

**EVALUATION OF PUMPKIN (*Cucurbita moschata* Duchesne ex
poir.) GENOTYPES BASED ON PHENOTYPIC TRAITS**

MD. ABDULLAH AL MASUD



**DEPARTMENT OF GENETICS AND PLANT BREEDING
SHER-E-BANGLA AGRICULTURAL UNIVERSITY
DHAKA-1207**

JUNE, 2016

**EVALUATION OF PUMPKIN (*Cucurbita moschata* Duchesne ex
poir.) GENOTYPES BASED ON PHENOTYPIC TRAITS**

BY

MD. ABDULLAH AL MASUD

REGISTRATION NO. 10-03944

A Thesis

Submitted to the Faculty of Agriculture,
Sher-e-Bangla Agricultural University, Dhaka,
in partial fulfillment of the requirements
for the degree of

**MASTER OF SCIENCE (MS)
IN
GENETICS AND PLANT BREEDING
SEMESTER: Jan -June, 2016**

Approved by:

(Prof. Dr. Mohammad Saiful Islam)
Supervisor

(Prof. Dr. Naheed Zeba)
Co-supervisor

(Prof. Dr. Md. Jamilur Rahman)
Chairman
Examination Committee



Dr. Mohammad Saiful Islam

Professor

*Department of Genetics and Plant Breeding
Sher-e-Bangla Agricultural University*

Dhaka 1207, Bangladesh

Mob: +8801742843195

E-mail: saiful_sau@yahoo.com

CERTIFICATE

*This is to certify that the thesis entitled, "EVALUATION OF PUMPKIN (*Cucurbita moschata* Duchesne ex Poir) GENOTYPES BASED ON PHENOTYPIC TRAITS" submitted to the Faculty of Agriculture, Sher-e-Bangla Agricultural University, Dhaka, in partial fulfillment of the requirements for the degree of **MASTER OF SCIENCE** In Genetics And Plant Breeding, embodies the result of a piece of bona fide research work carried out by MD. ABDULLAH AL MASUD; REGISTRATION NO. 10-03944, under my supervision and guidance. No part of this thesis has been submitted for any other degree or diploma.*

I further certify that any help or sources of information, as has been availed of during the course of this investigation have been duly acknowledged.

*Dated: June, 2016
Dhaka, Bangladesh*

*(Professor Dr. Mohammad Saiful Islam)
Supervisor*



*Dedicated to
My
Beloved Parents*

ACKNOWLEDGEMENTS

First of all, the author expresses his gratitude to the “Omnipotent Creator” for his never-ending blessing, it is a great pleasure to express profound thankfulness to his respected parents, who entiled much hardship inspiring for prosecuting his studies, thereby receiving proper education.

The author expresses his sincere gratitude, heartfelt respect and immense indebtedness to his reverend Supervisor, Prof. Dr. Mohammad Saiful Islam , Department of Genetics and Plant Breeding, Sher-e-Bangla Agricultural University, Dhaka for his sincere guidance, scholastic supervision, constructive criticism and constant inspiration throughout the course and in preparation of the manuscript of the thesis.

The author wishes to extend his deepest gratitude and sincere appreciation to his Co-supervisor, Prof. Dr. Naheed Zeba Department of Genetics and Plant Breeding, Sher-e-Bangla Agricultural University, Dhaka for his utmost cooperation and constructive suggestions to conduct the research work as well as preparation of the thesis.

He is grateful to Prof. Dr. Kamal Uddin Ahmed, Honourable Vice Chancellor, Sher-e-Bangla Agricultural University, Dhaka for providing me with all possible help during my studies.

He expresses his sincere respect to the Chairman, Prof. Dr. Jamilur Rahman and my honourable teachers Prof. Dr. Md. Shahidur Rashid Bhuiyan, Prof. Dr. Md. Sarowar Hossain, Prof. Dr. Firoz Mahmud , Prof. Dr. Md. Ashaduzzaman Siddiquee and Associate Prof. Dr. Abdur Rahim, and all other staffs of Department of Genetics and Plant Breeding, Sher-e-Bangla Agricultural University, Dhaka for their teaching, direct and indirect advice, encouragement and co-operation during the study period.

He would like to thank Dr. Md. Abdul Malek, honourable Principle Scientific Officer of Bangladesh Agricultural Research Institute (Plant Genetic Resources Centre, BARI) for his help to supply 37 advanced breeding line of pumpkin

The author also would like to thank all the staff of Bangladesh Agricultural Research Institute of BARI, Gazipur for their immense support.

The author also would like to thank all of his family members who have helped him with technical support to prepare this thesis paper. He also thanks all of his friends and well wishers to help him in his research specially Dewan Mahmudul Hoque, Shah Jalal, Rasel, Raja Masum, Najmul Khan, Shamima Khan, Samsun Naher.

He found no words to thank my mother, brother and sister-in-law Aleya Parvin, Md. Abdulla Al Mamun and Kaberi Mamun for their unquantifiable love, sacrifice and never ending affection for bringing my dream to proper shape. They are my constant source of zeal, inspiration and enthusiasm in the critical moment of my studies.

June, 2016

The Author

SAU, Dhaka

LIST OF CONTENTS

CHAPTER	TITLE	PAGE NO.
	ACKNOWLEDGEMENT	I-II
	LIST OF CONTENTS	III
	LIST OF TABLES	VIII
	LIST OF FIGURES	IX
	LIST OF PLATES	X
	LIST OF APPENDICES	XI
	ABBREVIATIONS	XII
	ABSTRACT	XIII
I	INTRODUCTION	01
II	REVIEW OF LITERATURE	04
2.1	Variability, heritability and genetic advance	04
2.2	Correlation co-efficient	15
2.3	Path coefficient analysis	18
2.4	Genetic diversity	21
III	MATERIALS AND METHODS	27
3.1	Experimental site	27
3.2	Geographical location	27
3.3	Climate	27
3.4	Characteristics of soil	28
3.5	Planting materials	28
3.6	Design and layout of the experiment	28
3.7	Polybag preparation and raising seedling	30
3.8	Land preparation	30

LIST OF CONTENTS (Cont'd)

3.9	Pit preparation	30
3.10	Application of manure and fertilizers	30
3.11	Transplanting of seedling	33
3.12.1	Thinning and gap filling	33
3.12.2	Weeding and mulching	33
3.12.3	Irrigation and after care	33
3.12.4	Pesticide application	33
3.13	Harvesting	34
3.14	Data recording	34
3.14.1	Plant characteristics	34
3.14.1.1	Leaf length (cm)	34
3.14.1.2	Leaf breadth (cm)	34
3.14.1.3	Internode distance (cm)	34
3.14.2	Inflorescences characteristics	34
3.14.2.1	Days to first male flowering	34
3.14.2.2	Days to first female flowering	35
3.14.2.3	Pedicel length of male flower	35
3.14.2.4	Pedicel length of female flower	35
3.14.3	Fruit characteristics	35
3.14.3.1	Fruit length (cm)	35
3.14.3.2	Fruit breadth (cm)	35
3.14.3.3	Fruit weight (kg)	35
3.14.3.4	Fruit yield per plant (kg)	35
3.15.1	Statistical analysis	36
3.15.1.1	Estimation of genotypic and phenotypic variances	36

LIST OF CONTENTS (Cont'd)

3.15.1.2	Estimation of genotypic and phenotypic correlation co-efficient	36
3.15.1.3	Estimation of genotypic and phenotypic co-efficient of variation	37
3.15.1.4	Estimation of heritability	38
3.15.1.5	Estimation of genetic advance	38
3.15.1.6	Estimation of genetic advance mean's percentage	38
3.15.1.7	Estimation of path co-efficient	39
3.15.2	Multivariate analysis	42
3.15.2.1	Principal component analysis (PCA)	42
3.15.2.2	Cluster analysis (CA)	42
3.15.2.3	Canonical vector analysis (CVA)	43
3.15.2.4	Calculation of D^2 values	43
3.15.2.5	Computation of average intra-cluster distances	43
3.15.2.6	Computation of average inter-cluster distances	44
3.15.2.6	Cluster diagram	44
3.15.2.7	Selection of varieties for future hybridization program	44
IV	RESULTS AND DISCUSSION	46
4.1	Genetic variability	46
4.1.1	Leaf length without petiole	46
4.1.2	leaf breadth	48
4.1.3	Internode distance	52
4.1.4	Days to first male flowering	52
4.1.5	Days to first female flowering	52
4.1.6	Male flower pedicel length	53

LIST OF CONTENTS (Cont'd)

4.1.7	Female flower pedicel length	57
4.1.8	No of male flower	57
4.1.9	No of female flower	58
4.1.10	Fruit length	58
4.1.11	Fruit breadth	59
4.1.12	Fruit weight	63
4.1.13	Yield per plant	64
4.2	Correlation Co-efficient	64
4.2.1	Leaf length without petiole	65
4.2.2	leaf breadth	65
4.2.3	Internode distance	65
4.2.4	Days to first male flowering	67
4.2.5	Days to first female flowering	67
4.2.6	Male flower pedicel length	67
4.2.7	Female flower pedicel length	68
4.2.8	No of male flower	68
4.2.9	No of female flower	68
4.2.10	Fruit length	69
4.2.11	Fruit breadth	69
4.2.12	Fruit weight	69
4.3	Path co-efficient Analysis	70
4.3.1	Internode distance	72
4.3.2	Days to first female flowering	72
4.3.3	Male flower pedicel length	72
4.3.4	Female flower pedicel length	73

LIST OF CONTENTS (Cont'd)

4.3.5	No of male flower	73
4.3.6	No of female flower	73
4.3.7	Fruit length	74
4.3.8	Fruit breadth	74
4.3.9	Fruit weight	74
4.4	Genetic diversity Analysis	75
4.4.1	Principal component analysis (PCA)	75
4.4.2	Non-hierarchical clustering	77
4.4.3	Canonical variate analysis	80
4.4.4	Contribution of phenotypic traits toward divergence of the genotypes	83
4.4.5	Selection of parents for future hybridization	85
V	SUMMARY AND CONCLUSION	86
	REFERENCES	89
	APPENDICES	96

LIST OF TABLES

Table No.	Title of the Tables	Page No.
1	Name of 24 pumpkin genotypes used in the study	29
2	Doses of manure and fertilizers used in study	32
3	Estimation of genetic variability for yield contributing characters related to yield of pumpkin	47
4	Coefficients of phenotypic and genotypic correlation among different yield components of twenty pumpkin genotypes	66
5	Path coefficient analysis showing direct and indirect effects of different characters on yield	71
6	Eigen value, % variance and cumulative (%) total variance of the principal components	76
7	Distribution of genotypes in different clusters	78
8	Cluster mean for thirteen yield and yield characters of pumpkin Genotypes	79
9	Intra (Bold) and inter cluster distances (D^2) for 20 genotypes	81
10	Relative contribution of thirteen characters towards divergence of the genotypes	84

LIST OF FIGURES

Figure No.	Title of the figures	Page No.
1	Cluster diagram of twenty four genotypes of pumpkin	82
2	Scattered diagram of twenty four genotypes of pumpkin	82

LIST OF PLATES

PLATE No.	NAME OF PLATES	Page No.
1	Raising of seedling in polybag	31
3a	Showing morphological variation in leaf among different pumpkin genotypes (G ₁ -G ₈)	49
3b	Showing morphological variation in leaf among different pumpkin genotypes (G ₉ -G ₁₆)	50
3c	Showing morphological variation in leaf among different pumpkin genotypes (G ₁₇ -G ₂₄)	51
4a	Showing morphological variation in flower among different pumpkin genotypes (G ₁ -G ₈)	54
4b	Showing morphological variation in flower among different pumpkin genotypes (G ₉ -G ₁₆)	55
4c	Showing morphological variation in flower among different pumpkin genotypes (G ₁₇ -G ₂₄)	56
5a	Showing morphological variation in fruits among different pumpkin genotypes (G ₁ -G ₈)	60
5b	Showing morphological variation in fruits among different pumpkin genotypes (G ₉ -G ₁₆)	61
5c	Showing morphological variation in fruits among different pumpkin genotypes (G ₁₇ -G ₂₄)	62

APPENDICES

Appendix No.	Title of Appendix	Page No.
1	Map showing the experimental site under the study	28
2	Monthly average temperature, relative humidity and total rainfall and sunshine of the experimental site during the period from March 2016 to July 2016	29
3	The mechanical and chemical characteristics of soil of the experimental site as observed prior to experimentation (0-15 cm depth)	39
4	Analysis of variance for different phenotypic plant characters of 24 pumpkin genotypes	40
5	Mean performance of 24 pumpkin genotypes based on different morphological traits related to yield	41

SOME COMMONLY USED ABBREVIATION

Full word	Abbreviations
Agro-Ecological Zone	AEZ
And others	<i>et al.</i>
Bangladesh Agricultural Research Institute	BARI
Bangladesh Bureau of Statistics	BBS
Centimeter	cm
Co-efficient of variation	CV
Days after sowing	DAS
Degree Celsius	⁰ C
Degrees of Freedom	d.f
Etcetera	etc.
Food and Agriculture Organization	FAO
Figure	fig
Genetic Advance	GA
Genotypic co-efficient of variation	GCV
Genotypic variance	δ^2_g
Mean sum of square	MSS
Gram (s)	g
Heritability in broad sense	h^2_b
Hectare	ha
Kilogram	kg
Genotypic variance	δ^2_g
Journal	j.
Phenotypic variance	δ^2_p
Meter	m
Randomized Complete Block Design	RCBD
Meter Square	m^2
Phenotypic co-efficient of variation	PCV
Sher-e-Bangla Agricultural University	SAU
percent	%
Standard Error	SE
Tons per hectare	t/ha
University	Univ.
Variety	var.

**EVALUATION OF PUMPKIN (*Cucurbita moschata* Duchesne ex Poir.) GENOTYPES
BASED ON PHENOTYPIC TRAITS**

By

MD. ABDULLAH AL MASUD

ABSTRACT

A field experiment was conducted during March 2016 to September 2016 using 24 pumpkin (*Cucurbita moschata* Duchesne ex Poir.) genotypes based on phenotypic traits in a randomized complete block design with three replications. The experiment conducted at the research farm of Sher-E-Bangla Agricultural University, Dhaka. The objectives of the study were to evaluate the variability of genotypes and select suitable genotypes for further breeding program. Experiment showed significant differences among the genotypes. Phenotypic variance was higher than that of genotypic variance for all the characters. High genotypic co-efficient of variation (GCV) was found for fruit yield, fruit weight and number of female flowers per plant. Low GCV was observed for days to first male flowering, days to first female flowering and leaf breadth. High heritability with high genetic advance in percent of mean was observed in fruit length, fruit breadth, fruit weight and pedicel length of female flower which indicated that these traits would be effective for genetic improvement. High heritability with low genetic advance in percent of mean was observed in days to first male flower, female flower and leaf breadth. Correlation studies showed that positive and significant correlation of fruit yield per plant with fruit length, fruit breadth and fruit weight. Path co-efficient analysis showed that days to first female flowering had the highest positive direct effect on yield per plant followed by internode distance, pedicel length of male flowers, pedicel length of female flowers, number of female flower per plant, fruit breadth and fruit weight. The highest intra cluster distance was found in cluster IV and the lowest was found in cluster III. Among five clusters the highest inter cluster distance was found in between cluster III and IV and the lowest between cluster I and IV. Considering group distance and phenotypic performances the inter genotypic crosses between G₁ (BD 10149) and G₂₄; G₁ (BD 10149) and G₂₃ (BD 4393); G₂₃ (BD 4393) and G₂₄ (PRITY F1) may be suggested for future hybridization program.

CHAPTER I

INTRODUCTION

Pumpkin (*Cucurbita moschata* Duchesne ex Poir.) or sweet gourd belongs to the family cucurbitaceae is a common year round climbing habit vegetable all over Bangladesh. This vegetable have great source of vitamin-A nutrient which have potentiality to solve the night-blindness particularly of the vulnerable groups of people. According to its nutritional and consumptive values it is grown widely from homestead to commercial field and marketed all over the country. Pumpkin also well grown in Europe, America, Canada, China, India, Turkey, Italy and Egypt.

There is debate about the taxonomy of the genus as the number of the accepted species varies from 13 to 30. The five domesticated species are *C. moschata*, *C. mixta*, *C. maxima*, *C. ficifolia* and *C. pepo* . The origin of cucurbits genus from central and south America with twenty pairs of chromosome ($2n=40$). Archaeological investigation have found evidence of domestication of cucurbits going back over 8000 year B.C. After the discovery of the new world, the cultivated cucurbits were introduced into the old world (Grubben, 2004).

Pumpkin vegetable is a monoecious plant which have both male and female flower in a single plant. It is mainly seed propagated, day neutral and well grown in tropical and sub-tropical region of the world. The typical cultivated species has five-lobed leaves with long petioles with the leaves alternately arranged on the stem. The stems are angular, spring like tendrils grow from each node. Egg shaped to heart shaped leaves, leaves may or may not have white spots. Fruits size and color are various according to species. Cucurbitaceae male flower generally have five stamens, but in *Cucurbita* there are only three, and their anthers are joined together so that there appears to be one (Mabberley, 2008). Female flowers have thick pedicels, and an inferior ovary with 3–5 stigmas that each have two lobes (Lu and Jeffery, 2015). The

calyx of *Cucurbita moschata* male flowers is comparatively short (Saade *et al.*, 2013).

According to the USDA National Nutrient Database, one cup of Pumpkin cooked, boiled, drained and without salt contains 49 calories, 1.76 gram of protein, 0.17 gram of fat, 0.00 gram of cholesterol, 12 gram of carbohydrates (including 2.7 gram of fibre and 5.1 g of sugar). Consuming one cup of cooked, canned pumpkin would provide well over 100 percent of our daily needs for vitamin-A, 20 percent vitamin-C, 10 percent or more for vitamin-E, riboflavin, copper, potassium and manganese and at least 5% for thiamin and pantothenic acid. It is also rich source of natural poly-phenolic flavonoid compounds such as, β carotenes, crypto xanthine, lutein and zeaxanthin. The fruit also available source of B-complex group of vitamins like folates, niacin, vitamin B-6 (pyridoxine). Seeds are an excellent source of dietary fiber and mono-unsaturated fatty acids, which are good for heart health.

Pumpkins have anti-diabetic, antioxidant, anti-carcinogenic, and inflammatory pharmacological properties (Yadav *et al.*, 2010). Pumpkin seeds have high levels of crude protein, calcium, iron, potassium, phosphorus, magnesium, zinc, (Mansour *et al.*, 1993) and beta-carotene.

Pumpkin are very versatile in their uses for cooking. Most part of the pumpkin are edible including the fleshy shell, the seeds, the leaves and even the flower. Ripen pumpkin is used to making soup, pie, purees in Europe, North America and Canada. Seeds are often roasted and eaten as a snack. In Middle East pumpkin is used for sweet dishes, a well-known sweet delicacy is called 'halawa yaqtin'. In south Asian countries like India pumpkin is cooked with butter, sugar and spices in a dish called 'kadu ka halwa'. Pumpkin can be used to flavor both alcoholic and non-alcoholic beverages. Pumpkin seed oil contains fatty acid (Bavec *et al.*, 2007). Raw vegetative part used as substitute for poultry feed.

The soil and agro climates of Bangladesh are highly conducive for growing numerous vegetables. Among them the cucurbit occupied 66 percent of the land under vegetables production in Bangladesh and contributes 11 percent of total vegetable production in the country (IPM CRSP, 2004). The total production area is about 27,377 acre where total yield 100493 MT (BBS, 2015).

Pumpkin is the perfect crop for maintaining our daily demand where the daily requirement of a full grown person need 285 g. Vegetable (Ramphall and Gill, 1990). Pumpkin also exportable vegetable crop which have good storability up to 6 months maintaining nutritional levels during the long dry seasons (Mendlinger *et al.*, 1991). So it has great potentiality to earn foreign currency. It is also low management practice and congenial to the environment rather than other vegetables. For this reason farmers are encouraged to cultivate this vegetable day by day.

As the pumpkin has wide species and numerous genotypes, it is grown in many places. Proper variety and genotypes would not be selected which is desire to our farmers. Being monoecious and cross-pollinated it has great potentiality for diseases and insect infestation. So it's a challenge to our breeder to develop high yielding quality varieties from existing genotypes or land races or from the segregates of a cross. For this we have to gather much knowledge about its phenotypic characters, character association and yield contributing characters. Genetic resource may helpful for its improvement. Therefore, the present research was under taken with the following objectives:

- To evaluate the phenotypic traits among different genotypes,
- To find out the relationship among the genotypes phenotypic traits towards yield and
- To select suitable genotypes of pumpkin and for future breeding program.

CHAPTER II

REVIEW OF LITERATURE

Pumpkin is a monoecious summer vegetables in our country. Because of its low cost of production and nutritious values, it is cultivated all over the world. But it is pessimistic issue that there is not available research work done to its improvement in our country. Meanwhile, for the crop improvement research work should be done on the molecular level, genetic resources, genetic diversity, correlation, path co-efficient analysis, heritability and genetic advance seem to be emaciated. So in this section, pumpkin and other some cucurbit crops information are reviewed.

2.1 Variability, Heritability and Genetic Advance

Mangal *et al.*, (1981) suggested that in bitter gourd significant variation for fruit length and diameter present and high heritability in bitter gourd for vine length. Mondal *et al.*, (1989) observed the genetic variability of 31 watermelon genotypes and found a wide range of variability for days to first fruit harvest, fruit length, and fruit diameter, number of fruits per plant and fruit yield per plant. Abusaleha and Dutta (1990) found out a study with 65 genetic stocks to assess the genetic variation and heritability in ridge gourd. Significant variability was observed for all the characters at phenotypic as well as genotypic level with a very wide range of values. Rahman *et al.*, (1991) noticed that male flower were earlier than female flower in several genotypes of bottle gourd, ribbed gourd and sweet gourd. They reported significant variations for that character among the genotypes of bitter gourd, sweet gourd, ribbed gourd and bottle gourd. Significant variation for fruit length and diameter were also observed. They also reported that bitter gourd, sweet gourd, ribbed gourd and bottle gourd genotypes differed significantly for fruit breadth and weight per fruit.

Saha *et al.*, (1992) studied the variability, character association and path analysis of pumpkin and reported that phenotypic coefficient of variation (PCV) was higher than the genotypic coefficient of variation (GCV). High genotypic variance and phenotypic variance were found for fruit length (30.34 and 3176), fruit weight (39.55 and 41.00) and low for fruit diameter (8.87 and 10.23) among the pumpkin genotypes. They also reported high heritability estimate, for both length (91.27) and diameter (75.07) of fruits indicating effectiveness of selection based on good phenotypic performances in pumpkin.

