

Generation Mean Analysis for Fruit Yield and Component Traits in Tomato (*Solanum lycopersicum* L.)

Pragati J. Prajapati^{1*}, J.N. Patel², Parthik Patel³ and N.A. Patel⁴

¹Ph.D. Scholar, Department of Genetics and Plant Breeding, Anand Agricultural University, Anand (Gujarat), India.

²Research Scientist & Head, Bidi Tobacco Research Station, Anand Agricultural University, Anand (Gujarat), India.

³Agriculture Officer, C/o Deputy Director of Agriculture (Ext.), Anand (Gujarat), India.

⁴Assistant Research Scientist, Main Vegetable Research Station, Anand Agricultural University, Anand (Gujarat), India.

(Corresponding author: Pragati J. Prajapati*)

(Received: 27 February 2023; Revised: 18 April 2023; Accepted: 22 April 2023; Published: 20 May 2023)

(Published by Research Trend)

ABSTRACT: Information of gene action governing the yield and its components is very crucial for formulating sound breeding programmes. In order to generate this information for tomato lines, the present investigation was carried out at Main Vegetable Research Station, Anand Agricultural University (AAU), Anand during *kharif-rabi* 2022-23. The experimental material comprised of six families developed from ten diverse lines. Each family is composed of six generations *viz.*, P₁, P₂, F₁, F₂, B₁ & B₂ evaluated with three replications in compact family block p; design (CFBD) to perform generation mean analysis. Results of simple scaling tests and joint scaling tests revealed adequacy of three parameter model in one out of eighty-four possible cases and adequacy of six parameter model in eighty two out of eighty-four possible cases.

Keywords: Tomato, Generation mean analysis, Scaling test, Gene action, Gene interaction.

INTRODUCTION

Tomato is the world's second-most widely cultivated vegetable crop trailing potato. Total area under tomato cultivation in the world was 5.05 million hectares, with production of 186 million tonnes and the average productivity of 37.0 tonnes/ha in 2020. China stands first in the major tomato growing countries followed by India, Turkey, Egypt, Iran, USA, Mexico, Italy, Brazil and Spain (Anonymous, 2020). Tomato [*Solanum lycopersicum* L.], ($2n = 20x = 24$) is self-pollinated, day-neutral, extensively cultivated and globally consumed vegetable crop in world (Sikder *et al.*, 2013). As yield is not simply an inherited trait, many genes' action and interaction determine inheritance and potential yield. Many yield attributing characters have positive and/or negative association, it is very hard to isolate line possessing all desirable traits. Hence, estimation of components of gene action and genetic variance is very essential to formulate robust breeding programmes. Knowledge of the nature and magnitude of gene effects controlling inheritance of yield and its attributing traits would aid in the choice of efficient breeding methods, ultimately aid in accelerating the pace of its genetic improvement and breaking the yield barriers. Most of the yield attributing traits generally show continuous variation and they are influenced by environment (Lecomte *et al.*, 2004).

Amount of genetic variability present in the breeding material and knowledge of genetic control of commercial traits is very crucial for breeding programmes in order to isolate improved cultivars and hybrids through proper breeding methodology. Even though tomato is self-pollinated crop, easy emasculation, better pollen dispersal, profuse flowering and higher seed multiplication ratio lead to better adoption and commercialization of hybrids (Damor *et al.*, 2021). Considerable important work has been done in this crop, but better information in the genetics of fruit yield, yield attributing traits and quality parameters of this crop, grown in middle Gujarat agro-climatic condition is still needed.

Generation mean analysis (Mather and Jinks 1982) is a useful tool for determining the nature of gene effects (additive, dominance and their digenic interaction) involved in the expression of traits such as yield and its associated traits. The scaling test examines generation means to determine presence or absence of epistasis as well as complementary (additive × additive) or duplicate (additive × dominance) and (dominance × dominance) interaction at digenic level. The present study was carried out to study the nature of gene action involved for the inheritance of yield and associated traits in six tomato crosses.

MATERIAL AND METHODS

The six generations of six crosses (2012/TODVAR-1 × AVTOV 1007, GAT-5 × 2015/TOLCV RES-1, 2014/TODVAR-5 × AVTOV 1002, 2016/TODVAR-12 × AVTOV 1005, 2017/TODVAR-8 × 2015/TOLCV RES-1, 2017/TODVAR-8 × 2015/TOLCV RES-4) comprising of P₁, P₂, F₁, F₂, B₁ and B₂ were developed from ten diverse parents *viz.*, 2012/TODVAR-1, AVTOV 1007, GAT-5, 2015/TOLCV RES-1, 2014/TODVAR-5, AVTOV 1002, 2016/TODVAR-12, AVTOV 1005, 2017/TODVAR-8 and 2015/TOLCV RES-4. The crosses were made in *kharif-rabi* 2020-21 and 2021-22 and evaluation of the experimental material was done during *kharif-rabi* 2022-23.

