.3 \pm 0.87 \mathrm{~A}$ |  | $32.7 \pm 0.19 \mathrm{CD}$ |  |
| $\mathrm{P}_{13} \times \mathrm{P}_{6}$ | $110.53 \pm 0.38 \mathrm{~A}$ | $112.24 \pm 0.91 \mathrm{~B}$ | $12.3 \pm 0 . .55 \mathrm{~A}$ | $12.3 \pm 0.57 \mathrm{~A}$ | $5.4 \pm 0.06 \mathrm{~B}$ | $5.0 \pm 0.63 \mathrm{~B}$ | $22.0 \pm 0.76 \mathrm{~A}$ | $34.2 \pm 0.08 \mathrm{~A}$ | $47.7 \pm 0.28$ A | $32.6 \pm 0.76 \mathrm{D}$ |
| $\mathrm{P}_{13}$ | $88.63 \pm 0.65$ D |  | $8.1 \pm 0.44$ B |  | $5.6 \pm 0.74$ B |  | $19.7 \pm 0.56$ B |  | $20.0 \pm 0.75 \mathrm{~F}$ |  |
| $\mathrm{P}_{6}$ | $121.42 \pm 0.89 \mathrm{~A}$ |  | $12.3 \pm 0.19 \mathrm{~A}$ |  | $4.1 \pm 0.24 \mathrm{C}$ |  | $17.7 \pm 0.32 \mathrm{~B}$ |  | $41.1 \pm 0.56 \mathrm{~A}$ |  |
| Mean | $110.92 \pm 0.35$ | $112.13 \pm 0.13$ | $10.32 \pm 0.02$ | $10.18 \pm 0.89$ | $6.2 \pm 0.02$ | $6.1 \pm 0.73$ | $20.5 \pm 2.26$ | $27.5 \pm 0.45$ | $37.4 \pm 0.54$ | $32.8 \pm 0.82$ |

$\mathrm{P}_{1}=$ First parent $; \mathrm{P}_{2}=$ Second parent; $\mathrm{PH}=$ Plant height; $\mathrm{SL}=$ Spike length; $\mathrm{NS}=$ Number of spike perplant; NSP $=$ Number of spikelet per spike; $\mathrm{NG}=$ Number of grain per spikelet; MP $=$ Mid-parent heterosis; HP $=$ High-
 Köse $220 / 39, \mathrm{P}_{8}=$ T. spelta, $\mathrm{P}_{9}=$ Penjamo $62, \mathrm{P}_{10}=$ Sivas $111 / 33, \mathrm{P}_{11}=$ Sürak $1593 / 51, \mathrm{P}_{12}=$ Sertak $52, \mathrm{P}_{13}=$ Yektay 406

Table 4. Heterosis percentages (\%) in the $\mathrm{F}_{1}$ and $\mathrm{F}_{2}$ for the MP and HP according to the examined traits in the combinations.

