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5 ABSTRACT

6 Most of the tomato varieties in Bangladesh are of inbred type and produced low yield indicating need to 7 develop high yielding variety through the hybridization. Heterosis breeding is used to improve yield and quality of tomato because traditional methods cannot be used to achieve this goal. A half diallel design 8 9 was employed to develop F_1 s from seven parents of winter tomato. 21 F_1 s along with their parents were evaluated for yield and quality traits. Heterosis analysis revealed that heterotic vigor was present for 10 11 growth and yield characters among hybrids. Heterosis for better parent was negative for days to 12 flowering, days to harvest, harvest duration, number of locules, and number of seeds per fruit but positive 13 for fruit set, number of fruits per plant, yield per plant, pericarp thickness and TSS. None of the hybrid was 14 heterotic for all characters simultaneously. The hybrids G5, G13, G16, G17, G18, and G20 had 25.73, 19.92, 39.20, 36.49, 53.77, and 50.31% higher heterosis compared to the better parent, respectively, for 15 16 fruit yield per plant as well as for many other yield contributing traits. High heterosis for yield appears to be the consequence of heterosis of yield attributing traits; therefore, these hybrids offer scope of 17 18 developing improved commercial lines through heterosis breeding.

Studies on Character Improvement in Tomato (Solanum

lycopersicum L.) by Heterosis

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20 Keywords: Heterosis breeding, quantitative trait, tomato, yield

22 1. INTRODUCTION

23 Tomato (Solanum lycopersicum L.) is one of the most popular and extensively consumed vegetable over 24 the world. Currently, tomato is grown around the globe for either fresh market or processing [1] and 25 considered as a high value crop. As a cash crop, it has a great demand in local as well as the 26 international market. Unfortunately, the production of tomato in Bangladesh is limited due to the scarcity 27 of high yielding varieties. As a result, a huge quantity of tomato is imported every year from the 28 neighboring countries to meet up the local market demand. Recently, the crop has received more 29 attention to the policy makers and researchers. As the development of hybrid varieties with higher yield 30 has been thought to be an effective strategy increasing tomato production, a number of projects have 31 been implemented recent years developing new hybrids in Bangladesh. On the other hand, heterosis 32 breeding is predicted to be the most powerful genetic approach developing hybrids with higher yield [2]. 33 Heterosis, which is the superiority in performance of hybrid individuals compared with their parents [3]. 34 has been reported for a wide range of crop species including both self and cross-pollinated crops. 35 Therefore, the estimation of heterosis is one of the goals to assess the hybrid vigor selecting promising 36 hybrids.

37 Heterosis was first observed by Hedrick and Booth [4] in tomato for higher yield. Afterwards a numerous 38 studies have been done in relation to heterosis for yield, its components and quality traits [3,5,6,7,8]. 39 However, the exploitation of heterosis is a quick and an effective way of selecting hybrids for high yield 40 potential, earliness and quality attributes. Unfortunately, a very few attempts in this regard has been taken 41 in the past in Bangladesh. The present study was therefore, executed to estimate the level of percent 42 better and mid parent heterosis among F_1 hybrids of tomato. This information would be useful to 43 investigate the performance and relationship of F_1 hybrids with their parents and to select suitable parents 44 and/or population for designing an effective breeding programme.

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46 2. MATERIALS AND METHODS

48 **2.1 Planting materials**: Seven inbred lines of tomato namely VRT001 (P1), VRT007 (P2), VRT008 (P3), 49 C11 (P4), C41 (P5), LE02 (P6) and TLB133 (P7) were used in the hybridization. A half diallel mating 50 fashion was followed in developing $F_{1}s$ in winter 2009-10 (Table 1). Twenty one $F_{1}s$ along with the seven 51 parents were evaluated in winter 2010-11. Parental genotype denoting VRT is virus tolerance, LE is 52 *Lycopersicon esculentum*, TLB is tolerance to late blight and C is heat tolerance.

2.2 Experimental site: The experiment was conducted at the Vegetable Research Field of Horticulture Research Centre (HRC), Bangladesh Agricultural Research Institute (BARI) Bangladesh from October 2010 to March 2011. The climate of the experimental site is subtropical characterized by heavy rainfall from May to September and scanty rainfall rest of the year. The soil of the experimental site was sandy loam in texture and acidic in nature with a pH around 6.0. This area belongs to the "Shallow red-brown terrace" soil of Madhupur tract as reported by Haider *et al.* [9]. The land was prepared and fertilized as described by Salim *et al.* [10].

