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# Genetic Diversity and Combining Ability in Maize (*Zea mays* L.)

Haydar, F. M. Ali

University of Rajshahi

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# **GENETIC DIVERSITY AND COMBINING ABILITY IN MAIZE (*Zea mays* L.)**



**Ph.D. Thesis**

**By**  
**F. M. Ali Haydar**  
**M.Sc, M.Phil.**

**JULY 2013**

**BIOMETRICAL GENETICS LAB.  
DEPARTMENT OF BOTANY  
FACULTY OF LIFE AND EARTH SCIENCE  
UNIVERSITY OF RAJSHAHI  
RAJSHAHI-6205, BANGLADESH**

# GENETIC DIVERSITY AND COMBINING ABILITY IN MAIZE (*Zea mays* L.)



*A thesis submitted for the degree  
of  
Doctor of Philosophy  
in the  
Department of Botany  
University of Rajshahi, Bangladesh*

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*Under the guidance of*

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**Ph.D.  
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**GENETIC DIVERSITY AND COMBINING  
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**F.M. Ali Haydar**

**2013**

**Professor Nishit Kumar Paul**  
B.Sc. (Hons.), M.Sc.  
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## *CERTIFICATE*

*I hereby certify that the research work, entitled “GENETIC DIVERSITY AND COMBINING ABILITY IN MAIZE (Zea mays L.)” submitted for the degree of Doctor of Philosophy in the subject of Botany is a bonafide research work carried out by F.M. Ali Haydar under my supervision in the University of Rajshahi, Rajshahi-6205, Bangladesh. The results of investigation, which embodied here are original and have not been submitted before in substance for any other degree of this or any other university.*

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**(Dr. Nishit Kumar Paul)**  
Professor and Supervisor  
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*DEDICATED  
TO  
LATE PROFESSOR M. A. KHALEQUE  
AND  
MY DEPARTED PARENTS*





## **DECLARATION**

I hereby declare that the whole research work submitted as a thesis which is the result of my own original investigation for the fulfillment of the degree of Doctor of Philosophy in Botany, Faculty of Life and Earth Science at the University of Rajshahi, Bangladesh.

.....  
**(F. M. Ali Haydar)**  
Candidate

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## ABSTRACT

The present study was carried out during 2008/2009, 2009/2010 and 2010/2011 to obtain information about the performance of maize inbred lines, genetic diversity, gene action and assessment of the combining ability of parental lines and their F<sub>1</sub>s by using diallel fashion. Genetic variability analysis revealed that days to maturity, plant height and number of rows/cob had higher variability, heritability and genetic advance in percentage of mean. Broad-sense heritability estimates of the characters were higher in magnitude (66.08 to 93.41%) indicating greater genetic impact on these characters. Positive significant phenotypic and genotypic correlations were found for days to maturity, number of kernels/row and number of grains/cob. The path analysis revealed that number of rows/cob, cob length and number of kernels/row had the highest direct effect on grain yield, while plant height and ear height and number of grains/cob had the highest moderate indirect negative effects on grain yield. Selection indices were constructed through the discriminate function using nine characters. From the results, the highest relative efficiency was observed with the selection index based on four characters viz., ear height, number of rows/cob, number of kernels/row and grain yield/plant. Cob length, number of kernels/row and number of grains/cob could be the important selection criteria in the improvement of maize lines and hybrids for higher grain yield. The average inter-cluster was always higher than the average intra-cluster distance suggesting wider genetic diversity among the inbred lines of the groups. Cluster III had the highest intra and inter-cluster distance was maximum between clusters I and II. So, the inbred lines chosen from these clusters would give broad spectrum of variability in the segregating generation.

A half diallel set of six maize inbred lines were utilized to evaluate combining ability and heterosis for yield and its components characters. General and specific combining ability effects were significantly different among the parental lines. The parents P<sub>5</sub> and P<sub>4</sub> were considered suitable according to their yield capacities

and general combining ability effects. The variances for general (gca) and specific (sca) combining ability for plant height, cob diameter and number of rows/cob were highly significant indicating the presence of additive as well as non-additive gene effects for controlling the characters.

In the present study the values of F for all the characters except cob diameter were positive and greater than zero, which expressed that dominant alleles were more frequent than recessive alleles. The estimate of additive genetic variance (D) was significant for days to silking, days to maturity, ear height and number of rows/cob indicating the importance of additive gene effect in their inheritance. Thus highly significant values of the components D, H<sub>1</sub> and H<sub>2</sub> indicated the importance of both additive and dominant gene effect for the characters under study. The ratio of  $[4DH_1]^{1/2} + F / [4DH_1]^{1/2} - F$  estimates the relative proportion of dominant and recessive alleles in the parents. From W<sub>r</sub>-V<sub>r</sub> graph it has been noticed that expression of dominant and recessive alleles in the parents was influenced by environment as the same parent showed different positions on graphs. From this study, it is concluded that parents with recessive and dominant genes can also contribute towards high yield.

The mid parent heterosis values ranged from 1.73% (P<sub>1</sub>×P<sub>6</sub>) to 5.21% (P<sub>1</sub>×P<sub>2</sub>) whereas, the useful heterosis values varied between -40.36% (P<sub>1</sub>×P<sub>5</sub>) and 4.94% (P<sub>1</sub>×P<sub>2</sub>), and only five crosses had higher grain yield. Of these crosses, P<sub>1</sub>×P<sub>2</sub>, P<sub>2</sub>×P<sub>5</sub>, P<sub>4</sub>×P<sub>5</sub> and P<sub>5</sub>×P<sub>6</sub> were considered promising hybrids and will be tested in yield trials for further evaluation. Taking the lines of these promising crosses into account, parents P<sub>5</sub> and P<sub>4</sub> may be used as parents in hybrid maize programs. In addition to these parents, P<sub>3</sub> with dominant genes, high yield and general combining ability may be recommended as another parent.



## ACRONYMS

|   |   |                                   |
|---|---|-----------------------------------|
| BBS   | = | Bangladesh Bureau of Statistics   |
| cm  | = | centimeter                        |
| df  | = | degrees of freedom                |
| <i>et al.</i>   | = | and others                        |
| FAO   | = | Food and Agriculture organization |
| Fig.  | = | Figure                            |
| g   | = | Gram                              |
| ha  | = | hectare                           |
| i.e   | = | that is                           |
| m   | = | Meter                             |
| m <sup>2</sup>  | = | Square Meter                      |
| mm  | = | Millimeter                        |
| S   | = | Significant                       |
| R (R <sub>1</sub> , R <sub>2</sub> , R <sub>3</sub> ) | = | Replication                       |
| NS  | = | Not Significant                   |
| t   | = | Ton                               |
| viz.  | = | Namely                            |
| %   | = | Percentage                        |
| /   | = | Per                               |

## INTRODUCTION

Maize (*Zea mays* L.;  $2n=20$ ) is an important cereal crop with high yield potential. Maize stands third among the cereal crops in Bangladesh after rice and wheat (BBS, 2009). It can be grown throughout the year because of its photo-insensitiveness. World wide maize ranks first in terms of production and yield but third in terms of area, next to wheat and rice (FAO, 2009). In Bangladesh, maize has emerged as a third important cereal due to its versatile uses. On global front, maize has gained tremendous importance due to rising demand from diversified sectors like food, feed and ethanol production. Maize is considered the third cereal crop after rice and wheat all over the world for production and consumption. In addition to its use as a human food, it is also utilized as a poultry and livestock feed and also as a fodder (White and Johnson, 2003). Maize is used as staple food in many countries of the world although its uses as human food are very limited in Bangladesh. Maize plays a significant role in human and livestock nutrition world wide (Bantle and Prasanna, 2004). In Bangladesh, area, production and yield of maize decreased by 2.9%, 3.59% and 0.69%, respectively from the year 1967-68 to 1986-87 due to utilization of traditional variety (Mohiuddin, 2003).

Introduction of hybrid varieties and appropriate management practices increased area, production and yield by 19.83%, 34.40% and 14.56%, respectively from the year 1987-88 to 2003-2004 (Moniruzzaman *et al.*, 2007). Now maize has become an important cereal in terms of yield (Maize: 5.36., wheat: 2.21; and rice: 2.15 ton/ha., Anonymous, 2003) but in terms of area and production, it could be good source of nutrients for under-nourished and mal-nourished populations in Bangladesh. From the trends of its increasing demand, yield, acreage and production in recent years it seems that maize is going to hold the second position next to rice in a few years.

Although maize research started in the early 1970's, its acreage and production did not increase much until the mid 1990's. With the introduction of hybrid varieties and recent growth of poultry industries, maize cultivation has expanded faster than any crop in the past. The present production is above 23 million tons from around 4 million hectares of land producing at a rate of 6 tons per hectare (Rashid *et al.*, 2010). With the introduction of hybrid maize varieties in the country the prospect of maize cultivation has become bright. Maize is a unique crop because of its versatile use and low cost per unit production. Maize is consumed either directly or indirectly by millions of people. About 75% of maize is fed to animal, thus, indirect consumption is greater than direct consumption. Demand of maize is likely to progressively increase in near future. In order to fulfill the demand of additional food and to maintain self-sufficiency in food of Bangladesh, maize can be considered as a supplementary food to rice.

The development of hybrid varieties and production of hybrid maize seeds is therefore, very important. Hybrid seed production requires development and selection of suitable inbred parents. Moreover, due to industrialization, urbanization and river erosion, 221 hectares of crop land is losing every day (Banik *et al.*, 2009). So, production of more food from limited land is essential and it is not possible to get required quantity of food from rice and wheat only. Maize is the crop which can fulfil the demand. Bangladesh is a rice consuming country. People of Bangladesh have changed their food habit to some extent and consuming wheat also. The climatic condition of Bangladesh is suitable for maize cultivation. That is why the acreage, production and yield of maize is increasing steadily and maize is becoming important crop in Bangladesh.

In working towards this goal, particular attention is paid to grain yield as the most important agronomic characteristic. Grain yield is a complex quantitative trait that depends on a number of factors. It is under great influence of environmental

conditions, has complex mode of inheritance and low heritability (Bocanski *et al.*, 2009). Most of the yield components are less complex, and because of that using some other traits which are highly correlated with grain yield and has higher heritability, would make the selection of the best progenies more reliable (Vasic *et al.*, 2001; Bekavac *et al.*, 2007, 2008). Because of that during selection of grain yield, in order to select the best individuals, we need to determine the mean values, components of variance and heritability of the studied traits. Besides, knowing the correlations between the characters is also of great importance for success in selection to be conducted in breeding programs, and analysis of correlation coefficient is the most widely used one among numerous methods that can be used (Yagdi and Sozen, 2009). Because correlation coefficient measures the mutual association only between a pair of variables, when more than two variables are involved, the correlations *per se* may not provide a clear picture of the importance of each component in determining grain yield. The appropriate knowledge of such interrelations between grain yield and its contributing components can significantly improve the efficiency of breeding programme through the use of appropriate selection indices (Mohammadia *et al.*, 2003). Assuming yield is a contribution of several characters which are correlated among themselves and to the yield, path coefficient analysis was developed (Wright, 1923; Dewey and Lu, 1959). Unlike the correlation coefficient which measures the extent of relationship, path coefficient measures, the magnitude of direct and indirect contribution of a component character to a complex character and it has been defined as a standardized regression coefficient which splits the correlation coefficient into direct and indirect effects. Path coefficient analysis provides more information among the variables that do correlation coefficients since this analysis provides the direct effects of specific yield components on yield, and indirect effects via other yield components (Garcia del Moral *et al.*, 2003). Because yield is a quantitative character and is

associated with other component characters which are influenced to varying degree by the fluctuations in the environmental conditions (Chaugale, 1967).

A complete satisfactory criterion based on discriminant function selection would be more desirable when a combination of two or more characters with yield is studied in a selection index. The use of selection index technique would serve a two-fold purpose: (1) to bring about the genetic progress simultaneously in several characters and (2) to improve the yield through selection for relatively more heritable auxiliary characters. The technique of discriminant function analysis was first developed by Fisher (1936) and adopted for plant selection by Smith (1936). Later on, different workers constructed selection indices for different crops, such as Robinson *et al.* (1951) worked on corn; Paroda and Joshi (1970) on wheat; Joarder *et al.* (1978), Samad (1991) on rapeseed, Hussain (1997) on chilli and Ferdous *et al.* (2010) on wheat. The demand for food is on the rise due to the growth of the human population. The global maize stocks that have been shrinking uninterruptedly over the last 5 years already reflected the increased demand (Dias, 2005).

The development of improved varieties with high yield potential can be seen as a possibility to increase production. Such varieties with qualitative and/or quantitative superior traits over previously recommended varieties are developed by genetic improvement, which represents one of the most successful modern technologies in agriculture, and accounts for approximately 50 % of the yield increments of most crops (Fehr, 1987). To obtain genetic gains in different traits there are some methodologies of simultaneous selection (Cruz and Carneiro, 2003). Of these, the selection index proposed by Smith (1936) and Hazel (1943) has been well-accepted in maize improvement programmes. This index associates the information of different traits of agronomic interest, based on economic weights, genotypic and phenotypic variances of each trait and the respective covariances. Construction of selection

indices and their analysis would give the most appropriate weightage to the phenotypic values of each of two or more characters to be used simultaneously for selection (Ferdous *et al.*, 2010).