| Combinations | PH (cm) |  |  |  | SL (cm) |  |  |  | NS (no) |  |  |  | NSP (no) |  |  |  | NG (no) |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | MP |  | HP |  | MP |  | HP |  | MP |  | HP |  | MP |  | HP |  | MP |  | HP |  |
|  | $\mathrm{F}_{1}$ | $\mathrm{F}_{2}$ | $\mathrm{F}_{1}$ | $\mathrm{F}_{2}$ | $\mathrm{F}_{1}$ | F ${ }_{2}$ | $\mathrm{F}_{1}$ | $\mathrm{F}_{2}$ | $\mathrm{F}_{1}$ | $\mathrm{F}_{2}$ | $\mathrm{F}_{1}$ | F ${ }_{2}$ | $\mathrm{F}_{1}$ | $\mathrm{F}_{2}$ | $\mathrm{F}_{1}$ | $\mathrm{F}_{2}$ | $\mathrm{F}_{1}$ | $\mathrm{F}_{2}$ | $\mathrm{F}_{1}$ | $\mathrm{F}_{2}$ |
| $\mathrm{P}_{1} \times \mathrm{P}_{2}$ | $16.3{ }^{* *}$ | - | $11.7{ }^{\text {*** }}$ | - | $24.6{ }^{* *}$ | - | $9.5 * *$ | - | $24.7{ }^{* *}$ | - | -1.7 | - | 0 | - | -3.0 | - | $24.6{ }^{* *}$ | - | $18.2{ }^{* *}$ | $-9.3{ }^{*}$ |
| $\mathrm{P}_{1} \times \mathrm{P}_{3}$ | 6.2 | 6.1 | 3.7 | 4.6 | 7.5 | $10.2{ }^{* * *}$ | $9.2{ }^{* * *}$ | 7.0 | 84.0*************) | 124.0** | 53.3 *** | $86.7{ }^{* * *}$ | 1.0 | $55.1{ }^{* *}$ | 0.5 | 54.3** | $24.5{ }^{* * *}$ | -4.4 | $16.4{ }^{* *}$ | $32.7{ }^{* *}$ |
| $\mathbf{P}_{4} \times \mathbf{P}_{3}$ | 6.7 | -2.5 | 6.3 | -0.5 | $19.7{ }^{* * *}$ | -44.2** | $12.9{ }^{* * *}$ | 5.1 | -21.4***********) | 24.0***********) | -29.8***********) | 10.6********* | $12.4{ }^{* * *}$ | 8.4 | $12.1{ }^{* *}$ | 8.1 | $20.1{ }^{* *}$ | -30.6** | $14.2{ }^{* *}$ | -31.6***********) |
| $\mathbf{P}_{4} \times \mathbf{P}_{2}$ | $11.4{ }^{* *}$ | 3.0 | 4.1 | -1.1 | $24.5{ }^{* *}$ | -0.1 | $32.3{ }^{* *}$ | $-22.5{ }^{* *}$ | 13.6 ** | 18.5***********) | -13.2**********) | -9.4**********) | 6.1 | -2.0 | 5.0 | -3.0 | 1.6 | $-9.0{ }^{*}$ | -2.1 | -9.8** |
| $\mathrm{P}_{5} \times \mathrm{P}_{3}$ | 5.4 | 3.0 | 4.0 | -7.5 | 0.6 | $-37.0{ }^{* * *}$ | -5.8 | -5.0 | 2.4 | -47.6** | 0 | $-48.8{ }^{* *}$ | -9.2* | 4.3 | 11.0 ** | 2.4 | $18.6{ }^{* *}$ | -29.0 ** | -2.4 | 2.4 |
| $\mathbf{P}_{5} \times \mathbf{P}_{6}$ | 0.4 | -6.0 | -1.6 | -13.0 ** | 18.3 ${ }^{* * *}$ | 4.8 | $23.9{ }^{* *}$ | $9.7{ }^{*}$ | 17.1** | -2.9 | -18.0 ********) | $-32.0{ }^{* *}$ | $11.2{ }^{* *}$ | $81.0{ }^{* * *}$ | 1.5 | 65.2*******) | 7.9 | 13.2*********** | -3.3 | 2.4 |
| $\mathbf{P}_{7} \times \mathbf{P}_{6}$ | 8.7 | $12.3{ }^{* *}$ | 1.6 | -19.3** | $51.8{ }^{* * *}$ | 52.0********) | 7.5 | 7.5 | 5.2 | $-17.0^{* *}$ | -19.3 *** | -36.4*********) | 14.6**********) | $91.6{ }^{* * *}$ | 10.0 ** | 83.0 ** | -3.3 | $-19.6{ }^{* *}$ | $44.0{ }^{* *}$ | 24.1*************) |
| $\mathbf{P}_{7} \times \mathbf{P}_{8}$ | -6.3 | 2.9 | -7.3 | -7.3 | $6.1{ }^{* *}$ | $-36.5{ }^{* *}$ | -1.6 | -41.1** | 152.6*********) | $148.4{ }^{* * *}$ | $135.3^{* * *}$ | $131.4 * *$ | $10.3{ }^{*}$ | 0.6 | -2.9 | 0.4 | $65.3{ }^{* *}$ | 43.0** | 5.8 | -71.2** |