Narayan *et al.*, (1996) conducted genetic variability, heritability in broad sense, genetic advance in 25 diverse populations of bottle gourd. Wide range of variation was observed in most of the characters. The high value of GCV and heritability estimates associated with greater genetic advance was observed for number of primary branches per plant and yield per plant indicated that these two characters had additive gene effect and, therefore, they are more reliable for effective selection. Rumarán *et al.*, (1997) studied 30 pumpkin genotypes in a field trial and reported that genotypic co-efficient of variation was smaller than phenotypic coefficient of variation for most of the traits studied. However, GCV was high for mean fruit weight, number of fruits per plant, number of seeds per fruit, yield per plant and fruit, total soluble solids content. High heritability coupled with high genetic advance were observed for vine length, mean fruit weight, number of fruits per plant, number of seeds per fruit, fruit yield per plant and total soluble solids content of fruits.

Mathew and Khader (1999) conducted an experiment on genetic studies in snake gourd (*Trichosanthen anguina*) and observed the genetic variability and heritability of 12 traits in 34 *Trichosanthen anguina* in Kerela, India and reported that the genotypic co-efficient of variation (GCV) and phenotypic coefficient of variation (PCV) were almost equal for all characters. The highest GCV and PCV were recorded for mean fruit weight, seed per fruit, fruit yield

per plant and fruit length. High heritability was observed for mean fruit weight, seeds per fruit, fruit length, days to first male flower and fruit yield per plant.

Miah *et al.*, (2000) evaluated 30 genotypes of bitter gourd and observed the highest genotypic as well as phenotypic co-efficient of variation were found for fruit length followed by days to female flowering, fruit yield per plant, fruit weight and nodes per vine. Sharma *et al.*, (2000) conducted ten cucumber lines and testers under different environmental conditions and reported that day to first female flower, nodal position of fruits per plant, marketable yield per plant, fruit length and fruit diameter had wide range of variation. Chowdhury and Sarma (2002) studied genetic variation, heritability, genetic advance, and correlation for yield and yield components (vine length, number of nodes, node on which the first flower appeared, number of fruits per plant, fruit length, fruit girth, and fruit weight) were studied in 12 *Luffa acutangula* cultivars (AAUJ-1, AAUJ-2, AAUJ-3, Mangaldoi, Tezpeu, Tihu, Mirza Short, Rangamati Long, Borpeta Long, Tiniali Long, Pusa Nazder, and HRS C-2) grown in Gwahati, Assam, India. The genetic coefficient of variation (GCV) was higher than the phenotypic coefficient of variation (PCV) for all characters. High values of heritability, PCV, GCV, and genetic advance were recorded for vine length, yield per hectare, and fruit weight, indicating that these traits were characterized by additive gene effects. The correlation coefficients revealed that yield per hectare can be improved through selection for greater fruit number per plant, fruit length and girth, and individual fruit weight.

Banik (2003) conducted an experiment on variability and genetic advance of 26 genotypes of snake gourd with respect of 15 quantitative yield contributing characters and found significant difference among the characters like vine length at harvest (2.197 to 3.87 m), number of primary branches (5.23 to 11.88), days to first male flowering (41.67 to 68.67 days), days to first female flowering (48.67 to 71.33 days), node number of first male flower (6.33 to 17.67 days), fruit length (20.67 to 71.17 cm), seeds per fruit (39.03 to 69.50).

Banik also found that significant differences in first female flower, node number (mean value 19.28) and fruits per plant. The highest phenotypic coefficient of variation was observed for fruiting node on main vine, fruit yield per plant, fruit length and first male flower node. The PCV was lowest for days to maturity, 100 seed weight and days to first male flower opening. The GCV along with heritability was high for the above characters. High heritability coupled with high genetic advance was noticed for fruit yield per plant (GCV and PCV 30.75 and 30.96; h^2b 98.64%), fruit length (GCV and PCV 29.92, and 30.04; h^2b 99.19%) and first female flower node number (GCV and PCV 25.87 and 26.59; h^2b 94.63%) and number of fruits per plant (GCV and PCV 19.82 and 20.59; h^2b 92.67%). Dora *et al.*, (2003) eleven pointed gourd (*T. dioica*) selections were assessed to estimate genetic variability and correlation for yield and its attributes. High genetic coefficient of variation (GCV) estimate was observed for the characters such as node at which first female flower appears, length of vine, number of nodes per plant, and number of fruits per plant. The heritability estimate was high for all the characters. The characters having high GCV also exhibited high genetic advance. Yield per plant had a significant positive correlation with number of fruits per plant, fruit set and fruit retention. Banik (2003) studied a field experiment, to study the nature and extent of combining ability of parents and crosses and the mode of gene action in controlling the individual characters in 6×6 diallel including reciprocals in snake gourd. The significant mean sum of squares due to general and specific combining ability (GCA and SCA) for these characters indicated both additive and as well as non-additive type of gene actions were involved for the expression of these characters.

The experiment was conducted at the experimental farm of Bangabandhu Sheikh Mujibur Rahman Agricultural University (BSMRAU), Salna, Gazipur, Bangladesh during May 2001 to September 2002. Narayanankutty *et al.*, (2006) evaluated genetic parameters of 36 snake gourd (*Trichosanthes cucumerina*) genotypes indicated a good amount of genetic variation in the

germplasm collections. Characters such as fruit yield, fruit weight and seeds per fruit exhibited high values of heritability and genetic gain indicating additive gene effects are important in determining these characters. The character association analysis revealed that yield was strongly correlated with fruit weight, fruits per plant, fruit girth, fruit length, days to first harvest, flesh thickness and days to first female flower opening. Fruit weight and fruits per plant have the maximum positive direct effects on yield and the indirect contribution of other characters was mainly through days to first harvest, seeds per fruit and 100 seed weight. Bharathi *et al.*, (2006) assessed genetic variability for 10 characters (days to flowering, vine length, number of nodes on which first flower appears, internode length, fruit length, girth, weight and volume, number of fruits, and yield per plant) in 32 genotypes of spine gourd (*Momordica dioica*) in Bhubaneswar, Orissa, India. Analysis of variance revealed significant differences among the genotypes studied. Phenotypic coefficient of variation (PCV) ranged from 15.26% for fruit girth to 34.28% for fruit weight, while genotypic coefficient of variation (GCV) ranged from 14.38% for fruit girth to 33.52% for fruit weight. High heritability coupled with high genetic advance were recorded for fruit weight, fruit volume and number of fruits per plant, indicating the preponderance of additive gene effects for these characters and their potential use in selection programmes to improve spine gourd productivity. Masud *et al.*, (2006) conducted a field experiment with seven inbred lines and their twenty-one hybrids of bottle gourd. Result showed significant variation in seven characters of the twenty eight populations. Variabilities were high in all seven characters indicating the possibilities of improvement through selection. Specific combining ability variance were significant for all characters while general combining estimates were significant for days to anthesis, fruit length, fruit diameter and yield per plant which indicated the presence of dominance for all the characters but additive is for only few characters. Parent-two showed good GCA for earliness and fruit length, Parent-five showed good GCA for fruit length only and parent-seven showed good GCA for fruit diameter and fruit yield per plant. The

cross involving parent-three and parent- five, which is the best for earliness, fruit length (53.5%) and; fruit yield per plant (106.8%). Rajkumar (2007) *et al.*, studied a field experiment in Tamil Nadu, India, from 2003 to 2005, to determine the genetic variation including the mean, genotypic coefficients of variation (GCV) and phenotypic coefficients of variation (PCV), heritability and genetic advance with 30 genotypes of snake gourd (*Trichosanthes cucumerina*). Significant differences among genotype for all the characters were noted. All the characters exhibited less difference between GCV and PCV values. The characters flesh thickness, fruits per plant, days to fruit maturity and 100-seed weight showed equal GCV and PCV values indicating less influence of environment in their expression. The heritability estimate was high for all the characters except days to first female flower. The maximum heritability was observed for ascorbic acid content of the fruit, followed by the crude fiber content and nodes for first female flower. The genetic advance as a percentage of mean was high for fruits per plant and fruit length. High heritability coupled with high genetic advance was observed for fruits per plant and fruit length. They are governed by additive genes and could be effectively improved through selection.

Quamruzzaman *et al.*, (2009) assessed heterosis in bottle gourd in a set of 13 F₁, with 26 parents. Results indicated highly significant differences for all the character among the materials studied. Heterosis was higher for yield per plant, number of fruits per plant and individual fruit weight, medium in fruit length and fruit diameter, and lower in days to 1st harvest. Hybrids F₁ 10 x 17 and 19 x 26 manifested highest heterosis over mid parent (73.1%) and better parent (61.8%), respectively, for yield per plant. Naik *et al.*, (2012) studied an experiment to study of genetic variability, heritability and genetic advance for fruit quality characters in Teasle gourd. Higher phenotypic coefficients of variation were observed for all the characters except fruit length at marketable stage. Total in mesocarp, total sugar in exocarp, reducing sugar in mesocarp, ascorbic acid in exocarp, ascorbic acid in mesocarp, total soluble solids (TSS)

in exocarp, β -carotene in exocarp, acidity in mesocarp, β -carotene in mesocarp, TSS in mesocarp, acidity in exocarp showed high heritability coupled with high genetic advance indicating that these traits were under the additive gene control and simple selection can be used for further improvement in these traits of teasle gourd. The experiment were carried out at the research field of All India Coordinated Project on Vegetable Crops situated at C Block farm, Bidhan Chandra Krishi Viswa vidyalaya, Nadia during 2007 to 2008.

2.1.1 Leaf Length (cm)

Gaffar (2008) studied an experiment with fifteen genotypes of sponge gourd in Sher-e Bangla Agricultural University. He found that the genotypic and phenotypic variances of leaf length were 24.13 and 25.55, respectively. The GCV (20%) was slightly lower than PCV (20.58%). Heritability for this trait was 97% with moderate genetic advance (9.83) and genetic advance in percent of mean (40.03) was considerable for this trait indicating apparent variation was due to genotypes.

Ahamed *et al.*, (2011) conducted an experiment to assess morphological and yield attributes of pumpkin (*Cucurbita moschata*) in northern area of Bangladesh during kharif season. The range of variability was distinct for leaf length ranged from 30.6-47.2 cm in different genotypes.

2.1.2 Leaf breadth

Asmaul Husna (2009) found GCV (22.87) was lower than PCV (23.04) for this character in bottle gourd. Gaffar (2008) observed GCV (20.94%) was slightly lower than the PCV (23.31%) heritability in broad sense was high (94%) with moderate genetic advance (7.81) for this character in sponge gourd.

2.1.3 Days to first flowering

In Bitter gourd, Mannan (1992) estimated considerable variability among eight lines for days to first male flower (66.7-81.6 days) and female flower (72.80-

85.67 days) opening. Ramchandran and Gopal krishnan (1979) also reported significant variability among 25 diverse genotypes of bitter gourd

Rajkumar (2007) *et al.*, found significant differences among genotype for all the characters in snake gourd. The heritable estimate was high for all the characters except days to first female flower. Banik *et al.*, (2010) found in his experiment the parent P4 was the best general combiner for fruits per plant, first male and female flower. Quamruzzaman *et al.*, (2008) studied experiment the genetic divergence among thirty genotypes of ridge gourd (*Luffa acutangula*) at the farm of Olericulture Division, HRC and in different RARS, BAR] during the summer season of 2005. The genotype RGNO5, RGNO6, RGNO7, RGNO8, RGN 13, RGN 17, RGN 18, RGN27, RGN29 recorded highest cluster mean values for days to 1st male flower open (56.0 days) and single fruit weight (141.0 g) and RGNO3, RGN 12 lowest mean values for days to 1st female flower open (27.0 days) and single fruit weight (85.0 g). The role of days to 1st male flower open, days to 1 female flower open, fruit diameter, single fruit weight and fruit number in PCA indicates their importance in genetic divergence. Sureshbabu (1989) studied 50 genotypes of pumpkin and observed considerable variability for days to first male flower anthesis (41.0-73.0 days) and days to first female flower opening (41.0-84.5 days). Lowest PCV was observed for days to first male flower anthesis (13 .08). Ahamed *et al.*, (2011) found the range of first flowering among twenty genotypes of pumpkin was at 52.0-73.7 days.

2.1.4 Number of male and female flowers per plant

Akter *et al* (2013) studied experiment among thirty accessions of pumpkin genetic divergence at the Research Farm of the Bangabandhu Sheikh Mujibur Rahman Agricultural University (BSMRAU), Salna, Gazipur, during the growing season 2011-12. High genotypic coefficient of variation (GCV) and high heritability coupled with high genetic advance in percent of mean were observed for beta-carotene followed by non-reducing sugar, number of male

flowers per plant and number of female flowers per plant which indicated that these characters were under additive gene control and selection for genetic improvement for these traits might be effective.

2.1.5 Pedicel length of flower (cm)

Asmaul Husna (2009) found that in bottle gourd male flower pedicel length is 3.5-21 cm and in female flower pedicel length is 3.13-9 cm. Rashid (1993) reported that in bottle gourd, male flower pedicel length is longer than female flower pedicel length. Grubben (2004) stated that male flowers have 7-31 cm long pedicel and female flowers have 2-10 cm long pedicel in bottle gourd.

2.1.6 Fruit length and breadth (cm)

Mathew and Khader (1999) noticed the highest GCV and PCV for fruit length in snake gourd. Banik (2003) found high heritability coupled with high genetic advance for fruit length (GCV and PCV 29.92, and 30.04; h^2_b 99.19%) in snake gourd. Rahman *et al.* (1986) indicated high GCV and PCV for both length (31.73 and 33.75) and diameter (39.23 and 41.96) of fruits in bottle gourd. They also observed minimum difference between GCV and PCV. Characters having high GCV indicate high potentiality for effective selection (Burton and de Vane, 1953). Saha *et al.*, (1992) observed high GCV and PCV for fruit length (30.34 and 31.76) and low for fruit diameter (8.87 and 10.23) in pumpkin. They estimated high h^2_b for both length (11.27 %) and diameter (75.07 %). They also found high genetic advance for fruit length (59.72) but low for fruit diameter (15.82)

2.1.7 Fruit weight (Kg)

Correlation studies revealed that highest significant association of yield per plant with reproductive characters number of fruit per plant followed by fruit weight at genotypic and phenotypic level. Path co-efficient analysis revealed maximum direct contribution towards yield per plant with of number of fruit per plant followed by fruit weight. Mathew and Khader (1999) recorded the

highest GCV and PCV were for mean fruit weight. They observed high heritability for mean fruit weight in snake gourd. High GCV and PCV were reported (39.55 and 41.00) by Saha *et al.*, (1992); (30.2 and 36.4) by Doijode and Sulladmath (1986) for fruit weight in pumpkin. Rana *et al.*, (1986) also obtained high value for this trait in pumpkin. Mannan (1992) reported narrow difference between GCV and PCV for this trait in bitter gourd indicating less environmental influence on this character. High h^2 coupled with genetic advance for average fruit weight was noticed in pumpkin (82.9% and 49.6) by Doijode and Sulladmath (1986); (93.03% and 78.58) by Saha *et al.*, (1992). Prasad and Singh (1992) also obtained similar results for this trait in snake gourd and cucumber. On the other hand, low heritability (45.1%) and very high genetic advance (133.05) was recorded for this trait in ribbed gourd by thakur and Choudhury (1965). Vashistha *et al.*, (1983) and Vijay (1987) noticed low GCV and PCV for fruit weight in water melon (0.28 and 0.41) and musk melon (0.01 and 0.02), respectively, whereas Mangal *et al.*, (1981) found high value (291.89 and 318.47) in bitter gourd.

2.1.8 Number of fruits per plant

Rahman *et al.*, (1986) recorded the value of genotypic and phenotypic variances for number of fruits per vine per plant in bottle gourd (1.43 and 3.10), whereas Prasad and Singh (1989), Abusaleha and Dutta (1990), Mangal *et al.*, (1981) reported the value in ribbed gourd (202.26 and 475.98), muskmelon (1.71 and 1.90), cucumber (1.15 and 1.24) and bitter gourd (9.02 and 10.45).

Mathew and Khader (1999) noted the highest GCV and PCV were for fruit yield per plant and fruit length. High heritability was observed for fruit yield per plant in their experiment. Banik (2003) also found that significant differences in fruits per plant. The highest phenotypic co-efficient of variation was observed for fruit yield per plant. High heritability coupled with high genetic advance was noticed for number of fruits per plant (GCV and PCV

19.82 and 20.59; h^2b 92.67%). Akter *et al.*, found that (2013) Correlation coefficient between yield per plant with number of fruits per plant and single fruit weight was positive and highly significant. Path coefficient analysis revealed that the maximum direct contribution towards yield was obtained through number of fruits per plant followed by days to first female flower and single fruit weight indicated that these traits should be considered as primary components of yield.

2.1.9 Yield per plant (kg)

Banik (2003) also studied that significant differences in fruits per plant. The highest phenotypic co-efficient of variation was observed for fruiting node on main vine, fruit yield per plant, fruit length and first male flower node. High heritability coupled with high genetic advance was noticed for fruit yield per plant (GCV and PCV 30.75 and 30.96; h^2b 98.64%). The variation for yield per plant was recorded in bottle gourd (Rahman *et al.* 1991), water melon (Chezhiyan, 1984), musk melon (Swamy *et al.*, 1984) and pumpkin (Rana *et al.*, 1986; Shaha *et al.*, 1992). Mangal *et al.*, (1981) found high value (47759.63 and 55149.80) in bitter gourd while, low GCV and PCV were recorded for this character in water melon (0.44 and 1.15) and musk melon (0.04 and 0.07) by Vashistha *et al.*, (1983) and Vijay (1987). Singh and Prasad (1989) and Saha *et al.*, (1992) recorded high GCV and PCV for yield per plant in pointed gourd (46.50 and 64.10) and pumpkin (28.82 and 31.21). High h^2 associated with high genetic advance for yield per plant was reported by Saha *et al.*, (1992).

Husna *et al.*, (2012) studied Variability, correlation and path analysis among different characters of thirty one bottle gourd genotype. High genotypic co-efficient of variation (GCV) was observed for yield per plant, fruit weight whereas low genotypic co-efficient of variation was observed fruit breadth. Path co-efficient analysis resulted maximum direct contribution towards yield per plant with number of fruit per plant followed by fruit weight.

2.2 Correlation Co-efficient

Singh *et al.*, (1986) yield was significantly correlated with fruits per plant ($r = 0.60$) and days to flowering, days to fruit set and days to ripeness were negatively correlated with all the other characters with the exception of a positive correlation between days to flowering and fruit weight in pointed gourd. Reddy and Rao (1984) observed negative and non-significant correlation between male flower pedicel length, female flower pedicel length traits ($r = 0.222$) in ribbed gourd. Mandal (1987) operated a study on cucumber 30 genotypes and found maximum positive correlation at the genotypic and phenotypic levels between yield per plant with number of fruits and female flowers per plant, fruit length and weight. Abusaleha and Dutta (1989) observed that cucumber yield is positively correlated with vine length ($r = 0.35$), branches per vine ($r = 0.29$), fruits per vine ($r = 0.48$), fruit length ($r = 0.60$) and fruit girth ($r = 0.43$). Days to first male and female flowering, nodal position female flower, percentage of misshapen fruits and non-marketable yield were negatively correlated with yield. Narayan *et al.*, (1996) estimated correlation analysis on 25 genotypes of bottle gourd. Correlation coefficient revealed that fruit yield per plant can be successfully improved by making selection or greater fruit number, higher fruit weight, greater number of primary branches and genotypes with lesser number of days to anthesis of first male flower. Li *et al.*, (1997) recorded that the cucumber genotypes were positively correlated to yield in respect of number of fruits per plant, average fruit per plant, average fruit weight, fruiting rate and leaf area. Days to flowering and vine length were negatively correlated.

Kumaran *et al.*, (1998) conducted an experiment on pumpkin in respect of correlation and path analysis. They found that positive and significant correlation of vine, length, mean fruit weight, number of fruit per plant and number of seeds per fruit with fruit yield per plant. Sarker *et al.*, (1999) found that 16 divergence types of pointed gourd positively and significantly correlated with yield per plant at genotypic and phenotypic levels. Miah *et al.*,

(2000) pointed that fruit yield in bitter gourd showed significant positive association with average fruit weight, fruit breadth and number of nodes per vine in genotypic and phenotypic correlation with days to male flowering. Badade *et al.*, (2001) studied an experiment to find out the correlation of 20 bottle gourd (*Lagenaria vulgaris*) genotypes. Significant and positive correlation of yield was found with the number of branch per vine, number of fruits per vine and negatively correlated with days to first male and female flower appearance and weight of deformed fruits per vine at both phenotypic and genotypic levels. Fruit length showed positive but insignificant correlation with fruit yield. Shah and Kale (2002) studied an experiment on ridge gourd of 55 yield components genotypes in correlation co-efficient analysis. The fruit weight per vine was positively and significantly correlated with number of fruits per vine, average fruit weight, number of female flower per vine and vine length, indicating the close association and dependency of yield these characters. The fruit length was negatively correlated with fruit diameter and fruit number per vine, while it was positively correlated with average fruit weight. Singh *et al.*, (2002) noted out 98 hybrids of cucumber derived from crosses involving fourteen male and seven female parents and found that fruit weight, fruit girth and fruit length had high correlations with fruit yield. Genotypic correlation coefficient were higher than phenotypic co-efficient which indicated strong association among these traits. Prasana *et al.*, (2002) estimated the correlation between the yield and yield components of ridge gourd (*Luffa acutangula*) in Bangalore, Karnataka, India, during the Rabi season of 1999. Fruit yield per hectare was positively associated with vine length at 90 days after sowing (DAS), number of leaves at 90 DAS, number of female flowers, total dry weight of plant, number of fruits, and fruit girth and weight. Hazra *et al.*, (2003) conducted sixty-eight diverse female clones of pointed gourd (*Trichosanthes dioica*). These were grown at the Horticultural Research Station, Mondouri, West Bengal, India to evaluate growth, morphological, yield and quality characters and their relationship through correlation and path analysis. The magnitude of genotypic correlation

coefficients was higher than phenotypic correlation coefficients for all the pairs of characters, and in most cases, a wide gap was recorded between the two estimates of correlation coefficients, indicating the influence of environment on the correlated response of the pair of characters. Most of the character pairs showed negligible or insignificant correlation that might have resulted due to simultaneous vegetative and reproductive growth in the plant. Only fruit number per plant had significant positive correlation with yield, whereas fruit weight showed highest positive direct effect on yield. However, from the overall study most of the fruit characters, viz. fruit weight, pulp content of fruit, fruit number per plant and fruit volume, and growth traits, such as leaves per plant and leaf length, were identified as important yield contributors. Singh and Ram (2003) conducted an experiment on 28 musk melon genotypes to determine the correlation among fruit characters. The simple correlation among fruit traits showed that polar diameter, latitudinal diameter, flesh thickness and seed cavity size were positively correlated with fruit weight. Kumaresan *et al.*, (2006) conducted field, experiments in Madurai, Tamil Nadu, India, during the 2000 rabi season, to determine correlations among different economic parameters and their direct and indirect effects on fruit yield in 6 snake gourd (*Trichosanthes cucumerina*) cultivars and their 30 hybrids. Yield per vine in snake gourd was positively associated with main vine length, number of fruits per vine, fruit weight, number of seeds per fruit, seed weight per fruit and ascorbic acid content of the fruits. However, negative association was observed with days to first female flower opening, days to first male flower opening, fruit length, fruit girth and acid content of the fruit. This indicated that the selection for the characters would simultaneously result in improving the yield per vine. Kumar *et al.*, (2007) studied an experiment to study the correlation coefficient of 20 bottle gourd (*Lagenaria vulgaris*) genotypes. Fruit yield per vine in bottle gourd is the result of interaction of number of inter-related characters. Therefore, selection should be based on these components character after assessing their correlation with fruit yield per vine. The fruit yield per vine showed positive and significant correlation with number of branches per

vine length, nodes number of first male flower, nodes number of first female flower, length of edible fruits, and number of fruits per vine, number of seeds per fruits and 100-seed weight at genotypic and phenotypic levels. This indicated that fruit yield can be improved by making selection on the basis of no. of branches per vine, vine length, nodes no. of first female flower, length of edible fruit and no. of fruit per vine. Khule *et al.*, (2011) conducted field experiments to determine correlation and path coefficient analysis in sponge gourd *Luffa cylindrica* (Linn.) at Vegetable Research Station Jagudan, (Gujarat) with thirty sponge gourd genotypes. They found genotypic correlation coefficients were higher than the corresponding phenotypic correlation coefficients suggesting that the environmental influence reduces the relationship between yield and yield contributing characters of sponge gourd. Path coefficient analysis showed that number of fruits per plant, days to appear first female flower, fruit length, fruit diameter, and number of seeds per fruit had direct positive effects on marketable fruit yield per plant which indicates that this character was the major contributor to fruit yield.