2.3 Seedling raising and transplanting: Seeds were sown thinly in a raised seed bed on October 15, 2010. Seed bed was shaded partially with black net after sowing the seeds. Young seedlings were also covered by a fine mesh white net to protect them from insect attack. 7-days old seedlings were transplanted to a second seed bed at the spacing of 5 x 5 cm for hardening. Thirty days old seedlings were transplanted in the main field on November 15, 2010. Light irrigation was given to each seedling immediately after transplanting for their better establishment.

2.4 Experimental design and plot layout: Tomato seedlings were grown in a raised seed bed and 30days old seedlings were transplanted in the main field following randomized complete block design with three replications. Each genotype with spacing of 60 cm x 40 cm represented double row having 12 plants per row accommodating in total 24 plants per plot. The unit plot was separated by 50 cm irrigation drain, while blocks were separated by 75 cm drain. Recommended cultural practices as well as plant protection measures were followed.

2.5 Data collection and statistical analysis: Data for different characters (Table 2) were recorded from
 10 randomly selected plants of parents and F₁s. Analysis of variance (ANOVA) was performed as
 suggested by Gomez and Gomez [11]. Heterosis was estimated using basic formula described by

Falconer [12]. Usually, the magnitude of heterosis depends on the accumulation of favorable dominant alleles in the F_1 population. If the parental populations differ from each other for favorable dominant alleles, the magnitude of heterosis supposed to be proportionally higher. This relationship was estimated by the basic formula 1. Where; d = magnitude of dominance, y = difference between the parental

79 population for allelic frequencies at the locus.

Heterosis in F1=
$$\sum dy^2$$
.....(1)

For estimation of heterosis in each character the mean values of the 21 F₁'s have been compared with better parent (BP) for heterobeltiosis and with mid parent (MP) for heterosis over mid parental value.

83 Percent heterosis was calculated by the formula 2 and 3.

Heterosis (BP) =
$$\frac{(FI-BP)}{BP} \times 100$$
.....(2)

84

85 86 Heterosis (MP) = $\frac{(FT-MP)}{MP} \times 100$(3)

Where, F₁ = mean performance of F₁ hybrid, BP = mean performance of better parent and MP = mean
performance of mid parent.

The test of significance for heterosis was done by using standard error of the value of better parent and mid parent as suggested by Turner [13]. Mean error variance from the combined analysis of variance of parents and F1's were used for calculating the standard error (SE) of difference. The mean values over replications were used for the comparison. Finally, critical difference (CD) was calculated by the formula 4 and 5 for heterosis over better and mid parent respectively. Note that the difference between F_1 and the parent used for the estimation of heterosis were taken into account cross wise. While the difference between F_1 and the parent was greater than CD it was considered significant and vice versa.

$$CD (BF) = \sqrt{\frac{2}{r}} EMS \times t \dots (4)$$

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$$CD (MP) = \sqrt{\frac{3}{2r}} EMS \times t \dots (5)$$

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- 99 Where, EMS = error mean square from ANOVA table, r = number of replications and t = tabulated value
- 100 either at 5% or 1% level of probability.

101 3. RESULTS AND DISCUSSION

- 103 **3.1 Analysis of variance:** Analysis of variance (ANOVA) for the genotypes *i.e.* parents and F_{1S} showed 104 highly significant differences (P = 0.05 or P = 0.01) for the maximum characters studied except fruit set
- 105 percentage (Table 2). The estimation of percent heterosis observed in F_{15} over mid and better parent was

106 presented in Table 3 to Table 5.