Genetic diversity is one of the useful tools to select appropriate lines for hybridization. Precise information on the nature and degree of genetic diversity helps the plant breeder in choosing the diverse parents for purposeful hybridization (Samsuddin, 1985). The genetic diversity between the lines is important as the genetically diverged parents are able to produce high heterotic effects (Falconer, 1981; Arunachalam, 1981; Ghaderi *et al.*, 1984; Mian and Bahl, 1989). Maize breeders are consistently emphasizing the importance of diversity among the parental genotypes as a significant factor contributing to heterotic hybrids (Ahloowalia and Dhawan, 1963).  $D^2$  analysis is a useful tool for quantifying the degree of divergence between biological population at genotypic level and in assessing relative contribution of different components to the total divergence both intra and inter-cluster level. Genetic divergence analysis estimates the extent of diversity existed among the selected genotypes (Murty and Arunachalam, 1966; Ram and Panwar, 1970; Sachan and Sharma, 1971 and Mondal, 2003).

The concept of general combining ability and specific combining ability was introduced by Sprague and Tatum (1942) and its mathematical modeling was set about by Griffing (1956) in his classical paper in conjunction with the diallel crosses. The value of any population depends on its potential *per se* and its combining ability in crosses (Vacaro *et al.*, 2002). The usefulness of these concepts for the characterization of an inbred in crosses have been increasingly popular among the maize breeders since the last few decades. Combining ability is a powerful tool in identifying the best combiners for hybridization especially, when a large number of advanced inbred lines are available and most promising ones are to be selected on the basis of their ability to give superior quality maize hybrids. Information on heterotic patterns and

combining ability among the maize germplasms is essential in maximizing the effectiveness of hybrid development (Beck *et al.*, 1990). Development of commercial maize hybrid usually requires a good knowledge of combining ability of the breeding materials to be used. The success in commercial production of hybrid maize depends on the availability of productive diverse quality maize inbred lines and clear knowledge of gene action for specific characters. The nature and magnitude of gene action is an important factor in developing an effective breeding program. Combining ability analysis is useful to assess the potential inbred lines and also helps in identifying the nature of gene action involved in various quantitative characters. A series of combining ability studies have been made by many workers from the International Maize and Wheat Improvement Centre (CYMMIT) to establish heterotic patterns among several maize check populations and gene pools, and to maximize their yield for hybrid development (Beck *et al.*, 1990, 1991; Crossa *et al.*, 1990; Vasal *et al.*, 1992a). Likewise, the variances of general and specific combining ability are related to the type of gene action involved. Variance for GCA includes additive portion while that of SCA includes non-additive portion of the total variance arising largely from dominance and epistatic deviations (Rojas and Sprague, 1952).

Diallel crosses were devised, specifically, to investigate the combining ability of the parental lines for the purpose of identification of superior parents for use in hybrid development programmes (Malik *et al.*, 2004). Analysis of diallel data is usually conducted according to the methods of Griffing (1956) which partition the total variation of diallel data into GCA of the parents and SCA of the crosses (Yan and Hunt, 2002).

Combining ability describes the breeding values of parental lines to produce hybrids. Sprague and Tatum (1942) used the term GCA to designate the average performance of a line in hybrid combinations, and used the term SCA to define those cases in which certain combinations do relatively better or

worse than would be expected on the basis of the average performance of the lines involved. In many studies, GCA effects for parents and SCA effects for crosses were estimated in maize (Dehghanpour *et al.*, 1996; San-Vicente *et al.*, 1998; Konak *et al.*, 1999; Chaudhary *et al.*, 2000; Araujo and Miranda, 2001; Kalla *et al.*, 2001).

Heterosis breeding using best combiners is one of the methods to improve upon the existing lines. Information on the genetic structure of a set of parents and mode of gene action governing yield and its attributes could be useful in designing suitable breeding procedures. For genetic studies various workers had used different biometrical methods but amongst them the approach of Hayman (1954a) and Mather and Jinks (1971) had been followed frequently. Genetic analysis of some economic traits showed different pattern of inheritance. The combining ability analysis helps in classifying the parents in terms of their hybrid performance and in gaining greater understanding of the nature of quantitatively inherited trait (Abd El-Aty and Katta., 2002; Ahmed and Saleem, 2003; El- Borhamy, 2004 and Ahmed *et al.*, 2011).

The most limiting factors of maize research in Bangladesh are the development, improvement and maintenance of parental/inbred lines. On the other hand, the problems of imported hybrid seed are the introduction of high price and uncontrolled quality. Moreover, the farmers can not get the seeds timely. One important approach to improve this situation is the development of inbred lines which can produce high yielding hybrid varieties. Before hybrid development, prospective parent (inbred line) selection is a pre-requisite. Several studies on maize have shown that inbred line from diverse stocks tend to be more productive than crosses between inbred lines from the same variety (Vasal, 1992b). Development of suitable inbred parents based on the genetic variability, diversity and combining ability for the production of hybrid seeds have been used as an important breeding approach in maize improvement.



A diallel analysis provides good information on the genetic identity of genotypes especially on dominance-recessive relations and some other genetic interactions. Diallel crosses have been used in genetic research to determine the inheritance of a trait among a set of genotypes and to identify superior parents for hybrid or cultivar development (Weikai Yan and Manjit Kang, 2003).

Heterosis and combining ability is one of the powerful tools in identifying the best combiner that can be used in crosses either to exploit heterosis or to accumulate fixable genes. Genetic diversity and combining ability of lines are important to obtain high heterosis values in the formation of maize hybrids (Sallahuddin, 2008). For developing desirable hybrids, information about combining ability of the parents and the resulting crosses is essential. (Banik, 2008).

One important approach to improve this situation is the development of inbred lines which can produce high yielding hybrid varieties. Before hybrid development, prospective parent (inbred line) selection is a pre-requisite. An inbred line is a “pure line” developed by self-pollination and selection until apparently homozygous plants are obtained. This usually requires five to seven generations of inbreeding. After five to seven generations of inbreeding and vigorous selection, vigorous inbred lines, uniform in appearance, are developed. Each inbred will have a different combination of genes. The main goal of maize breeding is to develop potential lines that ensure highest and stable production in a range of environments.

Therefore, we require stable inbred lines, which can help in the development of stable hybrids/varieties.

Keeping these points in view the present study was planned and executed with the following objectives.

- i. To evaluate and screen out the suitable inbred lines for yield parameters for their *per se* performance.

- ii. To study the nature and magnitude of genetic variability, heritability and genetic advance as per cent of mean for grain and yield contributing traits.
- iii. To study the association between quantitative characters and grain yield, between grain and yield component characters.
- iv. To estimate the contribution of yield component characters to the grain through the path coefficient analysis.
- v. To study the genetic divergence exists among the 25 maize inbred lines
- vii. To study the combining ability to identify good combining inbreds as well as their high heterotic hybrid combination.
- viii. To know the mode of gene action in governing the characters.
- ix. Study of inheritance pattern and mode of gene action for various yield contributing characters and positions of parents along the regression line on graph
- x. To select the superior parent/line suitable for commercial cultivation in Bangladesh.

## REVIEW OF LITERATURE

In maize, a lot of work has been done all over the world on genetic parameters. However, only selected reports on genetic variability, character association, selection index, divergence, combining ability, heterosis and genetic components of variation in maize inbred lines of some traits which are relevant to this work, are included in this review.

### GENETIC VARIABILITY

In designing a breeding programme it is essential to have a critical survey of genetic variability of inbred lines available. Burton (1952) suggested that a genetic coefficient of variation together with a heritability estimate would likely to give the best picture of the amount of genetic advancement to be expected from selection. On the other hand, Johnson *et al.* (1955) reported that heritability estimates along with genetic gain were more useful in predicting the effect of selection of the best individual. If heritability is mainly owing to the non-additive gene effect, the expected gain would be low; but if it is owing to additive gene effect, a high genetic advance may be expected.

Swamy *et al.* (1970) reported moderate to high heritability estimates of 55.55 and 86.04 percent for ear diameter, whereas Patil *et al.* (1972) noticed moderate heritability of 68.88 per cent for plant height trait and a value of 27.45 per cent for ear length. Johnson (1981) reported moderate to high range of heritability for 100 grain weight. However, Shahi and Singh (1985) reported high heritability for days to flowering, plant height and ear height. The expected genetic gain was about 17 per cent for plant height and 12 per cent for ear height in one of the locations. Bhalla *et al.* (1986) in their study reported high heritability estimates associated with high genetic advance for grain yield per plant, ear height, number of kernel rows per cob and plant height.

Debnath (1987) found that heritability was high for plant height and ear height, but low for the remaining characters. Expected genetic advance was high for grain yield, plant height and ear height. High heritability estimate of 70 per cent for plant height in 3-way cross hybrid maize was noticed by Debnath *et al.* (1988) and Mahmoud *et al.* (1990) noticed high heritability for ear diameter and number of kernels row per ear in maize. Arha *et al.* (1990) and Mani and Bisht (1996) found moderate heritability for grain yield, ear height and moderate for plant height. High heritability estimates for ear height was also noticed by Reddy and Agrawal (1992). Stem thickness showed the highest narrow sense heritability followed by ear position, thousand grain weight and ear girth. El-Hosary *et al.* (1994) noticed moderate to broad sense heritability values for ear height.

Ali *et al.* (1994) observed the highest genotypic variations for 1000-grain weight but minimum genetic advance in 32 genotypes of maize. Genotypic coefficient of variation, heritability and genetic advance are high for grain yield in maize as reported by most of the previous workers (Ali *et al.*, 1994; Singh *et al.*, 1995; Mani and Bisht, 1996; Satyanarayana and Saikumar, 1996; Ali *et al.*, 1997; Akanda *et al.*, 1997 and 1998; Tiwari and Verma, 1999; Singha and Prodhan, 2000 and Alam, 2009). On the contrary, Satyanarayana and Saikumar (1995) reported low genotypic coefficient of variation estimates combined with medium heritability and low genetic advance for grain yield.

Altinbos (1995) reported low heritability estimates and suggested that selection for ear length and ear diameter in early generations. Saxena *et al.* (1996) noted high heritability for plant height, ear height, ear girth and kernel rows per ear and low for grain yield, ear length and number of kernels per row. Maximum genetic gain was reported for ear height. The estimation of broad sense heritability is useful to predict the response even though it has got some

limitations. Mani and Bisht (1996) also reported high genetic advance for plant height. But Arha *et al.* (1990) found moderate heritability and genetic advance for plant height.

However, Mani and Bisht (1996) reported that genotypic coefficient of variation revealed low genetic variability for the ear girth in 38 local germplasms of maize. High genotypic coefficient of variation and heritability coupled with high genetic advance for grains/row has been reported by Akanda *et al.* (1997), Ali *et al.* (1997) and Singh *et al.* (1998).

Ali *et al.* (1997) reported high genotypic and phenotypic coefficient of variation for ear length. As an important yield attributing character, variability of ear girth has been studied. They also reported high genetic advance accompanied by high heritability for ear breadth.

Akanda *et al.* (1998) reported that genotypic and phenotypic coefficient of variation, heritability and genetic advance were moderate for ear girth. Contrarily, They reported comparatively low genotypic coefficient of variation and heritability and genetic advance for the same character.

Genotypic coefficient of variation, heritability and genetic advance are high for 1000-kernel weight reported by Ali *et al.* (1997), Singh *et al.* (1998) and Akanda *et al.* (1997 and 1998). Akanda *et al.* (1997, 1998) and Singha and Prodhan (2000) reported that heritability along with genetic advance and genotypic coefficient of variation was high for ear size and ear length.

Singh and Dashi (2000) reported high heritability and genetic advance for plant height. Singha and Prodhan (2000) reported high heritability along with high genetic advance and high genotypic coefficient of variation for ear height in 34 genotypes of maize. Rafiq *et al.* (2010) reported that grain yield, ear length, ear height, 100-seed weight and ear diameter had high GCA estimates with high heritability. Genetic advance was higher for plant height, ear length, grains per row and grain yield.

Shamim *et al.* (2010) studied that broad sense heritability estimates for plant height, cob length, grains/row, 1000-grain weight and harvest index were higher in magnitude (61.0 to 99.0%) indicating greater genetic impact on these traits.

Wannows *et al.* (2010) showed that high narrow sense heritability estimates were detected for leaf area index, number of kernel per row, plant height, ear height, physiological maturity, number of rows per ear, ear length and ear diameter and emphasizing that the additive genetic variance was the major component of genetic variation in the inheritance of these traits and would likely be in selection of improving these traits.

