

Genetic Variability and Traits Association Analysis of Tomato (*Lycopersicon esculentum* L.) Genotypes for Yield and Quality Attributes

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Abstract The experiment was conducted at the experimental farm of the department of Genetics and Plant Breeding, Bangladesh agricultural university, Mymensingh, Bangladesh, with 30 different tomato (*Lycopersicon esculentum* L.) genotypes to study the genetic variability and characters association for yield and yield-contributing traits along with quality traits. The experiment was laid out in a Randomized Complete Block Design with three replications. Analysis of variances showed high degree of variation existed among the genotypes of the studied traits. Yield contributing traits showed higher phenotypic co-efficient of variation as compared to their corresponding genotypic co-efficient of variation indicating considerable environmental influences on them. Individual fruit weight showed high heritability (99.71) with high genetic advance (85.4) followed by fruits per plant (99.65 & 81.26) suggested a good scope for the improvement of these characters through selection. High heritability but low genetic advance showed in number of primary branches (75.87 & 3.67), fruit cluster per plant (99.49 & 6.12), fruit diameter (98.83 & 5.59), total soluble solid (80.51 & 1.85) and ascorbic acid content (90.75 & 9.52) indicated limited scope for the improvement of these traits through selection. Correlation coefficient analysis showed that yield per plant had significant positive correlations with individual fruit weight (0.609***) and number of fruit cluster per plants (0.396*) whereas significant negative correlation was observed with number of primary branches per plant (-0.421*). Path

co-efficient analysis revealed that the maximum positive direct contribution towards yield was through individual fruit weight (0.704) whereas plant height (-0.038) showed negative direct contribution towards yield. Individual fruit weight and number of fruit cluster per plant could be crucial for improving tomato yield.

Keywords Tomato, Genetic Diversity, Heritability, Genetic Advance, Correlation co-efficient, Path co-efficient, Yield, Quality

1. Introduction

The cultivated tomato (*Lycopersicon esculentum* L.) occupies the prime position among different vegetables and grown in a variety of climatic condition all over the world. It belongs to the family Solanaceae with chromosome number of 24 and is a self-pollinated, annual crop [1]. The position of tomato in world vegetable production is second only after potato [2]. It is a rich source of vitamin A, C (ascorbic acid, AsA), glutathione (GSH) and mineral nutrients with preponderance of Ca, P and Fe [3]. Tomato fruit and its products are the main source of lycopene and other antioxidants in the human diet [4]. The level of lycopene in tomato fruit increases 500 times during ripening [5]. High antioxidant capacity in both fresh and processed tomatoes associated with the higher capacity to eliminate reactive oxygen species (ROS) and helps in lowering the incidence of

certain forms of human cancer [6]. In many countries it is considered as “poor man’s orange” because of its improved nutritional values [7]. Recent epidemiological studies have shown that their consumption helps to prevent cardiovascular disease [8,9] and some types of cancers, such as prostate cancer [10,11].

Tomato like other crops is the final product attributed by a complex chain of interrelating effects of different characters [12,13]. The degree of association between characters as indicated by the correlation coefficients has always been a helpful instrument for the selection of desirable characters in a breeding program. Correlation coefficient analysis measures the mutual relationship between various plant characters and determines the component characters on which selection can be based for yield improvement. Correlation coefficient is not adequate to anticipate traits interrelationship leading to yield. Under such circumstances, path coefficient analysis is an additional informative tool [12,14]. Path analysis splits the correlation coefficients into direct and indirect effects of a set of dependent variables on the independent variables which help in selecting elite genotypes. The first and foremost prerequisite for crop improvement is the availability of wide genetic variability in the form of germplasm [15]. The influence of environment in expression of characters and the extent to which improvement is possible after selection can be determined by heritability and genetic advance [16]. Heritable variation can be effectively studied in conjunction with genetic advance. High heritability alone is not enough to make efficient selection in segregating generation and needs to be accompanied by a substantial amount of genetic advance [17]. Amounts of variability present in germplasm can be detected by phenotypic and genotypic coefficients of variation.

In Bangladesh there are nearly 30 released tomato varieties and 5 tomato hybrids developed by Bangladesh Agricultural Research Institute (BARI) and Bangladesh Institute of Nuclear Agriculture (BINA) and other private seed companies. However, information on the quality attributes like antioxidant compounds in the released varieties is unexpectedly rare. Additionally, in Bangladesh majority of the Asian Vegetable Research and Development Center (AVRDC) germplasm has not been characterized for their yield and antioxidant properties. Importantly, we need to characterize them under the climatic conditions of Bangladesh for determining potential genotypes that can be used for future plant breeding program to improve yield and quality. The present investigation was therefore undertaken to study the genetic variability and character association among 30 diverse tomato genotypes for yield and quality attributes.