Field Experiment. The six generations of these six crosses were raised in Compact Family Block Design (CFBD) at Main Vegetable Research Station (MVRS), Anand Agricultural University (AAU), Anand during *kharif-rabi* 2022-23 with three replications. An individual replication had six families as blocks and each block consisted of one row of each P₁, P₂ and F₁ generation, four rows of each F₂ generation and two rows of each B₁ and B₂. 25 days old seedling then transplanted to well-prepared field keeping inter and intra-row spacing of 90 × 45 cm. The observations for different characters were recorded on randomly selected plants from each experimental unit from each replication. Observations were recorded on five plants each for P₁, P₂ and F₁, on twenty plants for F₂ and on ten plants for B₁ and B₂, respectively.

RESULTS AND DISCUSSION

Analysis of variance between the family comparison depicted significant differences among the families for all the traits. For estimation of components of gene effect, Simple Scaling Tests (Hayman and Mather, 1955) were applied. The non-significance test of all the Simple Scaling Tests suggest adequacy of additive dominance model; hence, principle gene effects *i.e.* additive and dominance were estimated as suggested by Jinks and Jones (1958) three parameters model (m, d and h). For the families and characters, wherein any of the Simple Scaling Test was significant, six parameters model (m, d, h, i, j and l) as suggested by Hayman (1958) was applied to partition the gene effect into epistatic components including principle gene effects. However, for confirmation of adequacy of additive dominance model, and to realize presence of higher order interallelic interactions, Joint Scaling Test as suggested by Cavalli (1952) was also applied. Further, the results were confirmed by significance of χ^2 test. The opposite sign of [h] and [l] indicated duplicate gene interaction. While, equal sign of [h] and [l] indicated complimentary gene interaction.

Days to Flowering. Additive dominance model was found to be inadequate since estimates of individual simple scaling tests, A, B, C and D found significant for the all six families except, family II and VI. In family II and VI scales A and D found significant. Further, presence of higher order epistasis was indicated and confirmed by significance of χ^2 test value of joint scaling test, which indicated inadequacy of the additive-

dominance model in all the six families for days to first flowering. Significant additive and dominance components of gene action were reported for the families I, II, III and IV indicating importance of both components. While, in family IV, additive component was found significant. All the families except, family I possessed significant estimates of additive × additive [i] and additive × dominance [j] type of epistatic gene interaction. Dominance × dominance [l] epistatic interaction found significant for all the families. The results were in accordance with the findings of Damor *et al.* (2021).

Branches per Plant. Additive dominance model found inadequate since simple scaling tests, A and D found significant in family I, B in family II, A and B in family III, A in family IV, C and in family V and B and C in family VI. Further the results were confirmed by significance of χ^2 test for all families. Additive component of gene action was recorded significant for family III and IV; whereas dominance component was significant for family I and V. Additive × additive [i] type of epistasis found significant in family I, V and VI. While, additive × dominance [j] interaction reported significant in family II and IV. Dominance × dominance [l] type of epistatic interaction found significant in family I, II, III and VI. From the signs of [h] and [l], complimentary epistasis reported for branches per plant except, in family I. Duplicate epistasis reported in family I was in accordance with findings of Das *et al.* (2020) who reported both duplicate and complementary type of epistasis. Analysis contradicted the findings of Parida *et al.* (2021); Kumar and Srivastava (2021) who reported adequacy additive dominance for the inheritance of branches per plant.

Plant Height. For plant height, simple additive dominance was found inadequate as scaling testes were found significant in all families. Significance of χ^2 test value from the joint scaling for all families confirmed the results. Significance of additive component of gene action was reported for family I, II and III; whereas only family I exhibited significant estimate of dominance component. Predominant role principal component of gene action (*i.e.* additive and dominant) recorded in family I. Additive × additive [i] type of epistatic interaction found significant in family I and III. Additive × dominance [j] interaction was found significant only in family II; whereas higher estimates of dominance × dominance [l] epistatic gene interaction reported in in family IV and V which indicates role of non additive gene effects. Complementary epistasis was reported to govern the traits in all families except family I. The results were in accordance with the study of Negi *et al.* (2013).

Fruit Length. Significant values of individual scaling tests were reported for fruit length in all families. Further, it is supported by significance of χ^2 test values. Significant values for additive genetic component were observed for all families under study for fruit length. While, significant estimates of dominance component were reported in family III and IV. Significant estimates of additive × additive [i] and dominance × dominance [l] were reported in family I with higher

magnitude of dominance \times dominance [I] inter allelic gene interaction. On the other hand, additive [I] gene action with additive \times dominance [j] epistatis interaction was reported in family II. In family III and IV, both principle gene action components with all three epistatis interaction *viz.* additive \times additive [i], additive \times dominance [j], dominance \times dominance [I] with higher magnitude of dominance \times dominance [I] type of gene interaction found to govern the fruit length. Four out of six families (*viz.* I, II, III and IV) showed duplicate type of gene interaction indicating complex inheritance of fruit length. Magnitude of all type epistatis was higher in all families, which governs the character fruit length. Importance of additive gene effect and duplicate epistatis were in accordance with the findings of Chauhan *et al.* (2019).