## **CORRELATION AND PATH COEFFICIENT**

Grain yield in maize as in other crop is associated with a number of yield components. A study of the nature and degree of association of these components with yield assumes greater importance for fixing up characters that play a decisive role in influencing yield. Selection would therefore be more effective if it is based on component characters rather than directly on grain yield. According to Appadurai and Nagarajan (1975), grain weight per ear and grain numbers per row had little effect on yield, while ear length and ear circumference had positive correlation with yield. Kim (1975) and Hallauer *et al.* (1989) reported correlation coefficients among the characters were generally positive and significant for plant height and ear height.

Probecky (1976) reported that grain yield primarily depends on the number of grains per plant; which in turn depend mainly on the number of grains in the rows. Utkhede and Shukla (1976) revealed highly significant positive genotypic and phenotypic correlation between yield and number of grain row per ear, weight of ear, ear height and ear length. Ear height and dry ear weight contributed substantially to yield. Singh and Nigam (1977) found that grain yield was positively and significantly correlated with five yield components

*i.e.*, 100 grain weight, number of kernel rows per cob, ear weight, ear diameter and number of grain per rows. Sharma *et al.* (1982) reported that grain yield was positively correlated with grains per ear, hundred grain weight, plant height and ear height. Saha and Mukherjee (1985) observed that grain yield per plant was significantly correlated with ovules per ear, grains per ear and 100-grain weight.

Malhotra and Khehra (1986) recorded positive correlation between grain yield and yield components like ear length, ear circumference, number of rows per ear, 1000-grain weight, shelling percentage, ear height and plant height. Tyagi *et al.* (1988) opined that grain yield was influenced more by ear weight, ear length, plant height, kernels per row and 100 grain weight. Maharajan *et al.* (1990) concluded that grain yield was positively correlated with ear length, number of kernels per row and plant height. Singh *et al.* (1991) noted that grain yield per plant had significant positive correlations with plant height and ear height in F<sub>1</sub> and F<sub>2</sub> generations under alkaline soil and with leaf area in both the generations under normal soil. Debnath and Khan (1991) revealed that plant height, number of kernels per row and 1000-grain weight had strong positive contributions to grain yield.

Boraneog and Duara (1993) observed that plant height and ear height exhibited significant positive correlation with grain yield. Saha and Mukherjee (1993) reported positive significant correlation between grain yield per plant with 100-grain weight, ear length, ear circumference, number of kernel rows per ear and number of kernels per row, the ear circumference and number of grains per row. Krishnan and Natarajan (1995) obtained high positive association between grain yield and plant height, ear length, ear weight and number of kernels per row. Rahman *et al.* (1995) reported that grain yield was significantly and positively correlated with plant height, ear height, number of grains per ear and 1000 grain weight. According to Satyanarayana and Saikaman (1996), grain

yield was positively correlated with number of kernel rows per ear, ear length, ear circumference and 100 grain weight. Kumar and Kumar (1997) reported that values of genotypic correlation were slightly higher than the corresponding phenotypic values. Significant positive correlation was recorded for plant height, ear length and ear height with yield per plant. Annapurna *et al.* (1998) reported that seed yield was positively and significantly correlated with plant height, ear circumference, number of seeds per row, number of seed rows per ear.

Khakim *et al.* (1998) noticed that grain yield was positively correlated with plant and ear insertion height, leaf area, ear number, ear length, number of kernel rows per cob, number of grains per row and grain weight per cob, ear weight and 1000-grain weight.

Gautam *et al.* (1998) reported maximum correlation between grain yield and number of kernels per row, leaf area, plant height and cob length. Dutu (1999) indicated that at phenotypic and genotypic levels, growth period was strongly correlated with plant height and leaf numbers. He also reported correlations using the date of flowering, plant height and the number of leaves which were used as indirect selection criteria and resulted in positive correlated response in earliness and yielding potential.

Basheruddin *et al.* (1999) reported that correlation coefficients had highly positive significant influence on plant height, number of leaves, leaf area per plant and stem girth. Nawar *et al.* (1999) observed that additive components were significant for number of kernel rows per cob. Highly significant positive correlation coefficients were detected among yield per plant, components of ear and plant height. Kumar (1999) revealed that the number of grains per row, number of rows per ear, ear circumference and ear length had direct effect on grain yield.



Mani *et al.* (2000) reported that grain yield per plant indicated that highly significant positive correlation with all the attributes and was highest with ear weight per plant. Umakanth *et al.* (2000) observed that grain yield per plot showed significant and positive correlations with ear circumference, ear length, plant height and 100-seed weight. Path analysis revealed that plant height followed by number of seeds per row, 100-seed weight, ear length and ear circumference showed maximum positive direct genotypic effects as well as indirect contribution through other characters on grain yield.

Vaezi *et al.* (2000) noticed that grain yield was significantly and positively correlated to ear weight, ear circumference, number of kernel rows per cob and number of kernels per row. Netaji *et al.* (2000) reported that yield per plot was significantly and positively correlated with all the characters except days to 50 per cent tasseling, silking and dry husk. Maximum variability was observed for plant height, followed by ear height and test weight. Singh and Dashi (2000) revealed high positive correlation of green fodder yield with plant height, leaf length and leaf stem ratio. Prodhan and Rai (2000) reported that significant positive correlation of popping expansion was found with popping percentage, tenderness and pericarp thickness, while popping expansion showed significant negative association with grain weight and non-significant negative association with grain yield. On the other hand, grain yield was strongly associated with grain weight.

Geetha and Jayaraman (2000) observed number of grains per row exerted a maximum direct effect on grain yield. Vaezi *et al.* (2000) showed that 300-kernel weight and kernel depth had the highest positive effect on grain yield whereas ear diameter had a negative indirect effect on grain yield through some traits. Path analysis, for grain yield showed that kernel weight and kernel depth had the highest positive effect on grain yield. Kumar and Satyanarayana (2001) concluded that grain yield was positively associated with plant height, ear

height, ear length, ear circumference, number of seed rows per ear and test weight. Swarnalatha and Mohammad (2001) indicated that the plant height, days to 75 per cent silking and maturity, ear length, number of seeds per row and 100-grain weight positively influenced the yield directly and also indirectly through several yield components. Guang Cheng *et al.* (2002) showed that importance of eight yield components to grain yield and suggested that more attention should be paid to cob length, cob diameter and kernel percentage. They also noticed that grain yield per plot had significant positive correlation with 100-kernel weight.

Venugopal *et al.* (2003) indicated that plant height, ear height, ear length, ear girth, 100-seed weight and number of seeds per row were positively associated with grain yield. Although the character number of seed rows per ear had a direct positive contribution towards grain yield, but it had indirect negative influence through ear length, 100 seed weight and number of seeds per row.

Ahmad and Saleem (2003) reported that vegetative phase had the highest positive direct contribution to grain yield per plant followed by growing degree days to tasseling and growing degree days to maturity. Growing degree days to the reproductive phase had the highest negative direct effect on grain yield. Viola *et al.* (2003) revealed that early silking and harvesting of fresh cobs, greater plant height, cob length, cob weight, cob height and number of cobs per plant and lesser cob girth directly contributed to increased cob yield.

Singh *et al.* (2003) observed that ear leaf area had the highest positive direct effect on green fodder yield per plant at genotypic and phenotypic levels followed by dry matter yield per plant, ear length and days to 50 per cent silking. Ear length had the maximum direct effect on grain yield followed by 500-kernel weight and ear leaf area. Number of leaves per plant, leaf: stem ratio and girth of basal internodes had also highly positive direct effect on grain yield per plant.

Srivas and Singh (2004) observed that dry fodder yield per plant, a dependent trait was significantly and positively associated with green fodder yield and its contributing traits such as plant height, days to 50 per cent silking, number of leaves per plant, stem girth, leaf blade length, leaf width and sheath length.

Kumar and Singh (2004) reported that cob length had maximum positive direct effect on grain yield at genotypic and phenotypic levels, respectively. Path analysis revealed that six characters had positive contribution to grain yield via number of other characters both at genotypic and phenotypic levels. Whereas, days to 50% tasseling and days to maturity had maximum negative effect on grain yield.

Bao Heping *et al.* (2004) reported that maize yield was mainly influenced by ear length, followed by number of kernels per row, ear width, number of rows per ear, growth period and 1000-seed weight. Kernel percentage per ear and number of pointless ears had minimum effect on maize yield.

Rafique *et al.* (2004) reported that grain yield was positively and significantly associated with all parameters studied. Number of kernels per row was positively correlated with grain yield followed by plant height, ear height, ear length and its diameter

Srivas and Singh (2004) observed that characters such as plant height, days to 50 per cent silking, stem girth, leaf length, leaf width and number of leaves per plant had positive direct effect on dry fodder yield at phenotypic levels. Patel *et al.* (2005) reported that dry matter yield per plant, number of leaves per plant, days to 50 per cent silking and plant height had positive direct effects on green fodder yield. Shelake *et al.* (2005) noticed that grain yield was positively and highly correlated with number of grains per cob, biological yield per plant, harvest index, 100-grain weight, cob length, number of grain rows per cob and cob girth. The number of days to 50 per cent tasseling, number of days to 50

per cent silking and harvest index showed higher genotypic direct effect. Biological yield per plant had the highest negative genotypic direct effect on grain yield. Ei-Shouny *et al.* (2005) showed that grain yield per plant correlated positively and significantly with ear diameter, ear length, number of kernels per row, 100-kernel weight, number of rows per ear, ear height, plant height and days to silking. Under normal planting date and with number of kernels per row, ear diameter, 100-kernel weight, ear length, number of rows per ear, ear height and days to silking under late planting date.

Sumathi *et al.* (2005) genotypic correlation studies indicated that ear weight, number of rows per ear, number of kernels/row, and total number of kernels/ear were positively associated with grain yield. Path coefficient analysis revealed that number of kernels per row showed high direct effect on grain yield followed by 100- seed weight, number of rows per ear and total number of kernels per plant.

Kumar *et al.* (2006) observed that days to 50% tasseling, anthesis silking interval, ear height and 100-seed weight had highest direct effect on grain yield. The days to 50 % silking exhibited negative direct effect on grain yield.

Harjinder *et al.* (2006) reported significant positive correlations for grain yield with days to 75 % husk, plant height, ear height, and number of ears.

Tan Heping *et al.* (2006) noticed that grain yield was significantly correlated with plant height, ear diameter, ear length, rare ear length, 100-kernel weight and grain production rate. Grain yield was most highly correlated with ear diameter, followed by 100-kernel weight, plant height, ear length and grain production rate. Wang Dachun (2006) reported that kernel weight per ear mainly affected by ear length and ear diameter and the ear length with bearing kernel played an important role on kernel weight per ear in high yielding combinations.

Wali *et al.* (2006) observed that yield was positively associated with plant height, ear length, ear circumference, number of kernels per row, fodder yield per plot and 100-grain weight, but was negatively correlated with number of days to 50 per cent silking at the phenotypic and genetic levels. The grain yield per plant was positively associated with plant height, ear length, ear circumference, number of kernels per row, fodder yield per plot and 100-grain weight at the phenotypic and genetic levels. Abirami *et al.* (2007) indicated that grain yield showed positive association with oil content and protein content. Path analysis showed that the weight of the cob contributed to the maximum direct effect to grain yield. It implied that selection for weight of the cob will be highly effective for the improvement of grain yield.

Bhoite *et al.* (2007) reported that dry matter and crude protein yields showed positive and significant correlation with green forage yield and had positive direct influence on their correlation with green forage yield.

Sofi and Rather (2007) reported that the genotypic correlation coefficient revealed that ear diameter, 100-seed weight, ear length, number of kernel rows per ear and number of kernels per row showed the greatest correlation with grain yield. Path analysis indicated that 100-seed weight had greatest direct effect on grain yield, followed by number of kernels per row, number of kernel rows per ear, ear length and ear diameter.

Xie Zhen Jiang *et al.* (2007) showed that kernels per plant was arranged for the top position among the many agronomic traits that contributed to the yield enhancement of a single plant and was followed by kernels per row, 1000-kernel weight and leaf orientation value.

Akbar *et al.* (2008) noticed that plant height had highly significant genotypic and phenotypic association with cob height and days to 50% tasseling with days to 50% silking. All traits had significant genotypic association but not

significant phenotypic association with grain yield. They showed that all traits exerted positive direct effect on grain yield per plant except days to 50% silking.

Bello *et al.* (2010) reported that positive and phenotypic and genotypic correlation coefficients had found for days to 50% tasseling with plant and ear height, and grain yield with plant height, number of grains/ear and ear weight. They also reported that days to 50% silking, ear weight and number of grains/ear had the highest direct effect on grain yield at genotypic level. Ferdous *et al.* (2010) reported that grain yield per plant indicated that significant and positive correlation with days to maturity grains/spike, 100-grain weight and harvest index in wheat.

Nastasic *et al.* (2010) reported that grain yield was positively and significantly genotypic correlation coefficients with kernel row number, ear length, kernel depth and 1000-kernel weight. They also reported that the direct effects, obtained in path coefficient analysis, in both studied population (S1, HS) indicated that grain yield at most depended upon 1000-kernel weight.