2.3 Path Co-efficient

Rahman *et al.*, (1986) studied variability, correlation and path coefficients in four lines of bottle gourd. Path coefficient analysis revealed that fruit diameter and fruit length had high positive direct effect on fruit weight per plant. Number of fruits per plant also had considerable positive direct effect on fruit weight per plant.

Chaudhury and Mandal (1987) conducted a study on 30 diverse cucumber genotypes and Path co-efficient analysis revealed that the number of fruits, female flowers per plant, fruit length, fruit weight and fruit diameter were the most important characters determining yield. Mondal *et al.*, (1989) studied path co-efficient in 31 genotypes of watermelon and observed that the number of fruits per plant and fruit diameter affected fruit yield directly. Path co-efficient analysis revealed that for increasing fruit yield selection should be based on plant having more number of fruits with larger diameter. Abusaleha and Dutta

(1989) carried out an experiment on correlation and path analysis studies in cucumber. Path coefficient analysis revealed that fruits per vine and fruit length had the greatest direct effects on yield. Parhi *et al.*, (1995) studied correlation and path co-efficient of thirteen genotypes of bitter gourd. Path analysis revealed that fruit breadth, days to opening of first male and female flower, vine length and number of seeds per fruit had the maximum positive direct effect on yield in bitter gourd. The characters like fruit weight and fruit length though have significant positive correlation with yield, exhibited low direct effect. Besides direct selection for yield, indirect selection through number of seeds per fruit and fruit weight would prove worth for further improvement in yield of bitter gourd. Kumaran *et al.*, (1998) carried out an experiment on correlation and path analysis studies in pumpkin. They found that number of fruit per plant exhibited the highest direct effect on yield. High positive indirect effects were exerted by number of fruit per plant and mean fruit weight. Sarker *et al.*, (1999) studied path co-efficient of 16 divergence types of pointed gourd. The path analysis revealed that fruit volume followed by fruit weight and fruit diameter had maximum positive direct effects on yield. Li *et al.*, (1997) conducted an experiment on cucumber genotypes. From path analysis, they concluded that fruits per plant and average fruit weight affected the yield directly. Rao *et al.*, (2000) conducted an experiment on the segregating population of ridge gourd for correlation and path coefficient analysis. Path analysis revealed that yield improvement could be achieved by direct selection for days to 50% flowering, girth of fruit, fruits per plant or vine, fruit per branch and length of the vine of ridge gourd. Miah *et al.*, (2000) conducted an experiment on bitter gourd for correlation and path coefficient analysis. Path analysis revealed that average fruit weight, number of fruits per plant, days to male flowering and fruit length had positive direct effect on fruit yield. Singh *et al.*, (2002) were noted out 98 hybrids of cucumber derived from crosses involving fourteen male and seven female parents. Path coefficient analysis indicated that fruit weight had the highest direct effect on fruit yield. Rao *et al.*, (2000) conducted an experiment on the segregating population of ridge gourd

for correlation and path coefficient analysis. Path analysis revealed that yield improvement could be achieved by direct selection for days to 50% flowering, girth of fruit, fruits per plant or vine, fruit per branch and length of the vine of ridge gourd. Miah *et al.*, (2000) conducted an experiment on bitter gourd for correlation and path coefficient analysis. Path analysis revealed that average fruit weight, number of fruits per plant, days to male flowering and fruit length had positive direct effect on fruit yield. Prasanna *et al.*, (2002) studied the correlation between the yield and yield components of ridge gourd (*Luffa acutangula*) in Bangalore, Karnataka, India, during the rabi of 1999. Fruit yield per hectare was positively associated with vine length at 90 days after sowing (DAS), number of leaves at 90 DAS, number of female flowers, total dry weight of plant, number of fruits, and fruit girth and weight. Path coefficient analysis showed that vine length at 90 DAS, number of female flowers per vine, number of branches per vine, number of fruits per vine, fruit girth, and fruit weight had direct positive effects on fruit yield, whereas the number of leaves at 90 DAS, total dry weight of the plant, and fruit length had negative direct effects on fruit yield. The fruit yield of ridge gourd can be enhanced through the improvement of vine length at 90 DAS, number of female flowers, number of branches, number of fruits per vine, fruit girth, and fruit weight. Umamaheswarappa *et al.*, (2004) conducted an experiment on the effect of various rates of nitrogen (0, 60 and 120 kg/ha), phosphorus (0, 50 and 100 kg/ha) and potassium (0, 30 and 60 kg/ha) on bottle gourd (*Lagenaria siceraria*), conducted in Bangalore, Karnataka, India, in 1999 showed that fruit yield/ha had strong positive association with vine length, number of leaves per vine, number of female flowers per vine, number of branches per vine, vine girth, total chlorophyll content in leaf, total dry weight of plant, number of fruits per vine, fruit weight, fruit length and fruit girth. Path coefficient analysis revealed that number of fruits per vine had maximum direct effect on fruit yield followed by fruit weight. Kumaresan *et al.*, (2006) conducted field experiments in Madurai, Tamil Nadu, India, during the 2000 rabi season, to determine correlations among different economic parameters and their direct and indirect

effects on fruit yield in 6 snake gourd (*Trichosanthes cucumerina*) cultivars and their 30 hybrids. Path coefficient analysis revealed that it would be highly rewarding to lay emphasis on the number of fruits per vine and fruit weight to increase the yield per vine directly. Kumar *et al.*, (2007) conducted an experiment to study the path coefficient of 20 bottle gourd (*Lagenaria vulgaris*) genotypes. Path analysis revealed that number of branches per vine, vine length, nodes number of first female flower and number of fruit per vine had positive direct effect on fruit yield per vine. Narayan *et al.*, (1996) studied path-coefficient analysis in 25 diverse populations of bottle gourd. Path coefficient analysis revealed that maximum weight age should be given primarily to days to first harvest followed by average weight of edible fruit, number of fruits per plant and days to anthesis of first female flower while formulating selection indices for improvement of yield in bottle gourd.

2.4 Genetic Diversity

For the self and cross-pollinated crops Genetic diversity is very important elements (Griffing and Lidstorm, 1954; Murty and Arunachalam, 1966; Guar *et al.*, 1978). Being a successful hybridization programme genetically diverse plants selected by the quantification of genetic diversity in the biometrical procedure (Rao, 1952). D^2 analysis (originally outlined by Mahalanobis, 1936 and extended by Rao, 1952) is one of potential methods of estimating the degree of genetic diversity. Genotypic diversity can be shown by the cluster analysis from same geographical regions. To understand the usable variability, grouping or classification of genotypes based on suitable scale. Multivariate analysis formulated by Mahalanobis (1936) is a powerful tool in quantifying the degree of divergence among biological population based on multiple characters. Studies on genetic diversity in bottle gourd carried out so far are presented as follows: Masud *et al.*, (1995) conducted an experiment among 27 genotypes of pumpkin (*Cucurbita moschata*) to study the genetic divergence collected from eight districts of Bangladesh was grouped into seven clusters. No relationship was found between genetic divergence and geographic distribution

of the genotypes. Maximum inter cluster distance was observed between cluster II and VII and was minimum between V and VI. Number of fruits per plant and yield per plant showed maximum contribution to the total divergence. The results obtained by D^2 analysis were confirmed by principal component analysis. According to Dora (2001) eleven genotypes of *Trihosanthes dioica* and the genotypes were grouped into four clusters based on Mahalanobis's D^2 statistics and found that inter cluster distances were greater than intra cluster distances, indicating considerable genetic diversity among genotypes. The highest D^2 value (984.3) was recorded between cluster II and IV. Genetic divergence using Mahalanobis D^2 statistics was studied for seven quantitative characters including yield per vine in a collection of twenty diverse cultivars of bottle gourd by Badade *et al.*, (2001). The cultivars differed significantly for almost all of the characters and were grouped into 10 clusters based on the similarities of D^2 value. Considerable diversity within and between clusters was noted and it was observed for the characters viz. vine length, number of branches, fruit per vine, length and diameter of fruit and yield per vine. Banik (2003) carried out an experiment about 26 genotypes of snake gourd using multivariate analysis and the genotypes were grouped into seven distinct cluster. No relationship was found between genetic divergence and geographical distribution of genotypes. The highest inter genotypes distance was observed between genotypes SO 026 and SO 010 (1.897). The inter cluster distance was maximum between cluster II and IV (17.74). Main vine length, first female flower node number, nodes on main vine, fruit length and number of seeds per fruit had the highest contribution towards the divergence. Harshawardhan and Ram (2003) conducted an experiment for yield and its components on severity germplasms of musk melon lines to elucidate genetic divergence. The genotypes were grouped into 11 clusters irrespective of geographic and genetic diversity. Group VIII contained the largest number of 11 genotypes. The maximum genetic distance occurred between cluster II and X. Islam (2004) evaluated genetic divergence among 42 bottle gourd (*L. siceraria*) accessions from Bangladesh was estimated in Japan during 2000

using D² and canonical analysis. The accessions were grouped into five clusters. No clear relationship was observed between geographic origin and genetic diversity. The maximum inter cluster distance was between clusters I and cluster II, and the minimum was between cluster III and cluster IV. Primary branches per plant, fruit length and weight, number of fruits and yield per plant contributed the most to the total genetic divergence. The results obtained by D² analysis were also confirmed by canonical analysis. The accessions included in the most divergent clusters I and II, are promising parents for a hybridization programme for obtaining high heterosis and thus, better segregates in bottle gourd. Bharathi *et al.*, (2005) The genetic divergence among 32 genotypes of spine gourd (*Momordica dioica*) for 12 traits (vine length, number of days to flowering, node on which the first flower appeared, internode length, mature leaf size, pedicel length, petiole length, fruit weight, fruit length, fruit girth, number of fruits per plant, and yield per plant) was evaluated in Orissa, India. The analysis of variance revealed significant variation among the genotypes for all traits. The genotypes were grouped into 7 clusters based on D² values. Cluster III had the highest number of genotypes (11), followed by clusters IV (9) and VI (4). The intra cluster distance ranged from 30.34 (cluster I) to 371.56 (cluster III). The inter cluster distance was greatest between clusters VI and VII (864.75). Genotypes included in cluster II were characterized by early flowering, and presence of the longest vines and internodes. Cluster VI recorded the greatest number of fruits, pedicel length and yield. Cluster VII was superior with regard to the node on which the first flower appeared. Cluster III had the greatest fruit weight, fruit length and fruit girth. Yield per plant, number of fruits, fruit weight, internode length, fruit length and pedicel length accounted for 93.5 5% of the diversity. Thus, selection for divergent parents based on these traits is recommended. Karuppaiah *et al.*, (2005) evaluated genetic divergence in 12 genotypes of bitter gourd (*Momordica. charantia*) grown in Annamalai, Tamil Nadu, India, during June-July 2001. Using Mahalanobis D² technique, the genotypes were grouped into clusters I (4 genotypes), II (one genotype), III (3 genotypes) and

IV (four genotypes). Among the four clusters, cluster IV (LA-7, LA-9, LA- 10 and LA-12) registered the highest mean values for vine length (6.2 m), number of male flowers per plant (79.3), number of female flowers per plant (23.2), yield per plant (5.2 kg), single fruit weight (242.2 g), fruit length (29.4 cm), number of fruits per plant (24.1), number of seeds per fruit (52.3), fruit size index (173.2), and 100-seed weight (18.6 g). Hence, it is desirable to involve LA-7, LA-9, La- 10 and LA- 12 of cluster IV in breeding programmes. Kabir (2007) reported that genetic divergence studied 24 accessions of pointed gourd. The accessions were grouped into five clusters. The cluster I and III had the highest number of accessions (6) followed by cluster V (5), cluster 11(4) & Cluster IV (3). The highest intra cluster distance was computed for cluster IV (35.80) followed by cluster I (28.12) and Cluster V (26.63). The minimum intra cluster distance was found in III (18.87).’ Quamruzzaman *et al.*, (2008) studied the genetic divergence among thirty genotypes of ridge gourd (*Luffa acutangula*) using D^2 and principal component analysis. The genotypes were grouped into six clusters. The highest intra cluster distance was noticed for the cluster II (0.882) and the lowest for the cluster III (0.220). The highest inter-cluster distance was observed between cluster I and II (15.045) whereas the lowest was observed between cluster IV and V (3.402). Gaffar (2008) conducted an experiment with 15 sponge gourd genotypes at the experimental farm of Sher-e-Bangla Agricultural University, during April, 2007 to October 2007. The genotypes were grouped into five clusters. The highest intra cluster distance was noticed for the cluster III (0.999) and the lowest for the cluster IV (0.43 9). The highest inter-cluster distance was observed between cluster IV and V (7.163) whereas the lowest was observed between cluster I and IV (2.258). Khan *et al.*, (2008) assessed the genetic diversity among 64 pointed gourd genotypes through multivariate analysis from an experiment conducted in Regional Agricultural Research Station, Ishurdi, Pabna during the growing season 2002-2003. The genotypes were grouped into twelve clusters. The cluster V consisted of highest number of genotypes and it was nine, the cluster VI and cluster VIII contained the lowest number of genotypes and it was two in

each. The clustering pattern of the genotypes under this study revealed that the genotypes collected from the same location were grouped into different clusters. The genotypes of Jessore were distributed in different clusters. The highest inter genotype distance as 366.3 observed between the genotypes P0022 and P0007 and the lowest 2.6 as observed between the genotypes P0043 and P0044. Cluster V had the highest cluster mean value for internodes length, fruit weight per plant and yield. the highest inter-cluster distance was noticed between cluster III and II (45.71) and the lowest between cluster VII and VI (3.33). The highest intra cluster distance was computed for cluster III and that was lowest for the cluster II. The first five axes accounted for 77.65% of the total variation among the 13 characters describing 64 pointed gourd genotypes. Fruit weight, seeds per fruit and fruit weight per plant contributed maximum to the total divergence. BARI annual report 2008-09 revealed that Genetic divergence among 30 snake gourd genotypes was estimated using Mahalanobis's D^2 statistic. Cluster V contained the highest number of genotypes (13) and cluster III & IV contained the lowest (3). The highest intra- cluster distance was observed in cluster III (1.665) and the lowest in cluster V (0.430). The highest inter- cluster distance was observed between cluster I and III (26.954) and the lowest in cluster II and I (5.693). BARI annual report 2008-09 revealed that Genetic divergence among 30 snake Khatun *et al.*, (2010) conducted at the field and laboratory of the Department of Horticulture, Bangladesh Agricultural University, Mymensingh during the period from April 2004 to September 2004 to study the nature and magnitude of genetic diversity of 38 snake gourd genotypes collected from different regions of the country. Based on D2 analysis, the genotypes were grouped into four different clusters, where the cluster I possessed maximum number (21) of genotypes followed by the cluster II (8), III (7), and IV (2). Clustering pattern revealed that geographical diversity was not associated with genetic diversity i.e., genotypes collected from same location were grouped into different clusters. The maximum inter-cluster distance was observed between the clusters III and IV and that of minimum in between the clusters I and II. In case of intra-cluster

distance, the maximum distance was observed in the cluster IV and that of minimum was observed in the cluster III. Considering cluster mean, the genotypes of cluster IV could be selected for yield per plant and other yield contributing characters. Islam *et al.*, (2010) studied genetic divergence of twenty bitter gourd genotypes through Moahalanobis's D^2 and principal component analysis in Pakistan. The genotypes under study fall into four clusters. The cluster I contained the highest number of genotypes and it was 10. Cluster IV contained the lowest number of genotypes. Cluster II produced the highest mean value for weight per fruit. The inter cluster distances were much higher than the intra cluster distances. Cluster I exhibit the highest intra cluster distance while the lowest distance was observed in cluster III. The highest inter cluster distance was observed between I and H while the lowest distance was observed between the cluster II and IV. The highest intra cluster means for weight per fruit and five important yield contributing characters were obtained from cluster II. Therefore, more emphasis should be given on the cluster for selecting genotypes as parents for crossing with the genotypes of cluster II which may produce new recombination with desired traits. Considering all the characters the 01 (Shaparan), G₅, (Rampaligaj), G₉ (Nabil), G₁₂ (Nandita) G₁₄ (Eureca), G₁₆ (Tia) and G₁₉ (Maharaj) were selected for future breeding programme. Preeti *et al.*, (2010) observed wide range of genetic diversity among twenty three germplasm lines of ash gourd collected from different parts of U.P. and Uttarakhand. Genotypes PAG-50, Pant Petha-1, PAG-64, PAG-12, PAG-14 and PAG-09 were high yielding lines while considering both the season's summer and kharif 2006. Based on Mahalanobis D^2 analysis all germplasm lines were grouped into 5 clusters. The clustering pattern indicated that geographical distribution need not necessarily be related to the genetic diversity. Cluster I was very large containing 14 genotypes (summer) and 10 genotypes (kharif) season. The commercially released cultivar Pant Petha-1 was grouped in cluster II along with other genotypes in both the seasons. The inter-cluster distance was found maximum between cluster III and cluster IV (summer) and cluster II and cluster V in Kharif seasons.

CHAPTER III

MATERIALS AND METHODS

The research programme was carried out at the experimental field of Sher-e-Bangla Agricultural University, Dhaka, Bangladesh during the period from March 2016 to July, 2016 to study on evaluation of pumpkin genotypes based on phenotypic traits. In this section a brief description about the locations of the experimental site, characteristics of soil, climate, materials, layout and design of the experiment are given whereas land preparation, manuring and fertilizing, transplanting of seedlings, intercultural operations, harvesting, data recording procedure, economic and statistical analysis etc. are presented as follows:

3.1. Experimental site

The present experiment was conducted at the experimental farm of Sher-e-Bangla Agricultural University, Dhaka-1207.

3.2 Geographical location

The research area was situated at 23°74'N latitude and 90°35'E longitude at an altitude of 8.6 meter above the sea level (Anon., 2004). The experimental field belongs to the Agro-ecological zone of The Modhupur Tract, AEZ-28. This was a region of complex relief and soils developed over the Modhupur clay, where floodplain sediments buried the dissected edges of the Modhupur Tract leaving small hillocks of red soils as 'Islands' surrounded by floodplain. The experimental site was shown in the map of AEZ of Bangladesh in (Appendix D).

3.3 Climate

Area has subtropical climate, characterized by high temperature, high relative humidity and heavy rainfall in Kharif season (March-July) and scanty rainfall associated with moderately low temperature during the Kharif season (March-

July). Meteorological information regarding temperature, relative humidity, rainfall and sunshine hours prevailed at the experimental site during the study period was presented in Appendix II.

3.4 Characteristics of soil

Soil of the experimental site belongs to the general soil type, Shallow Red Brown Terrace Soils under Tejgaon Series. Top soils were clay loam in texture, olive-gray with common fine to medium distinct dark yellowish brown mottles. Soil pH ranged from 5.47 to 5.63, organic matter 0.82%. Experimental area was flat having available irrigation and drainage system and above flood level. Soil samples from 0-15 cm depths were collected from experimental field. The analyses were done by Soil Resource and Development Institute (SRDI), Dhaka. Physicochemical properties of the soil are presented in (Appendix III).

3.5 Planting materials

Twenty four genotypes of pumpkin were used as experimental materials for the present research work. The purity and germination percentage were leveled as around 100 and 80, respectively. The genetically pure and physically healthy seeds of these genotypes were collected from Plant Genetic Resources Centre (PGRC) of Bangladesh Agricultural Research Institute (BARI) Gazipur. The name and origin of these genotypes are presented in (Table 1).

3.6 Design and layout of the experiment

The experiment was laid out Randomized Complete Block Design (RCBD) with three replications. The genotypes were distributed into the pit of each block of the prepared layout of the experiment. The twenty four genotypes of the experiment were assigned at random into pits of each replication. The distance maintained spacing pit to pit 1 m.

Table 1. List of twenty four pumpkin genotypes with their accession and source

Sl. No.	Genotype NO.	BARI ACC Number	Source
1	G ₁	(BD 10149)	PGRC,BARI
2	G ₂	(BD 10141)	PGRC,BARI
3	G ₃	(BD 4361)	PGRC,BARI
4	G ₄	(BD 4353)	PGRC,BARI
5	G ₅	(BD 10137)	PGRC,BARI
6	G ₆	(BD 10058)	PGRC,BARI
7	G ₇	(BD 4349)	PGRC,BARI
8	G ₈	PRONROY-F1	AGRO SERVICE Pvt.Ltd
9	G ₉	(BD 4371)	PGRC,BARI
10	G ₁₀	(BD 4357)	PGRC,BARI
11	G ₁₁	(BD 4350)	PGRC,BARI
12	G ₁₂	(BD 4352)	PGRC,BARI
13	G ₁₃	(BD 10078)	PGRC,BARI
14	G ₁₄	(BD 10366)	PGRC,BARI
15	G ₁₅	(BD 4363)	PGRC,BARI
16	G ₁₆	BARMASI	AGRO SERVICE Pvt.Ltd
17	G ₁₇	(BD 4372)	PGRC,BARI
18	G ₁₈	(BD 4391)	PGRC,BARI
19	G ₁₉	(BD 4389)	PGRC,BARI
20	G ₂₀	(BD 4354)	PGRC,BARI
21	G ₂₁	(BD 4393)	PGRC,BARI
22	G ₂₂	(BD 4370)	PGRC,BARI
23	G ₂₃	(BD 10134)	PGRC,BARI
24	G ₂₄	PRITY-F1	AGRO SERVICE Pvt.Ltd

Here, PGRC = Plant Genetic Resources Centre, BARI = Bangladesh Agricultural Research Institute, Agro Service Pvt.Ltd

3.7 Poly bag preparation and raising seedling

Due to uncertain rainfall during the period of the study, the seeds were dibbled in poly bag for higher germination percentage and to get healthy seedlings. When the seedlings become 12 days old those were transplanted in the main field in the pit. Seeds were sown 28th March, 2016 before sowing seeds were treated with Bavistin for 5 minutes. Plate 1 showing raising of seedling in polybag.