2. Materials and Methods

The experiment was conducted during the period from November 2014 to April 2015 in the experimental farm of

the Department of Genetics and Plant Breeding, Bangladesh Agricultural University, Mymensingh. The experiment was laid out in RCBD with three replications. This study involved 14 exotic (V1007282, V1006136, V1006422, V1006186, V1006484, V1045786, V1063607, WP7, WP8, WP10, C11, C41, C51, C71), 2 indigenous advance line (Phili-1, Phili-2) and 14 varieties (BARI tomato-9, BARI tomato-11, BARI tomato-14, BARI tomato-15, BARI tomato-8, BINA tomato-2, BINA tomato-7, BINA tomato-3, BINA tomato-4, BINA tomato-5, BINA tomato-8, BINA tomato-9, BINA tomato-10) of tomato. The exotic tomato lines were collected from AVRDC, Taiwan. The indigenous tomato lines and variety were collected from Bangladesh Agricultural Research Institute (BARI), Gazipur and Bangladesh Institute of Nuclear Agriculture (BINA), Mymensingh. Data on 14 different characters i.e. i) plant height (cm), ii) number of primary branches per plant, iii) days to first flowering, iv) days to 50% flowering, v) number of fruit cluster per plant (no.), vi) days to first harvest, vii) individual fruit weight (g), viii) fruit diameter (cm), ix) fruits per plant (no.), x) Days to last harvest, xi) Yield per plant (kg), xii) dry matter content (%), xiii) determination of total soluble solid (TSS) of tomato (%), xiv) ascorbic acid (AsA) content (mg/100g) were recorded for the study. Data were collected from three replications for each genotype and five randomly selected plants from each replication. Mean values for each character were assembled and used for statistical analysis. AsA content was determined according to the method described by Plummer [18].

Analysis of variance was performed using the statistical software SAS. Genotypic and phenotypic variances were estimated according to the formula given by Johnson *et al.* [17]. Heritability in broad sense (h^2_b) was estimated according to the formula suggested by Johnson *et al.* [17] and Hanson [19]. Genotypic and phenotypic co-efficient of variations were estimated according to Burton [20] and Singh *et al.* [21]. Genetic advance was calculated following formula given by Johnson *et al.* [17] and Allard [22]. Genetic advance in percent of mean was calculated by the formula of Comstock *et al.* [23]. The phenotypic correlations were estimated by the formula suggested by Miller *et al.* [24]. Correlation coefficient were further partitioned into components of direct and indirect effects by path coefficient analysis originally developed by Wright [25] and later described by Dewey and Lu [26].

3. Result and Discussion

The analysis of variance of different tomato genotypes for yield and yield contributing traits are shown in Table 1. Results showed a significant ($p < 0.001$) variation among the genotypes for different traits indicating presence of significant variability in the materials which can be exploited through selection. Importantly, the success of breeding

program depends upon quantum of variability present in the available germplasm. Significant variability among the genotypes for various characters except number of primary branches per plant was also reported [27]. The highest plant height (149.17 cm) was observed in BARI Tomato-11 which was statistically significant and different from all other genotypes. The lowest value (58.13 cm) was recorded in V1006422. The genotype V1006484 bears maximum number of primary branches (13.6) and the genotype Binatomato-10 bears minimum number of primary branches (5.93). The average numbers of primary branches were (9.42). C11 took maximum days (75.6) to flower and Binatomato-5 required the minimum number of days to first flowering (57.8). The average value of days to first flowering was (68). The genotype Binatomato-5 took minimum days (62) to 50 % flowering and the genotype WP8 required the maximum number of days (88) to 50% flowering. Binatomato-10 bears maximum number (17.57) of fruit cluster and BARI Tomato-15 bears minimum number (3.54) of fruit cluster. The genotype Binatomato-3 was early maturing which took (90.33) days for first fruit maturity. The genotype Binatomato-8 required highest number of days (128.67) for first fruit maturity. The genotype Bahar gave the highest mean value of individual fruit weight (217.91g) which was significantly superior to all other genotypes. The lowest fruit weight (8.09g) was observed in V1045786. The genotype Bahar had highest (10.50 cm) fruit diameter on the other hand Binatomato-10 had lowest (1.78 cm) fruit diameter. The highest number of fruits (177.67) recorded in BARI Tomato-11 and the lowest number of fruits (13.67) was recorded in V1006136. Fruit harvesting duration was maximum (154.67) days for the genotype V1006484. Minimum harvesting duration (131.00) days were recorded for Binatomato-8. The genotype Bahar had highest (17.89%) dry matter percentage. The lowest dry matter content was recorded for the genotype C71. Binatomato-10 had highest (7.10%) TSS and the lowest (2.53%) TSS content was recorded for V1045786. Highest AsA content was recorded in C71 (38.87mg %). The genotype WP8 contain lowest amount (16.13mg %) of AsA among the selected genotypes. The highest yield (6.43) kg per plant was obtained from

Binatomato-9 and the lowest yielding (0.37 kg per plant) genotype was V1006136 (Table 2).