Fruit Girth. Additive dominance model was found to be inadequate since individual scaling tests had significant estimates for the fruit girth. It was further supported by χ^2 test value of joint scaling test at three degree of freedom for all families under study. Significant estimates of additive gene effect were reported for family II, III and V; whereas family I, III and IV exhibited significant estimates of dominance gene effect for the trait under study. Family I possessed significant estimate of additive \times additive [i] type of gene interaction along with significant dominance gene effect. Additive gene effect along with additive \times dominance [j] epistatis interaction found to govern the character in family II. All three types of epistatis gene interactions (*viz.* additive \times additive [i], additive \times dominance [j], dominance \times dominance [I]) were found significant along with significant in families III and IV; however, family III also exhibited significant estimate of dominance component of principle gene effect. In family IV, dominance gene effect with additive \times dominance [j] found significant. Only additive \times dominance [j] gene interaction reported significant in family VI. All families except, family II had opposite signs of [h] and [I] stating presence of duplicate type of gene interaction. The findings were partially in accordance with Patel *et al.* (2010).

Average Fruit Weight. Significant estimates of the individual simple scaling test as well as significance of χ^2 test value from joint scaling test revealed presence of digenic interaction in the all six families under study. Significant estimates of principle gene effect components *i.e.* additive and dominance components were reported only in family I and VI but in negative direction. Family I and VI exhibited significant estimates of additive \times additive [i] and dominance \times dominance [I] epistatis interaction with estimates of dominance \times dominance [I] in positive direction revealing preponderance of non-additive gene action in the inheritance of average fruit weight in these families. Additive \times dominance [j] gene interaction was found significant in family II and III along with significant dominance \times dominance [I] interaction in family III. Family IV and V epistatis interaction additive \times additive [i] and dominance \times dominance [I] with higher estimates of dominance \times dominance [I] interaction. Here, both duplicate (4 families) and complimentary (2

families) gene interaction reported in the analysis of this trait. Similar findings with more prominent dominance gene effects were reported by Patel *et al.* (2010); Chauhan *et al.* (2019). Duplicate epistatis for this trait was also reported by Mawasid *et al.* (2019); Kumar and Srivastava (2021); Parida *et al.* (2021).

Pericarp Thickness. The failure of additive-dominance model was observed due to significance of scaling tests in all the families. The χ^2 test value of joint scaling tests were significant for the all families under study. Significant additive gene effect was recorded for family V only; whereas significant estimates of dominant gene effect were recorded in family IV, V and VI. Only additive \times dominance [j] and dominance \times dominance [I] gene interaction was found significant in family I and III, respectively. While, heterozygous [additive \times dominance] inter allelic interaction along with dominance \times dominance [I] found significant in family II. All three types of intergenic interactions *viz.* additive \times additive [i], additive \times dominance [j] and dominance \times dominance [I] were found significant in families IV and V. Significant additive \times additive [i] and additive \times dominance [j] epistatis interactions were reported for the family VI. Duplicate type of epistatis was observed in all the families except, family I, as [h] and [I] had opposites sign which reveals complex inheritance for the character under study.

Fruit Yield per Plant. Inadequacy of additive dominance to explain the inheritance of fruit yield per plant was revealed by significance of individual scaling test for all the families. The outcomes were supported by significance of χ^2 test values from the joint scaling test for all families. This revealed presence of digenic interaction for the character under study. Significant estimates of both additive and dominance gene action components were found for the families IV, V and VI. Although, significant value of dominance gene action was reported for family III. Only additive \times dominance [j] and dominance \times dominance [I] epistatis interaction found significant in family II and III, respectively. Additive \times dominance [j] and dominance \times dominance [I] type of epistatis interactions were found significant in family VI and V, along with significant dominance \times dominance [I] interaction in family V. Significant estimate of dominance \times dominance [I] interaction component with higher magnitude was reported in family VI. Duplicate type of epistatis was reported in all families. Similar findings with presence of non-allelic interaction along with were reported by Patel *et al.* (2010) reported higher non additive gene action comparable to additive gene actions responsible for fruit yield per plant. Duplicate epistatis for this trait was reported by Chauhan *et al.* (2019); Parida *et al.* (2021).

Locules per Fruit. The estimates of the individual simple scaling tests as well as χ^2 test value of joint scaling test were significant for all the families under the study. Both additive and dominance components of gene action were found significant for the family I, III and IV. While, additive gene action found significant in family V and dominance gene action in family VI. All three types of gene interaction *viz.* additive \times additive [i], additive \times dominance [j] and dominance \times

dominance [l] were found significant I, III, IV and VI. Significance of additive \times additive [i] gene interaction was found significant in family II and additive \times additive [i] and additive \times dominance [j] in family V. Duplicate type of epistasis reported in families I, II, IV and VI; while complimentary epistasis recorded for families III and IV. Higher magnitude of dominance \times dominance [l] gene interaction recorded for families I, III and IV indicating role of non-additive effects for inheritance of this trait. Due to presence of non-allelic interaction selection should be postponed to later generations.