3.8 Land preparation

The experiment plot was prepared by several ploughing and cross ploughing followed by laddering and harrowing with tractor and power tiller to bring about good tilth in the first week of March, 2016. Weeds and other stables were removed carefully from the experimental plot and leveled properly.

3.9 Pit preparation

After final land preparation, pits of 55 cm x 55 cm x 50 cm were prepared in each block with spacing of 3 m x 3 m. Pits were kept open in the sun for 7 days to kill harmful insect and microorganisms. To control field cricket 5 mg Furadan was also mixed with the soils of each pit before making it ready for dibbling.

3.10 Application of manure and fertilizers

Total cowdung, half of TSP and one third MOP were applied in the field during final land preparation Remaining TSP and one third MOP and whole gypsum and zinc oxide and one third of urea were applied in pit one week prior to transplantation Remaining urea and MOP were applied as top dressing in four instalments at 20, 40, 60 and 75 days after transplanting Doses of manure and 11 fertilizers used in the study are shown in Table 2.



Plate 1. Raising of seedling in polybag

Table 2. Doses of manure and fertilizers used in the study

SL No.	Fertilizers/Manures	Dose
1	Cowdung	10 ton/ha
2	Urea	125 kg/ha
3	TSP	125 kg/ha
4	MOP	150 kg/ha
5	Gypsum	75 kg/ha
6	Zinc Oxide	10 kg/ha

3.11 Transplanting of seedlings

Within 10 days germination of seeds was completed and the seedlings of different accessions were planted in the pit on 09 April, 2016. In each pit two seedlings were planted and the soil around the plant was firmly pressed by hand. Field view of plants after transplanting of seedling is presented in Appendix vi.

3.12 Intercultural operations

3.12.1 Thinning and gap filling

Only one healthy seedling was kept per pit for the proper development and for avoiding crowd environment. For this whenever need thinning and gap filling was done.

3.12.2 Weeding and mulching

Several weeding and mulching were done as per requirement. At the very first stage, weeding was done for ease of aeration and less competition seedling growth and mulch was provided after an irrigation to prevent crust formation and facilitate good aeration.

3.12.3 Irrigation and after-care

In the early stage irrigation was done twice daily by water cane. In mature stage flood irrigation was done when ever it's necessary.

3.12.4 Pesticide application

At the seedling stage red pumpkin beetle attacked tender leaves for this Malathion and Ripcord was sprayed in the field. In mature stage cucurbit fruit fly caused, severe damage to the fruit. For a protection from fruit fly, MSGT, (Mashed Sweet Gourd Trap) and Pheromone bait was used along with ripcord, sevin powders.

3.13 Harvesting

The fruit takes about 7-10 days from setting to reach marketable stage. Fruits were picked on the basis of horticultural maturity, size, color and age being determined for the purpose of consumption as the fruit grew rapidly and soon get beyond the marketable stage, frequent picking was done throughout the harvesting period. Fruits were picked with sharp knife and care was taken to avoid injury of the vine.

3.14 Data recording

Data were recorded on following parameters from the studied plants during the experiment. The details of data recording are given below on individual plant basis.

3.14.1 Plant characteristics

3.14.1.1 Leaf length (cm)

Leaf length was measured in three to five leaves in each germplasm in cm and average data was recorded.

3.14.1.2 Leaf breadth (cm)

Leaf breadth was measured in three to five leaves in each germplasm in cm and average data was recorded.

3.14.1.3 Internodes distance (cm)

Internodes distance was measured in three to five Internodes in each germplasm in cm and average data was recorded.

3.14.2 Flower characteristics

3.14.2.1 Days to first male flowering

The number of days required for first male flower flowering was counted for three replications separately and average data was recorded.

3.14.2.2 Days to first female flowering

The number of days required for first female flower flowering was counted for three replications separately and average data was recorded.

3.14.2.3 Pedicel length of male flower (cm)

Pedicel length of male flower was measured in three to five flowers in each germplasm in cm and average data was recorded.

3.14.2.4 Pedicel length of female flower (cm)

Pedicel length of female flower was measured in three to five flowers in each germplasm in cm and average data was recorded.

3.14.3 Fruit characteristics

3.14.3.1 Fruit length (cm)

Fruit length was measured in three to five fruits in each germplasm in cm and average data was recorded during fruit harvest for vegetable use.

114.3.2 Fruit breadth (cm)

Fruit diameter was measured in three to five fruits in each germplasm in cm, then the data was divided by two and average data was recorded during fruit harvest for vegetable use.

3.14.3.3 Fruit weight (kg)

Weight of three to five fruits in each germplasm during harvest for vegetable use was measured in kilogram.

3.14.3.4 Fruit yield per plant (kg)

Weight of edible fruits of selected plants from each accession was weighed in kilogram (kg).

3.15.1 Statistical analysis

Mean data of the characters were subjected to multivariate analysis. Univariate analysis of the individual character was done for all characters under study using the mean values (Singli and Chaudhury, 1985) and was estimated using MSTAT-C computer program. Duncan's Multiple Range Test (DMRT) was performed for all the characters to test the differences between the means of the genotypes. Mean, range and co-efficient of variation (CV %) were also estimated using MSTAT-C. Multivariate analysis was done by computer using GENSTAT 5.13 and Microsoft Excel 2000 software through four V techniques viz., Principal Component Analysis (PCA), Principal Coordinate Analysis (PCO), Cluster Analysis (CA) and Canonical Vector Analysis (CVA).

3.15.1.1 Estimation of genotypic and phenotypic variances

Genotypic and phenotypic variances were estimated according to the formula given by Johnson *et al.*, (1955).

$$\text{Genotypic variance } (\sigma^2_g) = \frac{\text{GMS} - \text{EMS}}{r}$$

Where,

GMS = Genotypic mean sum of squares

EMS = Error mean sum of square

r = number of replications

$$\text{Phenotypic variance } (\sigma^2_p) = \sigma^2_g + \text{EMS}$$

Where,

σ^2_g = Genotypic variance

EMS = Error mean sum of square

3. 15.1.2 Estimation of genotypic and phenotypic correlation co-efficient

For calculating the genotypic and phenotypic correlation co-efficient for all possible combinations the formula suggested by Miller *et al.*, (1958), Johnson *et al.*, (1955) and Hanson *et al.*, (1956) were adopted. The genotypic co-variance component between two traits and have the phenotypic covariance component were derived in the same way as for the corresponding variance

components. The co-variance components were used to compute genotypic and phenotypic correlation between the pairs of characters as follows:

$$\text{Genotypic correlation } (r_{gxy}) = \frac{\sigma_{gxy}}{\sqrt{\sigma^2_{gx}\sigma^2_{gy}}}$$

Where,

r_{gxy} = Genotypic co-variance between the traits x and y

σ^2_{gx} = Genotypic variance of the trait x

σ^2_{gy} = Genotypic variance of the trait y

$$\text{Phenotypic correlation } (r_{pxy}) = \frac{\sigma_{pxy}}{\sqrt{\sigma^2_{px}\sigma^2_{py}}}$$

Where,

σ_{pxy} = Phenotypic covariance between the traits x and y

σ^2_{px} = Phenotypic variance of the trait x

σ^2_{py} = Phenotypic variance of the trait y

3.15.1.3 Estimation of genotypic and phenotypic co-efficient of variation

Genotypic and phenotypic co-efficient of variation were calculated by the formula suggested by Burton (1952)

$$\text{Genotypic co-efficient of variation (GCV \%)} = \sqrt{\frac{\sigma^2_g}{\bar{x}}} \times 100$$

Where,

σ^2_g = Genotypic variance

\bar{x} = Population mean similarly,

The phenotypic co-efficient of variation was calculated from the following

$$\text{Phenotypic co-efficient variation (PCV)} = \sqrt{\frac{\sigma^2_{ph}}{\bar{x}}} \times 100$$

Where,

σ^2_{ph} = Phenotypic variance

\bar{x} = Population mean

3. 15.1.4 Estimation of heritability

Broad sense heritability was estimated (Lush, 1943) by the following formula, suggested by Johnson *et al.*, (1955).

$$h^2_b \% = \frac{\sigma^2_g}{\sigma^2_{ph}} \times 100$$

Where,

h^2_b = Heritability in broad sense

σ^2_g = Genotypic variance

σ^2_{ph} = Phenotypic variance

3.15.1.5 Estimation of genetic advance

The expected genetic advance for different characters under selection was estimated using the formula suggested by Lush (1943) and Johnson *et al.*, (1955).

$$\text{Genetic advance (GA)} = K \cdot h^2_b \cdot \sigma_{ph}$$

$$GA = K \cdot \frac{\sigma^2_g}{\sigma^2_{ph}} \cdot \sigma_{ph}$$

Where,

K = Selection intensity, the value which is 2.06 at 5%

Selection intensity

σ_{ph} = Phenotypic standard deviation

h^2_b = Heritability in broad sense

σ^2_g = Genotypic variance

σ^2_{ph} = Phenotypic variance

3.15.1.6 Estimation of genetic advance mean's percentage

Genetic advance as percentage of mean was calculated from the following formula as proposed by Comstock and Robinson (1952):

$$\text{Genetic advance (\% of mean)} = \frac{\text{Genetic advance (GA)}}{\text{Population mean } (\bar{x})} \times 100$$

3.15.1.7 Estimation of path co-efficient

Path coefficient analysis was done according to the procedure employed by Dewey and Lu (1959) quoted in Singh and Chaudhary (1985), using phenotypic correlation coefficient value. In path analysis, correlation coefficients between yield and yield contributing characters were partitioned into direct and indirect effects of yield contributing characters on grain yield per hectare. In order to estimate direct and indirect effects of the correlated characters, i. e. 1, 2, 3..... and 13 on yield y, a set of simultaneous equations (eight equations in this example) is required to be formulated as shown in below:

$$r_{1.y} = P_{1.y} + r_{1.2} P_{2.y} + r_{1.3} P_{3.y} + r_{1.4} P_{4.y} + r_{1.5} P_{5.y} + r_{1.6} P_{6.y} + r_{1.7} P_{7.y} + r_{1.8} P_{8.y} + r_{1.9} P_{9.y} + r_{1.10} P_{10.y} + r_{1.11} P_{11.y} + r_{1.12} P_{12.y}$$

$$r_{2.y} = r_{1.2} P_{1.y} + P_{2.y} + r_{2.3} P_{3.y} + r_{2.4} P_{4.y} + r_{2.5} P_{5.y} + r_{2.6} P_{6.y} + r_{2.7} P_{7.y} + r_{2.8} P_{8.y} + r_{2.9} P_{9.y} + r_{2.10} P_{10.y} + r_{2.11} P_{11.y} + r_{2.12} P_{12.y}$$

$$r_{3.y} = r_{1.3} P_{1.y} + r_{2.3} P_{2.y} + P_{3.y} + r_{3.4} P_{4.y} + r_{3.5} P_{5.y} + r_{3.6} P_{6.y} + r_{3.7} P_{7.y} + r_{3.8} P_{8.y} + r_{3.9} P_{9.y} + r_{3.10} P_{10.y} + r_{3.11} P_{11.y} + r_{3.12} P_{12.y}$$

$$r_{4.y} = r_{1.4} P_{1.y} + r_{2.4} P_{2.y} + r_{3.4} P_{3.y} + P_{4.y} + r_{4.5} P_{5.y} + r_{4.6} P_{6.y} + r_{4.7} P_{7.y} + r_{4.8} P_{8.y} + r_{4.9} P_{9.y} + r_{4.10} P_{10.y} + r_{4.11} P_{11.y} + r_{4.12} P_{12.y}$$

$$r_{5.y} = r_{1.5} P_{1.y} + r_{2.5} P_{2.y} + r_{3.5} P_{3.y} + r_{4.5} P_{4.y} + P_{5.y} + r_{5.6} P_{6.y} + r_{5.7} P_{7.y} + r_{5.8} P_{8.y} + r_{5.9} P_{9.y} + r_{5.10} P_{10.y} + r_{5.11} P_{11.y} + r_{5.12} P_{12.y}$$

$$r_{6.y} = r_{1.6} P_{1.y} + r_{2.6} P_{2.y} + r_{3.6} P_{3.y} + r_{4.6} P_{4.y} + r_{5.6} P_{5.y} + P_{6.y} + r_{6.7} P_{7.y} + r_{6.8} P_{8.y} + r_{6.9} P_{9.y} + r_{6.10} P_{10.y} + r_{6.11} P_{11.y} + r_{6.12} P_{12.y}$$

$$r_{7.y} = r_{1.7} P_{1.y} + r_{2.7} P_{2.y} + r_{3.7} P_{3.y} + r_{4.7} P_{4.y} + r_{5.7} P_{5.y} + r_{6.7} P_{6.y} + P_{7.y} + r_{7.8} P_{8.y} + r_{7.9} P_{9.y} + r_{7.10} P_{10.y} + r_{7.11} P_{11.y} + r_{7.12} P_{12.y}$$

$$r_{8,y} = r_{1.8} P_{1,y} + r_{2.8} P_{2,y} + r_{3.8} P_{3,y} + r_{4.8} P_{4,y} + r_{5.8} P_{5,y} + r_{6.8} P_{6,y} + r_{7.8} P_{7,y} + P_{8,y} + r_{8.9} P_{9,y} + r_{8.10} P_{10,y} + r_{8.11} P_{11,y} + r_{8.12} P_{12,y}$$

$$r_{9,y} = r_{1.9} P_{1,y} + r_{2.9} P_{2,y} + r_{3.9} P_{3,y} + r_{4.9} P_{4,y} + r_{5.9} P_{5,y} + r_{6.9} P_{6,y} + r_{7.9} P_{7,y} + r_{8.9} P_{8,y} + P_{9,y} + r_{9.10} P_{10,y} + r_{9.11} P_{11,y} + r_{9.12} P_{12,y}$$

$$r_{10,y} = r_{1.10} P_{1,y} + r_{2.10} P_{2,y} + r_{3.10} P_{3,y} + r_{4.10} P_{4,y} + r_{5.10} P_{5,y} + r_{6.10} P_{6,y} + r_{7.10} P_{7,y} + r_{8.10} P_{8,y} + r_{9.10} P_{9,y} + P_{10,y} + r_{10.11} P_{11,y} + r_{10.12} P_{12,y}$$

$$r_{11,y} = r_{1.11} P_{1,y} + r_{2.11} P_{2,y} + r_{3.11} P_{3,y} + r_{4.11} P_{4,y} + r_{5.11} P_{5,y} + r_{6.11} P_{6,y} + r_{7.11} P_{7,y} + r_{8.11} P_{8,y} + r_{9.11} P_{9,y} + r_{10.11} P_{10,y} + P_{11,y} + r_{11.12} P_{12,y}$$

$$r_{12,y} = r_{1.12} P_{1,y} + r_{2.12} P_{2,y} + r_{3.12} P_{3,y} + r_{4.12} P_{4,y} + r_{5.12} P_{5,y} + r_{6.12} P_{6,y} + r_{7.12} P_{7,y} + r_{8.12} P_{8,y} + r_{9.12} P_{9,y} + r_{10.12} P_{10,y} + r_{11.12} P_{11,y} + P_{12,y}$$

Where,

r_{1y} Genotypic correlation coefficients between y and I th character (y = Grain yield)

P_{iy} = Path coefficient due to ith character (i= 1, 2, 3,.....,13)

1 = Days to first male flowering

2 = Days to first female flowering

3 = Leaf length (cm)

4 = Leaf breadth (cm)

5 = Internode distance (cm)

6 = Pedicel length of male flower (cm)

7 = Pedicel length of female flower (cm)

8 = Number of male flower per plant

9 = Number of female flower per plant

10 = Fruit weight (kg)

11 = Fruit length (cm)

12 = Fruit breadth (cm)

Total correlation, say between 1 and y i. e., r_1 is thus partitioned as follows:

$P_{1,y}$ = the direct effect of 1 on y

$r_{1.2} P_{2,y}$ = indirect effect of 1 via 2 on y

$r_{1.3} P_{3,y}$ = indirect effect of 1 via 3 on y

$r_{1.4} P_{4,y}$ = indirect effect of 1 via 4 on y

$r_{1.5} P_{5,y}$ = indirect effect of 1 via 5 on y

$r_{1.6} P_{6,y}$ = indirect effect of 1 via 6 on y

$r_{1.7} P_{7,y}$ = indirect effect of 1 via 7 on y

$r_{1.8} P_{8,y}$ = indirect effect of 1 via 8 on y

$r_{1.9} P_{9,y}$ = indirect effect of 1 via 9 on y

$r_{1.10} P_{10,y}$ = indirect effect of 1 via 10 on y

$r_{1.11} P_{11,y}$ = indirect effect of 1 via 11 on y

$r_{1.12} P_{12,y}$ = indirect effect of 1 via 12 on y

Where,

$P_{1,y}, P_{2,y}, P_{3,y}, \dots, P_{8,y}$ = Path coefficient of the independent variables 1, 2, 3,, 12 on the dependent variable y, respectively.

$r_{1,y}, r_{2,y}, r_{3,y}, \dots, r_{12,y}$ Correlation coefficient of 1, 2, 3,, 12 with y, respectively.

After calculating the direct and indirect effect of the characters, residual effect (R) was calculated by using the formula given below (Singh and Chaudhary, 1985)

$$P^2_{RY} = 1 - (r_{1,y} + r_{2,y} P_{2,y} + \dots + r_{12,y} P_{12,y})$$

Where,

$$P^2_{RY} = R^2$$

Hence residual effect, $R = (P^2_{RY})^{1/2}$

$P_{1,y}$ = Direct effect of the 1st character on yield y.

$r_{1,y}$ = Correlation of the 1st character with yield y.

3.15.2 Multivariate analysis

The genetic diversity among the genotypes was assessed by Mahalanobis's (1936) general distance (D^2) statistic and its auxiliary analyses. The parents' selection in hybridization programme based on Mahalanobis's D^2 statistic is more reliable as requisite knowledge of parents in respect of a mass of characteristics is available prior to crossing. Rao (1952) suggested that the quantification of genetic diversity through biometrical procedures had made it possible to choose genetically diverse parents for a hybridization programme. Multivariate analysis viz. Principal Component analysis, Principal Component analysis, Cluster analysis and Canonical Vector analysis (CVA), which quantify the differences among several quantitative traits, are efficient method of evaluating genetic diversity. These are as follows:

3.15.2.1 Principal component analysis (PCA)

Principal component analysis, one of the multivariate techniques, is used to examine the inter-relationships among several characters and can be done from the sum of squares and products matrix for the characters. Thus, PCA finds linear combinations of a set variate that maximize the variation contained within them, thereby displaying most of the original variability in a smaller number of dimensions. Therefore, principal components were computed from the correlation matrix and genotypes scores obtained for first components (which has the property of accounting for maximum variance) and succeeding components with latent roots greater than unity. Contribution of the different morphological characters towards divergence is discussed from the latent vectors of the first two principal components.

3.15.2.2 Cluster analysis (CA)

Cluster analysis divides the genotypes of a data set into some number of mutually exclusive groups. Clustering was done using non-hierarchical classification. In Genstat, the algorithm is used to search for optimal values of chosen criterion proceeds as follows. Starting from some initial classification

of the genotypes into required number of groups, the algorithm repeatedly transferred genotypes from one group to another so long as such transfer improved the value of the criterion. When no further transfer can be found to improve the criterion, the algorithm switches to a second stage, which examines the effect of swooping two genotypes of different classes and so on.

3.15.2.3 Canonical vector analysis (CVA)

Canonical vector analysis (CVA) finds linear combination of original variabilities that maximize the ratio of between group to within group variation, thereby giving functions of the original variables that can be used to discriminate between the groups. Thus, in this analysis a series of orthogonal transformations sequentially maximizing of the ratio of among groups, to the within group variations. The canonical vector are based upon the roots and vectors of WB, where W is the pooled within groups covariance matrix and B is the among groups covariance matrix.

3.15.2.4 Calculation of D² values

The Mahalanobls's distance (D²) values were calculated from transformed uncorrelated means of characters according to Rao (1952), and Singh and Chaudhury (1985). The D² values were estimated for all possible combinations between genotypes. In simpler form D² statistic is defined by the formula

$$D^2 = \sum_1^x d_1^2 = \sum_1^x (Y_i^j - Y_j^k)^2 \quad (j \neq k)$$

Where,

Y = Uncorrelated variable (character) which varies from i = 1----- to x

x = Number of characters.

3.15.2.5 Computation of average intra-cluster distances

Average intra-cluster distances were calculated by the following formula as suggested by Singh and Chuadhury (1985).

$$\text{Average intra-cluster distance} = \frac{\sum D_i^2}{n}$$

Where,

D_i^2 = the sum of distances between all possible combinations (n) of genotypes included in a cluster.

n = Number of all possible combinations between the populations in cluster.

3.15.2.6 Computation of average inter-cluster distances

Average inter-cluster distances were calculated by the following formula as suggested by Singh and Chuadhury (1985).

$$\text{Average inter-cluster distance} = \frac{\sum D_{ij}^2}{n_i \times n_j}$$

Where,

$\sum D_{ij}^2$ = The sum of distances between all possible combinations of the populations in cluster i and j.

n_i = Number of populations in cluster i.

n_j = Number of populations in cluster j.

3.15.2.7 Cluster diagram

Using the values of intra and inter-cluster distances ($D = \sqrt{D^2}$), a cluster diagram was drawn as suggested by Singh and Chuadhury (1985). It gives a brief idea of the pattern of diversity among the genotypes included in a cluster.