3.1. Genetic Parameter Analysis

A clear idea can be gained by comparing the relative amount of phenotypic coefficient of variation (PCV) and genotypic coefficient of variation (GCV) for the actual strength of variability. The PCV was higher than the GCV for plant height (PCV=543.48, GCV=521.24); number of primary branches per plant (PCV=58.62, GCV=44.48); days to first flowering (PCV=87.99, GCV=77.04); days to 50% flowering (PCV=100.80, GCV=88.93); days to first harvest (PCV=96.22, GCV=93.62); individual fruit weight (PCV=2425.67, GCV=2418.64); fruits per plant (PCV=3566.48, GCV=3554.12) and dry matter percentage (PCV=62.34, GCV=61.60) indicated the influence of environment for the expression of these characters (Table 3). Highest GCV (15.09%) and PCV (22.19%) for plant height in a study of 12 tomato genotypes received from AVRDC, Taiwan grown at BARI farm, Joydebpur [12]. High genotypic variance indicating more contribution of genetic component for the total variation. Mohamed *et al.* [28] reported GCV and PCV (16.41%) and (17.91%) respectively for number of primary branches for plant. Similar results were also found by Biswas and Mallik [29] and Kaushik *et al.* [30] for days to first flowering. Gadekar *et al.* [31] also reported maximum phenotypic (381.74) and genotypic (323.14) variance with highest GCV (42.18%) for individual fruit weight in tomato. This indicated the possibility of obtaining higher selection response in respect of these traits. Bahar had highest and C71 had the lowest dry matter percentage. Dutta *et al.* [32] studied six tomato varieties including two advanced lines Bahar and E-6 and found dry matter content 4.85%, 4.70% and 4.50% for Bahar, Oxheart and Ratan, respectively, which is similar to the present findings. Prodan [33] found early and tall varieties had higher dry matter content than that of later and dwarf varieties which support the present results.

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Table 1. Analysis of variance (mean square) for different characters of 30 genotypes of tomato

S.V	DF	PH	PB	DFD	50% F	FC	FH	IFW	FD	FP	LH	DM	TSS	AsA	YP
Rep.	2	10.72	1.88	0.20	3.61	0.14	5.41	10.29	0.11	11.20	0.68	0.08	0.66	6.29	0.15
Trt.	29	1482.17***	13.91***	101.66***	142.18***	26.64***	257.32***	5175.41***	9.58***	4689.80***	119.40***	22.42***	3.25***	72.96***	5.12***
Error	58	20.79	1.33	4.60	6.06	0.05	2.35	5.01	0.05	5.43	0.23	0.09	0.24	2.40	0.09

*** indicates significant at $p < 0.001$

Here, Trt.= treatment, PH= Plant height; PB= Number of primary branches per plant; DFF= Days to first flowering; 50%F= Days to 50% flowering; FC= Number of fruit cluster per plant; FH= Days to first harvest; IFW= Individual fruit weight; FD= Fruit diameter; FP= Fruits per plant; YP= Yield per plant; LH= Days to last harvest; DM= Dry matter percentage; TSS= Total soluble solid; AsA = Ascorbic acid content.

Table 2. Average performance for 14 different traits (morphological and biochemical) of 30 genotypes of tomato

Ge.	PH (cm)	PB (no.)	DFD (days)	50%F (days)	FC (no.)	FH (days)	IFW (g)	FD (cm)	FP (no.)	YP (kg)	LH (days)	DM (%)	TSS (%)	AsA (mg/100g)
1	74mn	11.07bcd	61gh	71jkl	5.50kl	110jk	83.96g	7.08de	29.67jkl	2.42gh	136o	10.95lm	3.63hij	24.93efg
2	92.8hij	12.20abc	65f	82bcde	4.60m	119gh	32.49r	5.06kl	13.67r	0.37p	152c	8.00s	4.57cdefg	16.87lm
3	58.13q	13.07a	60hi	71jkl	7.60f	120g	45.27q	6.76ef	15.00qr	0.73nop	154b	12.26hi	4.53cdefg	22.73ghi
4	71.53no	13.00a	69de	81cdef	6.74ghi	120fg	24.29s	6.93de	20.67op	0.61op	150e	9.58qr	3.67hij	20.53ijk
5	124.07c	13.60a	61ghi	68l	6.50hij	122def	45.31qp	7.05de	18.33opq	0.89no	155a	14.31e	3.50hijk	21.27hij
6	69.33nop	12.27abc	58hi	70kl	5.60k	108k	8.09t	2.82q	127.00c	0.98nom	151d	14.94d	2.53l	27.13cde
7	65.87op	7.33ghi	72bcd	84abc	4.60m	124cd	53.39o	4.86lm	17.00pqr	0.92nom	141k	8.43s	4.70cd	25.67def
8	112.8de	9.13efg	67ef	74ijk	6.23j	123cd	58.54n	6.38gh	35.67h	1.76jk	138n	11.29kl	5.57b	22.00hi
9	91.93hijk	8.40fgh	74ab	88a	6.87g	122def	75.88ij	4.64mn	29.33jkl	2.12hij	150e	8.27s	3.03ijkl	16.13m
10	84.53kl	12.93ab	59hi	68l	7.33f	111jk	104.52d	6.54fg	34.33hi	3.10ef	140kl	10.32nop	3.77ghi	23.47fgh
11	72.6no	8.60efg	76a	83bcde	5.24l	119gh	69.73kl	4.45no	30.67ij	1.74jk	139lm	9.37r	3.63hij	29.33b
12	80.4lm	8.13fgh	72bcd	85ab	8.43e	123de	43.21q	4.83lm	30.33jk	1.16lmn	144hi	9.99opq	2.93jkl	24.93efg
13	88.67ijk	11.87abc	61ghi	71jkl	9.87c	117hi	77.58ih	7.14d	43.33fg	3.25ef	142j	10.80mn	4.63cdef	27.87bcd
14	132.8b	9.53def	72bcd	83bcde	6.47hij	119ghi	66.36ml	4.57mn	30.67ij	1.87ijk	152c	7.96s	5.20bc	38.87a