Rafiq *et al.* (2010) reported significant correlation of grain yield with ear diameter, 100-grain weight, ear length, rows per ear and grains per row in maize. They also reported that the highest direct effect on grain yield was exhibited by 100-seed weight followed by grains per row, grain rows per ear, ear length and ear diameter. Sreckov *et al.* (2010) observed that grain yield had significant positive correlation coefficients through ear height and length. They showed that population NSU1×568/11 had high significant, undesirable direct influence on grain yield and ear height.

Wannows *et al.* (2010) reported that grain yield was positively and significantly correlated with number of kernels per row, ear length, and leaf area index. They also reported that each of leaf area index, ear diameter and

physiological maturity had high positive direct effects on grain yield at the phenotypic level. Path analysis revealed that three characters had positive contribution to grain yield via number of other characters at phenotypic level.

## **SELECTION INDEX**

Different selection indices were formulated using different combinations of yield and yield contributing characters and their expected genetic gains were estimated. The development of improved varieties with high yield potential can be seen as a possibility to increase production. Such varieties with qualitative and/or quantitative superior traits over previously recommended varieties are developed by genetic improvement, which represents one of the most successful modern technologies in agriculture, and accounts for approximately 50 % of the yield increments of most crops suggested by Fehr (1987).

To obtain genetic gains in different traits there are some methodologies of simultaneous selection (Cruz and Carneiro, 2003). Of these, the selection index proposed by Smith (1936) and Hazel (1943) has been well-accepted in maize improvement programs. This index associates the information of different traits of agronomic interest, based on economic weights, genotypic and phenotypic variances of each trait and the respective co-variances. Kumar and Kumar (2000) suggested that selection based on plant height with greater ear weight, number of seeds rows per ear and number of seeds per ear was desirable for grain yield.

Bergale *et al.* (2002) suggested that the number of spikes/plant, grains/spike and harvest index must be given preference in selection along with optimum plant height and days to flowering to selection the superior wheat genotypes. The demand for food is on the rise due to the growth of the human population. The global maize stocks that have been shrinking uninterruptedly over the last 5 years already reflect the increased demand reported by Dias (2005).

Shiv *et al.* (2008) reported that number of tillers/plant, numbers of spikelets/ear, number of grains/ear, grain weight/ear; 100-grain weight and biological yield could form effective selection indices for selection of high yielding genotypes of wheat.

Ferdous *et al.* (2010) reported that selection indices were constructed through the discriminate functions using eight respective characters. From that result, the highest relative efficiency was observed with the selection index based on three characters; plant height, grains/spike and grain yield/plant in spring wheat.

## **GENETIC DIVERGENCE**

Wide range of variation was observed in cluster mean performance for most of the characters studied. All the genotypes were grouped into 5-clusters, indicating the presence of diversity for different characters. Williams and Hallaver (2000) reported that the cluster II (82) had the highest number of genotypes followed by cluster III (36) and cluster IV (19). The clusters IX and X were mono-genotypic reported by Williams and Hallaver (2000).

Yin ZhiTong *et al.* (2004) studied cluster analysis for various plant traits (including plant height, ear height, tassel length, stem diameter, ear length, ear diameter, number of rows, number of grains per row, 100-grain weight, yield per plant, dried ear weight and maturity period) classified some 110 maize inbreds into 5 groups. The genetic diversity among the inbred groups was greater than that within the same group, and heterosis among the groups was greater than that within the same group. To breed outstanding crosses, the parents should be selected from the various groups.

Singh *et al.* (2005) estimated  $D^2$  analysis using 23 genotypes of maize in an experiment conducted in Karnal, Haryana, India, during kharif 1998. Observations were recorded for 50 per cent tasselling, 50 per cent silking, plant height, cob height, days to maturity, cob girth, cob length, number of rows per



cob, number of grains per row, 100-grain weight and grain yield per plant. The genotypes fell into 6 clusters. The inter-cluster distances were higher than intra-cluster distances, suggesting wide genetic diversity among the genotypes of different groups. The inter-cluster  $D^2$  values indicated the maximum distance between clusters III and VI and the lowest distance between clusters I and IV. The cluster means were higher for 50% tasselling, 50% silking, plant height, cob height, cob length, number of grains per row and 100-grain weight in cluster IV; for cob girth, days to maturity and number of rows per cob in cluster II; and for grain yield per plant in cluster III followed by cluster II. The genotypes of these clusters would offer a good scope for the improvement of this crop through natural selection and hybridization. The genotypes included in the diverse clusters can be used as promising parents for hybridization to obtain high heterotic response and thus better segregates in maize. Based on genetic divergence and mean performance of yield and other characters, genotypes Vijay composite, NC-300, K-614, K-679, K-771, K-801, K-808, K-621 and CML-326 were selected.

Yuai *et al.* (2006) studied 24 varieties introduced from the Crop Institute, Chinese Academy of Agricultural Sciences to Hohhot, Nei Menggu [Nei Mongol], China and reported that cumulative contribution percentage of 7 principal components (morphology and yield factor, growth duration factor, rows/ear factor, low yield factor, oil content factor, protein content factor and ear length factor) to variation reached 83.149 per cent. Based on the analysis for each principal component vector, the introduced 24 varieties were grouped into 6 clusters.

More *et al.* (2006) reported that 45 diverse genotypes of forage maize for genetic diversity and identify the suitable genotypes for hybridization programmes based on clustering pattern. The genotypes were grouped into 7 clusters using Mahalanobis  $D^2$  statistics. Cluster II was the largest with 25 genotypes followed by cluster III with 11 genotypes and cluster I with 5

genotypes. The clusters IV, V, VI and VII were mono-genotypic. The maximum inter-cluster distance was observed between clusters I and VI followed by distance between clusters I and IV and clusters I and V. Clusters V and VI exhibited the minimum inter-cluster distance.

Chen FaBo *et al.* (2007) reported that 186 maize hybrids could be classified into ten clusters, with 88.2 per cent of the hybrids included in Cluster 4, Cluster 8 and Cluster 10. The analysis of pedigree sources of 51 hybrids showed that 36 hybrids had close genetic relationships with the hybrids of Pioneer Company developed in late 1980s and early 1990s in the United States, such as "Y78599", "Y7865", "and Y78698", accounting for 70.58 per cent. Meanwhile, 13 hybrids had close genetic relationship with "Y78599", accounting for 8.66 per cent. The cluster analysis showed that 88.2 per cent of the 51 hybrids were in Cluster-4, Cluster-8, and Cluster-10. It was indicated that the similarity was high and the genetic diversity was narrow among the 186 hybrids. It is necessary to broaden the genetic basis of breeding germplasm in maize.

Ivy *et al* (2007) and Hoque *et al* (2008) reported that 24 maize inbred lines were grouped into 4 different clusters by using clustering techniques. The maximum inter-cluster distance was observed between clusters I and III and the lowest in II and IV. The highest mean values for kernel yield/plant, 1000-kernel weight, number of kernels per row and ear were observed in the same cluster III.

Azad *et al.* (2012) revealed that 30 maize inbred lines were grouped into 6 different clusters, based on medium to high inter-cluster distances, 6 lines were selected for hybrid program. Cluster VI showed the highest mean values for kernel yield and all the yield contributing characters except days to 50% tasseling and 50% silking. Days to maturity and ear diameter showed maximum contribution towards total divergence among different characters. These characters should be given importance, for selecting diverse parents for breeding programme.

## **DIALLEL AND COMBINING ABILITY**

The concept of combining ability was originally evolved through the work of Davis (1927) who suggested that use of inbred cross to test the combining ability of inbreeds in maize. It may be defined as the ability of a strain to produce superior progeny upon hybridization with other strains. Combining ability provides information about the nature and magnitude of gene action controlling various quantitative characters. General combining ability (GCA) is the average performance of a line in hybrid combination and specific combining ability (SCA) is the deviation of crosses on the basis of average performance of the lines involved.

Diallel analysis is used to estimate GCA and SCA effects and their implications in breeding (Griffing, 1956; Gardner and Eberhart, 1966; and Baker, 1984). Griffing (1956) proposed an analysis for diallel mating systems that estimate the general and specific combining abilities of lines and hybrids. Combining ability analysis is important in identifying the best parents or parental combinations for a hybridization program. General combining ability is associated to additive genetic effects while specific combining ability is associated to non-additive genetic effects (Falconer and Mackay, 1996). Combining ability is an important aspect of hybrid breeding program. The proper identification of the genetically superior parents is dependent on the information obtained from the analysis of the combining ability.

The concept of combining ability was put forward by Sprague and Tatum (1942). According to them, general combining ability is the average performance of a strain in series of cross combinations estimated from the performance of F<sub>1</sub> is from the crosses, whereas specific combining ability is used to designate those cases in which certain combinations do relatively better or worse than would be expected as the basis of average performance of lines involved. The general combining ability is associated with genes which are additive in effects and specific combining ability is attributed primarily to

deviation from the additive scheme caused by dominance and epistasis (Rojas and Sprague, 1952).

Griffing (1956) has shown relationship between various heritable variance components and GCA and SCA variances. Thus GCA variance is due to additive variation and additive  $\times$  additive interaction variance, while SCA variance is due to dominance variance, additive  $\times$  additive variance, additive variance is due to dominance variance, additive  $\times$  additive variance, additive  $\times$  dominance variance components. Estimates of the variances due to GCA and SCA provide an appropriate diagnosis of the predominant role of additive or non-additive variance. Among the different biometrical methods employed to study combining ability, line  $\times$  tester is the most widely used method.

Allard (1960) reported that the combining ability analysis is highly useful technique for the plant breeder which provides indication of the genetic behavior of the parental material and is therefore desirable to select the parents for hybridization on the basis of their combining ability. Ratio of additive and non-additive gene action is to be considered in order to decide the predominance of the kind of genetic variation for a given character. If the ratio of additive to non-additive gene action is more than unity, indicates the major role of additive variance in controlling the expression of a character, whereas less than unity indicates the importance of non-additive variance (Gardner, 1963).

Muthiah (1989) reported that the proportion of GCA variance was higher than the SCA variance showing preponderance of additive genetic effects for all the characters studied. Crossa (1990) noted high general combining ability effects for high grain yield.

Results of Crossa *et al.* (1990) suggested that both additive and non-additive effects are important in controlling the expression of grain yield and days to silk; however, non-additive effects are more important in controlling grain

yield than in determining days to silk in CIMMYT's tropical late yellow maize germplasm.

Pal and Prodhan (1994) have also showed a greater influence of additive component of gene action in the expression of maturity. However, equal importance of both additive and non-additive gene action for days to maturity has been reported by Roy *et al.* (1998). Preponderance of additive genetic effects in the inheritance of ear girth has been reported by Das and Islam (1994), Pal and Prodhan (1994). On the other hand, specific combining ability is found to be very important for ear diameter (El-Hosary *et al.*, 1994 and Singh and Singh, 1998). More importance of additive gene action for days to maturity is to be found in 15 F<sub>1</sub> hybrids and their parents (Zelleke, 2000).

Pal and Prodhan (1994) have also showed a greater influence of additive component of gene action in the expression of maturity. However, equal importance of both additive and non-additive gene action for days to maturity has been reported by Roy *et al.* (1998).

Satyanarayana and Saikumar (1995) recorded low genotypic coefficient of variance combined with low to medium heritability and low genetic advances for grain yield.

But According to Altinbas (1995), GCA and SCA variances for grain yield per plant and other yield components indicated that screening parental lines and crosses based on combining ability effects for 100-grain weight and ear length should be effective. In case of other agronomic characters, genotypic coefficient of variance estimates were high combined with high heritability and genetic advance indicating the inheritance of additive gene action controlling these traits.

Dass *et al.* (1997) reported that non-additive gene action played major role in the inheritance of grain yield and majority of ear traits.

Joshi *et al.* (1998) observed both additive and non-additive gene effects in their early maturing inbred materials. However, the ratio of additive/non-additive

genetic variance revealed that there was preponderance of non-additive gene action in the expression grain yield/plant, protein and starch content, while for oil content and 100-grain weight there was preponderance of additive gene action.

Dutu (1998) observed that plant height was controlled by additive and non-additive gene actions and cytoplasmic inheritance, whereas number of leaves per plant was controlled by additive gene effects only.

Lou-Xiang Yang (1998) concluded that most plant and ear characters were improved with additive and dominance effects of the female parents. GCA variance is found to have significant for ear girth (Mathur *et al.*, 1998). General combining ability variance (Tulu and Ramachandrappa, 1998; Beck *et al.*, 1990) and both general and specific combining ability variances (Spaner *et al.*, 1996; Dahlan *et al.*, 1997; San-Vicente *et al.*, 1998; Lemos *et al.*, 1999; Talleci and Kochaksaraei, 1999 and Alam, 2009) are significant for plant height. Both additive and non-additive effects are to be found in genetically control of plant height reported by Choukan (1999). But additive gene effect is more important for plant height observed by (Crossa *et al.*, 1990; Mahajan and Khehra, 1991; Odongo and Bockholt, 1995; Paul and Debnath, 1999 and Zelleke, 2000). On the other hand, Suneetha *et al.* (2000) found preponderance of non-additive gene action for plant height. Both GCA and SCA effects are significant for ear height reported by Beck *et al.* (1990) and Talleci and Kochaksaraei (1999).