Lycopene Content. The individual estimates of simple scaling tests as well as χ^2 test value of the joint scaling test indicated the non-adequacy of additive-dominance model due to significance of individual scaling tests for all six families. Significant estimates of additive gene effect were reported in family I and III; while significant dominance gene effect was reported in family III only. Significant additive \times dominance [j] and dominance \times dominance [l] type of epistasis interaction were reported in family I, II, V and VI. Whereas, significant dominance \times dominance [l] and additive \times dominance [j] epistatic interactions were found in family III and IV, respectively. Duplicate epistasis reported in the family I, III, IV and VI; whereas complimentary epistasis reported in the family II and V. Higher magnitude of dominance gene effect with additive \times additive [i] type of gene interaction reveals the role of both fixable and non-fixable gene effects. Higher magnitude of dominance \times dominance [l] gene interaction was profound in family II and V. Similar findings were reported by Kumar and Srivastava (2017).

Total Soluble Solids. Additive and dominance model was inadequate for all the six families, as significant values of individual scaling tests were reported for seeds per fruit. In addition of this, significant estimates of χ^2 test of joint scaling test indicated presence of non-allelic interactions. Significant values of dominance gene effect as well as dominance \times dominance [l], additive \times additive [i] and additive \times dominance [j] gene interaction effect was reported in family I. All six parameters in family II found significant, revealing the predominant role of dominance, additive genetic effects and additive \times dominance [j], dominance \times dominance [l] and additive \times additive [i] type of epistasis gene interaction in the expression of the trait under study. Significant additive and dominant gene effects along with significant additive \times additive [i] and dominance \times dominance [l] interactions was reported in family III and V with higher magnitude of dominance \times dominance [l] interaction. However, additive gene effect coupled with additive \times dominance [j] and dominance \times dominance [l] interaction found to govern the character in family IV. Duplicate type of gene interaction in five out of six families. The similar findings were reported by Kumar and Srivastava (2017).

Moisture Content. Additive dominance was found inadequate to explain since the values of individual scaling tests were found significant for all families

along with significant estimates of χ^2 test value. Significant estimates of additive gene effect were reported for family I and II; whereas significant dominance effect was reported in family III. Significant additive effect along with significant additive \times dominance [j] interaction epistasis reported in family I. In family II, significant additive gene effect along with all three types of gene interaction *viz.* additive \times additive [i], additive \times dominance [j] and dominance \times dominance [l] found significant with higher magnitude of additive \times additive [i] type of interaction revealing. Significant dominance effect coupled with significant additive \times additive [i] and dominance \times dominance [l] type of gene interaction reported in family III. Whereas, only significant dominance \times dominance [l] and additive \times additive [i] gene interactions were reported in family IV and V, respectively. Only mean value was found significant in family VI and other parameters were found non-significant due to difference in comparable environmental or difference in fertility and viability. Abundance of inter genic interactions suggests postponement of the selection for the character under study.

1000 Seed Weight. Significance estimates of individual simple scaling tests revealed inadequacy of additive dominance model to explain inheritance of 1000 seed weight. The results of simple scaling test were confirmed by significance of χ^2 test value of joint scaling test except, family VI. Even though, scale C found significant in family VI, non-significant χ^2 test was reported. This lack of congruence may be due to differential fertility and viability of individuals of different segregating generations or may be due to sampling error. Significant estimates of additive gene effect were reported in all six families under study. Whereas, significant estimates of dominance gene effects were reported for the family IV and VI. In family I, only additive \times dominance [j] gene interaction found significant. All three epistatic interactions *viz.* additive \times additive [i], additive \times dominance [j] and dominance \times dominance [l] along with significant additive gene effect reported in family II and V. In family III, additive \times dominance [j] and dominance \times dominance [l] epistasis interaction were found significant. In family IV, significant additive \times additive [i] and additive \times dominance [j] were reported along with significant principle gene effects. From the sign of [h] and [l], duplicate epistasis depicted to govern the character in all the families except, family III. The findings were partially in accordance with Damor *et al.* (2021) as they also recorded importance inter allelic interactions.