Ge.	PH (cm)	PB (no.)	DFF (days)	50%F (days)	FC (no.)	FH (days)	IFW (g)	FD (cm)	FP (no.)	YP (kg)	LH (days)	DM (%)	TSS (%)	AsA (mg/100g)
15	75.73mn	9.33def	74ab	84abcd	6.73ghi	128ab	71.10k	4.38no	26.67klm	1.79jk	140k	16.33c	3.47hijk	17.60lm
16	149.17a	6.20i	70cde	78fghi	5.27kl	113j	8.26t	2.70q	177.67a	1.40klm	137o	10.40on	4.83bc	24.93efg
17	118.87cd	8.27fgh	73abc	86ab	5.45kl	123de	98.3e	5.47jl	26.33lm	2.55gh	152c	16.03c	3.43hijk	27.13bcde
18	100.27fg	8.33fgh	73ab	83bcde	3.54n	119ghi	63.07m	7.13d	40.67g	2.55gh	149f	14.77de	3.63hij	18.33klm
19	63.8pq	8.13fg	74ab	80cdef	7.63f	103l	80.24h	6.24gh	35.00h	2.75fg	144h	11.99ij	3.63hij	18.33klm
20	89.33ijk	8.20fgh	70cde	79efgh	4.60m	105l	67.16l	6.12h	25.00mn	1.48kl	142j	12.56gh	4.67cde	18.33klm
21	109.13e	8.40fgh	74ab	86ba	3.77n	108k	101.41ed	8.07b	22.00no	2.20hij	153bc	11.84ij	4.57cdefg	20.53ijk
22	90.8hijk	8.67efg	73abc	83bcde	6.40ij	124cd	217.91a	10.50a	20.33op	4.12bc	143i	17.89a	3.53hijk	19.07jkl
23	85.07kl	10.40cde	65fg	68l	12.60b	95m	48.97p	4.32no	45.67f	2.17hij	141k	12.49gh	3.83fghi	16.87lm
24	87.2jkl	8.20fgh	72abcd	80defg	4.76m	123de	138.76b	7.65c	21.33no	2.79fg	143i	11.66jk	3.53hijk	20.53ijk
25	88.93ijk	8.20fgh	61gh	70l	6.80gh	90n	94.49f	5.16jkl	26.67klm	2.30ghi	139mn	16.90b	2.77kl	19.07ijk
26	90.93hijk	7.73fghi	71bcd	75ij	5.57kl	125bc	83.93g	3.90p	47.00f	3.83cd	137o	12.98g	3.03ijkl	28.60bc
27	107.07ef	8.47fgh	58i	62m	12.60b	116i	73.27kj	4.15po	64.67d	4.41b	147g	14.81d	6.53a	25.67def
28	97.93gh	10.47cde	69cde	75hi	9.53de	129a	56.83on	5.24jk	55.67e	3.06ef	131p	11.75jk	4.00defgh	27.13bcde
29	95.27ghi	6.67hi	74ab	76ghi	9.20d	121efg	123.20c	5.62i	53.33e	6.43a	144hi	13.80f	3.87efgh	26.40cde
30	134.73b	5.93i	61hi	69l	17.57a	124cd	22.15s	1.78r	154.33b	3.36de	138on	9.87pq	7.10a	22.73ghi
Min.	58.13	5.93	57.8	62.33	3.54	90.33	8.09	1.78	13.67	0.37	131.00	7.96	2.53	16.13
Max.	149.17	13.60	75.6	87.67	17.57	128.67	217.91	10.50	177.67	6.43	154.67	17.89	7.10	38.87
Avr.	93.46	9.42	67.99	77.02	7.12	116.77	71.26	5.58	43.93	2.30	144.18	12.08	4.08	23.10
SE	4.06	0.39	1.06	1.26	0.54	1.69	7.58	0.33	7.22	0.24	1.15	0.50	0.19	0.90
SD	22.23	2.15	5.82	6.88	2.98	9.26	41.53	1.79	39.54	1.31	6.31	2.73	1.04	4.93
LSD _{0.05}	7.45	1.89	3.50	4.02	0.34	2.50	3.65	0.35	3.80	0.49	0.78	0.48	0.80	2.53

Here, Ge 1= V1007282, 2= V1006136, 3= V1006422, 4= V1006186, 5= V1006484, 6= V1045786, 7= V1063607, 8= WP7, 9= WP8, 10= WP10, 11= C11, 12= C41, 13= C51, 14= C71, 15= BARI Tomato-9, 16= BARI Tomato-11, 17= BARI Tomato-14, 18= BARI Tomato-15, 19= BARI Tomato-8, 20= Phili-1, 21= Phili-2, 22= Bahar, 23= Binatomato-2, 24= Binatomato-7, 25= Binatomato-3, 26= Binatomato-4, 27= Binatomato-5, 28= Binatomato-8, 29= Binatomato-9, 30= Binatomato-10.