- 107 **3.2 Days to 1st flowering:** All the F_1 s showed highly significant differences (P = 0.05 or P = 0.01) heterosis for flowering time, ranging from -9.89 to -0.09% over mid parent and -11.59 to -2.22% over 108 109 better parent (Table 3). Out of 21 F₁ combinations, the highest heterobeltiotic effect of -11.59% was found 110 in cross G4 followed by G15 (-11.50%), and G20 (-11.44). The entire cross combinations produced negative heterosis indicating early flowering in hybrids when compared with their parents. Earliness 111 112 actually leads to the early production and early supply in the market, resulting good price for the 113 producers. Thus the heterosis for flowering time is considered to be an economic parameter for this 114 study. The negative heterosis for flowering time was also reported in earlier studies [5,6,14,15].
- **3.3 Days to 50% flowering:** The significant differences (P = 0.05 or P = 0.01) were also observed among the F₁ crosses for the heterosis over mid and better parent (Table 3). Positive heterosis was shown for mid parent whereas negative heterosis ranging from -4.45 to -14.82% was shown for better parent. Negative heterosis showed in flowering indicating earliness by the hybrids as compared to their parents. As the farmers prefer to get a high price from the early supply, therefore, negative heterosis for this trait is preferable. This study is in accordance with the findings of Patwary *et al.* [16], Islam *et al.* [17] and Baishya *et al.* [18], those who reported negative heterosis for this trait over better parent in their studies.
- 3.4 Days to 1st harvest: Out of 21 cross combinations, 20 exhibited significant different (P = 0.05 or P = 122 123 0.01) negative heterosis over better parent ranging from -3.05 to -11.92% whereas 18 combinations 124 showed negative heterosis over mid parent (Table 3). The results were very similar to Sharma et al. [19] 125 who reported heterosis ranged of -2.90 to -11.20% over better parent in tomato. More than 10% negative 126 heterosis over better parent was observed from three F₁s viz. G5 (-11.92%), G1 (-10.38%), and G12 (-127 10.18%), which was superior to the previous study -7.14% of heterosis over better parent, reported by 128 Sharma et al. [20]. Negative heterosis here is suggesting early harvest of tomato fruits. Therefore, those 129 genotypes can further be utilized to develop inbred lines toward a variety development program.
- 130 3.5 Harvest duration (days): Harvest duration showed significant negative better parent heterosis in 131 fourteen F_{1s} whereas negative mid parent heterosis was showed in thirteen F_{1s} (Table 3). The highest 132 significant negative heterosis over better parent was estimated from the cross combination G1 (-6.77%) 133 followed by G12 (-6.68%). On the other hand, the highest negative heterosis over mid parent was also 134 estimated from the cross G1 (-6.50%). In contrast, four crosses produced significant positive heterosis 135 over better parent viz. G18 (5.58%), G16 (4.72%), G8 (3.87%), and G17 (3.05%), which also showed 136 positive heterosis over their mid parent (Table 3). Positive heterosis suggests longer harvest period 137 whereas negative heterosis suggests shorter harvest period. Generally, longer and shorter harvest 138 duration is preferred by the homestead and commercial growers, respectively. Positive heterosis for the 139 trait was also reported by Kumari and Sharma [14] and Khan and Jindal [21]. Therefore, these genotypes

would be the effective combination in exploiting heterosis for the homestead and commercial growers astheir desire.

- 142**3.6 Plant height at 1**st harvest: Significant negative heterosis for better parent was manifested by five143 F_{15} viz. G11 (-15.32%), G20 (-10.56%), G13 (-10.25%), G1 (-9.76%) and G19 (-7.74%). Only two F_{15} viz.
- G11 (-11.85%), and G20 (-6.76%) produced significant negative heterosis for their mid parent (Table 3).
- Significant positive heterosis for better parent was also found from the crosses G14 (16.60%) and G17
- 146 (8.87%). This result is similar to that of Baishya et al. [18] and Padma et al. [22]. Patwary et al. [16]
- reported both positive and negative heterosis for their study whereas Fageria *et al.* [23] reported only positive heterosis. So, these genotypes can further be used to develop inbred lines toward developing of both taller and dwarf varieties.
- **3.7 Fruit set (%):** Seventeen out of 21 F_{1S} produced significant different (P = 0.05 or P = 0.01) positive heterosis over their better parent whereas 16 produced significant positive heterosis over their mid parent (Table 4). Ten cross combinations *viz*. G20 (25.57%), G8 (17.00%), G18 (14.82%), G9 (10.29%), G19 (4.71%), G16 (3.72%) and G11 (2.04%) produced significant positive heterosis either their mid or better parent indicating potential increment of fruit set. On the other hand, seven F_{1S} performed negative heterosis ranging from -1.68 to -22.11% indicating a reduction in fruit setting. Both positive and negative heterosis in respect of fruit setting was reported by El-Ahmadi and Stevens [24].
- 157 **3.8 Number of fruits per plant:** About 50% of the F_{1s} showed significant different (P = 0.05 or P = 0.01) 158 positive heterosis over their better parent ranging from 7.86 to 45.99% (Table 4). More than 40% 159 heterosis over their better parent was produced by four crosses viz. G3, G10, G13, G18. On the other 160 hand, about 76% of the F1s produced significant positive heterosis over their mid parent ranging from 161 12.05 to 63.55% (Table 4). This result suggested a potential increment of fruits number in the tomato 162 plant. This study showed a bit higher amount of heterosis for fruits number than the previous study by Patwary et al. [16]. It could be due to the variation of the parents used in the study. Our study also had an 163 164 agreement with the previous research [6,18,19,20,23].
- **3.9 Fruit length (cm):** Fourteen hybrids showed positive heterosis, of which 5 hybrids exhibited positive significant heterosis over better parents (Table 4). More than 10% heterosis was estimated from four crosses *viz.* G6, G4, G14, and G20. Only one hybrid G18 (-12.93%) produced the significant negative heterosis over better parent. Since, only a genotype out of twenty one showed significant negative heterosis over better parent, indicating character is mainly governed by non-additive gene effects. Islam *et al.* [18] also reported similar results for fruit length. Significant positive heterosis has been reported by
- 171 Ahmad *et al.* [6], and Sharma *et al.* [20]. These findings of significant positive heterosis over mid and 172 better parent are in line with the findings of Singh *et al.* [5] and Kumar and Singh [25] as well.
- 3.10 Fruit diameter (cm): About 62% hybrids exhibited with significant positive heterosis over better
 parent, whereas 76% produced significant positive heterosis over mid parent (Table 4). The highest value
 of positive heterotic effect was exhibited by the cross G4 (53.70 %) followed by G2 (48.46 %), G13 (46.54
- 176 %), G7 (42.50 %) and G14 (40.00 %). One-third of the hybrids produced significant negative heterosis for