| $\mathbf{P}_{9} \times \mathbf{P}_{8}$ | 7.8 | $-{ }^{* * *}$ | 7.6 | - | 0.9 | - ${ }^{* *}$ | -14.6** | $-{ }^{* * *}$ | $-8.9{ }^{*}$ | - | -28.1** | 5 | 5.0 | . | $-1.0$ | - | 6.2 | $-$ | $17.0^{* *}$ | - |
| $\mathrm{P}_{9} \times \mathrm{P}_{6}$ | 12.0 ** | $12.7{ }^{* *}$ | $11.9{ }^{* *}$ | 2.2 | $11.0{ }^{* *}$ | 26.7 ${ }^{* * *}$ | $-1.6$ | $12.3{ }^{* * *}$ | $10.1{ }^{*}$ | 83.5************) | 5.3 |  | 0.5 | 46.4*********) | -25.0 ** | 42.0**********) | 40.3 ** | 35.0** | -5.6 | -12.0** |
| $\mathrm{P}_{10} \times \mathrm{P}_{6}$ | 1.2 | 6.9 | -3.3 | -8.2 | 0.6 | -10.6********) | $-27.2^{* * *}$ | $-20.0{ }^{* * *}$ | 0 | -11.8**************) | -26.1***************) | $-34.8{ }^{* *}$ | 0.3 | 41.6*************) | -1.6 | 39.0***********) | 0 | 60.3*************) | 23.0**********) | $-51.3^{* *}$ |
| $\mathrm{P}_{10} \times \mathrm{P}_{8}$ | $-11.9{ }^{* * *}$ | 3.8 | -13.4** | $-0.4$ |  | -2.3 | -49.7******) | -30.3 ** | $41.0{ }^{* *}$ | $-24.0{ }^{* * *}$ | $39.3{ }^{* *}$ | $-24.6{ }^{* * *}$ | -12.9** | $42.1{ }^{* * *}$ | -23.0 ** | 25.7*************) | $33.7{ }^{* *}$ | -54.5***********) | - $9.9{ }^{* *}$ | $-58.3{ }^{* * *}$ |
| $\mathbf{P}_{11} \times \mathbf{P}_{3}$ | 9.0 * | $7.1{ }^{* *}$ | 4.1 | $12.8{ }^{* *}$ | $10.6{ }^{* * *}$ | $-3.8$ | 8.0 | 8.0 | 21.0**************) | -16.6** | $11.0{ }^{* *}$ | $-23.7{ }^{* * *}$ | -2.4 | $31.0{ }^{* * *}$ | $-9.1{ }^{*}$ | 22.0***********) | -7.7 | -45.9 **********) | $-3.0$ | $-46.7^{* * *}$ |
| $\mathbf{P}_{11} \times \mathbf{P}_{6}$ | 3.5 | $28.0{ }^{* *}$ | 2.1 | 8.2 | 15.2** | 65.5** | $-1.0$ | $42.3{ }^{* *}$ | $36.0{ }^{* *}$ | -3.1 | 8.8 | $-17.5^{* *}$ | 8.3 | 18.6******* | 5.0 | 15.0 ** | 0 | $86.3{ }^{* *}$ | - 50.3********) | -26.6 ** |
| $\mathbf{P}_{12} \times \mathrm{P}_{6}$ | 6.1 ** | ${ }^{-}$ | $-1.9{ }^{\text {* }}$ | ${ }^{-}$ | -6.3 | - ${ }^{* * *}$ | $-14.3^{* * *}$ | . | 2.0 | . | $-11.7^{* * *}$ | .7 | 20.0 ** | ${ }^{-}$ | $18.3^{* * *}$ | - ${ }^{\text {- }}$ | 69.0 ** | - | - $16.5{ }^{* * *}$ | - |
| $\mathrm{P}_{12} \times \mathrm{P}_{8}$ | -26.6 ** | $-11.8{ }^{* *}$ | $28.1{ }^{* *}$ | -14.9 ** | -0.5 | -43.0 ** | -28.4** | -59.0 ** | 0 | 42.0** | - 13.3 ** | $22.7{ }^{* * *}$ | 0 | $74.4 *$ | 12.9 ** | $52.0{ }^{* *}$ | 0 | $13.9{ }^{* *}$ | - 35.3 ** | -4.9 |
| $\mathbf{P}_{13} \times \mathrm{P}_{8}$ | 5.0 | 3.3 | 3.5 | 4.3 | -22.3 ** | -50.0 *** | -41.0** | -62.0 ** | 0 | -12.9** | -9.7** | $-13.0{ }^{* *}$ | 5.4 | -8.4 | 0.5 | $-12.7{ }^{* *}$ | $-33.1^{* * *}$ | -19.0 ** | $16.1^{* *}$ | $-30.2^{* *}$ |
| $\mathrm{P}_{13} \times \mathrm{P}_{6}$ | 5.2 | -9.0* | 6.9 | -12.0 ** | 20.6 ** | $-66.7^{* * *}$ | 0 | -38.2*** | $11.3{ }^{*}$ | 3.1 | -3.6 | $-10.7{ }^{* * *}$ | $17 .{ }^{* *}$ | 82.9 ** | $11.7{ }^{* * *}$ | -73.6** | $56.4 * *$ | 6.4 | -2.1 | -20.7*******) |
| Mean | 3.3 | 4.0 | 3.8 | 3.0 | 8.5 | 1.3 | -4.6 | -7.3 | 22.0 | 21.0 | 4.4 | 5.1 | 4.9 | 38.0 | -1.4 | -31.1 | 1.0 | 3.1 | 8.0 | -4.5 |