Ge= Genotype, PH= Plant height; PB= Number of primary branches per plant; DFF= Days to first flowering; 50%F= Days to 50% flowering; FC= Number of fruit cluster per plant; FH= Days to first harvest; IFW= Individual fruit weight; FD= Fruit diameter; FP= Fruits per plant; YP= Yield per plant; LH= Days to last harvest; DM= Dry matter percentage; TSS= Total soluble solid; AsA = Ascorbic acid content, Min.= Minimum, Max.= Maximum, Avr.= Average, Sd. Er.= Standard Error, Sd. Dev.= Standard deviation, LSD0.05= Least significant difference at 5% level of significance

Table 3. Genetic Parameters of different characters of 30 tomato genotypes

Traits	Genotypic variance	Phenotypic variance	Heritability (%)	GCV (%)	PCV (%)	GA (%)
PH	487.13	507.92	95.91	521.24	543.48	44.53
PB	4.19	5.52	75.87	44.48	58.62	3.67
DFF	32.35	36.95	87.56	77.04	87.99	10.96
50%F	45.37	51.43	88.22	88.93	100.80	13.03
FC	8.87	8.91	99.49	124.51	125.15	6.12
FH	84.99	87.34	97.31	93.62	96.22	18.73
IFW	1723.47	1728.48	99.71	2418.64	2425.67	85.40
FD	3.18	3.22	98.57	56.89	57.72	3.65
FP	1561.46	1566.89	99.65	3554.12	3566.48	81.26
YP	1.68	1.77	94.84	72.82	76.78	2.60
LH	39.72	39.95	99.43	33.61	33.81	12.95
DM	7.44	7.53	98.83	61.60	62.34	5.59
TSS	1.00	1.25	80.51	24.59	30.54	1.85
AsA	23.52	25.92	90.75	101.83	112.21	9.52

Here, PH= Plant height; PB.= Number of primary branches per plant; DFF= Days to first flowering; 50%F= Days to 50% flowering; FC= Number of fruit cluster per plant; FH= Days to first harvest; IFW= Individual fruit weight; FD= Fruit diameter; FP= Fruits per plant; YP= Yield per plant; LH= Days to last harvest; DM= Dry matter percentage; TSS= Total soluble solid; AsA= Ascorbic acid content; GCV= Genotypic co-efficient of variation; PCV=Phenotypic co-efficient of variation; GA= Genetic advance; GA (%)= Genetic advance in percentage of mean.

Determinate plants had a lower content of dry matter than the indeterminate [34], mainly as a result of reduced leaf surface area. Indeterminate plant types were best for use in breeding for a high content of dry matter [35]. Number of fruit cluster per plant (PCV=125.15, GCV=124.51); fruit diameter (PCV=57.72, GCV=56.89) and days to last harvest (PCV=33.81, GCV=33.61) showed relatively low difference between PCV and GCV. This indicates the low impact of environment on the expression of characters and hence, they could be improved by following different phenotypic selections like directional, disruptive and stabilized selections. Tasisa *et al.* [36] obtained slightly higher PCV (12.98%) than GCV (12.46) for number of fruit cluster per plant. Chernet *et al.* [37] recorded radial and polar diameter of fruit and recorded higher PCV (10.09 & 13.95) than GCV (10.04 & 13.90) for dry matter percentage. Similar result was also found by Osekita and Ademiluyi [38] for days to last harvest. Vinod *et al.* [39] reported PCV (8.14) and GCV (7.29) for TSS content. High values of GCV are an indication of high genetic variability among the germplasm and thus the scope for improvement of these characters through simple selection would be better. The differences between PCV and GCV was minimum for number of fruit cluster per plant, fruit diameter, days to last harvest and dry matter percentage suggesting that these traits were least affected by environment.

Heritability and Genetic advance

By GCV and PCV alone, it is not possible to determine the amount of variation which is heritable. The heritability along with genetic advance is more meaningful and helps in predicating the resultant effect of selection on phenotypic expression [17]. High heritability with high genetic advance

showed in plant height ($h^2_b=95.91$, GA=44.53); individual fruit weight ($h^2_b=99.71$, GA=85.40); fruits per plant ($h^2_b=99.5$, GA=81.26) indicating that these characters were under additive genetic effects and that this trait could be considered as reliable indices for selection (Table 3). Similar results were also reported by other researchers in tomato. Many researchers found high genetic advance for individual fruit weight [40-42] and plant height [43-45]. For number of fruits per plant Mehta and Asati [46] observed moderate heritability (54.70) with genetic advance in percentage of mean (27.86). Tasis *et al.* [47] reported high heritability (99%) with genetic advance (11.32) and genetic advance in percentage of mean (32.54%). Meitei *et al.* [27] observed high values of heritability (98.55) associated with high genetic advance (45.91) and high genetic advance in percentage of mean (132.30%) for number of fruits per plant. High heritability with low genetic advance observed in number of primary branches ($h^2_b=75.87$, GA=3.67); days to first flowering ($h^2_b=87.56$, GA=10.96); days to 50% flowering ($h^2_b=88.22$, GA=13.03); number of fruit cluster per plant ($h^2_b=99.49$, GA=6.12); days to first harvest ($h^2_b=97.31$, GA=18.73); fruit diameter ($h^2_b=98.57$, GA=3.65); yield per plant ($h^2_b=94.84$, GA=2.6); days to last harvest ($h^2_b=99.43$, GA=12.95); dry matter percentage ($h^2_b=98.83$, GA=5.59); total soluble solid ($h^2_b=80.51$, GA=1.85); AsA content ($h^2_b=90.75$, GA=9.52). This situation indicated limited scope for the improvement of this character through selection but hybridization followed by progeny selection will be effective. Kaushik *et al.* [30] also reported high amount of heritability (69.20%) but low genetic advance as percentage of mean (3.2%) for days to first flowering. Mohanty [48] reported that the range of days to 50% flowering was 51 to 54 days and low heritability and low