Additive effects are found in controlling ear height shown by Tulu and Ramachandrappa (1998), Choukan (1999) and Lemos *et al.* (1999). On the other hand, additive gene action is found more important than non-additive for ear height reported by Mahajan and Khehra (1991), Altinbas (1995), Odongo and Bockholt (1995) and Paul and Debnath (1999). Contrary to those, Dehghanpour *et al.* (1996) and Singh and Singh (1998) found more importance of non-additive effects in the expression of ear height. Significant GCA variance for ear length has been reported by Alike (1994), Spaner *et al.* (1996)

and Mathur *et al.* (1998). But Das and Islam (1994) reported equal importance of additive and non-additive variances for ear length. On the other hand, Zelleke (2000) observed more importance of additive gene action in controlling ear length and identified A1-175 x A1-178 as the best specific combiner. On the contrary, Pal and Prodhan (1994) showed more importance of non-additive gene effects in controlling of ear length. Similarly, Singh and Singh (1998) reported more importance of SCA for ear length.

Ramech *et al.* (2000) observed greater ratios of GCA to SCA mean squares for all traits except for number of seed rows per ear, indicating the importance of non-additive gene effects in their genetic control.

Rosa *et al.* (2000) obtained highest SCA values in 13 x 13 diallel crosses, viz., AS-910 x AS-4450 and PP-9538 x AS-948.

Desai and Singh (2001) reported significant difference in gca and sca effects for plant height, ear height and number of leaves per plant.

Kara (2001) observed significant gca effects for all the traits and significant sca effects for ear circumference, ear height and grain yield per unit area.

Konak *et al.* (2001) obtained non-additive gene effects for ear length and number of kernel rows per ear and additive gene effect for yield, 1000-kernel weight, plant height, ear height and days to silking.

Shabir and Saleem (2002) performed diallel analysis of six elite lines of maize and reported that all the characters being studied were under the control of over-dominance type of gene action, except protein percentage which showed additive type of gene action.

Ahmad and Saleem (2003) observed that both additive and non-additive gene effects played an important role in the genetic control of all the traits. The estimates of mean of squares due to general combining ability were significant for all the traits except growing degree days in both F<sub>1</sub> and F<sub>2</sub> generations. Mean squares due to specific combining ability were significant for all the traits in both the generations.

Abdel-Sattar (2003) observed that value of GCA and SCA were increased from 53 to 55 self generations. The ratios of GCA/SCA through all self generations showed greater role of dominance variance than that of additive variance. Lines contributed much more than the testers to the total genetic variation.

Srivastava and Singh (2004) reported that the specific combining ability variance was greater than general combining ability variance indicating the importance of non-additive gene effects for ear diameter, ear length, number of rows per ear and grain yield.

Ming *et al.* (2004) revealed that the inbreds had high combining ability for grain yield of ten quality protein maize (QPM) lines and showed high general combining ability (GCA) and it is possible to get high yielding crosses by using them as parents. The results from the yields of F<sub>1</sub> crosses suggest that lines with high GCA and specific combining ability (SCA) should be selected as parents.

Koinuma *et al.* (2004) noticed that new inbred line H049 shows high combining ability with dent inbred lines.

Reddy *et al.* (2004) reported that no association was found between mean *per se* and all other estimates. They noticed that general combining ability failed to exhibit closer association in grain yield per plant and ear length.

Malik *et al.* (2004) reported that GCA effects were highly significant for all the traits under study, but SCA effects were less significant in certain cases suggesting predominance of additive genes. High gca effects for grain yield were observed in the temperate material, i.e., QPM-1 (0.168), QPM-3 (0.169) and QPM-5 (0.485), while sca effects were remarkable for hybrids QPM-3 NCML-1078 (0.890), NCML-NCML-1084 (0.878) and NCML-1082 NCML-1083 (0.831).

Uddin *et al.* (2006) studied that significant differences for GCA and SCA indicated the presence of additive as well as non-additive gene effects were more prominent for all the characters studied except grain yield/plant.



Ojo *et al.* (2007) reported that GCA mean squares were however highly significant and higher than SCA mean squares for grain yield, indicating a predominance of additive gene action for grain yield.

Aliu *et al.* (2008) reported that both GCA and SCA effects are significant for ear weight, indicating that both additive and non-additive genetic actions were important combining of hybrids from the diallel crosses.

Uddin *et al.* (2008) reported that GCA and SCA variance for yield/plant, number of kernels/row and 100-kernel weight was observed significant, which indicated importance of additive type of gene action for these characters.

Alam *et al.* (2008) reported that almost equal role of additive and non-additive gene action was observed for days to maturity. Additive genetic variance was preponderant for grains per ear and 1000-grain weight and non-additive gene action was involved in plant height, ear height, days to silking and days to maturity.

Ahmad *et al.* (2011) reported that the relative magnitude of GCA and SCA variances indicated that additive gene effects were more prominent for days to heading and grain weight.

Singh *et al.* (2012) revealed that estimates of SCA variances were higher than GCA variances for number of kernels per row, cob height and 50% silking under study, indicating predominance of non-additive gene action of these traits. The parents HUZQPM 3-2, HUZQPM 6-2 and HUZQPM 5 were identified as good combiners for yield and its related traits.

Estakhr and Haidari (2012) reported that significant for GCA and SCA mean squares for plant height, ear height, ear diameter, ear length, grain number per ear, grain moisture and cob percentage which is an indication of the importance of both additive and non-additive genetic effects.

## GENE ACTION AND THEIR PROPORTIONS

The choice of a suitable breeding method depends to a large extent on the nature of gene action involved. Fisher (1918) and Wright (1935) defined three types of variances as additive genetic variance, variance due to dominance deviations and epistatic variance resulting from the interaction of non-allelic genes.

Hayman and Mather (1955) further showed that epistatic variance can be partitioned into genetic interactions of the additive x additive, additive x dominance and dominance x dominance types and higher order interactions.

The selection of suitable parents is one of the most important steps in hybridization program. Selection of the parents on the basis of phenotypic performance alone is not sound since phenotypical superior lines may yield poor recombination in the segregating generations. It is, therefore, essential that parents should be chosen on the basis of their genetic values. There are several techniques for the evaluation of varieties or strains in terms of their genetic makeup. Of these, diallel analysis technique (Hayman 1954a, b) is the popular method to study components of variation. It was developed by Jinks (1954), Hayman (1954a, b) and Jinks and Hayman (1957) using Mather's concept of D, H components of variation. While a negative F value indicates an excess of recessive alleles in the parents, a positive value shows more dominant alleles than recessive alleles of each gene are distributed equally among the parents, the F value will be equal to zero reported by Allard (1962). Over dominance as well as predominant role of non-additive genetic variance in the inheritance of kernel yield of maize was also reported by Gardner and Lonquist (1961), Gamble (1962a), Gardner (1963), Nawar *et al.* (1980, 1981), Genova (1984), Shahi and Singh (1985) and Genov (1987). Regarding 1000-kernel weight non-allelic interaction with over dominance was found by Gamble (1962b) and Debnath and Sarkar (1990b). As an indicator of the relative frequency of dominant and recessive alleles in the parents, the F value was found to be

positive but non-significant for grain yield, which means either that no allele's exhibit dominant or else that the dominant and recessive alleles are distributed equally among the parents observed by Verhalen and Murry(1967).

Mather and Jinks (1971) concluded that Hayman's analysis was the most useful to determine the significance of principal genetic components. The components D, H<sub>1</sub> were significant for ear length, ear diameter and kernel rows per ear. The ratio  $(H_1/D)^{1/2}$  was greater than one for kernel yield, kernel rows per ear and 1000-kernel weight indicating over dominance in the inheritance of these characters. All the characters except ear diameter exhibited presence of non-allelic interaction in their inheritance. Over dominance as well as importance of non-additive gene action in the inheritance of grain yield was observed by Gamble (1962a), Darrah and Hallauer (1972) and Genov (1987). On the other hand, Lonquist and Castro (1967) and Murthy (1978) reported partial dominance as well as predominant role of additive genetic variance for this trait. Epistasis was also observed in grain yield by Gamble (1962a) and Darrah and Hallauer (1972). Partial dominance as well as importance of additive gene effects in the inheritance of kernel rows per ear was reported by Singh (1979). Over dominance for predominant non-additive genetic variance was observed by Nawar *et al.* (1980) for kernel rows per ear and by Gamble (1962b) for kernel weight.

Debnath and Sarker (1989) reported complete dominance in the inheritance of ear diameter as the ratio  $(H_1/D)^{1/2}$  was very close to unity. Positive but insignificant values of F for all characters except ear length and ear diameter indicated almost symmetrical gene distribution with a small excess of dominant alleles in the parents. The ratio  $(h^2/H_2)$  suggested that the kernel yield be under control of highest number of genes or gene group. However, kernel weight was controlled by the lowest number of genes. They also reported that all characters were under polygene control.

Debnath and Sarker (1990b) undertook an experiment to characterize the nature and magnitude of genetic variability for grain yield and some of its components in nine maize inbreds. The results revealed evidence for existence of enough genetic variability in the parental materials, which is essential for the improvement of a crop. Over dominance as well as non-additive component of genetic variance was observed to be important for all traits except kernel rows per ear. Partial dominance as well as additive gene effect was found to be predominant in the inheritance of kernel rows per ear. Debnath and Sarker (1990b) observed epistasis along with over dominance in the inheritance of ear length. For ear breadth they reported absence of non-allelic interaction in the inheritance. They also reported over dominance for this trait.

Kara (2001) and Alam (2009) estimated that a non-additive gene effect was involved in maize grain yield. The estimated heritability degree of yield (narrow sense; 0.236) is consistency with other researcher's results by Dehghanpour *et al.* (1996), Singh *et al.* (1998), Chaudhary *et al.* (2000), Kara *et al.* (2001) and Muhammad and Muhammad (2002).

Onay *et al.* (2004) reported the letter alternative may apply since the variances for  $H_1$  and  $H_2$  were significantly different from zero. It may thus be concluded that the dominant and recessive alleles of the related genes are distributed equally among the parents. Since the mean dominance effect of the heterozygote locus ( $h^2$ ) was significant, high heterotic effect values would be expected for grain yield among crosses. The parameters, E, an estimate of the environmental variation and D, the additive genetic variation, were not different from zero. The parameter D, which may also include a portion of the additive  $\times$  additive epistatic variances as well as additive genetic variance itself, was non-significant for grain yield. Dominance variance ( $H_1$ ) and corrected dominance variance ( $H_2$ ) were significantly different from zero. It may thus be concluded that grain yield is under the dominance effect. This result was also supported

by the GCA/SCA ratio (0.65). They also showed that over-dominance from the  $W_r$ - $V_r$  graph for the inheritance of grain yield.

Ahmed *et al.* (2011) who found preponderance of recessive alleles of weight of spikes/plant in wheat genotypes. Adel and Ali (2013) had shown that dominant genes being to the origin for grain yield/plant in wheat.

## **HETEROSIS**

Hybrid maize has made a significant contribution to increasing productivity during the course of twentieth century both in the developed as well as the developing world. Exploitation of heterosis in maize can be achieved by using progenitors of different kinds, which may be inbreds, non-inbreds and even a combination of both (Leon and Vasal, 2000). The superiority of variety crosses over the parental varieties was established by Beal (1980). He advocated commercial cultivation of inter-variety hybrids. The basis of inbred-hybrid concept resulted from systematic researches done by East (1908), Shull (1908, 1909) and Jones (1918). Various theories have been pronounced to explain and understand the phenomenon for heterosis.

Jinks (1955) suggested that the non-allelic interaction might be the cause of heterosis rather than the special relation between genes at the same locus Mather (1955) considered heterosis as an expression of genetic balance, which might vary, with the breeding behaviour of species. Jinks and Jones (1958) stated that heterosis was a complex genetically phenomenon depending upon the balance of additive action, dominance and interaction of homozygous / homozygous and homozygous/heterozygous components as well as on the distribution of the genes in parental lines. Williams (1959) suggested that dominance or partial dominance of the alleles for favorable expression of the component characters could explain heterosis. Appreciable percentage of heterosis (112, 90, 43, 93 and 111 to 128%) for grain yield in maize was also reported by Lonquist and Gardner (1961), Ruckij (1963), Akhtar and Sing

(1981) and Gerrish (1983) respectively. Heterosis has been exploited profitably in many cross-pollinated crops by crossing highly selected inbred lines (Mian, 1985). According to him heterosis in  $F_1$  is the combined expression of genetically, cytoplasmic and physiological factors and might be attributed to stimulation resulting from the interaction of different heritable factors of the parents.