Seed to Pulp Ratio. Additive-dominance model found inadequate to explain the inheritance of this novel character since, estimates of simple scaling test were found significant for all the families except, family II and also confirmed by significance of χ^2 test for these families. Adequacy of additive-dominance model of family II confirmed by non-significant value of χ^2 test of joint scaling test for this family. Family I exhibited significant value of additive \times dominance [j] type of inter allelic interaction. Significant and higher value of

dominance component of gene action along with significant additive × additive [i] and additive × dominance [j] gene interaction was reported in family III. While, all six parameters of generation mean were found significant for family IV. Significant principle gene effect components viz. additive and dominance along with significant additive × dominance [j] and dominance × dominance [l] epistatis gene interaction

and additive × additive [i], additive × dominance [j] and dominance × dominance [l] reported for families V and VI, respectively. Four out of six families possessed opposite signs of [h] and [l] components indicating prevalence of duplicate gene interaction; whereas complimentary gene interaction reported for the remaining two families.

Table 1: Analysis of variance of generation means in six families of tomato for characters under study.

Source	Degree of freedom	Mean sum of square							
		Days to flowering	Branches per plant	Plant height	Fruit Length	Fruit girth	Average fruit weight	Pericarp thickness	Fruit yield per plant
Analysis of variance between families									
Replication	2	2.12	0.01	45.04	0.10	0.01	1.019	0.003	0.37
Families	5	17.77**	1.04**	2288.62**	0.21*	1.072**	32.29**	0.681*	3.33**
Error	10	0.65	0.13	16.05	0.03	0.15	3.65	0.009	0.59

Source	Degree of freedom	Mean sum of square					
		Locules per fruit	Lycopene content	Total soluble solids (TSS)	Moisture content	1000 seed weight	Seed to pulp ratio
Analysis of variance between families							
Replication	2	0.069	0.000*	0.032*	1.95**	0.123**	0.001*
Families	5	1.21**	0.015**	0.958**	5.13**	0.275**	0.027**
Error	10	0.018	0.000	0.003	0.32	0.005	0.000

Table 2: Estimates of simple scaling test and gene effects for characters under study.

Family	Scaling Test				χ^2 at 3 d.f.	Gene effect									Gene action
						Three parameter Model			Six Parameter Model						
	A	B	C	D	m	d	h	m	d	h	i	j	l		
Days to flowering															
I	-9.20**	-6.73**	23.46**	-3.76**	163.38**	-	-	-	45.01**	-4.00**	14.00**	7.53	-1.23	8.39**	C
II	10.53**	-2.00	-3.20	4.67*	45.63**	-	-	-	43.00**	-9.00**	-17.20**	-9.33*	-4.26*	21.86**	D
III	17.40**	-8.86**	15.40**	3.43*	127.37**	-	-	-	43.36**	11.83**	-15.63**	-6.86*	13.13**	-1.66	C
IV	19.60**	10.93**	20.40**	-5.07**	181.56**	-	-	-	46.11**	-1.63	-1.10	10.13**	4.33**	40.67**	C
V	9.33**	23.33**	14.53**	14.27**	319.17**	-	-	-	45.46**	10.67**	-34.73**	28.53**	16.33**	42.53**	D
VI	17.00**	-0.07	-5.00	6.03**	105.06**	-	-	-	46.00**	-6.96**	-2.96	12.06**	-8.46**	29.13**	D
Branches per plant															
I	-4.13**	-3.66**	-0.86	3.46**	187.09**	-	-	-	13.18**	0.83	-4.26**	-6.93**	-0.23	14.73**	D
II	-5.00**	-0.60	-6.46**	-0.43	36.56**	-	-	-	12.55**	0.53	2.66	0.86	-2.20**	4.73**	C
III	-4.20*	-5.33**	-7.13**	1.20	22.97**	-	-	-	12.85**	1.76*	0.59	-2.40	0.56	11.93**	C
IV	0.53	-3.33**	-2.66	0.06	13.82**	-	-	-	12.66**	1.53**	1.73	-0.13	1.93**	2.93	C
V	-0.53	-0.46	-4.46**	-1.73**	13.42**	-	-	-	11.68**	0.30	5.19**	3.46*	-0.03	2.46	C
VI	-1.40	-5.26**	-7.26**	-0.30	61.53**	-	-	-	13.05	0.39	2.33	0.59**	1.93	6.06**	C
Plant height															
I	-6.14	23.46**	-59.55	-38.43*	8.54*	-	-	-	172.38**	27.79**	153.71**	76.87**	-14.80	-94.19	D
III	-28.99*	41.44**	79.97**	-4.76	16.94**	-	-	-	147.74**	1.91*	33.06	9.53**	6.22	60.91	C
IV	57.13**	-40.10*	80.25**	8.49	16.55**	-	-	-	151.67**	-1.23	7.57	-16.98	-8.51	114.22*	C
V	50.34**	-20.37	77.48**	-3.41	10.53*	-	-	-	100.28**	-8.03	35.23	6.83	-15.01	63.81	C
VI	42.91**	45.27**	79.16**	4.51	31.72**	-	-	-	111.88**	4.76	14.77	-9.02	1.18	97.21**	C
Fruit length															
I	-1.38	-0.18	3.22*	2.39**	11.22*	-	-	-	9.60**	-1.27*	-2.10	-4.78**	-6.03	6.35*	D
II	3.64**	-0.98	-0.34	-1.50	12.18*	-	-	-	8.50**	2.02**	3.78	3.00	2.31**	-5.67	D
III	-6.14**	-4.26**	1.36	5.88**	159.52**	-	-	-	9.37**	1.52**	10.08**	11.76**	0.93**	22.16**	D
IV	3.29**	-4.13**	8.48**	4.6**	334.75**	-	-	-	10.54**	2.07**	-5.76**	-9.31**	3.71**	10.15**	D
V	6.50**	-0.56	9.20**	1.63**	244.92**	-	-	-	9.77**	2.68**	-1.84	-3.26**	3.53**	-2.68	C
VI	8.06**	3.50**	12.25**	0.34	280.80**	-	-	-	9.86**	1.05*	0.78	-0.69	2.27**	10.86**	C
Fruit girth															
I	0.23	-0.85	8.40**	4.51**	49.07**	-	-	-	17.71**	-0.90	-6.31**	-9.02**	0.54	9.65**	D
II	-0.72	-5.28**	-7.18**	-0.59	20.69**	-	-	-	14.09**	2.13*	1.97	1.17	2.28*	4.82	C
III	10.42**	-1.52	1.91	6.93**	245.24**	-	-	-	16.42**	4.70**	14.33**	13.86**	4.44**	25.81**	D
IV	2.76*	-4.04**	-5.83**	-2.27	21.71**	-	-	-	13.06**	1.11	7.65**	4.54	3.40**	-3.25	D