genetic advance as percentage of mean. Meena and Bahadur [45] found high heritability (100) with low genetic advance (16.85) and low genetic advance as percentage of mean (27.40%) for days to 50% flowering. Similar results were also found by Tasisa *et al.* [36] and Meitei *et al.* [27] for number of fruit cluster per plant. Many researchers reported high heritability with low genetic advance for days to maturity [49-51]. Osekita and Ademiluyi [38] found similar findings for days to last harvest. Meena and Bahadur [45] found high heritability (98%) and low genetic advance (2.16) and genetic advance in percentage of mean (51.59%) for TSS content. Mohamed *et al.* [28] also reported high heritability (80%) with low genetic advance (3.47) for AsA content. From the present study it is found that smaller fruit have more AsA than the larger one. Bahar had the highest fruit diameter (10.5 cm) and fruit weight (217.90 g) but the AsA content is very low (19.06 mg%) comparing other small sized fruit. Barooah and Mohan [52] studied 11 varieties and found that Sioux having largest fruit size had less AsA content and Chickugrande, the smallest fruit had highest ascorbic content (26.30 mg%), which were in agreement with this research findings. The better indicators of heritable proportion of variation estimation are heritability estimation. The high heritability indicates the effectiveness of selection based on phenotypic traits but does not necessarily mean a high genetic gain for a particular trait. So, consideration of both, heritability and genetic advance is more important for predicting effective selection than heritability alone. Johnson *et al.* [17] reported that heritability estimates along with genetic advance would be more rewarding than heritability alone in predicting the consequential effect of selection to choose the best individual. High values of heritability with high genetic advance were observed for individual fruit weight followed by fruits per plant and plant height indicated that these traits were under the control of additive gene action and directional selection for these characters in the genetically diverse material could be effective for desired genetic improvement. Similar findings were reported earlier by Mehta and Asati [46], Mohanty [48] in tomato.

3.2. Correlation Co-efficient Studies

A crop breeding program aimed at increasing the yield requires consideration not only of yield but also of its associate components that have direct or indirect impact on yield. Correlation analyses give an insight into the genetic variability present in populations. Correlation coefficient analysis measures the mutual relationship between various plant characters and determines the component characters on which selection can be based for improvement in yield. Correlation studies were made among 14 characters of which yield per plant were positively correlated with individual fruit weight (0.609) and number of fruit cluster per plant (0.396) (Table 4). The significant and positive association

between the traits suggested additive genetic model thereby less affected by the environmental fluctuation. Similar results have also been reported by Susic *et al.* [53] and Rani *et al.* [54] for individual fruit weight. Reddy *et al.* [55] also observed positive association between number of fruit cluster and fruit yield per plant. Similar findings were also reported by Kumar and Dudi [56] for fruit cluster. Yield per plant also showed positive but insignificant relationship with plant height (0.159), fruit diameter (0.105), number of fruits per plant (0.126), dry matter (%) (0.352), TSS (0.126) and AsA (0.163). Sharma and Singh [57] reported positive insignificant association with plant height, number of fruits per plant and TSS. Similar findings were also reported by Golani *et al.* [44] and Manna and Paul [34].

But Islam *et al.* [58] observed positive significant association between yield and fruit diameter, which is contradictory with the present research finding. On the other hand de Souza *et al.* [59] reported positive insignificant association with fruit diameter, which support the present results. Similar results also reported by Anitha *et al.* [60] for AsA. The positive and non-significant association referred information of inherent relation among the pairs of combination. Yield per plant showed negative significant relationship with number of primary branch per plant (-0.421) but previously it was reported by many researchers like Reddy *et al.* [55] and Sharma and Singh [57] that yield was positively significant with number of primary branches per plant, which are not in agreement with our findings. But Meena and Bahadur [61] reported negative significant association between yield and number of primary branches, which approves reports of the present investigation. Negative but insignificant association was found with days to 50% flowering (-0.240), days to last harvest (-0.318). The negative and non-significant association referred a complex linked of relation among the pair of combinations.

Character association revealed the mutual relationship between two characters, and it is important parameters for taking a decision regarding the nature of selection to be followed for improvement in the crop under study. The overall results indicated that yield per plant was positively associated with days to first flowering (0.099), number of fruit cluster per plant (0.396), individual fruit weight (0.609), number of fruits per plant (0.126) and negatively associated with number of primary branches per plant (-0.421) and days to last harvest (-0.318). The traits which do not show any significant association or very negligible amount of association can be discarded to reduce the number of traits to characterize. This correlation result can be used as basis for character discard if similar research is conducted in the future using additional morphological traits. Elimination of redundant traits will reduce the workload of researcher and will render characterization less cumbersome and more efficient.