Bhalla *et al.* (1979) observed that in general, the crosses between genetically diverse varieties showing superior heterosis values for yield also showed increased hybrid vigour in respect of ear length and ear diameter, but not for number of kernel rows per ear. Hallauer and Miranda (1981) summarized 47 independent reports and found the mean high-parent heterosis for yield to be 8.2% from 1394 varied crosses involving 611 parent varieties.

Debnath (1984, 1987) reported 37.4 to 245.0% heterobeltiosis for kernel yield in maize. Saha and Mukherjee (1985) studied heterosis manifest for number of grains in a set of inter-varietal crosses of maize. The investigation indicated the possibility of maximising heterosis for number of grains by identifying parents possessing the highest level of heterotic potential for number of ovules and per cent of grain conversion. Debnath (1987) studied heterosis in maize using nine maize inbreds and their 36  $F_1$ 'S and observed significant positive heterosis over mid and better parent. Among the component characters, highest percentages of heterosis were observed in number of kernels per row followed by ear length and 1000-kernel weight. Heterosis for ear diameter and kernel rows per ear was low. Seven crosses with highly significant desirable heterobeltiosis for grain yield and some of its attributes were selected for exploiting their heterotic performance. Debnath (1989) estimated 43.05 to 96.74% heterosis over better parent for kernel yield, 3.41 to 8.06% for days to silk and 6.62 to 36.98% for plant height in 18 hybrids from inbred lines. Misevic (1990) estimated 11.3 and 8.7% heterosis for grain yield and plant height, respectively.

Crossa *et al.* (1990) reported mid-parent heterosis ranging from 17.5% to 3.3% in tropical late-maturity yellow germplasm. High parent heterosis (9.6%) for grain yield among crosses was observed in CIMMYT's tropical early and intermediate maturity maize (Beck *et al.* 1990).

Han *et al.* (1991) conducted an experiment with CIMMYT inbred lines of maize and reported that low heterosis in the population crosses could be due mainly to insufficient genetic diversity between the two populations and great genetic variability within the populations and pools involved in their study.

Beck *et al.* (1991) also reported low estimate of high-parent heterosis (16% in U.S. and 9.9% in Mexican environment) in CIMMYT's subtropical and temperate intermediate-maturity maize germplasm, although in subtropical early-maturity germplasm moderate levels of heterosis (13%) was noticed by Vasal *et al.* (1992a). Ordas (1991) reported 32.7% mid-parent heterosis for yield in crosses between American and Spanish populations of maize.

Vasal *et al.* (1992b) reported that high x low yielding parent of tropical and subtropical germplasm exhibited the highest heterosis (23.7%). However they observed high-parent heterosis in crosses between tropical germplasms ranging from 12.7% to 3.1%. The magnitude of heterosis exhibited in any crop is extremely important in decision making process to initiate hybrid development efforts. Though optimum yield heterosis levels will differ for each crop, there is a consensus among most breeders that a heterosis level of at least 20% is desirable (Vasal, 2000).

Vasal *et al.* (1993a) reported 15.6% high-parent heterosis for grain yield in quality protein maize (QPM) populations. They stated that crosses between parents with divergent endosperm modifications resulted in higher heterosis for grain yield. Crosses between hard and soft endosperm parents were intermediate in their endosperm rating, indicating that polygenic system was involved in another study. Vasal *et al.* (1993b) reported that a cross combination involving QPM parent's population with white and yellow kernel

color showed 14.8% heterosis in Mexican environment and 19.7% in U.S. environment. They observed many high yielding crosses that involved parents with different kernel color.

Gomaa and Shaheen (1994) obtained heterotic effects for grain yield/plant of F1 hybrids ranging from 23.5 to 36.3% of the mid-parent and from 32.0 to 18.7% of the better parent.

Kim and Ajala (1996) reported that several factors could have influenced level of heterosis. They obtained highest yield in white x yellow cross. Relative distance between the two color groups might increase levels of heterosis; grain texture was another factor that might influence levels of heterosis. The advantage of dent x flint crosses over dent x dent or flint x flint was reported by Wellhausen (1978) and Kim *et al.* (1985).

Dass *et al.* (1997) conducted an experiment to identify bold seeded parental lines and to exploit them in the breeding program for developing high yielding hybrids/composite cultivars in maize. Combining ability analysis revealed that both additive and non-additive genetic variances were important in the expression of seed weight and yields in winter and summer seasons. Results on SCA effects indicated that most of the superior crosses were between high x low and high x medium parents and that involvement of one good general combiner was essential to get better specific combination. They suggested development of season specific hybrids.

Roy *et al.* (1998) obtained -16.42 to 71.82% heterobeltiosis for yield in crosses among six genetically diverse composite parents.

Koirala and Gurung (2002) observed that high parent heterosis for grain yield ranged from -17.8 to 23.9%.

Uddin *et al.* (2006) explained that the better performing four crosses (P<sub>1</sub>xP<sub>7</sub>, P<sub>6</sub>xP<sub>7</sub>, P<sub>1</sub>xP<sub>4</sub> and P<sub>4</sub>xP<sub>5</sub>) can be utilized for developing high yielding hybrid varieties as well as for exploiting hybrid vigor. Except cross P<sub>2</sub>xP<sub>5</sub>, rest of the crosses showed significant positive heterobeltiosis both in plant and ear height.



Alam *et al.* (2008) studied that the highest % of heterosis for grain/ear over mid parent and better parent were observed by the cross  $P_2 \times P_3$ . Crosses  $P_1 \times P_3$  and  $P_1 \times P_5$  showed significant negative heterosis for days to maturity.

Uddin *et al.* (2008) reported that the highest significant positive heterosis (28.41%) for grain yield was observed in the cross IPB911-11  $\times$  BM-7.

Amanullah *et al.* (2011) reported that 21 crosses had positive heterosis for grain yield/ha and ranged from +0.39% to 86%. The heterosis and heterobeltiosis for 1000-seed weight was 12.9% and 4.57%.

Singh *et al.* (2012) obtained highest heterosis in the crosses of HUZQPUM 3-2  $\times$  HUZQPM 4-2 and HUZQPUM 1-1  $\times$  HUZQPM 3-2 for grain yield and number of rows per ear.

Parihar *et al.* (2012) observed maximum economic heterosis in all the environments for grain yield per plant in the crosses  $P_9 \times P_{12}$  and  $P_4 \times P_{12}$ .

## MATERIALS AND METHODS

The details of materials used and methods followed in carrying out the present investigation are presented in this chapter.

### EXPERIMENT I

#### 3.1.1 MATERIALS

The experimental material consisted of 25 maize (*Zea mays* L.) inbred lines. They were IL1, IL2, IL3, IL4, IL5, IL6, IL7, IL8, IL9, IL10, IL11, IL12, IL13, IL14, IL15, IL16, IL17, IL18, IL19, IL20, IL21, IL22, IL23, IL24 and IL25. The maize inbred lines were received from Biometrical Genetics Lab, Department of Botany, University of Rajshahi, Bangladesh.

#### 3.1.2 METHODS

The methods followed to conduct the experiments and analysis of data is described below.

##### **Preparation of the Experimental Field**

The experiment was carried out in the Botanical Research Garden of Rajshahi University during the Rabi crop season of 2008, 2009 and 2011. The experimental land was first opened on the month of November in every experimental year and the operations were done by power tiller and harrows. The land was again ploughed and cross-ploughed with a power tiller followed by laddering during the course of final land preparation. All crop residues and weeds were removed completely before layout of the field and sowing of the inbred seeds. The field was pulverized and leveled properly. As the experimental field was sufficiently moist, no irrigation was given before sowing of the seeds. The recommended practices were provided to raise a good crop. Irrigation, weeding and other intercultural operations were done as and when necessary.

### **Soil and Climatic Condition of the Experimental Field**

The experimental field area was a medium high land. The texture of the soil was fertile and silty loam having pH 5.2-6.4. The topography of the field was medium high land above flood level. It was readily broken when pulverized. Well drained sandy loam and medium loam soils, rich in humus are most suitable for maize. The climate of the location is characterized by relatively low temperature and little rainfall during rabi or winter season (November to March) and high temperature and rainfall during kharif or summer season (April to October).

### **Experimental Layout and Size of the Experimental Field**

Layout of the experimental field and 25 inbred lines were sown in a Randomized Block Design with three replications. The field comprised total area of 12×18 square meter having 3 replications. Size of each replication was 16.5 m × 5 m having 25 rows (lines). Each line contained 10 plants. The distance was 100 cm from row to row and 30 cm from plant to plant and in each row ten hills was maintained. In each hill, one plant was maintained. The inbred lines were randomly assigned in each line of replication plot.

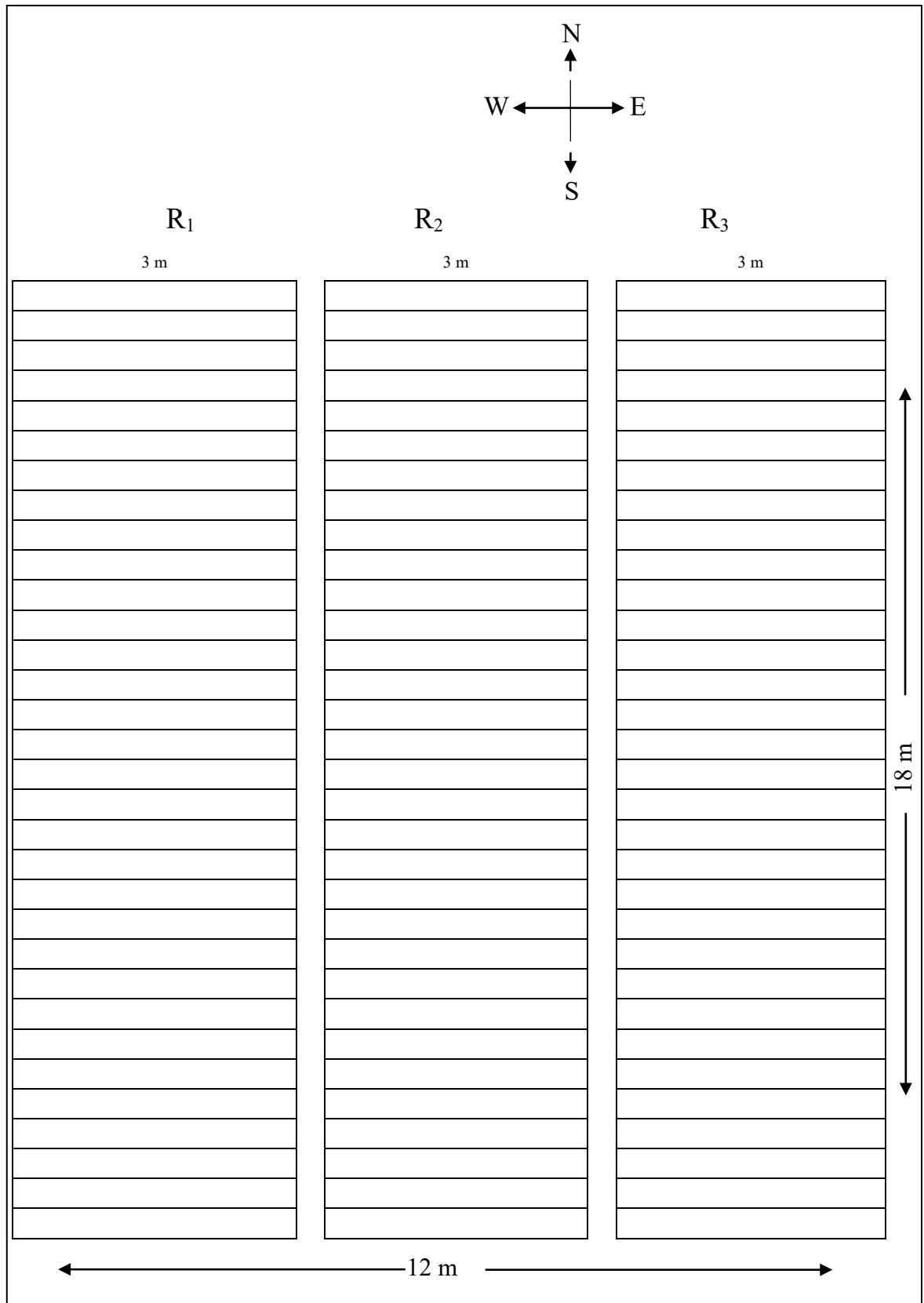


Figure 3.1.1 Design of the experimental field

### **Sowing of Maize Inbred Lines and Raising of Seedlings**

Twenty-five maize inbred lines were randomly assigned to the inner 25 lines in each replication. Before sowing, inbreds were treated with vitavex-200 (0.3%). After sowing, inbred lines were covered with soil. The sowing date in three years was 24 November 2008, 26 November 2009 and 28 November 2010, respectively. When seedling emergence was completed 10 plants were kept in each row (line) of each replication.

### **Intercultural Operation and Disease Control**

Weeding was done twice, one within 25-30 days after sowing and other within 40-45 days after sowing; the plants were 15-20 cm in height. Irrigation was done to the experimental plot as needed. The crops were almost free from diseases and insect pests. Only at the early stage of growth, seedlings were sprayed appropriated insecticide.