$\mathrm{PH}=$ Plant height, $\mathrm{SL}=$ Spike length, $\mathrm{NS}=$ Number of spike per plant; $\mathrm{NSP}=$ Number of spikelet per spike; $\mathrm{NG}=\mathrm{Number}$ of grain per spikelet; MP=Mid-parent heterosis; $\mathrm{HP}=\mathrm{High}$-parent heterosis,
$\mathrm{P}_{1}=$ Kunduru 414/44, $\mathrm{P}_{2}=$ T. dicoccum, $\mathrm{P}_{3}=$ T. carthlicum, $\mathrm{P}_{4}=$ Kunduru $1149, \mathrm{P}_{5}=$ Aköz 86, $\mathrm{P}_{6}=$ T. vavilovií, $\mathrm{P}_{7}=$ Köse 220/39, $\mathrm{P}_{8}=T$. spelta, $\mathrm{P}_{9}=$ Penjamo $62, \mathrm{P}_{10}=$ Sivas $111 / 33, \mathrm{P}_{11}=$ Sürak $1593 / 51, \mathrm{P}_{12}=$ Sertak 52 ,
ay 406
${ }^{*} \mathrm{P}<0.05,{ }^{* *} \mathrm{P}=0.01$

Table 5. Minumum and maximum mean values, heterosis percentages as MP and HP according to the combinations and generations, and phenotypic correlation coefficients.

\begin{tabular}{|c|c|c|c|c|c|c|c|c|c|c|c|c|c|}
\hline Traits \& Combinations \& Generations \& Means \& MP \& HP \& (P1-F1) \& (P1-F2) \& (P2-F1) \& (P2-F2) \& (MP-F1) \& (MP-F2) \& (HP-F1) \& (HP-F2) <br>
\hline PH (cm) \& $$
\begin{array}{r}
1 \\
\hline \\
\left(\mathrm{P}_{11} \times \mathrm{P}_{3}\right) \\
\left(\mathrm{P}_{7} \times \mathrm{P}_{8}\right) \\
\left(\mathrm{P}_{11} \times \mathrm{P}_{6}\right) \\
\left(\mathrm{P}_{12} \times \mathrm{P}_{8}\right)
\end{array}
$$ \& $\mathrm{F}_{1}$
$\mathrm{~F}_{2}$ \& $$
\begin{gathered}
(120.10 \pm 0.28) \\
(90.40 \pm 0.42) \\
(137.84 \pm 0.78) \\
(92.80 \pm 0.26)
\end{gathered}
$$ \& $$
\begin{gathered}
9.0 \\
-6.3^{* *} \\
3.5 \\
-26.6
\end{gathered}
$$ \& $$
\begin{array}{r}
4.1 \\
-7.3 \\
2.1 \\
-28.1
\end{array}
$$ \& $0.473^{* *}$ \& 0.331 ** \& $0.315^{* *}$ \& 0.220 ** \& 0.450 ** \& $0.386^{* *}$ \& $0.217^{*}$ \& 0.160* <br>
\hline SL (cm) \& $$
\begin{aligned}
& \left(\mathrm{P}_{13} \times \mathrm{P}_{8}\right) \\
& \left(\mathrm{P}_{1} \times \mathrm{P}_{3}\right) \\
& \left(\mathrm{P}_{11} \times \mathrm{P}_{6}\right) \\
& \left(\mathrm{P}_{5} \times \mathrm{P}_{3}\right)
\end{aligned}
$$ \& F

$\mathrm{F}_{2}$ \& \[
$$
\begin{array}{r}
(13.1 \pm 0.82) \\
(7.2 \pm 0.65) \\
(15.8 \pm 1.47) \\
(5.1 \pm 0.87)
\end{array}
$$

\] \& \[

$$
\begin{array}{r}
-22.3 \\
7.5 \\
15.2 \\
0.6
\end{array}
$$

\] \& \[

$$
\begin{array}{r}
-41.0 \\
9.2 \\
-1.0 \\
5.8
\end{array}
$$
\] \& $0.495^{* *}$ \& $0.129^{*}$ \& $0.456^{* *}$ \& 0.041 ns \& 0.570** \& $0.633^{* *}$ \& 0.120** \& 0.080 ns <br>