V	5.67**	-7.83**	3.14	2.65*	92.40**	-	-	-	16.05**	5.53**	-2.73	-5.30**	6.75**	7.46**	D
VI	3.83**	-1.02	4.16	0.67	21.05**	-	-	-	16.36**	0.22	2.50	-1.35	2.43**	-1.45	D
Average fruit weight															
I	-4.41	-3.16	9.66*	8.62**	19.09**	-	-	-	38.37**	-2.94*	-1.38**	-1.72**	-6.22	24.82**	D
II	12.29**	-4.27	9.92	0.95	12.83*	-	-	-	44.61**	4.62	-1.09	-1.90	8.28**	-6.10	C
III	18.00**	3.84	24.68**	1.41	56.12**	-	-	-	45.55**	3.25	8.42	-2.83	7.07**	-19.00*	D
IV	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
IV	21.09**	28.91**	24.77**	12.61**	46.69**	-	-	-	43.33**	-2.85	-6.55	25.23**	3.91	75.24**	D
V	-9.94*	-	-	6.00	41.55**	-	-	-	45.22**	-0.24	12.03	-	7.50	46.90**	C
VI	-	-	-2.19	12.04**	32.91**	-	-	-	43.41**	-3.53*	-	-	-1.65	50.35**	D
Pericarp thickness															
I	-1.04*	0.46	-0.45	0.06	9.72*	-	-	-	5.28**	0.20	0.30	-0.12	-	0.70	C
II	1.94**	0.23	1.65*	-0.26	27.40**	-	-	-	5.58**	0.37	0.11	0.53	0.85**	-2.71**	D
III	0.49	1.35**	1.19*	-0.32	18.36**	-	-	-	5.37**	-0.22	0.87	0.64	-0.42	-0.24*	D
IV	1.92**	0.46	-0.44	-	24.28**	-	-	-	3.86**	-0.44	3.07**	2.82**	0.73*	-5.21**	D
V	-	-	-	0.82**	51.00**	-	-	-	5.02**	0.74**	-1.73*	-1.64*	1.04**	5.79**	D
VI	1.79**	-	-	0.94**	31.99**	-	-	-	5.11**	-0.60	3.27**	1.88*	1.70**	-2.07	D
Fruit yield per plant															
I	3.61	-0.09	9.00**	2.74	10.20*	-	-	-	9.78**	1.34	-3.03	-5.48	1.85	1.96	D
II	5.39**	-0.09	5.22	-0.03	9.12*	-	-	-	9.49**	1.63	1.67	0.07	2.74**	-5.37	D
III	4.45**	3.70**	5.52*	-1.31	14.32**	-	-	-	8.43**	0.93	6.46**	2.63	0.37	-	D
IV	-3.00	2.61	-	-	23.84**	-	-	-	6.77**	-	11.59**	7.14**	-	-6.75	D
V	-1.63	-	9.14**	9.67**	92.04**	-	-	-	13.89**	2.93**	-	-	3.44**	29.51**	D
VI	6.19**	3.56*	6.21*	-1.76	14.13**	-	-	-	8.81**	3.16**	6.22*	3.53	1.31	-	D
Locules per fruit															
I	-0.86*	-	2.33**	2.80	115.08**	-	-	-	4.13**	1.00**	-4.56**	-5.59**	0.76**	8.86**	D
II	0.60	-0.33	1.73**	0.73*	11.51*	-	-	-	4.20**	0.33	-0.06	-1.46*	0.46	1.19	D
III	-	2.66**	1.46**	1.46**	98.61**	-	-	-	3.71**	-	3.96**	2.93**	1.93**	4.40**	C
IV	0.86*	-	4.80**	2.73**	86.74**	-	-	-	6.33**	0.53*	-3.20**	-5.46**	1.20**	6.13**	D
V	-0.46	1.26**	2.66**	0.93**	27.91**	-	-	-	4.90**	-	0.59	-1.86**	-	1.06	C
VI	1.13**	2.26**	1.00	-	31.41**	-	-	-	4.68**	-0.23	3.80**	2.40**	-0.56*	-5.80**	D
Lycopene content															
I	0.22**	0.55**	0.75**	-0.01	375.00**	-	-	-	0.45**	-	0.05	0.02	-	-0.80**	D
II	-	0.02	-0.13*	0.07	62.22**	-	-	-	0.53**	0.01	0.14	-0.14	-	0.43**	C
III	0.35**	0.34**	0.27**	-	88.19**	-	-	-	0.36**	0.08**	0.54**	0.42**	-0.00	-1.11**	D
IV	0.17**	-	0.00	-0.02	37.70**	-	-	-	0.36**	0.02	0.13	0.05	0.14**	-0.11	D
V	-	-	-	0.13*	63.65**	-	-	-	0.47**	0.01	0.06	-0.27	-0.11*	0.88**	C
VI	-0.01	0.21**	-0.20*	0.01	28.14**	-	-	-	0.43**	-0.02	-0.03	-0.02	0.09**	0.25*	D
Total soluble solids															
I	-	0.13	0.43	0.90**	52.83**	-	-	-	5.08**	0.10	-	-	-	3.19**	D
II	2.07**	0.76**	1.52**	-	111.80**	-	-	-	4.22**	1.04**	1.21**	1.31**	0.65**	-4.15**	D
III	-	-	0.50	1.47**	63.73**	-	-	-	4.43**	0.34*	-	-	-	5.39**	D
IV	0.43*	-	-	0.36	131.19**	-	-	-	3.43**	0.78**	-0.03	-0.73	1.45**	2.76**	D
V	-	-	-	0.76**	250.01**	-	-	-	4.06**	0.65**	0.12	-	-	7.31**	C
VI	2.10**	2.23**	1.13**	1.60**	98.82**	-	-	-	3.52**	-	-	-	0.06	7.54**	D
Moisture content															
I	-	0.17	-5.34	0.96	15.27**	-	-	-	90.52**	-	-4.52	-1.93	-	9.22	D
III	3.13	3.35	-2.45	-	16.99**	-	-	-	91.11**	-0.75	7.84**	8.94**	-0.10	-	D
IV	-	-	-	2.53	20.61**	-	-	-	91.57**	0.49	-4.55	-5.06	0.24	17.47**	D
V	-	-	-	0.30	5.87	-	-	-	91.26**	-0.14	-3.26	-0.61	-1.12	5.80	D
VI	-	-	-	-0.15	15.27**	-	-	-	91.11**	-0.44	-0.58	0.30	-1.03	6.62	D