3.1 5.2.8 Selection of varieties for future hybridization programme

Divergence analysis is usually performed to identify the diverse genotypes for hybridization purposes. The genotypes grouped together are less divergent among themselves than those, which fall into different clusters. Clusters separated by largest statistical distance (D^2) express the maximum divergence among the genotypes included into these different clusters. Variety (s) or line(s) were selected for efficient hybridization program according to Singh and

Chaudhury (1985). According to them the following points should be considered while selecting genotypes for hybridization program:

- i. Choice of cluster from which genotypes are selected for use as parent(s)
- ii. Selection of particular genotype(s) from the selected cluster(s)
- iii. Relative contribution of the characters to the total divergence
- iv. Other important characters of the genotypes performance

CHAPTER-IV

RESULTS AND DISCUSSION

The research experiment was carried out to evaluate the genetic variability, correlation, path coefficient analysis and genetic diversity of 24 pumpkin accessions. The results of the present study have been presented and discussed in this chapter under the following heading.

4.1. Genetic variability

The analysis of variance indicated that the existence of highly significant variation among the genotypes. The mean, range, mean sum of square, variance components, genotypic and phenotypic co efficient of variance, heritability, genetic advance, genetic advance in percent of mean are presented in Table 3.

4.1.1. Leaf length without petiole (cm)

Considerable variations were observed among 24 genotypes of pumpkin for leaf length. Significant mean sum of square for leaf length (6.03) indicated considerable variation was present among genotype studied (Table 3). The maximum leaf length was observed 17.50 in G₆ (BD 10058) and minimum was 11.67 recorded in G₁₇ (BD 4363) with mean value 15.01 (Appendix V). The phenotypic variance (4.74) appeared to be slightly higher than genotypic variance (1.29) suggested that less influence of environment on the expression

Table 3. Estimation of genetic variability for yield contributing characters related to yield of pumpkin

Traits	Min	Max	Mean	MS	CV (%)	σ_{g}^2	σ_{e}^2	σ_{P}^2	GCV	ECV	PCV	h^2_{b}	GA (5%)	GA(% mean)
LL	11.67	17.50	15.01	6.03	12.37	1.29	3.4449	4.74	7.58	12.37	14.50	27.31	1.22	8.16
LB	15.73	24.00	20.16	12.57	16.37	0.84	10.8859	11.73	4.55	16.37	16.99	7.18	0.51	2.51
ID	7.90	16.33	12.00	9.07	19.24	1.87	5.3301	7.20	11.40	19.24	22.36	25.98	1.44	11.97
DFMF	52.67	56.00	54.18	2.20	2.58	0.12	1.9541	2.08	0.65	2.58	2.66	5.97	0.18	0.33
DFFF	60.67	66.33	63.82	4.87	3.28	0.25	4.3726	4.62	0.78	3.28	3.37	5.37	0.24	0.37
PLM	15.67	28.33	22.41	16.71**	3.47	8.05	0.6031	8.66	12.66	3.47	13.13	93.03	5.64	25.16
PLF	6.17	11.83	9.60	7.54**	4.43	3.68	0.1806	3.86	20.01	4.43	20.49	95.32	3.86	40.24
NMF	6.33	12.00	8.93	8.32**	15.92	3.15	2.0211	5.17	19.87	15.92	25.46	60.90	2.85	31.94
NFF	4.33	12.00	6.83	8.09**	20.24	3.09	1.9112	5.00	25.73	20.24	32.74	61.77	2.85	41.65
FL	22.37	39.00	30.06	82.46**	3.08	40.80	0.8550	41.66	21.25	3.08	21.47	97.95	13.02	43.32
FB	41.37	76.60	54.60	390.48**	2.24	194.49	1.5024	195.99	25.54	2.24	25.64	99.23	28.62	52.41
FW	1.90	4.67	2.89	1.46**	4.20	0.72	0.01	0.74	29.44	4.20	29.73	98.01	1.73	60.03
YIELD	4.27	11.30	6.51	11.62**	15.59	5.29	1.03	6.32	35.34	15.59	38.63	83.71	4.34	66.62

LL= Leaf length, LB= Leaf breath, ID= Internode distance, DFMF= Days to first male flowering, DFFF= Days to first female flowering, PLM= Pedicel length of male, PLF= Pedicel length of female, NMF= Number of male flower, NFF= Number of female flower, FL= Fruit length, FB= Fruit breath, FW= Fruit weight (kg) and YIELD= Fruit yield (kg)

** = significant at 1%

*= significant at 5%

(7.58%) and phenotypic co-efficient of variation (14.50%) were close to each other. Asmaul Husna (2009) found 14.14 genotypic variance in bottle gourd. The GCV (22.63) and PCV (22.67) were close to each other respectively. This character showed heritability (27.31) and moderate genetic advance in percent of mean (8.16) which indicated that the character is controlled by additive genes and that selection based on this character would be effective. Gaffar (2008) found high heritability and moderate genetic advance in sponge gourd. Fayeun *et al.*, (2012) also found high heritability and moderate genetic advance in fluted pumpkin. Phenotypic variations among leaf length of different genotypes is presented in plate 3a and 3c.

4.1.2 Leaf breadth (cm)

Significant mean sum of square for leaf breadth (12.57) indicated considerable variation presented among the genotypes studied (Table 3). The maximum leaf breadth was observed 24.00 in G₆ (BD 10058) and minimum in 15.73 which was recorded in G₁₇ (BD 4363) with mean value 20.16 (Appendix V). The phenotypic variance (11.73) appeared to be higher than genotypic variance (0.84) suggests less influence of environment on the expression of this gene controlling the trait. The genotypic co-efficient of variation (4.55%) and phenotypic co-efficient of variation (16.99%) were far reach to each other. Husna (2009) found GCV (22.87) was lower than PCV (23.04) for this character in bottle gourd. This character showed heritability (7.78) and moderate genetic advance (0.51) and genetic advance in percent of mean (2.51) which indicated that the character was controlled by additive genes. Therefore the selection based on this character would be effective. Gaffar (2008) observed high broad sense heritability (94%) with moderate genetic advance (7.81) for this character in sponge gourd. Phenotypic variations among leaf breadth of different pumpkin genotypes is presented in 3a to 3c.



Plate 3a: Showing morphological variation in leaf among different pumpkin genotypes G₁(BD 10149), G₂(BD 10141), G₃(BD 4361) G₄(BD 4353), G₅(BD 10137) G₆(BD 10058) G₇(BD 4349) G₈(Pronoy-F1)



Plate 3b: Showing morphological variation in leaf among different pumpkin genotypes G₉(BD 4371), G₁₀(BD 4357), G₁₁(BD 4350), G₁₂(BD 4352), G₁₃(BD 10078), G₁₄(BD 10366), G₁₅(BD 4363), G₁₆(BARMASI)

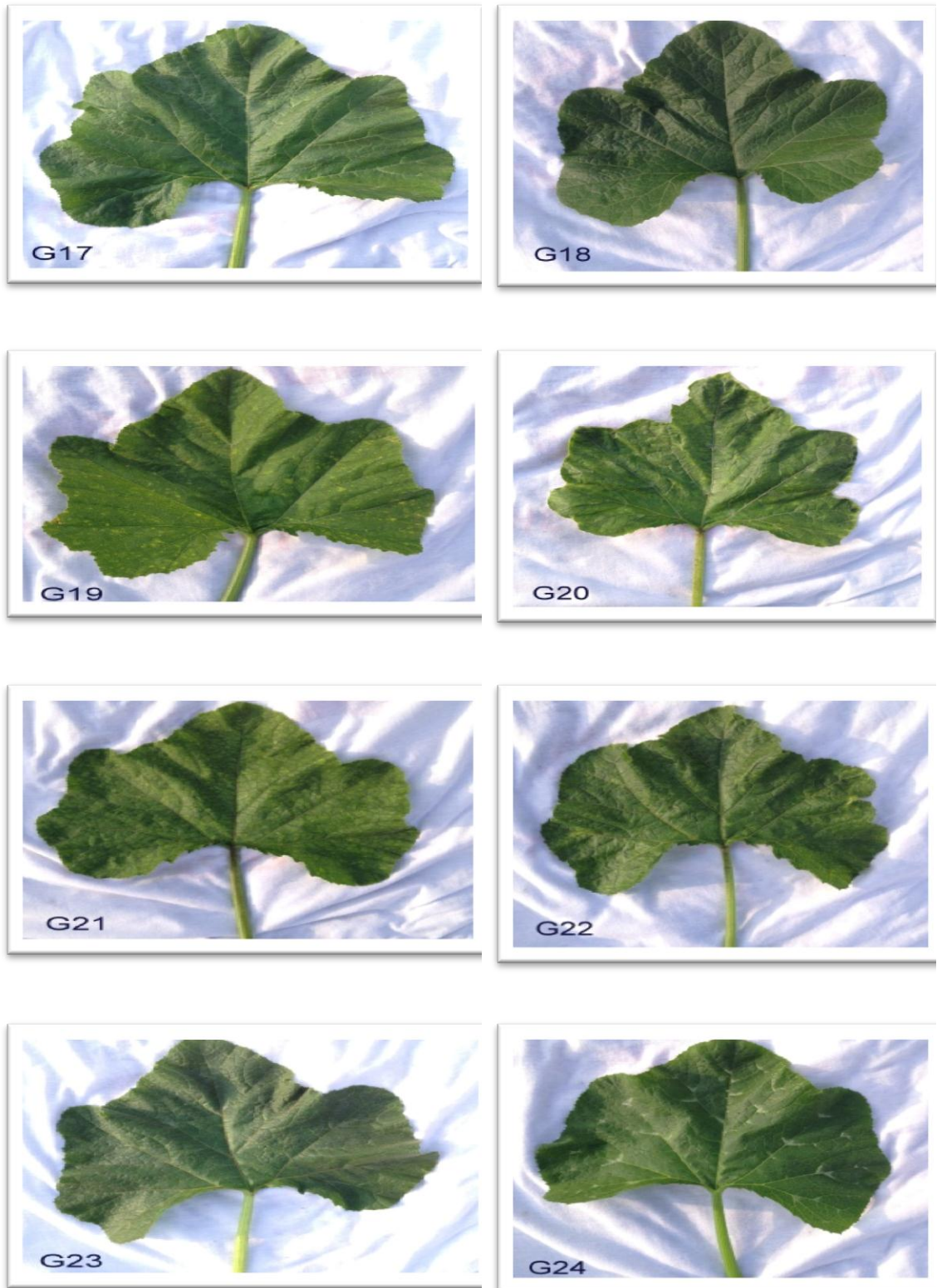


Plate 3c: Showing morphological variation in leaf among different pumpkin genotypes G₁₇(BD 4372), G₁₈(BD 4391), G₁₉(BD 4389), G₂₀(BD 4354), G₂₁(BD 4393), G₂₂(BD 4370), G₂₃(BD 10134), G₂₄(PRITY-F1)

4.1.3 Internode distance (cm)

Significant difference for internode distance observed among the pumpkin genotype studied (Table 3). Mean sum of square was significant (9.07). The maximum internode distance was observed (16.33cm) in G₂₁ (BD 4389) and minimum (7.90cm) which was recorded in G₁₇ (BD 4563) with mean value 12.00 (appendix V). The difference between phenotypic variance (7.20) and genotypic variance (1.87) was higher indicating less influence of environment on the expression of this character. The genotypic co-efficient of variation was 22.36%, and phenotypic co-efficient of variation was 11.40% (Table 3).

Heritability showed high (25.98) and moderate genetic advance (1.44) and genetic advance in percent of mean (11.97) revealed that character was controlled by additive genes the selection based on this character would be effective. Fayeun *et al.*, (2012) also found high heritability and moderate genetic advance in percent of mean for this trait in pumpkin.

4.1.4 Days to first male flowering

Days to first male flowering showed significant variation among genotype due to Mean square (2.20). The maximum duration(56.00)was observed in G₂₃ (BD 4393) and the minimum duration (52.67) was in G₂₄ (PRITY F1) with mean value of 54.18 (Table 3). The difference between phenotypic variance (2.08) and genotypic variance (0.12) was large which indicates presence of environmental influence for the expression of the characters (Table 3).

Heritability showed high (5.97) with low genetic advance in percent of mean (0.33) revealed that which indicated character was controlled by non-additive genes the selection based on this character would not be effective. Samsun.

4.1.5 Days to first female flowering

Significant difference was observed for days to first female flowering among the pumpkin genotypes (Table 3) as mean sum of square was significant. The maximum duration(66.33 days) was observed in G₁₁ (BD 4357) and the minimum duration(60.67 days) was in G₁ (BD 10149) with mean value 63.83.

The difference between phenotypic variance (4.62) and genotypic variance (0.25) was with large environmental influence. The genotypic co-efficient of variation and phenotypic co-efficient of variation for this character was 0.78% and 3.37% respectively (Table 3).

Heritability showed high (5.37) with low genetic advance in percent of mean (0.37) revealed that character was controlled by non-additive gene so the selection based on this character would not be effective. Singh and Lal (2005) also found similar result in their study.

4.1.6 Pedicel length of male flower (cm)

Mean sum of square for pedicel length was significant (16.71) among the genotypes of pumpkin (Table 3). The maximum pedicel length was observed (28.33) in G₁₆ (BARMASI) and minimum was (15.67) recorded in G₁₄ (BD 10078) with mean value of 22.41 (Appendix V). The phenotypic variance (8.66) appeared to be slightly higher than genotypic variance (8.05) suggested that less influence of environment on the expression of this gene controlling this trait. The genotypic co-efficient of variation (12.66%) and phenotypic co-efficient of variation (13.13%) were close to each other (Table 3).

Heritability showed high (93.03) and moderate genetic advance (5.64) and genetic advance in percent of mean (25.16) revealed that character was controlled by additive gene and the selection based on this character would be effective. Asmaul Husna (2009) also found high heritability (99.55%) and genetic advance for this trait in bottle gourd. Plate 4a to 4c showing variation of male flower among twenty four genotypes of pumpkin.



Plate 4a: Showing morphological variation in flower among different pumpkin genotypes G₁(BD 10149), G₂(BD 10141), G₃(BD 4361) G₄(BD 4353), G₅(BD 10137) G₆(BD 10058) G₇(BD 4349) G₈(Pronoy-F1)



Plate 4b: Showing morphological variation in flower among different pumpkin genotypes G₉(BD 4371), G₁₀(BD 4357), G₁₁(BD 4350), G₁₂(BD 4352), G₁₃(BD 10078), G₁₄(BD 10366), G₁₅(BD 4363), G₁₆(BARMASI)



Plate 4c: Showing morphological variation in flower among of different of pumpkin genotypes G₁₇(BD 4372), G₁₈(BD 4391), G₁₉(BD 4389), G₂₀(BD 4354), G₂₁(BD 4393), G₂₂(BD 4370), G₂₃(BD 10134), G₂₄(PRITY-F1)

4.1.7 Pedicel length of female flower

Mean sum of square for pedicel length was significant (7.54) in genotypes of pumpkin. The maximum pedicel length was observed 11.83 in G₁₆ (BARMASI) and minimum was 6.17 recorded in G₃ (BD 4361) with mean value 4.56(Appendix V). The phenotypic variance (3.86) appeared to be slightly higher than genotypic variance (3.68) suggested that less influence of environment on the expression of this gene controlling this trait. The genotypic co-efficient of variation (20.01%) and phenotypic co-efficient of variation (20.49%) were close to each other (Table 3). Asmaul Husna (2009) found similar result in bottle gourd.

Heritability showed high (95.32%) and high genetic advance in percent of mean (40.24) revealed that character is controlled by additive genes and the selection based on this character would be effective.

4.1.8 Number of male flowers per plant

Significant difference observed among number of male flowering in pumpkin genotypes studied (Table 3). Mean sum of square was significant (8.32). The maximum number of male flower was 12.00 observed in G₁ (BD 10149) and the minimum number was 6.33 in G₁₈ (BD 10134) with mean value 8.93. The phenotypic variance (5.17) appeared to be slightly higher than genotypic

Variance (3.15) suggested that less influence of environment on the expression of this gene controlling this trait. The genotypic co-efficient of variation was 19.87% and phenotypic co-efficient of variation was 25.46%, respectively. Husna *et al.*, (2011) found GCV 31.86 % and PCV 31.95% which was similar to the present study.

Heritability showed high (60.90) and with high genetic advance in percent of mean (31.94) revealed that character was controlled by additive gene so the selection based on this character would be effective.

4.1.9 Number of female flowers per plant

Significant difference was observed among number of female flowering in pumpkin genotypes studied (Table 3). Mean sum of square was significant (8.09). The maximum number of female flower was 12.00 observed in G₁ (BD 10149) and the minimum number was 4.33 in G₂₁ (BD 4389) with mean value 6.83. The phenotypic variance (5.00) appeared to be slightly higher than genotypic variance (3.09) suggested that less influence of environment acted on the expression of this gene controlling this trait. The genotypic co-efficient of variation was 25.73% and phenotypic co-efficient of variation was 32.74%, respectively. Husna *et al.*, (2011) found GCV 35.14% and PCV 38.08% which was similar to the present study.

High heritability showed (61.77) with genetic advance in percent of mean (41.65) revealed that character was controlled by additive gene so the selection based on the character would be effective. Husna *et al.*, (2011) also found high heritability (88.15) with high genetic advance in percent of mean (85.6).

4.1.10 Fruit length (cm)

Mean sum of square of fruit length was significant (82.46). The maximum fruit length was found 39.00 in G₂₄ (PRITY F1) and the minimum number was 22.37 in G₂ (BD 10141) with mean value 30.06. The phenotypic variance (41.66) appeared to be moderately higher than genotypic variance (40.80) suggested that moderate influence of environment acted on the expression of this gene controlling this trait. The genotypic co-efficient of variation and phenotypic co-efficient of variation were observed 21.25% and 21.47%, respectively. Banik (2003) found the highest phenotypic co efficient of variation for fruit length in snake gourd. Mathew and Khader (1999) also reported high heritability for fruit length in snake gourd. Rahman *et al.*,(1991) indicated minimum differences between GCV and PCV in bottle gourd for fruit length.

High heritability found (97.95) with moderately high genetic advance in percent of mean (43.32) revealed that character was controlled by additive gene so the selection based on this character would be effective. Devi and Mariappan (2013) found high heritability (99.99) with high genetic advance (97.13) which also revealed that the character was controlled by additive gene. Photographs showed variation in fruit length of sweet gourds. Variation of fruit length among twenty genotypes of pumpkin is presented in plate 5a to 5c.

4.1.11 Fruit breadth (cm)

Significant mean sum of square of fruit breadth was found (390.48). The maximum fruit breadth was found 76.60 in G₂₄ (PRITY F1) and the minimum number was 41.37 in G₁₂ (BD 4350) with mean value 54.60. The phenotypic variance (195.99) appeared to be higher than genotypic variance (194.49) suggested that influence of environment acted on the expression of this gene controlling this trait. The genotypic co-efficient of variation was 25.54% and phenotypic co-efficient of variation was 25.64% respectively. Devi and Mariappan (2013) found GCV Phenotypic co efficient of variation 24.67 was slightly higher than genotypic co efficient of variation 24.66.



Plate 5a: Showing morphological variation in fruits among different pumpkin genotypes G₁(BD 10149), G₂(BD 10141), G₃(BD 4361) G₄(BD 4353), G₅(BD 10137) G₆(BD 10058) G₇(BD 4349) G₈(Pronoy-F1)



Plate 5b: Showing morphological variation in fruits among different pumpkin genotypes G₉(BD 4371), G₁₀(BD 4357), G₁₁(BD 4350), G₁₂(BD 4352), G₁₃(BD 10078), G₁₄(BD 10366), G₁₅(BD 4363), G₁₆(BARMASI)

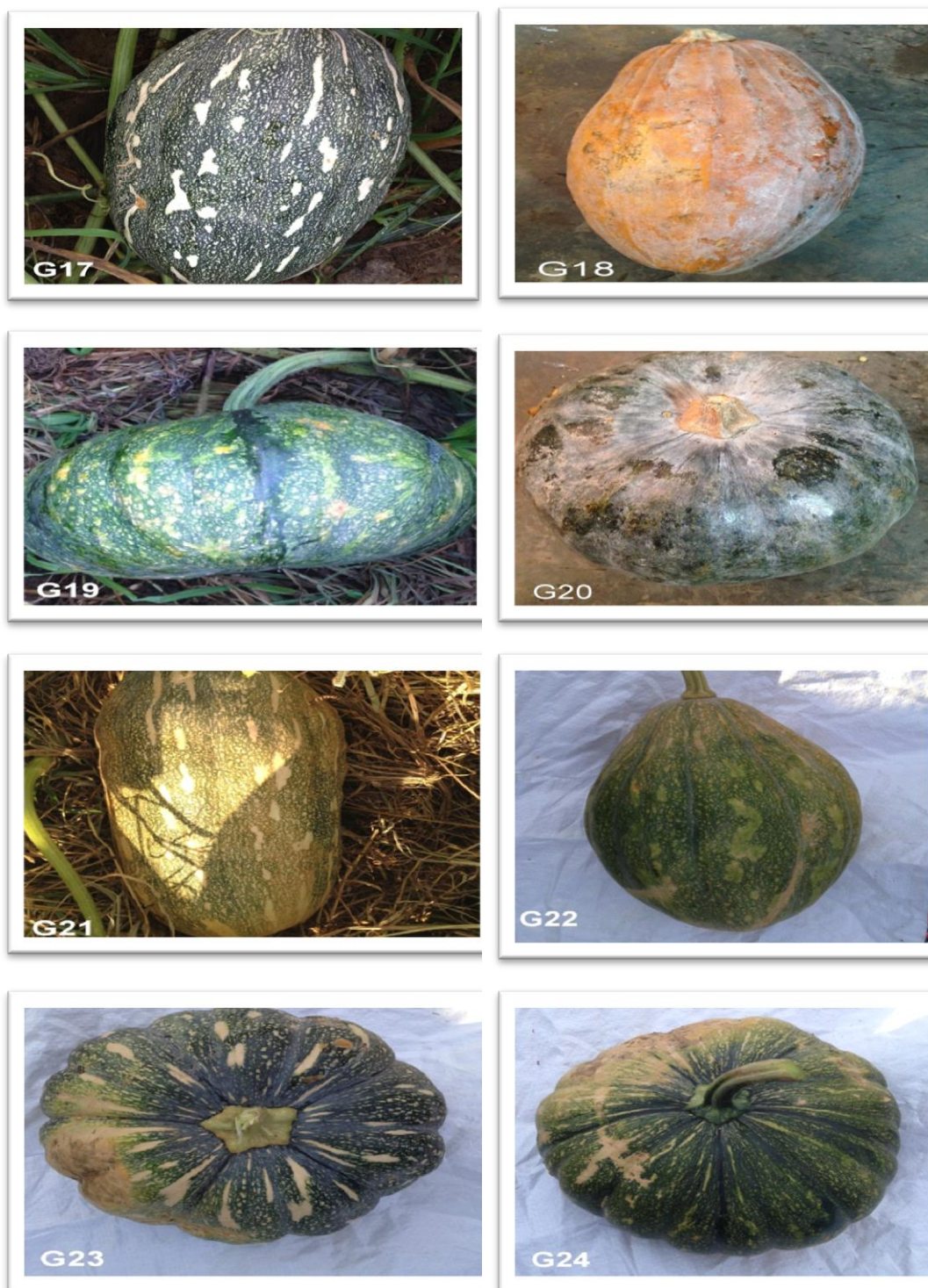


Plate 5c: Showing morphological variation in fruits of different pumpkin genotypes G₁₇(BD 4372), G₁₈(BD 4391), G₁₉(BD 4389), G₂₀(BD 4354), G₂₁(BD 4393), G₂₂(BD 4370), G₂₃(BD 10134), G₂₄(PRITY-F1)

Heritability was found high (99.23) with moderately high genetic advance in percent of mean (52.41) revealed that character is controlled by additive gene so the selection based on this character would be effective. Asmaul Husna (2009) reported GCV and PCV were 15.84 and 17.39 respectively in bottle gourd and heritability (82.93%) estimates for this trait was high along with moderately high genetic advance in percent of mean (38.08). Plate 5a to 5c showing variation of fruit breadth among twenty genotypes of pumpkin.