### **Collection of Data**

Five plants were tagged randomly for recording observations for each entry for all the quantitative characters. Mean of five plants for each entry in each replication was worked out for each character at each replication and used for statistical analysis. Data on the following quantitative characters were recorded at appropriate stages of plant growth.

#### **i). Days to tasseling**

The number of days from sowing up to the day on which 50% of the plants showed tassel emergence was recorded as days to 50% tasseling.

#### **ii). Days to silking**

The number of days from sowing up to the day on which 50% of plants showed silk emergence was recorded as days to 50% silking.

**iii). Days to maturity (DM)**

Maturity time was recorded in days from the date of planting to the date of yellowish layer formation of grain of 50% population.

**iv). Plant height (cm)**

Height of the plant from ground level up to the base of fully opened flag leaf was recorded in centimeters as plant height when plants were mature.

**v). Ear height (cm)**

Height from ground level up to the base of the upper most bearing internode was recorded as ear height in centimeters.

**Collection of Data on Harvesting and Cob Related Characters**

The maize inbred lines were harvested at 90 days (5 March 2009, 9 March 2010 and 12 March 2011). When the color of leaf turned yellow and dropped off then plants of individual orientation as tagged previously were separately harvested. Harvested inbred lines were cured and slotted properly. Data on different quantitative characters were collected on individual plant basis from five plants randomly selected in each line of each replication. All the measurements were taken in CGS system. Cob related data were measured and recorded on the following characters:

**vi). Cob length (cm)**

Length of the ear was measured and recorded in centimeters at the time of harvest as its total length (from the base to the tip of the ear).

**vii). Cob diameter (cm)**

Cob diameter was measured and recorded in centimeters as the thickness of the ear i.e., at the middle of the ear.

**viii). Number of kernel rows /cob**

Number of kernel rows per cob was counted and recorded.

**ix). Number of kernels /row**

Number of kernels per row was counted and average was recorded as number of kernels per row.

**x). Number of kernels /cob (number of grains/ear)**

The total number of kernels per cob was counted and recorded.

**xi). Grain yield/plant (g)**

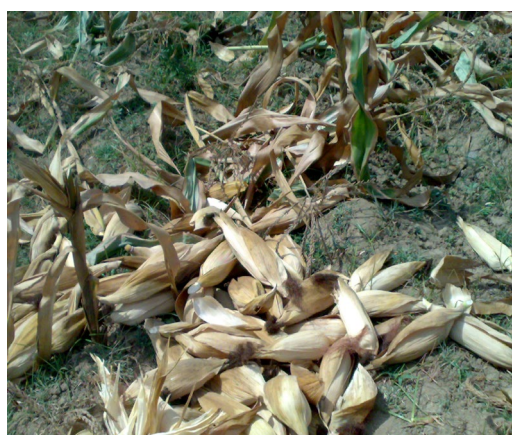
Grain yield per plant expressed in grams was recorded by weighing the grains obtained after shelling of cobs from individual plant.



Exp. I. Seedling stage of the maize inbreds



Exp.I. Maturity stage



Collected cob with husk



Collected cob of different inbreds



Exp.II Crossing block



Crossing block

**Plate No. 1: Different stages of inbreds and crossing block**





Collection of F<sub>1</sub> seed



Collection of F<sub>1</sub> seeds



Lay out of experiment II



Sowing period of F<sub>1</sub> seeds



F<sub>1</sub> plants



F<sub>1</sub> plants

**Plate No. 2: F<sub>1</sub> seeds and F<sub>1</sub> plants**

### Techniques of Statistical Analysis of Data

Mean data of three years were analyzed in the first experiment. The collected data were analyzed following the biometrical techniques of analysis as developed by Mather (1949) based on mathematical model of Fisher (1936).

Genotypic variance, phenotypic variance, genotypic coefficient of variation and phenotypic coefficient of variation were estimated as suggested by Singh and Chaudhary (1985).

Broad sense heritability was calculated as suggested by Johnson *et al.* (1955) and genetic advance was estimated using the formulae suggested by Johnson *et al.* (1955) and Hanson *et al.* (1956).

Genotypic and phenotypic covariances were estimated according to the formulae suggested by Singh and Chaudhary (1985).

Genotypic and phenotypic correlation coefficients were calculated using the formulae suggested by Miller *et al.* (1958).

Path coefficient analysis was done following the method as suggested by Dewey and Lu (1959). As per suggestion of Robinson *et al.* (1951) yield was also included as one of the independent characters.

The expected genetic advance from straight selection [GAS] and from the discriminant function [GAD] was first estimated and the expected gain from the discriminant function over straight selection was calculated for all the functions studied as follows: Expected gain (%) =  $[(GAD/GAS)-1] \times 100$ . Multivariate analysis was done based on Mahalanobis analysis,  $D^2$ - statistics (Mahalanobis, 1936).

General combining ability (GCA) and specific combining ability (SCA) were estimated by following Model-1, Method of Griffing (1956). The mean squares for GCA and SCA were tested against error variance desired. Mean data were used to estimate heterosis over mid parent and better parent according to Rai (1979).

The techniques used for analysis of data are described under the following sub-heads:

**(i) Mean**

Data on individual plant basis were added together and then divided by the total number of observations and the mean was obtained as follows:

$$\text{Mean } (\bar{x}) = \frac{1}{n} \sum_{i=1}^n X_i$$

Where,

X = The individual reading was recorded on each plant

n = Number of observations

i = 1, 2, 3.....n

$\Sigma$  = Summation

**(ii) Standard deviation (SD)**

Standard deviation is the average deviation of the individual observation from the mean. It was calculated as the square root of the variance as follows:

$$SD = \sqrt{\sigma^2}$$

Where,

SD = Standard deviation

$\sigma^2$  = Variance

**(iii) Standard error of mean (SE)**

If several samples are considered instead of taking one, it will be found that the standard deviations of the different samples also vary. This variation is measured by the standard error of mean, which was calculated as follows:

$$SE = \frac{SD}{\sqrt{n}}$$

Where,

SD = Standard deviation

SE = Standard error of mean

n = Total number of individual

**(iv) Coefficient of variability in percentage**

Coefficient of variability in percentage (CV %) was calculated according to the following formula:

$$CV\% = \frac{S}{\bar{x}} \times 100$$

Where,

CV% = Coefficient of variability in percentage

S = Standard deviation,

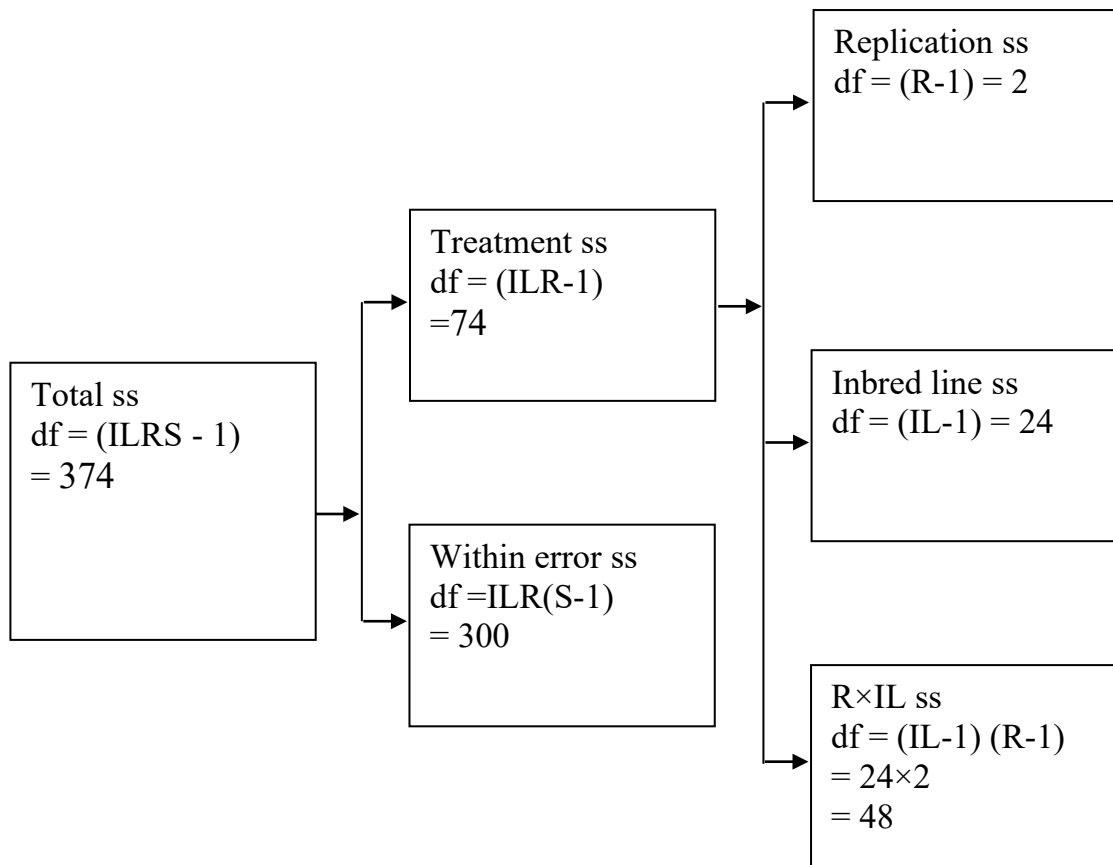
$\bar{x}$  = Genotypic mean.

**(v) Analysis of variance**

Variance is a measure of dispersion of a population. So, the analysis of variance is done for testing the significant differences among the populations. Variance analysis for each of the characters was carried out separately on individual plant value of a row.

The variances due to different sources such as; line (L), replication(R), interaction of R × L and within error (E) of a population were calculated as per the following skeleton of analysis.

The variance due to different sources such as inbred line (IL), replication (R), interaction of (IL×R) and within error (E) of population were calculated as per the following skeleton of analysis:



Where,

$$\text{Total ss} = \sum (\text{LRS})^2 - \text{CF}$$

$$\text{Treatment ss} = \frac{\sum_{ij} (\text{L}_i \text{R}_j)^2}{\text{S}} - \text{CF}$$

$$\text{Replication ss} = \frac{\sum_j \text{R}_j^2}{\text{SL}} - \text{CF}$$

$$\text{Line ss} = \frac{\sum_i \text{L}_i^2}{\text{SR}} - \text{CF}$$

$$(\text{L} \times \text{R}) \text{ ss} = \frac{\sum_{ij} (\text{L}_i \text{R}_j)^2}{\text{S}} - \text{CF} - \text{L}_{\text{ss}} - \text{R}_{\text{ss}}$$

$$\text{Error ss} = \text{Total ss} - \text{Treatment ss} - \text{Replication ss}$$

$\text{L}_i$  = The value of  $i^{\text{th}}$  line,

$\text{R}_j$  = The total of  $j^{\text{th}}$  replication

$L_iR_j$  = The value of  $i^{\text{th}}$  line in  $j^{\text{th}}$  replication

S = The value of the  $s^{\text{th}}$  replication

CF = Correction factor =  $(GT)^2 / N$

GT = Grand total,

N = Total number of observations = (SLR)

The analysis of variance of a mixed model was used, where line (L) was fixed and replication (R) effect is random.

The expectation of mean square (E.M.S) is derived as follows.

**Table 3.1.2.1 Analysis of variance (ANOVA)**

| Source of variation | df          | MS              | EMS   |
|---------------------|-------------|-----------------|---|
| Replication (R)     | R-1         | MS <sub>1</sub> | $\sigma_{we}^2 + S\sigma_R^2$                   |
| Inbred line(L)      | L-1         | MS <sub>2</sub> | $\sigma_{we}^2 + S\sigma_{LR}^2 + RS\sigma_L^2$ |
| R×L                 | (R-1) (L-1) | MS <sub>3</sub> | $\sigma_{we}^2 + S\sigma_{LR}^2$                |
| Within error        | RL(s-1)     | MS <sub>4</sub> | $\sigma_{we}^2$                                 |

Where,

L, R, S designated for line (genotype), replication and sib respectively.

MS<sub>1</sub>= Represents mean square of replication.

MS<sub>2</sub>= Represents mean square of line.

MS<sub>3</sub>= Represents mean square of R×L

MS<sub>4</sub>= Represents mean square of within error and

$S\sigma_R^2$  = Variance due to replication

$RS\sigma_L^2$  = Variance due to line

$S\sigma_{LR}^2$  = Variance due to L×R

$\sigma_{we}^2$  = Variance due to within error

\*, \*\* and \*\*\* denoted 1%, 5% and 0.1% level, respectively.