\hline NS (no) \& $$
\begin{aligned}
& \left(\mathrm{P}_{7} \times \mathrm{P}_{8}\right) \\
& \left(\mathrm{P}_{4} \times \mathrm{P}_{3}\right) \\
& \left(\mathrm{P}_{7} \times \mathrm{P}_{8}\right) \\
& \left(\mathrm{P}_{5} \times \mathrm{P}_{3}\right)
\end{aligned}
$$ \& F 1

$\mathrm{~F}_{2}$ \& \[
$$
\begin{array}{r}
(12.0 \pm 3.19) \\
(3.3 \pm 1.17) \\
(11.8 \pm 0.11) \\
(2.2 \pm 0.86)
\end{array}
$$

\] \& \[

$$
\begin{gathered}
152.6 \\
-21.4 \\
152.6 \\
-94
\end{gathered}
$$

\] \& \[

$$
\begin{array}{r}
135.3 \\
29.8 \\
135.3 \\
1.7
\end{array}
$$
\] \& $0.400^{* *}$ \& 0.050 ns \& 0.074 ns \& -0.250 ns \& 0.550 ** \& $0.625^{* *}$ \& 0.450 ** \& 0.001 ns <br>

\hline NSP (no) \& $$
\begin{aligned}
& \left(\mathrm{P}_{4} \times \mathrm{P}_{3}\right) \\
& \left(\mathrm{P}_{10} \times \mathrm{P}_{8}\right) \\
& \left(\mathrm{P}_{7} \times \mathrm{P}_{6}\right) \\
& \left(\mathrm{P}_{13} \times \mathrm{P}_{8}\right)
\end{aligned}
$$ \& F

$\mathrm{F}_{2}$ \& \[
$$
\begin{gathered}
(22.2 \pm 0.93) \\
(17.9 \pm 0.59) \\
(36.6 \pm 2.09) \\
(18.6 \pm 0.17)
\end{gathered}
$$

\] \& \[

$$
\begin{aligned}
& 12.4 \\
& 12.9 \\
& 14.6 \\
& 54.0
\end{aligned}
$$

\] \& \[

$$
\begin{gathered}
12.1 \\
-23.0 \\
10.0 \\
0.5
\end{gathered}
$$
\] \& 0.500** \& 0.330** \& 0.391 ** \& $0.200^{* *}$ \& $0.540^{* *}$ \& $0.113^{* *}$ \& 0.320 ns \& $0.380 * *$ <br>

\hline NG (no) \& $$
\begin{aligned}
& \left(\mathrm{P}_{13} \times \mathrm{P}_{6}\right) \\
& \left(\mathrm{P}_{13} \times \mathrm{P}_{8}\right) \\
& \left(\mathrm{P}_{11} \times \mathrm{P}_{6}\right) \\
& \left(\mathrm{P}_{10} \times \mathrm{P}_{8}\right)
\end{aligned}
$$ \& F1

$\mathrm{F}_{2}$ \& \[
$$
\begin{gathered}
(47.7 \pm 0.28) \\
(29.3 \pm 0.19) \\
(59.6 \pm 0.79) \\
(14.4 \pm 0.20)
\end{gathered}
$$

\] \& \[

$$
\begin{array}{r}
56.4 \\
33.1 \\
0 \\
-33.7
\end{array}
$$

\] \& \[

$$
\begin{array}{r}
-2.1 \\
-30.2 \\
-35.3 \\
-58.3
\end{array}
$$
\] \& -0.126 ns \& -0.050 ns \& 0.390 \& 0.110 ns \& 0.155 \& 0.125 \& -0.160 ns \& 0.150 <br>