4.1.12 Fruit weight (kg)

Significant mean sum of square of fruit weight was found (1.46). The maximum fruit weight found 4.67 kg in G₂₄ (PRITY F1) and the minimum fruit weight was 1.90 kg found in G₂ (BD 10141) with mean value 2.89. The phenotypic variance (0.74) appeared to be slightly higher than genotypic variance (0.72) suggested that less influence of environment acted on the expression of this gene controlling this trait. The genotypic co-efficient of variation was 29.44% and phenotypic co-efficient of variation was 29.73%, respectively. Saha *et al.*, (1992) found similar GCV and PCV for the fruit weight in pumpkin. Kumaran *et al.*, (1997) reported similar types of result which confirmed the present findings.

Heritability was found high (98.01) with moderately high genetic advance in percent of mean (60.03) revealed that the character was controlled by additive gene so the selection based on this character would be effective. Rahman *et al.*, (1986) also found the similar result in bottle gourd.

4.1.13 Fruit yield per plant (kg)

Mean sum of square of fruit weight was found was significant (11.62). The maximum fruit yield per plant found 11.30 kg in G₂₃ (BD 4393) and the minimum fruit yield per plant was 4.27 kg found in G₁₂ (BD 4350) with mean value 6.51. The phenotypic variance (6.32) appeared to be slightly higher than genotypic variance (5.29) suggested that less influence of environment acted on the expression of this gene controlling this trait. The genotypic co-efficient of variation was observed 35.34% and phenotypic co-efficient of variation was observed 38.63% respectively. Husna *et al.*, (2011) were found GCV (52.02 %) and PCV (54.35) for fruit yield per plant in pumpkin which confirmed the results of the present study. High heritability (83.71) with very high genetic advance in percent of mean (66.62) revealed that character was controlled by additive gene. So the selection based on this character would be highly effective. Narayankutty *et al.*, (2006) reported high values of heritability and genetic gain for fruit yield per plant indicating additive gene effects are important in determining this character.

4.2. Correlation co-efficient

Yield is a complex product being influenced by several inter-dependable quantitative characters. Thus selection for yield may not be effective unless the other yield components influence it directly or indirectly are taken in to consideration. When selection pressure is exercised for improvement of any character highly associated with yield, it simultaneously affects a number of other correlated characters. Hence knowledge regarding association of character with yield and among themselves provides guideline to the plant breeder for making improvement through selection provide a clear understanding about the contribution in respect of establishing the association by genetic and non-genetic factors (Dewey and Lu 1959). Result of genotypic and phenotypic correlation co-efficient analysis of thirteen yield and yield contributing characters of pumpkin were estimated separately as vegetative

character and reproductive character with yield shown in Table 4 which discussed character wise below:

4.2.1 Leaf length without petiole (cm)

Leaf length showed highly significant positive correlation with leaf breadth and internode distance at both genotypic and phenotypic level indicated that if leaf length increased these parameters will also be increased (Table 4). Non-significant and negative correlation was found with number of male flowers, number of female flowers and fruit yield. Positive but non-significant correlation was found for days to first male flowering, days to first female flowering, pedicel length of male, pedicel length of female, fruit breadth, fruit length and fruit weight. Husna *et al.*, (2014) also reported leaf length was positively non-significant correlated with fruit yield per plant. Li *et al.*, (1997) also found similar result in cucumber for this trait.

4.2.2 Leaf breadth (cm)

Highly significant and positive correlation was found between leaf breadth and internode distance. (Table 4). Non-significant and negative correlation was found with days to first male flowering, days to first female flowering, pedicel length of male flower, pedicel length of female flower and number of female flower both at genotypic and phenotypic level. Positive but non-significant correlation was found in number of male flower, fruit length, fruit breadth, fruit weight and fruit yield per plant. Husna *et al.*, (2014) also found leaf breadth has positive non-significant correlation with fruit yield per plant.

4.2.3 Internode distance (cm)

Positive and non-significant correlation was found in days to first male flowering, number of male flower and fruit weight per plant indicating that association among these traits is largely influenced by

Table 4. Coefficients of phenotypic and genotypic correlation among different yield components of twenty four pumpkin genotypes.

		LL	LB	ID	DFMF	DFFF	PLM	PLF	NMF	NFF	FL	FB	FW
LB	G	0.570											
	P	0.660**											
ID	G	0.764	0.650										
	P	0.555**	0.597**										
DFMF	G	0.675	0.157	0.048									
	P	0.063	-0.071	0.095									
DFFF	G	0.321	-0.135	-0.214	0.630								
	P	0.093	-0.074	-0.155	0.368								
PLM	G	0.106	0.015	-0.172	0.665	0.450							
	P	0.033	-0.023	-0.142	0.426*	0.382							
PLF	G	0.026	-0.243	-0.164	-0.231	0.390	0.322						
	P	0.015	-0.109	-0.075	-0.053	0.376	0.311						
NMF	G	-0.047	0.090	0.147	-0.452	-0.060	0.265	-0.016					
	P	0.044	0.120	0.161	-0.082	-0.040	0.234	-0.017					
NFF	G	-0.233	0.165	-0.456	-0.532	-0.392	-0.215	-0.332	0.526				
	P	-0.226	-0.051	-0.348	-0.493*	-0.378	-0.179	-0.300	0.337				
FL	G	0.525	0.607	-0.013	-0.613	-0.152	0.038	-0.264	0.035	0.012			
	P	0.343	0.247	-0.011	-0.216	-0.138	0.038	-0.259	0.029	0.008			
FB	G	0.297	0.493	-0.253	-0.569	-0.165	0.086	-0.156	0.003	0.035	0.872		
	P	0.193	0.191	-0.163	-0.204	-0.152	0.087	-0.153	0.004	0.032	0.868**		
FW	G	0.392	0.590	0.022	-0.283	-0.042	0.269	-0.001	0.171	-0.045	0.777	0.737	
	P	0.257	0.248	0.035	-0.107	-0.030	0.269	0.001	0.151	-0.054	0.773**	0.733**	
YIELD	G	-0.034	0.304	-0.346	-0.358	0.110	0.493	0.058	0.127	-0.015	0.525	0.663	0.770
	P	-0.022	0.140	-0.242	-0.060	0.081	0.461*	0.06	0.159	-0.052	0.499*	0.628**	0.733**

LL= Leaf length, LB= Leaf breath, ID= Internode distance, DFMF= Days to first male flowering, DFFF= Days to first female flowering, PLM= Pedicel length of male, PLF= Pedicel length of female, NMF= Number of male flower, NFF= Number of female flower, FL= Fruit length, FB= Fruit breath, FW= Fruit weight (kg) and YIELD= Fruit yield (kg).

environment (Table 4). Non-significant and negative correlation was found with days to first female flowering, pedicel length of male flower, pedicel length of female flower, number of female flower, fruit length, fruit breadth and fruit yield at both genotypic and phenotypic level. Positive and non-significant correlation between yield and internode distance showed the selection of genotypes with higher internode distance are expected to yield better.

4.2.4 Days to first male flowering

The character showed highly significant and positive correlation with days to first female flowering and pedicel length of male at both genotypic and phenotypic level indicated that the traits were governed by same gene and simultaneous improvement would be effective (Table 4). Negative correlation but significant was found with number of female flower. Negative correlation non-significant was found with number of male flower per plant, fruit breadth, fruit weight and fruit yield per plant both at genotypic and phenotypic level.

4.2.5 Days to first female flowering

Days to first female flowering showed highly insignificant and positive correlation with pedicel length of male, pedicel length of female and fruit yield both genotypic and phenotypic level indicated that if days to first female flowering increases fruit yield would be highly increased (Table 4). Negative correlation was found with number of male flower, number of female flower, fruit length, fruit breadth and fruit weight which suggested that delayed of first female flowering increases the number of male flower. Positive association was found with pedicel length of male flower, pedicel length of female flower and fruit yield per plant. Khan *et al.*, (2009) reported the similar result.

4.2.6 Pedicel length of male flower (cm)

Pedicel length of male flower showed positive and significant correlation with the yield per plant. Positive but non-significant correlation with pedicel length

of female flower, number of male flower per plant, fruit length, fruit breadth and fruit weight indicated the traits were governed by same gene and improvement would be effective. Highly negative correlation was found with number of female flower per plant.

4.2.7 Pedicel length of female flower (cm)

The character showed negative but insignificant relation with number of male flower per plant and number of female flower per plant indicating decreased pedicel length would increase the number of male and female flower. Positive but insignificant relation was found with fruit yield per plant indicating if pedicel length increased fruit yield would also be increased.

4.2.8 Number of male flowers per plant

Number of male flower per plant showed highly positive correlation with number of female flower per plant and positive but insignificant correlation with fruit yield per plant at both genotypic and phenotypic level. It indicated if number of male flower increased number of female flower would be highly increased and fruit yield would also be increased. Khan *et al* (2009) also found Number of male flower per plant has highly significant positive correlation with number of female flower per plant.

4.2.9 Number of female flowers per plant

This character showed highly positive correlation with fruit length and fruit breadth at both genotypic and phenotypic level indicating if number of female flower increased fruit breadth would be increased highly. Number of female flower had positive correlation with fruit yield per plant indicating that if the number of female flower increased number of fruit also increased. Mohanty (2001) reported similar trend of relationship.

4.2. 10 Fruit length (cm)

Fruit length showed positive and significant correlation with fruit breadth, fruit weight and fruit yield at phenotypic level (Table 4) that indicating if fruit length increased fruit weight and fruit yield per plant would be highly increased. Narayankutty *et al.*, (2006) reported that yield is strongly correlated with fruit length in snake gourd. Chowdhury and Sarma (2002) studied *Luffa acutangula* cultivars and observed that yield per hectare can be improved through selection of fruit length.

4.2.11 Fruit breadth (cm)

Positive and highly significant correlation was found with fruit weight and fruit length at both genotypic and phenotypic level indicating if fruit breadth may increase fruit weight and fruit length may also increase. Narayankutty *et al.*, (2006) reported that yield is strongly correlated with fruit breadth in snake gourd. Khan *et al.*, (2009) found fruit breadth is positively correlated with fruit weight.

4.2.12 Fruit weight (kg)

Fruit weight showed highly significant positive correlation with fruit yield per plant both at genotypic and phenotypic level (Table 4) indicated that if fruit weight increased, then the fruit yield and number of fruits also increased. Narayankutty *et al.*, (2006) reported that yield is strongly correlated with fruit weight in snake gourd. Khan *et al* (2009) also found fruit weight has positive high correlation with yield. Husna *et al.*, (2014) also found similar result in bottle gourd. Chowdhury and Sarma (2002) studied *Luffa acutangula* cultivars and observed that yield per hectare can be improved through selection of individual fruit weight. Prasana *et al.*, (2002) found in ridge gourd (*Luffa acutangula*) fruit yield per hectare was positively associated with fruit weight. Kumaresan *et al.*, (2006) yield per vine in snake gourd was positively associated with fruit weight.

4.3 Path analysis

Association of character determined by correlation co-efficient may not provide an exact picture of the relative importance of direct and indirect influence of each of yield components. In order to find out a clear picture of the inter relationship between yield per plant and other yield attributes, direct and indirect effects were worked out using path analysis at phenotypic level which also measured the relative importance of each component. Estimation of direct and indirect effect of path co-efficient analysis for pumpkin presented in (Table 5).

Table 5. Path coefficient analysis showing direct and indirect effects of different characters on yield

Characters	Direct effect	Indirect effect									Genotypic correlation with Yield
		ID	DFFF	PLM	PLF	NMF	NFF	FL	FB	FW	
ID	.137	-	-.29	-.23	-.22	.2	-.62	-.02	-.35	.3	-0.346
DFFF	0.80	-.1709	-	.36	.31	-.08	-.31	-.12	-.13	-.11	0.110
PLM	0.72	-.12	.32	-	.23	.19	-.15	.27	.62	.19	0.493
PLF	0.24	-.04	.92	.08	-	-.003	-.08	-.06	-.03	-.0002	0.058
NMF	-.125	-.18	.73	-.33	.2	-	-.66	-.04	-.003	-.214	0.127
NFF	.181	-.82	-.71	-.39	-.60	.95	-	.42	.63	-.08	-0.015
FL	-.126	-.16	.19	-.05	.33	-.04	-.02	-	-.11	-.98	0.525
FB	.159	-.4	-.26	.14	-.25	.05	.56	.14	-	.12	0.663
FW	0.76	.02	-.11	.204	-.0007	.13	-.03	.59	.56	-	0.770

ID= Internode distance, DFFF= Days to first female flowering, PLM= Pedicel length of male, PLF= Pedicel length of female, NMF= Number of male flower, NFF= Number of female flower, FL= Fruit length, FB= Fruit breath, FW= Fruit weight (kg) and YIELD= Fruit yield (kg)

Residual effect: 0.47

4.3.1 Internode distance (cm)

Internode distance showed a positive direct effect (0.137) on yield (Table 5). This character showed highest positive indirect effect through number of male flower per plant (0.2), fruit weight (0.3). The character also produced negative indirect effect on yield via days to first female flowering (-0.29), pedicel length of male (-0.23), pedicel length of female flower (-0.22), number of female flower (-0.62), fruit length (-0.02), fruit breadth (-0.35) which finally made insignificant negative correlation between internode distance and yield per plant (-0.346).

4.3.2 Days to first female flowering

The character showed a positive direct effect (0.80) on yield (Table 5). Days to first female flowering showed highest positive indirect effect on pedicel length of male flower (0.36), pedicel length of female flower (0.31). The negative indirect character via number of male flower per plant (-0.08), number of female flower per plant (-0.31), fruit length (-0.12) and fruit breadth (-0.13), fruit weight (-0.11) The cumulative effect produced a positive insignificant correlation with yield (0.110).

4.3.3 Pedicel length of male flower (cm)

Male flower pedicel length showed a positive direct effect (0.72) on yield (Table 5). The character showed highest positive indirect effect on days to first female flowering (0.32), pedicel length of female flower (0.23), number of male flower (0.19), fruit length (0.27), fruit breadth (0.62), fruit weight (0.19). The character also produced negative indirect effect on yield through internode distance (-0.12), number of female flower per plant (-0.15) which finally produced positive insignificant correlation with yield (0.493).

4.3.4 Pedicel length of female flower (cm)

Pedicel length of female flower showed a positive direct effect (0.24) on yield (Table 5). The character showed highest positive indirect effect on days to first female flowering (0.92), pedicel length of male flower (0.08). The character also produced negative indirect effect on yield through internode distance (-0.04), number of male flower per plant (-0.003), number of female flower per plant (-0.08), fruit length (-0.06), and fruit breadth (-0.03), fruit weight (-0.0002) which finally contributed positive insignificant correlation with yield (0.058).

4.3.5 Number of male flowers per plant

Number of male flowers per plant showed negative direct effect (-0.125) on yield (Table 5). The character showed the highest positive indirect effect via days to first female flower per plant (0.73), internode distance (0.18). The character also produced negative indirect effect on yield through pedicel length of male flower (-0.33), number of female flowers per plant (-0.66), fruit length (-0.04), fruit breadth (-0.003), fruit weight (-0.214). The cumulative effect produced a positive non-significant correlation with yield (0.127). Husna *et al.*, (2011) also found negative direct effect of number of male flower on yield.

4.3.6 Number of female flowers per plant

Number of female flowers per plant showed positive direct effect (0.181) on yield (Table 5). The character showed the highest positive indirect effect via number of male flower per plant (0.95), fruit length (0.42), fruit breadth (0.63). The character also produced negative indirect effect on yield through internode distance (-0.82), days to first female flowering (-0.71), pedicel length of male flower (-0.39), pedicel length of female flower (-0.60) and fruit weight (-0.08) which finally produced a negative insignificant yield (-0.015). Shamima Sultana (2011) found similar result in sweet gourd.

4.3.7 Fruit length (cm)

Fruit length showed negative direct effect (-0.126) on yield (Table 5). The character showed highest positive indirect effect via days to first female flowering (0.19) and pedicel length of female flower (0.33). The character also produced negative indirect effect on yield through internode distance (-0.16), pedicel length of male flower (-0.05), number of male flower per plant (-0.04), number of female flower per plant (-0.02), fruit breadth (-0.11), fruit weight (-0.98). The cumulative effect produced a highly insignificant positive correlation with yield (0.525). Husna *et al.*, (2011) also found negative direct effect of fruit length on yield.

4.3.8 Fruit breadth (cm)

Fruit breadth showed positive direct effect (0.159) on yield (Table 5). The character showed highest positive indirect effect through fruit weight (0.12), number of male flower per plant (0.05), pedicel length of male flower (0.14), number of female flower (0.56) and fruit length (0.14). The character also produced negative indirect effect on yield through internode distance (-0.4), days to first female flowering (-0.26), pedicel length of female flower (-0.25) which finally produced a positive insignificant yield (0.663).

4.3.9 Fruit weight (kg)

Fruit weight showed positive direct effect (0.76) on yield (Table 5). The character showed highest positive indirect effect through internode distance (0.02), pedicel length of male flower (0.204), number of male flower per plant (0.13), fruit length (0.59) and fruit breadth (0.56). The character also produced negative indirect effect on yield through days to first female flowering (-0.11) and number of female flower per plant (-0.03). The cumulative effect produced a highly insignificant positive correlation with yield (0.770). Husna *et al.*, (2011) also found negative direct effect of fruit weight on yield, and also found

highly significant positive correlation with yield. Kumaresan *et al.*, (2006) conducted an experiment in snake gourd and path coefficient analysis revealed that it would be highly rewarding to lay emphasis on the number of fruit per vine and fruit weight to increase the yield per vine directly. The result is similar with the findings of Asmaul Husna (2009) in bottle gourd.

4.4 Genetic Diversity Analysis

The genetic diversity of pumpkin advanced lines are presented in Table 6 to Table 10 and Figure 1 to 3.

4.4.1 Principal Component Analysis (PCA)

Principal component analysis was carried out with 24 genotypes of pumpkin. The computed Eigen values for the 13 variables subjected to principal component analysis together with the corresponding proportion and cumulative explained variance are given in Table 6. Following the Proportion of Variance Criterion, only one principal components was retained and these are the principal components whose cumulative explained variances was equal to or more than 99%. In conclusion, the principal component analysis resulted in the reduction of the 13 original variables to three independent linear combination, principal component of variables. The first principal component accounted for 26.54 % of the total variation while principal components two and three accounted for 18.56 % and 17.48 %, respectively (Table 6).

Table 6. Eigen value, % variance and cumulative (%) total variance of the principal components

Traits	Eigen values	Percent variation	Cumulative % of Percent variation
Leaf length	3.45	26.54	26.54
Leaf breath	2.41	18.56	45.10
Internode distance	2.27	17.48	62.58
Days to first male flowering	1.34	10.31	72.89
Days to first female flowering	1.06	8.13	81.02
Pediceal length of male	0.68	5.22	86.24
Pediceal length of female	0.53	4.10	90.34
Number of male flower	0.44	3.38	93.72
Number of female flower	0.25	1.90	95.62
Fruit length	0.23	1.74	97.36
Fruit breath	0.19	1.48	98.84
Fruit weight (kg)	0.10	0.76	99.60
Fruit yield (kg)	0.05	0.40	100.00

4.4.2 Non-Hierarchical Clustering

Twenty four pumpkin genotypes were grouped into five different clusters non-hierarchical clustering (Table 7). The clustering pattern of the genotypes obtained through principal component analysis. Kundu *et al.*, (2012) estimated 36 genotypes of bitter gourd and genotypes were grouped into six distinct clusters. Khatun *et al.*, (2010) conducted an experiment in 38 snake gourd genotypes and the genotypes were grouped into four different clusters. Asmaul Husna (2009) reported five clusters in bottle gourd. Gaffar (2008) reported similar number of clustering in fifteen sponge gourd genotype. In this study cluster III had the maximum number of genotypes eight, Cluster I and cluster II constitute equal number of five genotypes. Cluster IV and cluster V also had equal number 3 genotypes (Table 7).

Cluster III had G₂ (BD 10141), G₁₀ (BD 4371), G₁₁ (BD 4357), G₁₂ (BD 4350), G₁₉ (BD 4372), G₂₀ (BD 4391), G₂₁ (BD 4389) and G₂₂ (BD 4354). Cluster I and cluster II accumulated G₃ (BD 4361), G₄ (BD 4353), G₇ (BD 4349), G₉ (BD 4370) and G₁ (BD 10149), G₅ (BD 10137), G₆ (BD 10058), G₈ (PRONROY F1) and G₁₇ (BD 4363) respectively. Cluster IV consisted G₁₆ (BARMASI), G₂₃ (BD 4393) and G₂₄ (PRITY F1). Cluster V constituted by G₁₃ (BD 4352), G₁₅ (BD 10366) and G₁₈ (BD 10134).

Among the thirteen genotypes cluster V estimated the maximum cluster mean value for leaf length without petiole (16.10), leaf breath (22.59), and internode distance (14.22). (Table 8).

In cluster IV were highest mean value for days to first male flowering (54.45), pedicel length of male (25.78), number of male flower (9.67), fruit length (37.01), fruit breath (73.98), fruit weight (4.37), fruit yield (10.54). Cluster III produced maximum cluster mean for days to first female flowering (64.71), pedicel length of female (10.46). In cluster II produced highest mean value for number of female flowering (7.93).