- 177 either mid or better parent, which suggested that the character is possibly governed by non-additive gene
- action. Heterosis for fruit diameter in tomato was also reported by Ahmad et al. [6], Padma et al. [23], and
- 179 Sharma *et al.* [20].
- 180 **3.11 Average fruit weight (g):** The entire cross combinations except G18 and G4 exhibited with negative
- heterosis over mid and better parent, whereas two hybrids G18 (12.09%) and G4 (12.01%) showed
 significant positive heterosis over mid parent (Table 4). The best hybrid was G18, which showed the

 - 183 highest per se performance with the highest heterosis (12.09%) over mid parent. Positive heterosis for
 - fruit weight has been reported by Sharma et *al.* [19,20], whereas both positive and negative heterosis over better parent reported by Patwary *et al.* [16] and Ahmad *et al.* [6] in their studies. These findings of positive heterosis over mid parent and check co-relate with the findings of Kumari and Sharma [14] and Marbal *et al.* [26].
 - **3.12 Total soluble solid (TSS):** Significant positive heterosis over mid and better parent was observed in all the F_{1S} confirming additive gene effect for the trait (Table 4). The highest positive heterosis was observed in cross G20 (141.67%) followed by G17 (84.76%), and G16 (80.83%). Similar range bof heterosis was also noted by the previous studies [8,17,19,20,22,27]. Total soluble solid is responsible for the sweetness of tomato hereafter high TSS is a preferable character in processing tomatoes. So, these genotypes can further be advanced toward developing a processing variety.
- 194 **3.13 Fruit yield per plant (kg):** Off 21 crosses, six produced significant different (P = 0.05 or P = 0.01) 195 positive heterosis over better parent, whereas 15 produced significant positive heterosis over mid parent 196 (Table 5). More than 20% heterosis over better parent was observed in five F_{1s} viz. G18 (53.77%), G20 197 (50.31%), G16 (39.20%), G17 (36.49%), and G5 (25.70%). The cross combinations G18 (70.00%), G16 198 (58.74%) and G20 (55.63%) showed higher positive heterosis over mid parent. This result suggested a 199 potential yield increment by the heterosis, and is predicted to be the reason of high yielding parents used 200 in the hybridization [28]. Eight genotypes exhibited with significant negative heterosis over either mid or 201 better parent. Positive better parent heterosis ranging from 13.58 to 282.63% was reported in heat 202 tolerant tomato [16], which was higher than this study. Bhatt et al. [8.27] observed 2.92 to 54.17% better 203 parent heterosis for yield per plant in tomato, which is very identical to our findings. Similarly, 204 heterobeltiosis in tomato hybrids was also reported in many studies [3,6,14,25,26,29,30]. Therefore, 205 these genotypes may be selected as heterotic hybrids for yield and can further be advanced toward developing a high yielding variety. 206
- **3.14 Number of locules per fruit:** Seven cross combinations out of 21 showed positive heterobeltiosis but only two was significant. Positive heterosis for this trait ranged from 1.94 to 56.66% (Table 5). On the other hand, nine cross combinations produced significant negative heterosis over better parent ranging from -18.15 to -51.38%. More than 35 % negative heterosis was manifested by five F_{1S} namely G8 (– 51.38 %), G16 (– 46.03 %), G18 (– 46.03 %), G17 (– 40.02 %) and G15 (– 36.29 %). Similarly, eight F_{1S} showed significant positive heterosis over mid parent and five F_{1S} showed significant negative heterosis over mid parent. The hybrid G20 showed no heterosis regarding locule number in fruit (Table 5).