Genetic Variability and Traits Association Analysis of Tomato (*Lycopersicon esculentum* L.)
Genotypes for Yield and Quality Attributes

Table 4. Phenotypic correlation co-efficients between yield and yield contributing traits

	YP	PH	PB	DFF	50%F	FC	FH	IFW	FD	FP	LH	DM	TSS	AsA
YP	1.00	0.159	-0.421*	0.099	-0.240	0.396*	0.078	0.609***	0.105	0.126	-0.318	0.352	0.126	0.163
PH		1.000	-0.336	0.005	-0.047	0.184	0.146	-0.105	-0.239	0.476**	0.036	-0.022	0.488**	0.239
PB			1.000	-0.546**	-0.331	-0.100	-0.054	-0.209	0.300	-0.326	0.362*	-0.062	-0.221	-0.040
DFF				1.000	0.864***	-0.422*	0.312	0.322	0.157	-0.255	-0.027	-0.086	-0.259	-0.022
50%F					1.000	-0.554***	0.335	0.178	0.181	-0.322	0.222	-0.217	-0.271	-0.088
FC						1.000	-0.008	-0.163	-0.388*	0.406*	-0.245	-0.021	0.499*	0.047
FH							1.000	0.053	0.030	-0.058	0.025	-0.159	0.152	0.254
IFW								1.000	0.671***	-0.442*	-0.081	0.431*	-0.223	-0.093
FD									1.000	-0.653***	0.211	0.245	-0.222	-0.280
FP										1.000	-0.276	-0.029	0.336	0.199
LH											1.000	0.034	-0.065	-0.065
DM												1.000	-0.300	-0.228
TSS													1.000	0.172
AsA														1.000

*, ** and *** indicates significant at $p < 0.05$, $p < 0.01$ and $p < 0.001$

Here,

PH= Plant height; PB= Number of primary branches per plant; DFF= Days to first flowering; 50%F= Days to 50% flowering; FC= Number of fruit cluster per plant; FH= Days to first harvest; IFW= Individual fruit weight; FD= Fruit diameter; FP= Fruits per plant; YP= Yield per plant; LH= Days to last harvest; DM= Dry matter percentage; TSS= Total soluble solid; AsA = Ascorbic acid content.

3.3. Path Co-efficient Studies

Yield is the sum total of the many component characters which directly or indirectly contributed to it. Correlation studies give an idea about the positive and negative associations of different characters with yield and also among themselves. But the nature and extent of contribution of these characters towards yield is not obtained. Path coefficient analysis was used to make partition of the correlation coefficient of the different characters studied to know direct and indirect effects on yield. The information obtained helps in giving proper weightage to the various characters during selection or other breeding program so that the improvement of desirable traits can be achieved effectively. Path coefficient analysis revealed that individual fruit weight (0.704), days to first flowering (0.590), number of fruit cluster (0.259), number of fruits per plant (0.192), days to first harvest (0.107) and AsA (0.0087) had direct positive effect on yield per plot, indicating these are the main contributors to yield (Table 5). Dhankar *et al.* [62], Verma and Sarnaik [63], Mageswari [64], Yadav and Singh [65] found that fruits per plant had the highest positive direct effect on yield. The association of AsA content with yield is similar with the findings of Manna and Paul [34]. But days to last harvest (-0.318) was negatively correlated with yield per plant. This negative effect mainly was due to the fact that positive direct effect of days to last harvest on yield was nullified by its negative indirect effects via days to 50% flowering (-0.176), number of fruit cluster per plant (-0.053), number of primary branches (-0.061) and individual fruit weight (-0.057). However days to 50% flowering (-0.792), number of primary branches (-0.169), fruit diameter (-0.055), plant height (-0.038), dry matter (%) (-0.038) and TSS (-0.038) had direct negative effect on yield per plot. Similar results have also been reported by Tiwari and Upadhyay [66] for plant height. But plant height (0.159), fruit diameter (0.105), dry matter (%) (0.352) and TSS (0.126) were positively correlated with yield per plant. This positive effect

mainly was due to the fact that negative direct effect of plant height on yield was nullified by its positive indirect effects via fruits per plant (-0.091), number of primary branches (0.057) and number of fruit cluster (0.048). Negative effect of fruit diameter on yield was nullified by its positive indirect effects via individual fruit weight (0.472). Negative effect of dry matter (%) on yield was nullified by its positive indirect effects via individual fruit weight (0.303) and days to 50% flowering (0.172). Negative effect of TSS (-0.038) on yield was nullified by its positive indirect effects via days to 50% flowering (0.215) and number of fruit cluster (0.129). The residual effect determines how best the causal factors account for the variability of the dependent factor, the yield per plant in this case. In case of the present study the residual effect was 0.41 indicating that the fourteen traits explain only 59% of variability in yield per plant. The reason seems to be very low and non-significant correlation of some traits with yield. Besides, some other factors which have not been considered here need to be included to account fully for the variation in yield.

The above information revealed that highly significant positive correlation with highest positive direct effect was observed in individual fruit weight followed by number of fruit cluster per plant. Gorbtenko and Gorbtenko [67] and Kumar *et al.* [69] studied path analysis of economically useful characters of tomato and observed that single fruit weight had an appreciable direct effect on yield per plant. Similar results were also obtained by Alam *et al.* [68], Kumar *et al.* [69], Hidayatullah [70] and Islam and Khan [12] in tomato. Previously it was reported that average fruit weight and number of fruits per plant can be used as selection criteria for improvement in tomato [71-73]. So, from this research it may be recommended that the individual fruit weight and number of fruit cluster per plant could be considered as critical criteria for yield improvement in these genotypes of tomatoes.