Contd.

Family	Scaling Test				χ^2 at 3 d.f.	Gene effect										Gene action
	A	B	C	D		Three parameter Model			Six Parameter Model							
						M	d	h	m	d	h	i	j	l		
1000 seed weight																
I	-0.58	2.14*	1.46*	-0.04	50.74**	-	-	-	3.72*	0.61*	0.45	0.09	-	1.36*	-1.64	D
II	1.09*	0.31	1.35	1.06	54.58**	-	-	-	3.73*	0.79*	-1.08	-	2.13*	0.70*	2.92*	D
III	0.33	3.96*	2.99*	0.31	143.11*	-	-	-	3.56*	1.20*	0.16	-0.63	2.14*	4.26*	C	
IV	-0.54	0.92	3.30*	1.46*	20.39**	-	-	-	4.85*	1.58*	-3.01*	-2.92*	-0.73*	2.54	D	
V	1.23*	0.75	1.86	1.17*	23.02**	-	-	-	4.05*	-0.59*	-1.83	-2.35*	0.99*	2.83*	D	
VI	-0.62	-0.07	-1.61*	-0.45	5.01	-	-	-	3.53*	-0.60*	0.22*	0.91	-0.27	-0.20	D	
Pulp to seed ratio																
I	0.17*	0.03	0.12	-0.04	10.17	-	-	-	0.42*	0.03	0.17	0.08	0.06*	-0.03	C	
II	-0.07	-0.01	-0.01	0.03	3.59	0.67*	0.13*	0.25	0.68*	0.11*	-0.05	-0.07	-0.03	0.16	D	
III	0.21*	0.15*	0.41*	0.17*	60.45**	-	-	-	0.62*	-0.00	0.29*	0.35*	0.18*	0.29	D	
IV	0.71*	0.21*	0.11	0.30*	274.90*	-	-	-	0.59*	0.28*	0.49*	0.60*	0.46*	1.10*	D	
V	0.07	0.44*	0.28*	0.04	87.05**	-	-	-	0.60*	0.31*	0.11	-0.08	0.25*	0.45*	C	
VI	0.08	0.47*	-0.16	0.11*	94.37**	-	-	-	0.54*	0.24*	-0.03	-0.22*	0.27*	0.61*	D	

Note: * indicates non-significant values for scaling tests and/or ANOVA for particular character of particular family, C-Complementary and D-Duplicate gene action.