However, the estimation of negative heterobeltiosis from -4.50 to -51.39% was observed from the study, indicating the importance of non-additive gene action for the trait. As a result, heterosis breeding can be

- exploited very well to reduce the locule number in tomato fruits. This result supported by Duhan et al.
- 217 [31], Kurian et al. [7] and Dod et al. [32] in where identified heterotic hybrids for lower locule number in
- tomato. On the other hand, Ahmad *et al.* [6] reported significant positive heterosis for this trait. From the
- 219 quality point of view, less locule is desirable in tomato. This study is predicted the potential genotypes for
- 220 future breeding in reducing locule as we have seen negative estimation of heterosis.
- 221 **3.15 Pericarp thickness:** The highly significant different (P = 0.05 or P = 0.01) heterosis was estimated 222 by the majority of the hybrids towards positive heterosis over mid parent, whereas 12 hybrids produced 223 significant positive heterosis for better parent ranging from 26.67 to 109.06% (Table 5). More than 25% 224 heterosis exhibited by the 57% hybrids, indicating possibility of the enhancement of fruit quality by 225 improving pericarp thickness. Only a single hybrid G12 produced significant negative heterosis for both 226 mid and better parent. The results of the study in relation to pericarp thickness were agreed by the 227 previous studies [14,16,19,21,33,34]. Pericarp thickness usually contributes much for long storability. 228 Positive heterosis is the indicator of additive gene action for the trait, and is predicted to increase pericarp 229 thickness of tomato using these genotypes in a variety development program.
- **3.16 Number of seeds per fruit:** Significant negative heterosis was manifested by 19 hybrids varying from -10.30 to -67.56% for both mid and better parent (Table 5). The highest negative heterotic value was achieved by the hybrid G8 (-67.56) followed by G3 (-65.41), G21 (-59.51) and G9 (-59.39) whereas the lowest negative heterosis was provided by the hybrid G4. Ahmad *et al.* [6] and El-Ahmadi and Stevens [24] reported higher degree of heterosis for this trait. Negative heterosis is an indication of the reduction of seeds in tomato as the consumers expect. So, these cross combinations can be further used toward developing less seeded tomato varieties.
- **3.17 1000-seed weight:** The highly significant different (P = 0.05 or P = 0.01) positive heterosis was observed by 48% of the hybrids over better parent (Table 5) indicating seed quality can be improved through the hybridization. More than 10% positive heterosis was manifested by five hybrids *viz*. G4 (18.11%), G10 (15.81%), G9 (14.45%), G19 (11.39%), and G21 (12.13%). Nine hybrids provided significant negative heterosis ranging from -4.30 to -26.94%. This result is in accordance with the findings of Subburamu *et al.* [35].
- 243

244 4. CONCLUSION

None of the cross combinations was heterotic for all characters simultaneously. In this study, promising hybrids for yield per plant with significant over better parent in desirable direction and also revealed for other traits *viz*, days to flowering and harvesting, number of fruits per plant, fruit length, fruit diameter, pericarp thickness, number of locules per fruit, plant height, TSS, 1000-seed weight (Table 6). As a result, high heterosis for yield appears to be the consequence of heterosis of the yield attributing traits. Among the hybrids G5, G13, G16, G17, G18 and G20 were promising for yield per plant as well as for many

251	other y	ield contributing traits. Therefore, these hybrids can be used to develop high yielding varieties
252	along w	vith other quality traits.
253		
254 255	СОМР	ETING INTERESTS
256 257	Authors	s have declared that there was no competing interests exist
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Table 1 Developed F₁ **hybrids by a half diallel fashion**

Parent (P)	P 1	P2	P3	P4	P5	P6	P7
P1 (WP10)	-	P1× P2 (² G1)	P1 × P3 (G2)	P1 × P4 (G3)	P1 × P5 (G4)	P1 × P6 (G5)	P1 × P7 (G6)
P2 (VRT003)			P2 × P3 (G7)	P2 × P4 (G8)	P2 × P5 (G9)	P2 × P6 (G10)) P2 × P7 (G11)
P3 (VRT004)				P3 × P4(G12)	P3 × P5 (G13)	P3 × P6 (G14)) P3 × P7 (G15)
P4 (LE009)					P4 × P5 (G16)	P4 × P5 (G17)) P4 × P5 (G18)
P5 (TLB182)						P5 × P6 (G19)) P5 × P6 (G20)
P6 (WP02)							P6 × P7 (G21)
P7 (TLB111)							-
^z Hybrid							

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Table 2 ANOVA for various traits of 21 F₁s and seven parents of tomato

Characters	Mean Squares							
Characters	Replications (^Y df = 2)	Genotypes (df = 27)	Error (df = 54)					
Days to 1 st flowering	0.94	8.47**	0.77					
Days to 50% flowering	3.62	24.29**	0.58					
Days to 1 st harvesting	9.33	41.37**	10.54					
Harvest duration	52.27	95.44*	14.33					
Plant height at 1 st harvest (cm)	60.69	174.81**	11.253					
Fruit set percentage (%)	130.47	107.71	67.81					
Number of fruits per plant	7.20	291.75**	9.65					
Fruit length (cm)	1.48	1.44**	0.21					
Fruit diameter (cm)	0.65	4.33**	0.13					
Average fruit weight (g)	38.56	1829.54**	29.67					
Yield per plant (kg)	0.11	0.47**	0.07					
Total soluble solid (%)	0.35	6.09**	0.26					
Locules per fruit	0.81	5.55**	0.76					
Pericarp thickness (mm)	4.51	4.53**	0.44					
Seeds per fruit	1505.23	1063.47**	5.04					
1000-seed weight (g)	2.99	0.27**	0.001					