Table 5. Path-coefficient analysis showing the direct and indirect effect of different yield contributing traits on fruit yield. (Bold diagonal values are direct effects and off-diagonal values are indirect effects)

	PH	PB	DFP	50%F	FC	FH	IFW	FD	FP	LH	DM	TSS	AsA	Correlation with YP
PH	-0.038	0.057	0.003	0.037	0.048	0.016	-0.074	0.013	0.091	0.005	0.001	-0.019	0.021	0.159
PB	0.013	-0.169	-0.322	0.262	-0.026	-0.006	-0.147	-0.016	-0.063	0.045	0.002	0.008	-0.003	-0.421*
DFP	0.000	0.092	0.590	-0.684	-0.109	0.033	0.227	-0.009	-0.049	-0.003	0.003	0.010	-0.002	0.099
50%F	0.002	0.056	0.510	-0.792	-0.143	0.036	0.125	-0.010	-0.062	0.028	0.008	0.010	-0.008	-0.240
FC	-0.007	0.017	-0.249	0.438	0.259	-0.001	-0.115	0.021	0.078	-0.031	0.001	-0.019	0.004	0.396*
FH	-0.006	0.009	0.184	-0.265	-0.002	0.107	0.038	-0.002	-0.011	0.003	0.006	-0.006	0.022	0.078
IFW	0.004	0.035	0.190	-0.141	-0.042	0.006	0.704	-0.037	-0.085	-0.010	-0.016	0.009	-0.008	0.609***
FD	0.009	-0.051	0.093	-0.144	-0.100	0.003	0.472	-0.055	-0.125	0.026	-0.009	0.008	-0.024	0.105
FP	-0.018	0.055	-0.151	0.255	0.105	-0.006	-0.311	0.036	0.192	-0.034	0.001	-0.013	0.017	0.126
LH	-0.001	-0.061	-0.016	-0.176	-0.063	0.003	-0.057	-0.012	-0.053	0.125	-0.001	0.002	-0.006	-0.318
DM	0.001	0.011	-0.051	0.172	-0.006	-0.017	0.303	-0.013	-0.006	0.004	-0.038	0.011	-0.020	0.352
TSS	-0.019	0.037	-0.153	0.215	0.129	0.016	-0.157	0.012	0.065	-0.008	0.011	-0.038	0.015	0.126
AsA	-0.009	0.007	-0.013	0.070	0.012	0.027	-0.065	0.015	0.038	-0.008	0.009	-0.007	0.087	0.163

Residual effect: 0.41

*, ** and *** indicates significant at $p < 0.05$, $p < 0.01$ and $p < 0.001$

Here, PH= Plant height; PB= Number of primary branches per plant; DFP= Days to first flowering; 50%F= Days to 50% flowering; FC= Number of fruit cluster per plant; FH= Days to first harvest; IFW= Individual fruit weight; FD= Fruit diameter; FP= Fruits per plant; YP= Yield per plant; LH= Days to last harvest; DM= Dry matter percentage; TSS= Total soluble solid; AsA = Ascorbic acid content.

4. Conclusions and Recommendation

Plenty of variation was found among the genotypes for the studied traits. Among all the traits number of fruits per plant exhibited high estimates of PCV (3566.48), GCV (3554.12) followed by individual fruit weight (PCV=2425.67, GCV=2418.64), plant height (PCV=543.48, GCV=521.24) and number of fruit cluster per plant (PCV=125.15, GCV=124.51). The lowest PCV (30.54) and GCV (24.59) values were recorded for TSS contents. The studied traits expressed moderate to high heritability estimates ranging from 75.87 to 99.71%. In the present study, high heritability along with high genetic advance was noticed for the traits, individual fruit weight ($h^2_b = 99.71$, GA=85.40), fruits per plant ($h^2_b = 99.65$, GA=81.26) and plant height ($h^2_b = 95.61$, GA=44.53). These traits can be improved through simple or progeny selection methods. Other traits (i.e. number of primary branches, days to first flowering, days to 50% flowering, number of fruit cluster per plant, days to first harvest, fruit diameter, yield per plant, days to last harvest, dry matter percentage, total soluble solid and AsA content) showed high heritability along with moderate or low genetic advance which can be improved by inter-mating superior genotypes to develop segregating population for further selection and improvement. Yield per plant was significantly correlated with individual fruit weight (0.609***), number of fruit cluster per plant (0.396*) in positive direction and significantly negatively correlated with number of primary branch per plant (-0.421*). The findings suggested that the selection of genotypes having high individual fruit weight with reasonable compromises for higher number of fruit cluster, lower number of primary branches, higher fruit diameter, high number of fruits per plant and moderate plant height should be the priority of breeders to achieve higher yield. Based on our research findings, the genotypes Bahar, Binatomato-7, Binatomato-9, WP10 and Phili-2 can be used for future breeding program targeting yield and quality improvement because these genotypes have the suggested characteristics. The path coefficient analysis revealed that highly significant positive correlation, with highest positive direct effect was found in individual fruit weight (0.704) followed by number of fruit cluster per plant (0.259). Our findings suggest that individual fruit weight and number of fruit cluster per plant could be considered as important criteria for yield improvement in tomato genotypes.

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