\hline
\end{tabular}

$\mathrm{PH}=$ Plant height, $\mathrm{SL}=$ Spike length, $\mathrm{NS}=$ Number of spike per plant; NSP $=$ Number of spikelet per spike; NG $=$ Number of grain per spikelet; MP $=$ Mid-parent heterosis; HP $=$ High-parent heterosis, ${ }^{1)}$ Maximum, ${ }^{2)}$ Minimum,
$\mathrm{P}_{1}=$ Kunduru 414/44, $\mathrm{P}_{3}=$ T. carthlicum, $\mathrm{P}_{4}=$ Kunduru 1149, $\mathrm{P}_{5}=$ Aköz $86, \mathrm{P}_{6}=$ T. vavilovii, $\mathrm{P}_{7}=$ Köse 220/39, $\mathrm{P}_{8}=T$. spelta, $\mathrm{P}_{9}=$ Penjamo $62, \mathrm{P}_{10}=$ Sivas $111 / 33, \mathrm{P}_{11}=$ Sürak $1593 / 51, \mathrm{P}_{12}=$ Sertak $52, \mathrm{P}_{13}=$ Yektay 406
$\mathrm{F}_{1}=$ First generation, $\mathrm{F}_{2}=$ Second generation; P 1 : Mean values of the female parents, P 2 : Mean values of the male parents, ${ }^{\mathrm{P}}<0.05$. ** $\mathrm{P}<0.01$, ns $=$ non significant.
$\mathrm{MP}-\mathrm{Fn}=\mathrm{n}$ generation's mean values of the mid-parent heterosis values of the females, $\mathrm{HP}-\mathrm{Fn}=\mathrm{n}$ generation's mean values of the mid-parent heterosis values of the semi wilds.

Table 6. The parental and the combinational participations at the $\mathrm{F}_{1}$ and $\mathrm{F}_{2}$ for the MP and HP. (+ yes; - or blank no; s= significant; ns= non significant)