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Degree of freedom; *, ** = Significant difference at P = 0.05 and P = 0.01 respectively

Genotypes	Days to 1	I st flowering	Days to	Days to 50% flowering		Days to 1 st harvest		t duration	Plant height at 1 st harvest (cm)	
	MP	BP	MP	BP	MP	BP	MP	BP	MP	BP
G1	-9.890**	-10.87**	14.129**	-12.10**	-9.979**	-10.38**	-6.495**	-6.77**	-1.087	-9.76*
G2	-0.090**	-9.35**	0.268**	-7.64**	-0.086**	-9.89**	-0.056**	-6.49**	0.146**	6.12
G3	-2.571**	-4.32**	30.297**	-7.64**	-4.573*	-9.02**	-2.953**	-5.92**	-0.531	-6.394
G4	-9.290**	-11.59**	39.753**	-12.10**	-8.299**	-8.98**	-5.378**	-5.84**	5.172*	-4.18
G5	-8.834**	-10.15**	21.191**	-10.97**	-10.619**	-11.92**	-4.046**	-4.96**	10.810**	7.64
G6	-8.644**	-9.30**	28.177**	-14.82**	-5.660**	-7.78**	0.888	-0.58	8.778**	3.08
G7	-5.836**	-7.19**	11.644**	-12.10**	-6.899**	-7.85**	1.441*	0.76	13.370**	-3.47
G8	-2.222**	-2.22*	2.923	-8.91**	-2.877	-5.04*	5.400**	3.87**	0.971	-2.31
G9	-3.011**	-4.44**	14.573**	-9.56**	-5.703**	-6.82**	1.561*	0.78	-1.098	-1.25
G10	-3.357**	-3.71**	6.805**	-4.45**	-5.436**	-6.40**	-3.556**	-4.20**	9.213**	-2.94
G11	-3.284**	-5.01**	13.311**	-8.65**	-3.348	-5.93**	-2.161**	-3.87**	-11.851**	-15.32**
G12	-10.216**	-11.50**	1.038	-14.01**	-7.207**	-10.18**	-4.678**	-6.68**	8.902**	-4.61
G13	-5.933**	-8.63**	18.200**	-12.10**	-7.578**	-9.59**	-4.937**	-6.30**	5.535*	-10.25**
G14	-4.769**	-6.48**	19.816**	-5.73**	-6.977**	-6.98**	-4.775**	-4.78**	22.472**	16.60**
G15	-6.817**	-7.16**	21.154**	-12.35**	-3.774*	-7.27**	-0.686	-3.06**	9.953**	-3.18
G16	-3.755**	-5.18**	11.696**	-12.10**	-2.000*	-3.05*	5.442**	4.72**	1.355	-2.09
G17	-4.093**	-4.44**	3.398	-12.73**	-2.703**	-5.82**	5.263**	3.05*	18.877**	8.87*
G18	-1.102	-2.87**	7.447*	-4.94**	0.786	0.32	5.898**	5.58**	5.484*	4.72
G19	-5.660**	-6.72**	13.927**	-13.38**	-7.284**	-9.31**	-4.745**	-6.11**	3.951	-7.74*
G20	-8.501**	-11.44**	23.821**	-16.67**	-4.324**	-5.78**	-2.185**	-3.15**	-6.762**	-10.56**
G21	-8.036**	-10.01**	11.162**	-15.43**	-6.181**	-9.59**	-3.419**	-5.73**	7.221**	-1.15
SE	0.620	0.72	0.539	0.62	2.296	2.65	2.677	3.09	2.372	2.74
CD at 5%	0.507	0.83	0.442	0.72	1.879	3.07	2.192	3.60	4.766	7.78
CD at 1%	0.675	1.10	0.588	0.96	2.503	4.09	2.919	4.82	6.347	10.36

Table 3 Percent heterosis over mid parent (MP) and better parent (BP) for days to 1st flowering, days to 50 % flowering, days to 1st
 harvest, harvest duration and plant height at 1st harvest in winter tomato.

341 *, ** = Significant difference at P = 0.05 and P = 0.01 respectively

343Table 4 Percent heterosis over mid parent (MP) and better parent (BP) for fruit set (%), number of fruits, fruit length, fruit diameter,
average fruit weight, and TSS% in winter tomato.