Table 7. Distribution of genotypes in five different clusters

Cluster	Number of genotype	Genotype Number	Genotypes
I	5	G3, G4, G7, G9 and G14	4361, 4353, BD 4349, 4370 and 10078
II	5	G1, G5, G6, G8 and G17	10149, 10137, 10058, PRONROY f1 and 4363
III	8	G2, G10, G11, G12, G19, G20, G21 and G22	BD 10141, 4371, 4357, 4350, 4372, 4391, 4389 and 4354
IV	3	G16, G23 and G24	BARMASI, 4393 and PRITY f1
V	3	G13, G15 and G18	4352, 10366 and 10134

**Table 8. Cluster mean for thirteen yield and yield characters of pumpkin
Genotypes**

Traits	I	II	III	IV	V
Leaf length	15.31	14.42	14.53	15.69	16.10
Leaf breath	20.07	19.46	19.39	21.12	22.59
Internode distance	12.01	10.91	12.06	11.41	14.22
Days to first male flowering	53.87	53.87	54.42	54.45	54.33
Days to first female flowering	63.14	63.20	64.71	64.67	62.78
Pedicle length of male	20.00	22.67	22.27	25.78	23.00
Pedicle length of female	8.27	9.97	10.46	10.42	8.06
Number of male flower	8.47	9.13	8.83	9.67	8.89
Number of female flower	6.93	7.93	6.33	6.33	6.67
Fruit length	35.36	28.50	24.61	37.01	31.38
Fruit breath	65.20	54.60	42.82	73.98	48.95
Fruit weight (kg)	2.96	2.57	2.38	4.37	3.16
Fruit yield (kg)	5.74	6.87	5.33	10.54	6.34

4.3 Canonical Variant Analysis (CVA)

Canonical variant analysis was completed to compute the inter-cluster distances. The intra and inter-cluster distance (D^2) values were shown in Table 9. In this experiment, the inter-cluster distances were higher than the intra-cluster distances thus indicating broader genetic diversity among the genotypes of different groups. The highest inter cluster distance was observed between clusters III and IV (32.97), followed by between cluster IV and V (22.8), cluster I and cluster III (22.16), cluster II and IV (21.67), and between cluster I and IV (13.57). (Figure 1). Meanwhile, the maximum inter-cluster distance was observed between the Clusters III and IV (32.97), indicating genotypes from these two clusters, if involved in breeding program may produce a wide spectrum of segregating population. On the other hand, the highest intra-cluster distance was found in cluster V (4.58), which contained of 4 genotypes, while the lowest distance was found in cluster III (1.24) that comprises 2 genotypes. From figure 2 it can be concluded that different multivariate analysis was completed with suggested technique.

Following the scatter diagram all the genotypes were apparently distributed into five clusters. It is apprehended that the highest number of heterosis will be manifested in cross combination involving the genotypes belonging to most divergent clusters. It is suggested for future plant breeder to achieve higher level of production in addition to high heterosis.

Table 9. Intra (Bold) and inter cluster distances (D^2) for 24 genotypes

	I	II	III	IV	V
I	1.77				
II	10.42	1.88			
III	22.16	11.81	1.24		
IV	13.57	21.67	32.97	2.62	
V	14.45	5.83	10.78	22.80	4.58

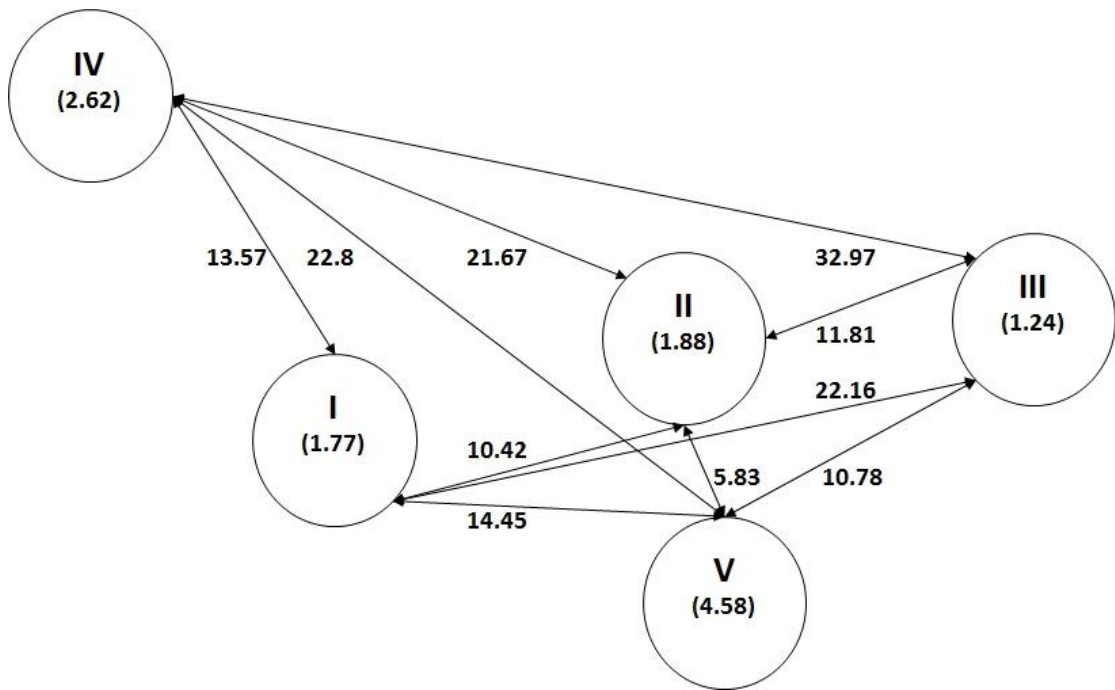


Fig 1. Cluster diagram showing the average intra and inter cluster distances ($D = \sqrt{D^2}$ Values) of 24 pumpkin genotypes.

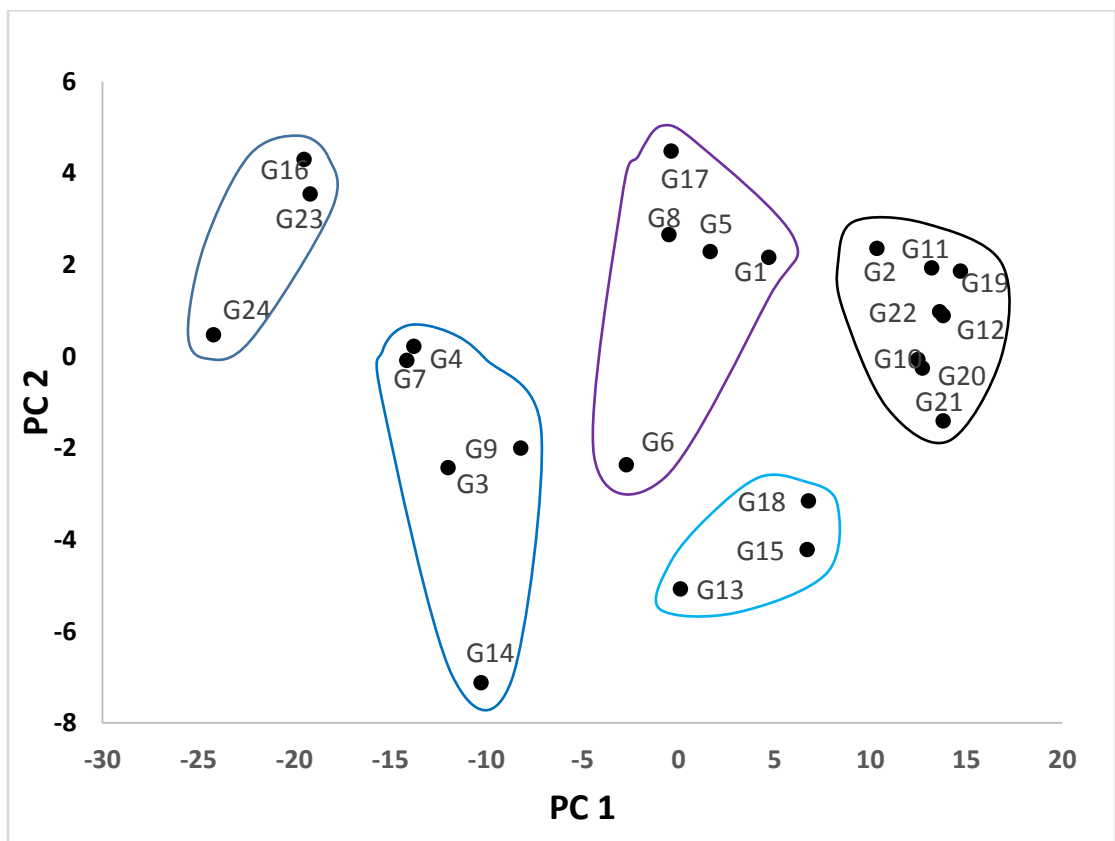


Fig 2. Scattered diagram of twenty four pumpkin genotypes

4.4.4 Contribution of Phenotypic traits towards divergence of the genotypes

In Table 10 the values of Vector I and Vector II are presented. Character assembled in vector-1 that were major contribution to the genetic divergence like leaf breath, internode distance, days to first female flowering, pedicel length of female. In vector-2 the important characters responsible for genetic divergence were internode distance, days to first male flowering, days to first female flowering, pedicel length of female flower (cm), number of male flower, number of female flower, fruit length (cm), fruit breadth (cm). Both vectors negative value determined the lower contribution of genetic divergence like leaf length, pedicel length of male flower, fruit weight and fruit yield (kg). Kundu *et al* (2012) found days to first male flowering and fruit yield per plant in both vectors is important components in genetic divergence of bitter gourd.

Table 10. Relative contribution of thirteen characters towards divergence of the genotypes

Traits	Vector-1	Vector-2
Leaf length	-0.8453	-0.0204
Leaf breath	0.1611	-0.4903
Internode distance	0.2824	0.0027
Days to first male flowering	-1.7596	0.4082
Days to first female flowering	0.7959	0.1056
Pedicle length of male	-0.1763	-0.7151
Pedicle length of female	0.2606	0.4066
Number of male flower	-0.241	0.3181
Number of female flower	-0.372	0.0767
Fruit length	-0.1314	0.2278
Fruit breath	-0.8021	0.2013
Fruit weight (kg)	-2.1504	-4.4036
Fruit yield (kg)	-0.0165	-0.1715

4.4.5 Selection of parents for future hybridization

This is very important and tough scheme to select genotypes for future breeding purposes. Meanwhile, considering the magnitude of genetic distance, contribution of character towards divergence, magnitude of cluster mean and phenotypic traits the genotype G_1 (BD 10149) for minimum days to first female flowering from cluster II, G_{23} (BD 4393) for maximum number of fruit yield and for maximum number of female flowering from cluster II, G_1 (BD 10149) for maximum fruit breadth from cluster IV, G_{24} (PRITY F1) for maximum fruit weight from cluster IV. So considering group distance and phenotypic performances the inter genotypic crosses between G_1 (BD 10149) and G_{24} ; G_1 (BD 10149) and G_{23} (BD 4393); G_{23} (BD 4393) and G_{24} (PRITY F1) may be suggested for future hybridization program.

CHAPTER V

SUMMARY AND CONCLUSION

The present study was carried out at the research field of Sher-e-Bangla Agricultural University, Sher-e-Bangla Nagar, Dhaka-1207, to evaluate the pumpkin genotypes based on phenotypic traits during the period from March 2016 to July 2016.

Randomized Complete Block Design (RCBD) with three replications was used to complete the field experiment. It was found that the existing genotypes have numerous significant variation among morphogenic traits. The highest value in respect to was recorded 17.50 cm G_6 (BD 10058) lowest was 11.67cm recorded in G_{17} (BD 4363). In case of leaf breadth, G_6 (BD 10058) recorded maximum 24.00 cm leaf breadth and minimum (15.73 cm) was recorded in G_{17} (BD 4363). Genotype G_{21} (BD 4389) recorded maximum internode distance (BD 16.33) and G_{17} (BD 4363) recorded minimum (7.90 cm). The highest value in respect to days to first male flowering was recorded as 56.00 in G_{23} (BD 4393) and the minimum duration was 52.67 in G_{24} (PRITY F1). Genotype G_{11} (BD 4357) recorded maximum duration of female flowering (66.33) and the minimum duration was 60.67 recorded in G_1 (BD 10149). Genotype G_{16} (BARMASI) recorded maximum pedicel length (28.33 cm) of male flower and minimum was 15.67 cm recorded in G_{14} (BD 10078). Genotype G_{16} (BARMASI) recorded 11.83 cm maximum pedicel length of female flower and minimum was 6.17 cm recorded in G_3 (BD 4361). Genotype G_1 (BD 10149) recorded maximum number of male flower (12) and the minimum number was 6.33 in G_{18} (BD 10134). The G_1 (BD 10149) recorded maximum number of female flower (12) and the minimum number was 4.33 in G_{21} (BD 4389). Genotype G_{24} (PRITY F1) recorded maximum fruit length (39.00 cm) and the minimum number was 22.37 cm in G_2 (BD 10141).

Genotype G₂₄ (PRITY F1) recorded the maximum fruit breadth (76.60 cm) and the minimum number was 41.37 cm in G₁₂ (BD 4350). In case of fruit weight G₂₄ (PRITY F1) was recorded maximum weight (4.67 kg) and the minimum fruit weight (1.90 kg) recorded in G₂ (BD 10141). Genotype number G₂₃ (BD 4393) recorded maximum average fruit yield (11.30 kg) per plant and the minimum fruit yield per plant was (4.27 kg) found in G₁₂ (BD 4350).

Here in this study the genotypic variance was lower than phenotypic variance in all the characters, indicating greater influence of environment on the expression of these characters. The maximum difference between phenotypic and genotypic co efficient of variation were 22.36% and 11.40%, which indicated that internode distance mostly dependent on environmental effect. The top most heritability estimates among thirteen yield contributing characters were 99.23%, 98.01%, 97.95%, 95.32%, 93.03%, in fruit breadth per plant, fruit weight per plant, fruit length per plant, pedicel length of female flower, pedicel length of male flower.

The lowest heritability was 5.37% in days to first female flowering. The maximum genetic advance was observed in fruit breadth (28.62), followed by fruit length of (13.02) among thirteen character of pumpkin genotypes. The maximum genetic advance in percent of mean was observed for fruit yield per plant (66.62%) and the lowest was in days to first male flowering (0.33 %).

Somehow the phenotypic correlation co-efficient were higher than their corresponding genotypic correlation co-efficient suggesting that both environmental and genotypic correlation in these cases acted in the same direction and finally maximize their expression at phenotypic level. The significant positive correlation with fruit yield per plant were found in fruit length ($G = 0.525$, $P = .499$), fruit breadth ($G = .663$, $P = .628$) and fruit weight ($G = 0.770$, $P = 0.773$).

According to the result of Path co-efficient analysis showed that days to first female flowering had highest positive direct effect (0.80) on yield per plant followed by internode distance (.137), pedicel length of male (0.72), pedicel

length of female (0.24), number of female flower per plant (.181), fruit breadth (.159) and fruit weight (0.76). So it was undoubtedly clear that improvement of pumpkin yield would be influenced by these character. On another view, number of male flower per plant (.125) and fruit length (.126) indicated the direct selection based on these characters would not be effective. Days to first female flowering via pedicel length of female flower had highest positive indirect effect (0.92). The highest negative indirect effect (-0.98) was fruit weight via fruit length. Residual effect was 0.47, which indicated that some other characters were responsible for contribution to yield per plant.

By using GENSTAT computer software program, pumpkin genotypes diversity was recorded through Principal Component Analysis (PCA), Canonical Variate Analysis (CVA), and cluster analysis. Twenty four pumpkin genotypes were grouped into five different clusters where cluster I and II both comprised five genotypes, cluster III comprised eight and cluster IV, V both comprised three genotypes. The maximum inter cluster distance was observed between clusters III and IV (32.97), followed by between cluster IV and V (22.8), cluster I and cluster III (22.16), cluster II and IV (21.67), and between cluster I and IV (13.57). On the other hand, the highest intra-cluster distance was found in cluster V (4.58), while the lowest distance was found in cluster III (1.24).

According to genetic distance, contribution of character towards divergence, magnitude of cluster mean and agromorphgenic traits the genotype G_1 (BD 10149) for minimum days to first female flowering from cluster II, G_{23} (BD 4393) for maximum number of fruit yield and for maximum number of female flowering from cluster II, G_1 (BD 10149) for maximum fruit breadth from cluster IV, G_{24} (PRITY F1) for maximum fruit weight from cluster IV. So considering group distance and agromorphgenic performances the inter genotypic crosses between G_1 (BD 10149) and G_{24} ; G_1 (BD 10149) and G_{23} (BD 4393); G_{23} (BD 4393) and G_{24} (PRITY F1) may be suggested for future hybridization program.

REFERENCES

- Abusaleha and Dutta, O. P. (1990). Studies on variability, heritability and scope of improvement in cucumber. *Haryana J. Hort. Sci.* **19**(3): 349-352.
- Ahamed, K.U., Akhter, B., Islam M. R., Ara N. and Humauan, M.R. (2011). An assessment of morphology and yield characteristics of pumpkin (*Cucurbita moschata*) genotypes in northern Bangladesh. *Tropic. Agril. Res. ext.* **14**(1).
- Akter, S., Rasul, M. G., Aminul, A. K. M. I and Hossain, M. M. (2013). Genetic variability, correlation and path coefficient analysis of yield and quality traits in pumpkin. (*Cucurbita moschata* Duch ex Poir). *Bangladesh J. Plant. Breed. Genet.* **26**(1): 25-33.
- Arora, S., Pandita M. L., Pratap P. S. and Sidhu A. S. (1983). Variability and correlation studies in sponge gourd (*Luffa cylindrica*). *Haryana Agric. Univ. J. Res.* **13**(1): 146-149.
- Badade, D. S., Warade, S. D. and Gaikwad, S. K. (2001). Correlation studies in bottle gourd. *J. Maharashtra Agric. Univ.* **26**(1): 20-22.
- Banik, B. R. (2003). Variability, gene action and heterosis in snake gourd (*Trichosanthes anguina* L.). Ph.D. Thesis, BSMRAU, Gazipur.
- BBS. (Bangladesh Bureau of Statistics). 2015. Statistical Year Book of Bangladesh 2014-2015. Bangladesh Bureau of Statistics Division, Ministry of Planning, Govt. of the People's Republic of Bangladesh, Dhaka.
- Bevec, f, Grobelnik, M. S., Rozman, C, Bovec, M. (2007). "Oil Pumpkins: Niche for Organic Producer", Purdue University Agriculture, Horticulture and Landscape Architecture.
- Bharathi, L. K., Naik G., and Dora, D. K. (2005). Genetic divergence in snake ground. *Veg. Sci.* **32**(2): 179-181.
- Bharathi, L. K., Naik, G., and Dora, D. K. (2006). Studies on genetic variability in gourd. *Indian J. Hort.* **63**(1): 96-97.
- Burton, G. W. (1952). Quantitative inheritance in Grasses. *Int. Grassland Cong.* **1**: 277-283.
- Chaudhury, M. L. and Mandal, G. (1987). Correlation and path analysis in cucumber (*Cucumis sativus* L.). *Haryana J. Hort. Sci.* **16** (3&4): 269-273.
- Chezhiyan, N. (1984). Studies on the performance of certain indigenous varieties of watermelon (*Citullus vulgaris*). *South Indian Hort.* **32**(3): 163-166.

- Chowdhury, D. and Sharma, K. C. (2002). Studies on variability, heritability, genetic advance and correlation in ridge gourd (*Luffa acutangula* Roxb.). *Hort. J.* **15**(3): 53-58.
- Dewey, D. R. and Lu, K. H. (1959). A correlation and path coefficient analysis of components of crested wheat grass seed production. *Agron. J.* **51**: 515-518.
- Dora, D. K., (2001). Genetic divergence in pointed gourd (*Trichosanthes dioica*). *Veg. Sci.* **28**(2): 170-171.
- Dora, D. K., Behera, T. K., Acharya, G. C., Motapatra, P., and Mishra, B. (2003). Genetic variability and character association in pointed gourd (*Trichosanthes dioica*). *Indian J. Hort.* **60**(2): 163-166.
- Fayeun, L. S. 1., Odiyi A. C., Makinde, S. C. O. and Aiyelari O. P. (2012). Genetic variability and correlation studies in the fluted pumpkin (*Telfairia occidentalis*). *J. Plant Breed. Crop Sci.* **4**(10): 156-160.
- Gaffar, A. (2008). Characterization and genetic diversity of sponge gourd (*Luffa cylindrica* L.). MS Thesis, SAU, Dhaka-1207.
- Gaur, P., Gupta, C. and Kishore, H. (1978). Studies on genetic divergence in potato. *Euphytica.* **27**: 361-368.
- Griffing, B. and Lindstone, E. W. (1954). A study of combining abilities of corn in breeds having varying proportion of corn germplasm. *Agron. J.* **46**(4): 454-552.
- Grubben, G. J. H. (2004). Vegetables, backhuys publishers, Leiden. pp: 354-357.
- Hanson, C. H., Robinson, H. F. and Comstock, R. E. (1956). Biometrical studies of yield in segregation population of Korean Lespedza. *Agron J.* **48**: 268-272.
- Harhsawardhan, C. and Ram, H. H. (2003). Genetic diversity studies in musk melon. *Ann. Agril. Res.* **24**(2): 345-349.
- Hazra, P., Ghosh, R. and Nath, S. (2003). Identification of important yield components in pointed gourd (*T. anguina* Roxb). *Crop Res. Hisar.* **25**(2): 244-252.
- Husna, A. Mahmud, F., Islam, M. R., Mahmud, M. A. A. and Ratna, M. M. (2011). Genetic Variability, correlation and path co-efficient analysis in bottle gourd (*Lagenaria siceraria* L.). *Adv. Biol. Res.* **5**(6): 323-327.
- Husna, A., Maih, M. A., Begum, S., Shilpi, S. Z. and Islam, M. R. (2014). Genetic variability, correlation and path co-efficient analysis based on vegetative characters in bottle gourd (*Lagenaria Siceraria* L.). *Adv. Agric. Biol.* **3**(1): 8-12.
- Husna, A. (2009). Genetic diversity, correlation and path co-efficient analysis in bottle gourd (*Lagenaria siceraria* L.). MS Thesis. SAU, Dhaka-1207

- IPM, CRSP, (2004). Technical Bulletin, Bangladesh Agricultural Research Institute, Gazipur, Bangladesh.
- Islam, M. R., Faruquei, M. A. B., Bhuiyan, M. A. R., Biswas, P. S. and Salam, M. A. (2004). Genetic diversity in irrigated rice. *Pakistan J. Biol. Sci.* **7**(2): 226-229.
- Islam, M. R., Hossain, M. S., Bhuiyan, M. S. R., Hasan, G. N., and Syed, A. (2010). Multivariate analysis of bitter melon (*Momordica charantia* L.). *J. Sci. Res.* **5**(2): 86-90.
- Johnson, H. W., Robinson H. F. and Comstock R. E. (1995). Estimation of genetic and environmental variability in soybean. *Agron. J.* **47**: 314-318.
- Kabir, M. E. (2007). Genetic variability, correlation and path analysis of pointed melon (*Trichosanthes dioica* Roxb.) MS Thesis. SAU, Dhaka-1207.
- Karuppaiah, P., Kavitha, R. and Senthilkumar, P. (2005). Divergent analysis in bitter melon (*Momordica charantia*). *Indian. J. Hort.* **46**(2): 314-319.
- Khan, M. A. S., Kabir, M. Y. and Alam, M. M. (2009). Variability, correlation path analysis of yield and yield components of pointed Melon. *J Agric Rural Dev.* **7**(1&2): 93-98.
- Khan, A S, M. R., Rabbani, M. G., Siddique, M. A. and Hossain, M. I. (2008). Study on genetic diversity of pointed melon using morphological characters. *Bangladesh J. Agril. Res.* **33**(3): 607-616.
- Khatun, R. M. and Rahnman. M. G. (2010). Estimation of genetic diversity in snake melon (*Trichosanthes cucumerina*). *Bangladesh J. Agril. Res.* **35**(1): 95-100.
- Khule, A. A., Tikka, S. B. S., Jadhav, D. J. and Kajale, D. B. (2011). Correlation and path coefficient analysis in sponge melon (*Luffa cylindrica*). *Int. J. Plant Sci.* **6**(2): 277-279.
- Kumar, S. Singh, R. and Pal, A. K. (2007). Genetic variability, heritability, genetic advance, correlation coefficient and path analysis in bottle melon (*Lagenaria siceraria* L.). *Indian J. Hort.* **64**(2): 163-168.
- Kumaran, S. S., Natarajan, S. and Thamburaj, S. (1998). Correlation and path analysis studies in pumpkin (*Cucurbita moschata* Poir.). *South Indian Hort.* **46**(3): 138-142
- Kumaresan, G. R., Makesh, S., Ramaswamy, N. (2006). Character association and path coefficient studies in snake melon (*Trichosanthes anguina* L.). *Res. Crops.* **7**(2): 510-513.
- Kundu, B. C., Hossain, M. M., Khaleque, M. A. and Mian, I. H. (2012). Genetic divergence in bitter melon (*Momordica charantia* L.) *J. Asiat. Soc. Bangladesh, Sci.* **38**(2): 125-134.