| Genotypes | PH (cm) |  |  |  | SL (cm) |  |  |  | NS (no) |  |  |  | NSP (no) |  |  |  | NG (no) |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | MP |  | HP |  | MP |  | HP |  | MP |  | HP |  | MP |  | HP |  | MP |  | HP |  |
|  | $\mathrm{F}_{1}$ | $\mathrm{F}_{2}$ | $\mathrm{F}_{1}$ | $\mathrm{F}_{2}$ | $\mathrm{F}_{1}$ | $\mathrm{F}_{2}$ | $\mathrm{F}_{1}$ | $\mathrm{F}_{2}$ | $\mathrm{F}_{1}$ | $\mathrm{F}_{2}$ | $\mathrm{F}_{1}$ | $\mathrm{F}_{2}$ | $\mathrm{F}_{1}$ | $\mathrm{F}_{2}$ | $\mathrm{F}_{1}$ | $\mathrm{F}_{2}$ | $\mathrm{F}_{1}$ | $\mathrm{F}_{2}$ | $\mathrm{F}_{1}$ | $\mathrm{F}_{2}$ |
| ( ${ }^{\text {( })}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| $\mathrm{P}_{1}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| $\mathrm{P}_{4}$ |  |  |  |  |  |  |  |  |  |  |  | (-) s | (+) s |  |  |  |  |  | $(-) \mathrm{s}$ |  |
| $\mathrm{P}_{5}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| $\mathbf{P}_{7}$ |  |  |  |  |  |  |  |  |  |  | (+) s | (+) s |  | (+) s |  | (+) s |  |  |  |  |
| $\mathrm{P}_{9}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| $\mathrm{P}_{10}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| $\mathrm{P}_{11}$ |  |  |  |  |  | (+) s |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| $\mathbf{P}_{12}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| $\mathbf{P}_{13}$ |  |  |  |  |  |  |  |  |  |  |  | (-) s |  |  |  |  | (+) s |  | $(-) \mathrm{s}$ |  |
| $\left(0^{\prime}\right)$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| $\mathrm{P}_{2}$ |  |  |  |  |  |  |  |  |  |  |  |  | $(-) \mathrm{s}$ |  |  |  |  |  |  |  |
| $\mathbf{P}_{3}$ |  |  |  |  |  |  |  |  |  |  |  |  | (+) s | (+) s | (+) s |  |  |  |  |  |
| $\mathrm{P}_{6}$ |  |  |  |  |  |  |  |  |  |  |  |  | (+) s | (+) s |  |  |  |  |  |  |
| $\mathrm{P}_{8}$ |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  | (+) s |  |  | (+) s |  |
| mbinations |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| $\mathrm{P}_{1} \times \mathrm{P}_{2}$ | (+) ns | No seed | (+) ns | No seed | (+) ns | No seed | (+) ns | No seed | (+) ns | No seed | $(-) \mathrm{ns}$ | No seed | (+) ns | No seed | $(-) \mathrm{ns}$ | No seed | (+) ns | No seed | (+) ns | $(-) \mathrm{ns}$ |
| $\mathrm{P}_{1} \times \mathrm{P}_{3}$ | $(+) \mathrm{ns}$ | (+) ns | $(+) \mathrm{ns}$ | (+) ns | $(+)$ ns | (-) ns | $(+) \mathrm{ns}$ | (+) ns | (+) ns | (+) ns | $(+) \mathrm{ns}$ | (+) ns | (+) ns | (+) ns | (+) ns | (+) ns | (+) s | $(-) \mathrm{ns}$ | $(+) \mathrm{ns}$ | (+) ns |
| $\mathbf{P}_{4} \times \mathbf{P}_{3}$ | $(+)$ ns | $(-) \mathrm{ns}$ | $(+) \mathrm{ns}$ | $(-) \mathrm{ns}$ | (+) s | $(-) \mathrm{ns}$ | $(+) \mathrm{s}$ | (+) s | $(-) \mathrm{ns}$ | (+) ns | $(-) \mathrm{ns}$ | $(+) \mathrm{ns}$ | $(+)$ ns | (+) ns | $(+) \mathrm{ns}$ | $(+)$ ns | $(+) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(+) \mathrm{ns}$ | $(-) \mathrm{ns}$ |
| $\mathbf{P}_{4} \times \mathbf{P}_{2}$ | $(+) \mathrm{ns}$ | (+) ns | $(+) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(+)$ ns | $(-) \mathrm{ns}$ | $(+) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(+)$ ns | (+) ns | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(+)$ ns | $(-) \mathrm{ns}$ | $(+) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(+) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ |
| $\mathrm{P}_{5} \times \mathrm{P}_{3}$ | $(+)$ ns | (+) ns | $(+) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(+)$ ns | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(+)$ ns | $(-) \mathrm{ns}$ | $(+)$ ns | $(-) \mathrm{ns}$ | $(+)$ ns | (+) ns | $(+) \mathrm{ns}$ | (+) ns | $(+) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | (+) ns |
| $\mathrm{P}_{5} \times \mathrm{P}_{6}$ | $(+)$ ns | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(+)$ ns | (+) ns | (+) ns | $(+) \mathrm{ns}$ | $(+)$ ns | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | (+) ns | $(+)$ ns | (+) ns | $(+) \mathrm{ns}$ | (+) ns | $(-) \mathrm{ns}$ | (+) ns |
| $\mathbf{P}_{7} \times \mathbf{P}_{6}$ | $(+) \mathrm{ns}$ | (+) ns | (+) s | $(-) \mathrm{ns}$ | $(+)$ ns | $(+) \mathrm{ns}$ | $(+) \mathrm{ns}$ | (+) s | $(+)$ ns | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(+)$ ns | (+) ns | (+) s | (+) ns | $(+) \mathrm{ns}$ | $(-) \mathrm{ns}$ | (+) s | (+) ns |