Genotypes	Frui	t set (%)		of fruits per lant	Fruit ler	ngth (cm)	Fruit dia	meter (cm)		fruit weight (g)		luble solid SS %)
	MP	BP	MP	BP	MP	BP	MP	BP	MP	BP	MP	BP
G1	17.873**	8.48**	16.130**	-0.79	14.129**	-5.10	18.182**	17.50**	-18.374**	-23.37**	43.992**	40.78**
G2	-0.209**	-22.1083**	-0.024	-14.73*	0.268**	3.47	0.493**	48.46**	-0.153**	-30.77**	0.487**	45.07**
G3	-14.062**	-22.1083**	4.847	40.47**	30.297**	6.03	0.242	-14.35**	1.402	-49.55**	60.432**	50.81**
G4	2.278	-11.3983**	14.251**	10.08	39.753**	26.91**	79.342**	53.70**	12.014**	-1.87	75.949**	41.26**
G5	-5.431**	-9.92**	45.655**	27.97**	21.191**	3.93	25.897**	25.53**	-15.462**	-26.28**	34.054**	26.02**
G6	-3.437*	-14.20**	23.217**	-3.43	28.177**	27.47**	37.422**	6.81*	-13.165**	-38.18**	89.394**	52.44**
G7	-1.727	-8.21**	-1.680	-24.83**	11.644**	9.03*	42.500**	42.50**	-14.049**	-33.07**	34.302**	34.04**
G8	18.968**	17.00**	18.833**	10.36*	2.923	0.17	-14.538**	-26.62**	-11.228**	-26.96**	51.266**	39.22**
G9	17.851**	10.29**	52.504**	26.34**	14.573**	3.83	31.680**	12.31**	-13.790**	-19.95**	74.170**	37.48**
G10	11.079**	7.12**	46.113**	41.42**	6.805**	2.91	24.397**	24.04**	-18.484**	-24.68**	58.228**	45.63**
G11	6.043**	2.04	38.380**	24.60**	13.311**	-6.19	17.764**	-8.85**	-18.163**	-39.32**	71.779**	35.92**
G12	1.652	-6.51**	30.239*	-5.34	1.038	0.69	8.434**	-6.90**	-23.550**	-48.02**	50.947**	38.69**
G13	1.689	-10.69**	58.967**	43.51**	18.200**	4.86	71.815**	46.54**	-9.009**	-32.77**	84.049**	45.07**
G14	-9.178**	-12.14**	0.562	-21.29**	19.816**	12.85**	40.405**	40.00**	-1.410	-27.41**	47.368**	35.40**
G15	-3.204*	-12.76**	-13.762	-38.61**	21.154**	-1.56	56.522**	21.15**	-1.887	-37.74**	83.599**	45.07**
G16	9.104**	3.72	63.554**	27.84***	11.696**	-1.21	0.366	-24.41**	-7.057**	-18.52**	114.227**	80.83**
G17	-2.192	-7.18**	39.286**	25.52**	3.398	-2.93	-11.594**	-24.28**	-1.117	-12.91**	84.758**	84.76**
G18	17.400**	14.82**	50.968**	45.99**	7.447*	-12.93**	-2.376	-32.00**	12.090**	-2.04	97.817**	67.44**
G19	15.743**	4.71	12.053**	-4.65	13.927**	6.88	38.009**	17.99**	-13.097**	-13.56**	103.010**	71.36**
G20	29.269**	25.57**	41.351**	7.86*	23.821**	11.88**	51.534**	34.61**	0.955	-21.08**	142.475**	141.67**
G21	5.802**	-1.68	22.581**	7.23	11.162**	-5.11	21.945**	-5.42	-6.990*	-27.01**	86.357**	57.74**
SE	5.823	6.72	2.197	2.54	0.326	0.38	0.255	0.29	3.851	4.45	0.254	0.29
CD at 5%	1.941	3.17	1.798	2.94	0.267	0.44	0.209	0.34	3.152	5.15	0.208	0.34
CD at 1%	2.586	4.22	2.394	3.91	0.356	0.58	0.278	0.45	4.198	6.86	0.277	0.45

345 *, ** = Significant difference at P = 0.05 and P = 0.01 respectively

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Table 5 Percent heterosis over mid parent (MP) and better parent (BP) for yield, number of locules, pericarp thickness, number of seeds and 1000-seed weight in winter tomato. 348 349