CONCLUSIONS

The present investigation revealed that main effects viz., additive and dominance along with additive × additive (i), additive × dominance (j) and dominance × dominance (l) were present at more or less extent indicating the importance of these interactions for the inheritance of the various traits. The result of epistatic gene effects for fruit yield and its related traits in different cross combinations revealed that recurrent selection and bi-parental mating between desirable segregants followed by selection would be profitable for development of desirable hybrids/lines/varieties.

FUTURE SCOPE

The segregating material can be advanced in order to produce advanced lines with superior traits. They can be employed further in breeding programmes.

Conflict of Interest. None.

REFERENCES

Anonymous (2020). FAOSTAT data retrieved from <https://www.fao.org/faostat/en>.

Cavalli, L. L. (1952). An analysis of linkage in quantitative inheritance. *An analysis of linkage in quantitative inheritance*.

Chauhan, V. B. S., Kumar, R., Behera, T. K., Yadav, R. K. and Verma, A. K. (2019). Inheritance of fruit weight and mode of gene action for yield contributing traits in tomato. *Research Journal of Biotechnology*, 14(4).

Damor, H. I., Acharya, R. R. and Patel, A. A. (2021). Genetic Analysis for Fruit Yield and its Component Traits in

Tomato (*Solanum lycopersicum* L.) Population. *Journal of Plant Development Sciences*, 13(1), 1-9.

Das, I., Hazra, P., Longjam, M., Bhattacharjee, T., Maurya, P. K., Banerjee, S. and Chattopadhyay, A. (2020). Genetic control of reproductive and fruit quality traits in crosses involving cultivars and induced mutants of tomato (*Solanum lycopersicum* L.). *Journal of Genetics*, 99(56).

Hayman, B. I. and Mather, K. (1955). The description of genic interactions in continuous variation. *Biometrics*, 11(1), 69-82.

Hayman, B. I. (1958). The separation of epistasis is from additive and dominance variation in generation mean. *Heredity*, 12, 371-390.

Jinks, J. L. and Jones, R. M. (1958). Estimation of the components of heterosis. *Genetics*, 43, 233-34.

Kumar, R. and Srivastava, K. (2021). Gene Effects and Heritability for Yield Traits in Tomato (*Solanum lycopersicum* L.). *Bangladesh Journal of Botany*, 50(3), 453-465.

Lecomte, L., Duffé, P., Buret, M., Servin, B., Hospital, F. and Causse, M. (2004). Marker-assisted introgression of five QTLs controlling fruit quality traits into three tomato lines revealed interactions between QTLs and genetic backgrounds. *TAG. Theoretical and applied genetics. Theoretische und angewandte Genetik*, 109(3), 658-668.

Mather, J. and Jinks, J. L. (1982). *Biometrical Genetics, The study of continuous variation*. Chapman and Hall, London, 2nd Ed.

Negi, P. K., Sharma, R. R., Kumar, R., & Chauhan, V. B. S. (2013). Genetic analysis for yield and its contributing traits in tomato under low temperature regime. *Vegetable Science*, 40(2), 189-194.

- Parida, A. P., Srivastava, A., Mathur, S., Sharma, A. K. and Kumar, R. (2021). Identification, evolutionary profiling, and expression analysis of F-box superfamily genes under phosphate deficiency in tomato. *Plant Physiology and Biochemistry*, 162, 349-62.
- Patel, U. J., Kathiria, K. B., Patel, J. S. and Saiyad, I. M. (2010). Heterobeltiosis and inbreeding depression in tomato (*Lycopersicon esculentum* Mill.). *International Journal of Plant Sciences*, 5(2), 636-638.
- Sikder, S., Biswas, P., Hazra, P., Akhtar, S., Chattopadhyay, A., Badigannavar, A. M. and D'Souza, S. F. (2013). Induction of mutation in tomato (*Solanum lycopersicum* L.) by gamma irradiation and EMS. *Indian Journal of Genetics and Plant Breeding*, 73(4), 392-399.

How to cite this article: Pragati J. Prajapati, J.N. Patel, Parthik Patel, and N.A. Patel (2023). Generation Mean Analysis for Fruit Yield and Component Traits in Tomato (*Solanum lycopersicum* L.). *Biological Forum – An International Journal*, 15(5): 230-237.