| $\mathbf{P}_{7} \times \mathbf{P}_{8}$ | $(-) \mathrm{ns}$ | (+) ns | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(+)$ ns | $(-) \mathrm{ns}$ | $(+) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(+)$ ns | (+) ns | (+) s | (+) ns | $(+)$ ns | (+) ns | $(-) \mathrm{ns}$ | (+) ns | $(-) \mathrm{ns}$ | (+) ns | (+) s | $(-) \mathrm{ns}$ |
| $\mathbf{P}_{9} \times \mathbf{P}_{8}$ | $(+) \mathrm{ns}$ | No seed | $(+) \mathrm{ns}$ | No seed | $(+)$ ns | No seed | $(-) \mathrm{ns}$ | No seed | $(-) \mathrm{ns}$ | No seed | $(-) \mathrm{ns}$ | No seed | $(+)$ ns | No seed | $(-) \mathrm{ns}$ | No seed | $(+) \mathrm{ns}$ | No seed | $(+) \mathrm{ns}$ | No seed |
| $\mathrm{P}_{9} \times \mathrm{P}_{6}$ | $(+)$ ns | (+) ns | $(+) \mathrm{ns}$ | (+) ns | (+) s | (+) ns | $(-) \mathrm{ns}$ | (+)ns | $(+)$ ns | (+) ns | $(+) \mathrm{ns}$ | (+) ns | $(+)$ ns | (+) ns | $(-) \mathrm{ns}$ | (+) ns | $(+) \mathrm{ns}$ | (+) ns | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ |
| $\mathbf{P}_{10} \times \mathbf{P}_{6}$ | $(+)$ ns | $(+) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(+)$ ns | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(+)$ ns | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | (+) ns | $(-) \mathrm{ns}$ | (+) ns | $(+) \mathrm{ns}$ | (+) ns | (+) ns | $(-) \mathrm{ns}$ |
| $\mathbf{P}_{10} \times \mathbf{P}_{8}$ | $(-) \mathrm{ns}$ | $(+) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(+)$ ns | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(+)$ ns | $(-) \mathrm{ns}$ | (+) ns | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(+)$ ns | $(-) \mathrm{ns}$ | $(+)$ ns | $(+) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ |
| $\mathrm{P}_{11} \times \mathrm{P}_{3}$ | $(+)$ ns | (+) ns | $(+)$ ns | $(+)$ ns | (+) s | $(-) \mathrm{ns}$ | $(+) \mathrm{ns}$ | (+) ns | $(+)$ ns | $(-) \mathrm{ns}$ | $(+)$ ns | $(-) \mathrm{ns}$ | $(+) \mathrm{ns}$ | $(+)$ ns | $(-) \mathrm{ns}$ | (+) ns | $(+) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(+) \mathrm{ns}$ | $(-) \mathrm{ns}$ |
| $\mathbf{P}_{11} \times \mathbf{P}_{6}$ | $(+)$ ns | (+) ns | $(+) \mathrm{ns}$ | (+) ns | (+) s | (+) ns | $(-) \mathrm{ns}$ | (+) ns | $(+)$ ns | $(-) \mathrm{ns}$ | $(+) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(+)$ ns | (+) ns | $(-) \mathrm{ns}$ | (+) ns | $(-) \mathrm{ns}$ | (+) ns | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ |
| $\mathbf{P}_{12} \times \mathrm{P}_{6}$ | $(+) \mathrm{ns}$ | No seed | $(-) \mathrm{ns}$ | No seed | $(+)$ ns | No seed | $(-) \mathrm{ns}$ | No seed | $(+)$ ns | No seed | $(-) \mathrm{ns}$ | No seed | $(+)$ ns | No seed | $(+) \mathrm{ns}$ | No seed | $(+)$ ns | No seed | $(-) \mathrm{ns}$ | No seed |
| $\mathbf{P}_{12} \times \mathbf{P}_{8}$ | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(+) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(+)$ ns | $(+) \mathrm{ns}$ | $(-) \mathrm{ns}$ | (+) ns | $(+)$ ns | $(+) \mathrm{ns}$ | $(+) \mathrm{ns}$ | (+) ns | (+) ns | $(+) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ |
| $\mathbf{P}_{13} \times \mathbf{P}_{8}$ | $(+) \mathrm{ns}$ | $(+) \mathrm{ns}$ | $(+) \mathrm{ns}$ | $(+) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(+)$ ns | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(+)$ ns | $(-) \mathrm{ns}$ | $(+) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(+) \mathrm{ns}$ | $(-) \mathrm{ns}$ |
| $\mathrm{P}_{13} \times \mathrm{P}_{6}$ | $(+) \mathrm{ns}$ | $(+) \mathrm{ns}$ | $(+) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(+)$ ns | $(-) \mathrm{ns}$ | $(+) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(+) \mathrm{ns}$ | $(+)$ ns | $(-) \mathrm{ns}$ | $(-) \mathrm{ns}$ | $(+)$ ns | $(+) \mathrm{ns}$ | $(+)$ ns | $(-) \mathrm{ns}$ | (+) ns | $(+)$ ns | $(-)$ ns | $(-) \mathrm{ns}$ |

$\mathrm{PH}=$ Plant height, $\mathrm{SL}=$ Spike length, $\mathrm{NS}=$ Number of spike per plant; $\mathrm{NSP}=$ Number of spikelet per spike; NG=Number of grain per spikelet; MP=Mid-parent heterosis; HP=High-parent heterosis
$\mathrm{P}_{1}=$ Kunduru 414/44, $\mathrm{P}_{2}=$ T. dicoccum, $\mathrm{P}_{3}=T$ carthlicum, $\mathrm{P}_{4}=$ Kunduru 1149, $\mathrm{P}_{5}=$ Aköz $86, \mathrm{P}_{6}=T$. vavilovii, $\mathrm{P}_{7}=$ Köse $220 / 39, \mathrm{P}_{8}=T$. spelta, $\mathrm{P}_{9}=$ Penjamo $62, \mathrm{P}_{10}=$ Sivas $111 / 33, \mathrm{P}_{11}=$ Sürak $1593 / 51, \mathrm{P}_{12}=$ Sertak $52, \mathrm{P}_{13}=$ Yektay 406