- Li, J. W., Sun, S. R. and Rer, Y. H. (1997). Study on genetic correlation, path analysis of the main agronomic characters of cucumber. *Acta. Agril. Univ.* **31**(3): 244-24.
- Lu, A. and Jeffrey, C. (2015). *Cucurbita linnaeus. Flora of China*. Retrieved.
- Mabberley, D. J. (2008). *The Plant Book : A portable dictionary of the vascular plants*. Cambridge: Cambridge University Press. pp.235.
- Mahalanobis, P. C. (1936). On the generalized distance in statistics. *Proc. Nat. Inst. Sci., India.* **2**: 49-55.
- Mangal, J. L., Dixit, J., Pandita, L. and Sidhu, A. S. (1981). Genetic variability and correlation studies in bitter gourd (*Momordica charantia* L). *Indian. J. Hort.* **38**: 94-98.
- Mannan, M. A. (1992). Studies on variability, genetic parameters and correlation in bitter gourd. M.S. Thesis, BAU.
- Mansour, E. H., Dworschak. E., Erno, L., Andrea., B. B. and Gergely, A. (1993). Nutritive value of pumpkin (*Cucurbita Pepo*) seed products. *J. Sci. Food Agric.* **61**(1): 73-78.
- Masud, M. A. T., Chauwdhury, M. A. Z., Hossain, M. A. and Hossain, S. M. M. (1995). Multivariate analysis in pumpkin (*Cucubita moschata*). *Bangladesh J. Plant. Breed. Genet.* **8**(1 &2): 45-50.
- Masud, M. A. T., Rashid, M. H., Chowdhury, M. A. Z. and Uddin, M. S. (2006). Combining ability and heterosis in bottle gourd (*Lagenaria siceraria*). *Bangladesh Plant. Breed. Genet.* **19**(1): 41-44.
- Mathew, S. S. and Khader, K. M. A. (1999). Genetic studies in snake gourd (*Trichosanthes anguina* L.). *J. Trop. Agril.* **37**(1-2): 71-72.
- Mendlinger, S, Chweya, J, Benzioni, A, Seme, A, Ventura, M, Lungaho, C, and Okoko, V, (1991). Collection, Evaluation and Breeding of African Edible Vegetables. BGUN-ARI-25-92. Annual Report (1991) on AID-CDR Program, Beer-Sheva, Israel.
- Miah, M. A., Rahman, M. M., Uddin, M. S., Rahman, A. K. M. M. and Ullah. M. H. (2000). Genetic association in bitter gourd *Bangladesh. J. Sci. Tech.* **2**(1): 2125.
- Miller, P.J., Williams, J. C., Robinson, H. F. and Comstock, R. E. (1958). Estimation of genotypic and environmental variance and co-variance in upland cotton and their implication in selection. *Agron. J.* **50**: 126-131.
- Mohanty, B. K. (2001). Studies on correlation and path analysis in pumpkin (*Cucurbita moschata*). *Haryana J. Hort. Sci.* **30**(1-2): 86-89.
- Mondal, SN., Rashid, A., Inouve, K., Hossain, A. K. M. A., and Hossain, M. A. (1989). Genetic variability, correlation and path co-efficient analysis in watermelon. *Bangladesh J. Plant Breed. Genet.* **2**(1): 31-35.

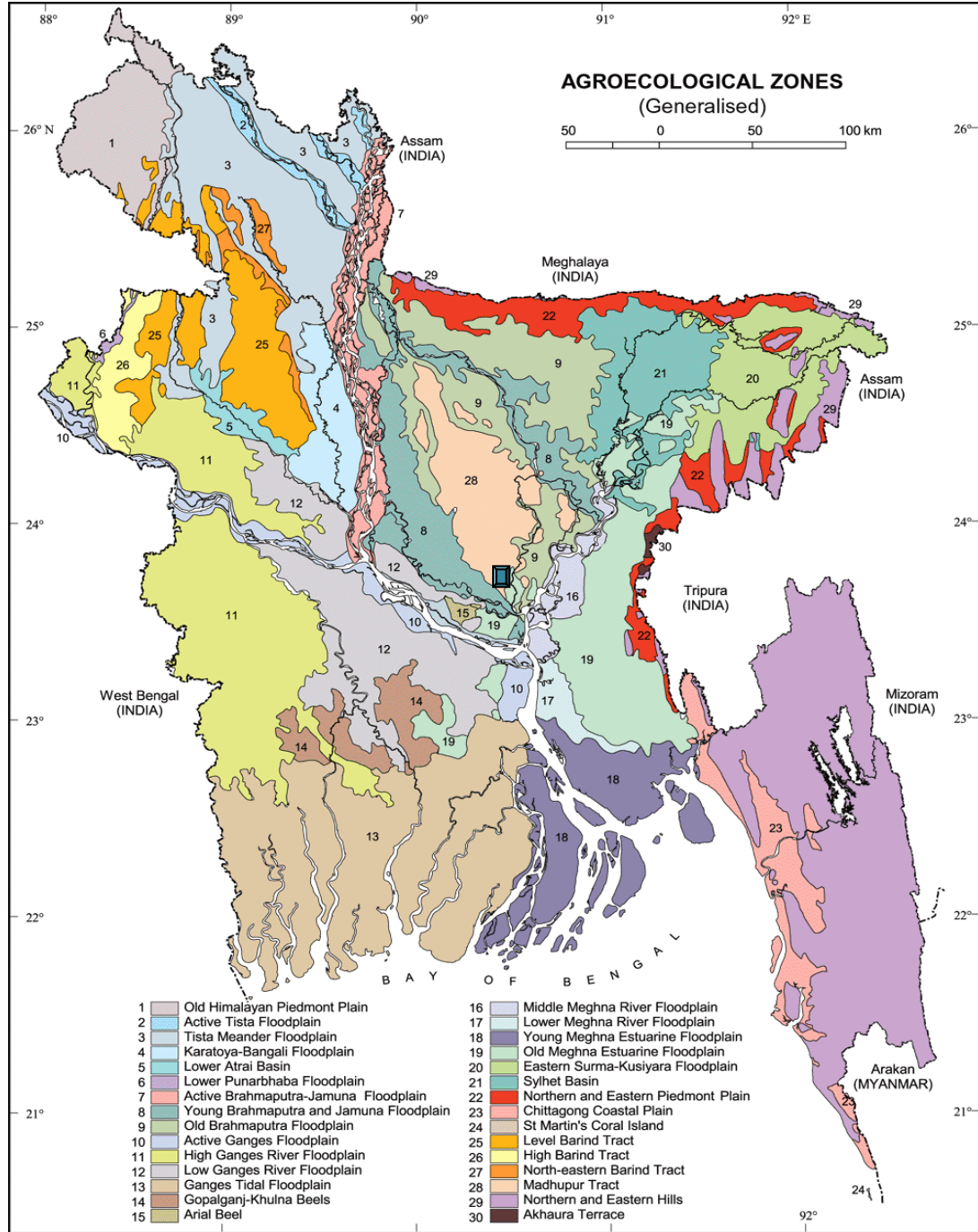
- Murty, B. R. and Arunachalam, I. J. (1966). The nature of genetic divergence in relation to breeding system in crop plants. *Indian J. Genet.* **26**: 188-198.
- Naik, A., Akhtar, S., Chattopadhyay, A. and Hazra, P. (2012). Study of genetic variability, heritability and genetic advance for fruit quality characters in Teasle gourd (*Momordica subangulata*). *African J. of Agric Res.* **7**(49): 650-652.
- Narayan, R., Singh, S. P., Sharna, D. K. and Rastogi, K. B. (1996). Genetic variability and selection parameters in bottle gourd (*Lagenaria siceraria* L.). *Indian J. Hort.* **53**(1): 53-58
- Narayanankutty, C., Sunanda, C. K., and Jaikumaran, U. (2006). Genetic variability and character association analysis in snake gourd. *Indian J. Hort.* **63**(4): 402-406.
- Parhi, G. G., Mishra, H. N. and Mishra, R. S. (1995). Correlation and path-coefficient studies in bitter gourd (*Momordica charantia* L.). *Indian J. Hort.* **52**(2):132-136.
- Prasad, V. S. R. K., and Singhi, D. P. (1990). Genetic variability and heritability studies in sponge gourd (*Luffa aegyptica*, Mill.). *Haryana J. Hort. Sci.* **19**(1- 2): 222-224.
- Prasanna, S. C., Krishnappa, K. S., and Reddy, N. S. (2002). Correlation and path coefficient analysis studies in ridge gourd. *Bangalore Agril. Univ. J. Res* **31**(9/10): 150-152.
- Preeti S., Singh, D. K., Damke, S. R., and Harshawardhan C. (2010). Genetic diversity in indigenous germplasm of ash gourd. *Indian J. Hort.* **67**: 208-213.
- Quamruzzaman, A. K. M., Rashid, M. A., Masud, M. A .T. and Uddin, M. N. (2009). Heterosis in bottle gourd. *Bangladesh J. Agril. Res.* **34**(3): 465-472.
- Rahman, M. M., Dey, S. K. and Wazuddin S. (1991). Study of yield, yield components and vine characters of some cucurbit genotypes. *BAU Res Progress* **5**: 75-85.
- Rajkumar, M., and Karuppaiah, P. (2007). Variability studies in snake gourd (*Trichosanthes anguina* L.). *Plant Archives.* **7**(2): 699-701.
- Ramchandran, C. and Gopalkrishnan. (1979). Correlation and regression studies in bitter gourd. *Indian J. Agric. Sci.* **49**: 850-854.
- Ramphall, N. and Gill H. S. (1990). Demand and Supply of Vegetables and Pulses in South Asia. Proc. Workshop Held at Islamabad, Pakistan on 24-29 September, 1990. AVRDC Publication No. 90. 331, AVRDC, Tainan, Taiwan.

- Rana, T. K., Vashistha, R. N., and Pandita. M. L. (1986). Genetic variability studies in Pumpkin (*Cucurbita moschata*). *Haryana J. Hort. Sci.* **15**(1-2): 71-75.
- Rao, M. N., Roy, P. V., and Reddy, B. M. M. (2000). Correlation and path analysis in segregating population of ridge gourd. *Crop Res. Hisar*, **20**(2): 338-342.
- Rashid, M. M. (1993). Vegetable Science (in Bangla). 1st ed., Bangla Academy. Dhaka. Bangladesh. pp.265-336.
- Saade, R., Hernandez L., Montes. S. 2013."Cucurbits". Purdue Horticulture pp.70-75.
- Saha, R. R., Mitra, B. N., Hossain, A. E., Jamaluddin, M. and Hoque, A.M.M.M. (1992).Genetic variability, character association and path coefficient analysis in pumpkin (*Cucurbita moschata* L). *Bangladesh Hort. J.* **20**(1): 59-62.
- Sarkar, S. K., Marity, T. K. and Som. N. G. (1999). Correlation and path coefficient studies in pointed gourd (*Trichosanthes dioica*). *Indian. J. Hort.* **56**(3): 252-255.
- Shah, S. R. and Kale, P. N. (2002). Yield component association analysis in ridge gourd. *J. Maharashtra Agril. Univ.*, **27**(2): 197-198.
- Sultana, S. (2011).Genetic diversity, correlation and path co-efficient analysis in sweet gourd. MS Thesis, Dept of Genet. Plant Breed., Sher-e-Bangla Agricultural University, Dhaka.
- Sharma, A., Vidyasagar, K. and Pathania, N. K. (2000). Studies on combining ability for earliness and marketable fruit yield in cucumber (*Cucumis sativus* L.). *Himachal. J. Agril. Res.* **26**(1 & 2): 54-61.
- Singh, G. and Lal, T. (2005). Correlation and path analysis of fruit yield and its component traits in muskmelon (*Cucumis melo* L.). *J. Crop improv.* **32**(1): 102-107.
- Singh, R. K. and Chaudhury, B. D. (1985). Biometrical methods of quantitative genetic analysis. *Haryana J. Hort. Sci.* **12**(2): 151-156.
- Singh, K. P., Panda, P. K. and Singh, A. K. (2002). Variability, heritability and genetic advance in ash gourd (*Benincasa hispida*). *Haryana J. Hort. Sci.* **31**(2): 139-140.
- Singh, R. V. and Ram, D. (2003). Correlation and path analysis in musk melon. *Indian J. Hort.* **50**(2): 124-128.
- Swamy, K. R. M., Dutta O. P., Ramachander, P. R. and Wahi, S. D. (1984). Interrelationships between yield and other quantitative characters in muskmelon (*Cucumis melo* L.) *South Indian Hort.* **32**(6): 334-339.
- Thakur, J. C. (1970). Genetic variability and correlation studies in watermelon. M. Sc. Thesis, Punjab Agricultural University, Ludhiana. India.

- Umamaheswarappa, P., Krishnappa, K. S., Murthy, P. V., Adivappar, N. and Muthu M. P. (2004). Correlation and path coefficient analysis studies in bottle gourd (*Lagenaria siceraria* (Mol) Standl.). *Env. Eco.* **22**(4): 636-640.
- Vashitha, R.N., Pratap, P. S. and Pandita, M. L. (1983). Studies on variability and heritability in watermelon (*Citrullus lanatus*). *Haryana Agric. Univ. J. Res.* **13**(2): 319-324.
- Vijay, O. P. (1987). Genetic variability, correlation and path-analysis in muskmelon (*Cucumis melo* L.). *Indian Hort.* **44**(4): 233-238.
- Yadav, M., Jain, S., Tomar, R., Prasad, G. B., and Yadav, H. (2010). Medicinal and biological potential of pumpkin: An Updated Review. *Nutri. Res. Rev.* **23**(2): 184–190.

APPENDICES

Appendix I. Map showing the experimental site under the study



The experimental site under study

Appendix II. Monthly records of air temperature, relative humidity, rainfall and sunshine hours during the period from March 2016 to July 2016

Month	Year	Monthly average air temperature (°C)		Average relative humidity (%)	Total rainfall (mm)	Total sunshine (hours)
		Maximum	Minimum			
March	2016	38.7	17.7	70	88	9.1
April	2016	34.9	22.3	72	207	7.9
May	2016	36.9	22.5	80	595	4.5
June	2016	35	23.7	83	625	3.7
July	2016	36	19.40	85	319	4.0

Source: Bangladesh Meteorological Department (Climate division), Agargaon Dhaka-1207.

Appendix III. The mechanical and chemical characteristics of soil of the experimental site as observed prior to experimentation (0 - 15 cm depth).

Mechanical composition:

Particle size	constitution
Sand	40%
Silt	40%
Clay	20%
Texture	Loamy

Chemical composition:

Soil characters	Value
Organic matter	1.44 %
Potassium	0.15 meq/100 g soil
Calcium	3.60 meq/100 g soil
Magnesium	1.00 meq/100 g soil
Total nitrogen	0.072
Phosphorus	22.08 µg/g soil
Sulphur	25.98 µg/g soil
Boron	0.48 µg/g soil
Copper	3.54 µg/g soil
Iron	262.6 µg/g soil
Manganese	164 µg/g soil
Zinc	3.32 µg/g soil

Source: Soil Resources Development Institute (SRDI), Khamarbari, Dhaka

Appendix IV. Mean performance of various growth parameter and yield components

Sl.	Genotype	LL	LB	ID	DFMF	DFFF	PLM	PLF	NMF	NFF	FL	FB	FW	YIELD
G1	10149	13.80	19.20	11.03	53.33	60.67	20.67	7.67	12.00	12.00	23.47	52.23	2.40	5.77
G2	BD 10141	12.93	21.50	10.83	53.67	64.00	21.33	8.67	7.00	7.33	22.37	46.40	1.90	7.60
G3	4361	16.83	20.00	11.67	54.33	64.67	21.00	6.17	7.33	5.67	35.47	65.47	2.70	5.43
G4	4353	14.67	19.77	11.67	53.67	62.67	21.33	10.17	7.67	6.00	34.03	68.13	2.73	5.43
G5	10137	14.43	19.07	10.47	54.67	64.33	24.33	10.83	7.33	8.00	29.40	53.17	2.53	5.90
G6	10058	17.50	24.00	13.00	53.67	64.33	21.67	10.67	9.00	7.33	30.67	57.10	2.67	6.87
G7	BD 4349	13.73	18.67	12.03	53.00	63.00	20.67	9.00	10.00	7.67	35.77	67.53	3.40	7.97
G8	Pronoy f1	14.70	19.30	12.17	54.00	63.00	23.33	11.33	9.00	5.33	27.67	56.13	2.37	7.10
G9	4370	14.73	20.23	11.20	55.33	62.67	21.33	7.33	7.33	8.00	34.43	61.93	2.53	4.30
G10	4371	15.07	19.83	13.00	55.33	65.33	22.67	10.67	10.00	4.67	25.57	43.13	2.17	5.00
G11	4357	14.27	19.90	10.07	54.67	66.33	21.33	10.83	8.00	7.33	23.40	43.23	2.57	4.97
G12	4350	14.20	18.70	12.00	53.67	65.00	25.00	10.17	12.00	8.67	26.37	41.37	2.53	4.27
G13	4352	14.53	21.97	13.90	53.33	62.67	21.00	7.67	8.33	7.67	33.83	52.93	3.30	5.33
G14	10078	16.57	21.70	13.50	53.00	62.67	15.67	8.67	10.00	7.33	37.10	62.93	3.43	5.57
G15	10366	17.07	23.73	13.50	54.67	63.67	25.00	7.67	12.00	7.67	32.13	46.00	2.90	7.77
G16	BARMASI	16.43	21.00	10.57	54.67	64.67	28.33	11.83	10.67	6.00	36.20	72.80	4.10	9.50
G17	4363	11.67	15.73	7.90	53.67	63.67	23.33	9.33	8.33	7.00	31.30	54.37	2.90	8.70
G18	10134	16.70	22.07	15.27	55.00	62.00	23.00	8.83	6.33	4.67	28.17	47.93	3.27	5.93
G19	4372	15.03	18.07	10.83	54.67	65.00	21.67	10.33	7.33	5.67	23.10	41.77	2.13	4.97
G20	4391	15.83	16.83	11.83	53.67	64.33	22.00	11.17	7.33	7.33	27.63	42.03	2.57	5.10
G21	4389	15.43	21.87	16.33	54.67	64.67	22.67	11.17	10.00	4.33	23.40	42.60	2.30	4.70
G22	4354	13.50	18.40	11.57	55.00	63.00	21.50	10.67	9.00	5.33	25.03	42.03	2.87	6.00
G23	4393	14.53	19.40	12.17	56.00	65.33	25.67	7.93	10.00	5.67	35.83	72.53	4.33	11.30
G24	PRITY f1	16.10	22.97	11.50	52.67	64.00	23.33	11.50	8.33	7.33	39.00	76.60	4.67	10.83

LL= Leaf length, LB= Leaf breath, ID= Internode distance, DFMF= Days to first male flowering, DFFF= Days to first female flowering, PLM= Pedicel length of male, PLF= Pedicel length of female, NMF= Number of male flower, NFF= Number of female flower, FL= Fruit length, FB= Fruit breath, FW= Fruit weight (kg) and YIELD= Fruit yield (kg)

Appendix V. Analysis of variance of 13 yield and yield contributing characters

Source	DF	Mean sum of square												
		LL	LB	ID	DFMF	DFFF	PLM	PLF	NMF	NFF	FL	FB	FW	YIELD
Genotype	23	6.03 ^{NS}	12.57 ^{NS}	9.07 ^{NS}	2.20 ^{NS}	4.87 ^{NS}	16.71 ^{**}	7.54 ^{**}	8.32 ^{**}	8.09 ^{**}	82.46 ^{**}	390.48 ^{**}	1.46 ^{**}	11.62 ^{**}
Replication	2	2.33 ^{NS}	2.09 ^{NS}	3.71 ^{NS}	2.06 ^{NS}	10.76 ^{NS}	0.05 ^{NS}	0.12 ^{NS}	2.18 ^{NS}	2.04 ^{NS}	1.44 ^{NS}	0.74 ^{NS}	0.21 ^{**}	0.56 ^{NS}
Error	46	3.4449	10.8859	5.3301	1.9541	4.3726	0.6031	0.1806	2.0211	1.9112	0.8550	1.5024	0.01	1.03

Z1-Z2 score

Genotype	PC1	PC2
G1	4.726	2.161
G2	10.357	2.359
G3	-11.987	-2.427
G4	-13.771	0.223
G5	1.665	2.292
G6	-2.695	-2.365
G7	-14.129	-0.083
G8	-0.474	2.657
G9	-8.197	-1.999
G10	12.493	-0.068
G11	13.203	1.934
G12	13.8	0.892
G13	0.121	-5.076
G14	-10.262	-7.12
G15	6.719	-4.214
G16	-19.491	4.3
G17	-0.358	4.485
G18	6.794	-3.15
G19	14.702	1.863
G20	12.725	-0.255
G21	13.81	-1.406
G22	13.617	0.976
G23	-19.166	3.55
G24	-24.203	0.473



Appendix vi : Field view of plants after transplanting of seedlings