Genotypes	Fruit yield per plant (kg)			Number of locules per fruit		Pericarp thickness (mm)		Number of seeds per fruit		1000-seed weight (g)		
	MP	BP	MP	BP	MP	BP	MP	BP	MP	BP		
G1	-3.967	-13.21**	52.624**	48.04**	100.667**	89.91**	-23.822**	-29.60**	10.891**	9.375**		
G2	-0.144**	-20.72**	0.254**	-4.50	0.254**	-4.75	0.066**	2.91*	0.017**	-9.907**		
G3	14.286**	-5.98	0.000	-18.96**	59.627**	26.67**	-65.341**	-65.41**	13.238**	-13.932**		
G4	26.923**	7.94	67.598**	56.66**	80.317**	79.18**	19.874**	-10.30**	19.284**	18.110**		
G5	26.588**	25.70**	22.549**	15.47	78.650**	77.81**	-30.740**	-34.59**	7.486**	2.941**		
G6	20.533**	5.61	20.950**	13.06	38.436**	34.07*	-14.287**	-23.42**	1.581**	0.000**		
G7	-7.364*	-9.81*	28.088**	-4.50	37.066**	0.33	-12.614**	-16.51**	-18.480**	-26.935**		
G8	7.759*	-5.66	-38.588**	-51.38**	66.221**	26.67**	-67.249**	-67.56**	-1.606**	-4.297**		
G9	29.639**	1.51	5.916	1.94	105.034**	95.21**	-42.850**	-59.39**	14.902**	14.453**		
G10	18.908**	6.79	26.103**	15.47	121.891**	109.06**	-37.866**	-38.93**	19.318**	15.809**		
G11	18.310**	-4.91	5.916	1.94	58.621**	54.88**	-41.260**	-43.38**	-13.450**	-13.619**		
G12	21.778**	9.16	3.704	-4.50	-23.130**	-27.54**	-32.397**	-34.45**	7.257**	-6.192**		
G13	50.125**	19.92***	12.570*	-18.15**	18.960*	-10.00	33.333**	-2.50	2.600**	-8.359**		
G14	10.390**	1.59	-8.576	-27.29**	37.849**	5.08	-27.156**	-28.82**	9.916**	1.238**		
G15	9.223*	-10.36*	-12.383*	-36.29**	10.915*	-17.54	-18.898**	-25.18**	-2.414**	-12.384**		
G16	58.739**	39.20**	-29.895**	-46.03**	41.149**	11.48	14.594**	-17.81**	-14.516**	-16.535**		
G17	40.488**	36.49**	-30.095**	-40.52**	19.070**	-5.19	-28.994**	-29.55**	7.782**	1.838**		
G18	70.000**	53.77**	-29.895**	-46.03**	9.677*	-15.00	-52.369**	-54.76**	5.812**	2.724**		
G19	-2.493	-16.59**	4.439	-7.62	97.788**	95.66**	-20.550**	-42.74**	15.209**	11.397**		
G20	55.627**	50.31**	0.000	0.00	86.230**	81.47**	-31.146**	-52.12**	0.587	0.000		
G21	18.280**	4.27	-13.055	-23.10*	45.543**	40.31**	-57.053**	-59.51**	15.312**	12.132**		
SE	0.184	4.27	0.616	0.71	0.470	0.54	1.587	1.83	0.022	0.03		
CD at 5%	0.151	0.25	0.505	0.82	0.385	0.63	1.299	2.12	0.018	0.03		
CD at 1%	0.201	0.33	0.672	1.10	0.512	0.84	1.730	2.83	0.024	0.04		

*, ** = Significant difference at P = 0.05 and P = 0.01 respectively

Table 6 Promising F_1 hybrids showing higher per se performance and better-parent heterosis (BPH) for yield per plant and significant BPH for other characters

Genotypes	Yield per plant (kg)	BPH (%) for yield	BPH for other characters
G18	3.06	53.77**	# of locule, # of seeds per fruit, 1000-seed weight, harvest duration, TSS
G20	2.42	50.31**	Days to 1 st flowering, days to 50% flowering, days to 1 st harvest, harvest duration, fruit length, fruit diameter, TSS, pericarp thickness, # of seeds per fruit, # of fruits per plant
G16	2.77	39.20**	Days to 1 st flowering, days to 50% flowering, days to 1 st harvest, harvest duration, fruit diameter, TSS, # of seeds per fruit, # of fruits per plant
G17	2.88	36.49**	Days to 1 st flowering, days to 50% flowering, days to 1 st harvest, harvest duration, plant height, fruit diameter, TSS, # of seeds per fruit, # of fruits per plant, 1000-seed weight
G5	2.67	25.70**	Days to 1 st flowering, days to 50% flowering, days to 1 st harvest, harvest duration, fruit diameter, TSS, # of seeds per fruit, # of fruits per plant, 1000-seed weight
G13	3.02	19.92**	Days to 1 st flowering, days to 50% flowering, days to 1 st harvest, harvest duration, fruit diameter, TSS, # of fruits per plant, # of locule