**(vi) Components of variation**

The components of variation were phenotypic ( $\sigma^2_p$ ), genotypic ( $\sigma^2_g$ ), Replication ( $\sigma^2_R$ ), interaction ( $\sigma^2_{LR}$ ) and error ( $\sigma^2_w$ ) variances. These were measured as follows:

$$\begin{aligned}\text{Step-I: } \sigma^2_R &= (MS_1 - MS_4) / sl \\ \sigma^2_L &= (MS_2 - MS_3) / rs \\ \sigma^2_{LR} &= (MS_3 - MS_4) / s \\ \sigma^2_{we} &= MS_4\end{aligned}$$

Step-I I:

- i) Phenotypic variance ( $\sigma^2_p$ ) =  $\sigma^2_L + \sigma^2_{LR} + \sigma^2_{we}$
- (ii) Line variance ( $\sigma^2_g$ ) =  $\sigma^2_L$
- (iii) Replication variance =  $\sigma^2_R$
- (iv) Line  $\times$  Replication variance =  $\sigma^2_{LR}$
- (v) Error variance =  $\sigma^2_{we}$

**vii) Estimation of genotypic and phenotypic coefficient of variation**

Genotypic coefficient of variation (GCV) and phenotypic coefficient of variation (PCV) were calculated as per Singh and Chaudhary (1985) by the following formulae.

a) Phenotypic coefficient of variability

$$PCV = \sqrt{\frac{\sigma^2_p}{\bar{x}}} \times 100$$

b) Genotypic (Line) coefficient of variability

$$GCV = \sqrt{\frac{\sigma^2_g}{\bar{x}}} \times 100$$

c) Genotype  $\times$  Replication coefficient of variability

$$G \times RCV = \sqrt{\frac{\sigma^2_{lr}}{\bar{x}}} \times 100$$

d) Error coefficient of variability

$$ECV = \sqrt{\frac{\sigma^2_e}{\bar{x}}} \times 100$$

Where,

$$\begin{aligned}\sigma^2_g &= \text{Genotypic variance} \\ \sigma^2_p &= \text{Phenotypic variance,} \\ \bar{x} &= \text{Population mean}\end{aligned}$$

### (viii) Heritability ( $h^2_b$ )

Heritability (in broad sense) estimates was computed by dividing the genotypic variance with phenotypic variance and then multiplying by 100 as suggested by Warner (1952).

$$h^2_b (\%) = \frac{\sigma^2_g}{\sigma^2_p} \times 100$$

Where,

$h^2_b$  = Heritability in broad sense

$\sigma^2_g$  = Genotypic variance

$\sigma^2_p$  = Phenotypic variance

### (ix) Genetic advance (GA)

Genetic advance was calculated by the following formula as suggested by Allard (1960):

$$GA = K \cdot \sigma_p (\sigma^2_g / \sigma^2_p)$$

Where,

K = the selection differential in standard units for the present study it was 2.06 at 5% level of selection (Lush, 1949).

$\sigma_p$  = Square root of the phenotypic variance

$\sigma^2_p$  = Phenotypic variance

$\sigma^2_g$  = Genotypic variance

### (x) Genetic advance in percentage of mean (GA %)

Genetic advance in percentage of mean was calculated by the following formula given by Comstock and Robinson (1952):



$$GA \% = \frac{GA}{\bar{X}} \times 100$$

Where,

$\bar{x}$  = Population mean

### (xi) Analysis of covariance

For the purpose of correlation coefficients and path-coefficient, the analysis of both variance and covariance are required (Miller *et al.*, 1958). Therefore, covariance was calculated between all possible pairs of characters.

Mean value per replication per genotype of three years were arranged in combined table and analyses of covariance were done as per following formula:

$$\text{Cov.} = \frac{\sum_{i=1}^n X_i Y_i - \left( \sum_{i=1}^n X_i \right) \left( \sum_{i=1}^n Y_i \right) / n}{n-1}$$

Where,

Cov. = Covariance

$\sum_{i=1}^n X_i Y_i$  = Sum of X and Y

$\sum_{i=1}^n X_i$  = Grand total of X

$\sum_{i=1}^n Y_i$  = Grand total of Y

n = Number of observation

n-1 = Degrees of freedom

i = 1,2,3 .....n

$\Sigma$  = Summation

The expectation of mean cross product (MCP) was derived as follows:

**Table 3.1.2.2. Analysis of covariance**

| Source of variation | df  | MS | EMS  |
|---------------------|-----|----|--|
| Replication (R)     | 2   |    | $\partial^2_{12} + G\partial^2R_{12}$                      |
| Inbred line (G)     | 24  |    | $\partial^2_{12} + \partial^2R_{12} + RL\partial^2_{12}$   |
| G×R                 | 48  |    | $\partial^2_{12} + R\partial^2GY_{12} + R\partial^2Y_{12}$ |
| Error (E)           | 300 |    | $\partial^2_{12}$  |

$G\partial^2R_{12}$  = Covariance due to replication.

$R\partial^2G_{12}$  = Covariance due to genotype

$R\partial^2GY_{12}$  = Covariance due to G×Y.

$\partial^2_{12}$  = Covariance due to error.

The phenotypic ( $\partial^2P_{12}$ ), genotypic ( $\partial^2g_{12}$ ), interaction ( $\partial^2GR_{12}$ ) and error covariance ( $\partial^2_{12}$ ) were determined as follows:

### Step-1

$$\partial^2g_{12} = (\text{MCP}_g - \text{MCP}_{g \times r})/R$$

$$\partial^2GR_{12} = (\text{MCP}_{g \times r} - \text{MCP}_e)/R$$

$$\partial^2_{12} = \text{MCP}_e$$

### Step-2

a. Phenotypic covariance ( $\partial^2P_{12}$ ) =  $\partial^2g_{12} + \partial^2_{12}$

b. Genotypic covariance ( $\partial^2g_{12}$ ) =  $\partial^2g_{12}$

### (xii) Correlation coefficient

The correlation coefficient at phenotypic ( $r_p$ ), genotypic ( $r_g$ ) and environmental ( $r_e$ ) levels were calculated as follows:

$$r_p = (\partial^2p_{12}) / (\partial^2p_{11} \times \partial^2p_{22})^{1/2}$$

$$r_g = (\partial^2g_{12}) / (\partial^2g_{11} \times \partial^2g_{22})^{1/2}$$

$$r_e = (\partial^2e_{12}) / (\partial^2e_{11} \times \partial^2e_{22})^{1/2}$$

Where,

$\partial^2 p_{12}$ ,  $\partial^2 g_{12}$  and  $\partial^2 e_{12}$  represent phenotypic, genotypic and environmental covariance of character 1 and 2.  $\partial^2 p_{11}$ ,  $\partial^2 g_{11}$  and  $\partial^2 e_{11}$  represent phenotypic, genotypic and environmental levels of character 1.  $\partial^2 p_{22}$ ,  $\partial^2 g_{22}$  and  $\partial^2 e_{22}$  indicate variance at phenotypic, genotypic and environmental levels of character 2.

**(xiii) Path coefficient**

The path coefficient analysis was carried out using the formula and Wright (1923) as illustrated by Dewey and Lu (1959). The path-coefficient analysis was done at both phenotypic and genotypic levels by solving the simultaneous equation using matrix method.

The form of equation is as follows:

$$r_{xy} = P_{xy} + r_{x2} P_{2y} + r_{x3} P_{3y} + \dots + r_{xn} P_{ny}$$

Where,

$r_{xy}$  = correlation between one components character and yield.

$P_{xy}$  = Path-coefficient between the same character and yield.

$r_{x2}, r_{x3} \dots r_{xn}$  = Represent correlation coefficient between that character and each of the other yield components in turn.

The above equation was written in a matrix form as:

$$\begin{matrix} \text{A} \\ \left[ \begin{array}{c} r_{1y} \\ r_{2y} \\ r_{3y} \\ r_{iy} \end{array} \right] \end{matrix} = \begin{matrix} \text{B} \\ \left[ \begin{array}{cccc} r_{11} & r_{12} & r_{13} & r_{1J} \\ r_{21} & r_{22} & r_{23} & r_{2J} \\ r_{31} & r_{32} & r_{33} & r_{3J} \\ r_{i1} & r_{i2} & r_{i3} & r_{iJ} \end{array} \right] \end{matrix} \times \begin{matrix} \text{C} \\ \left[ \begin{array}{c} P_{1y} \\ P_{2y} \\ P_{3y} \\ P_{iy} \end{array} \right] \end{matrix}$$

$$A = B \times C; \text{ Then } C = B^{-1} A$$

Where,

$P_{ry}$  = direct effect of the character  $i$  on the dependent trait  $y$  (yield).

The indirect effect of a particular character through other characters was obtained by multiplication of direct path and particular correlation coefficient between those two characters respectively.

$$\text{Indirect effect} = r_{ij} \times P_{iy}$$

Where,

$$i = 1, \dots, n,$$

$$j = 1, \dots, n,$$

$$P_{iy} = P_{1y} \dots P_{ny}$$

Where,  $r_{ij}$  = correlation coefficient between two independent characters.

The residual effect is assumed to be independent to the remaining variables. It was calculated from the formula as proposed by Wright (1923).

$$\text{Residual effect } (\chi) = 1 - R^2$$

$$R^2 = P_{1y} + P_{2y} r_{2y} + \dots + P_{ny} r_{ny}$$

$R^2$  is the required multiple correlation coefficient and is the amount of variation in yield that can be accounted for by the component characters.

**(xiv) Selection index**

The coefficients,  $b_1, b_2, \dots, b_n$  used in the discriminant function technique was obtained from the genotypic and phenotypic variances and covariances arranged in the matrix as follow:

$$\begin{bmatrix} X \\ X_{11} & X_{12} & X_{13} & X_{1j} \\ X_{21} & X_{22} & X_{23} & X_{2j} \\ X_{31} & X_{32} & X_{33} & X_{3j} \\ X_{i1} & X_{i2} & X_{i3} & X_{ij} \end{bmatrix} \begin{bmatrix} b \\ b_1 \\ b_2 \\ b_3 \\ b_n \end{bmatrix} = \begin{bmatrix} G \\ G_{11} & G_{12} & G_{13} & G_{1j} \\ G_{21} & G_{22} & G_{23} & G_{2j} \\ G_{31} & G_{32} & G_{33} & G_{3j} \\ G_{i1} & G_{i2} & G_{i3} & G_{ij} \end{bmatrix} \begin{bmatrix} a \\ a_1 \\ a_2 \\ a_3 \\ a_n \end{bmatrix}$$

The solution of this matrix gave the estimates of 'b' values in the following manner (Singh and Chaudhury, 1985).

$$b = X^{-1} GA$$

Where,

'b' is the column vector,  $X^{-1}$ , is the inverse of phenotypic variance and covariance matrix, 'G' is the genotypic variance and covariance matrix and 'a' is the column vector for economic weights. Assuming that all the characters are of economically equal importance, i.e.,  $a_1 = a_2 = a_3 = 1$ .

The values obtained for  $b_1, b_2, \dots, b_n$  were used in discriminant function selection technique. The phenotypic and genotypic variances and covariances as obtained were used for constructing the discriminant function using different character combinations according to the method as developed by Fisher (1925) and Smith (1936). Yield/plant was also included as one of the independent characters as suggested by Robinson *et al.* (1951). The expected genetic advance from straight selection {GA(S)} and from discriminant function {GA(D)} was calculated as follows:

$$GA(S) = (Z/P) \times (g_{yy}) / (t_{yy}) \frac{1}{2} \text{ and}$$

$$GA(D) = (Z/P) \times (b_1 g_{1y} + b_2 g_{2y}) \frac{1}{2}$$

Where,

$Z/P$  = the selection differential in standard units, for the present study it was 2.06 at 5% level of selection (Lush, 1949).

$g_{yy}$  and  $t_{yy}$  = the genotypic and phenotypic variances of character.

$b_1, b_2, \dots, b_n$  = the relative weights for character.

$g_{1y}, g_{2y}, \dots$  = the genotypic co variances of independent character with y.

The expected gain from the discriminant function over straight selection was calculated for all the functions as shown below:

$$\text{Expected gain (\%)} = [GA(D)/GA(S)] \times 100.$$

**(xv) Genetic divergence**

After analysis of variance and covariance, the data were  $D^2$  statistics. First,  $D^2$  values of all the individual population is (n-1) combinations were arranged in ascending order. After arranging the  $D^2$  values in this manner a method suggested by Rao (1952) was used for cluster formation.

After formation of the cluster on the basis of  $D^2$  values, the average intra-cluster  $D^2$  values were obtained by the formula  $\frac{\sum di^2}{n}$  where  $di^2 =$  sum of the distances between all possible combinations (n) of the populations included in a cluster. In this way, average inter-cluster  $D^2$  values were also obtained between any two groups. The square roots of the  $D^2$  values represented the distance between and within groups. A measure for group distance based on multiple characters was given by Mahalanobis (1936).

With  $x_1, x_2, x_3, \dots, x_p$  as the  $d_p$  as  $\bar{x}_1^1 - \bar{x}_1^2, \bar{x}_2^1 - \bar{x}_2^2, \dots, \bar{x}_p^1 - \bar{x}_p^2$ , respectively, being the difference in the means of two populations, Mahalanobis'  $D^2$  – statistics is defined as follows:

$${}_pD^2 = b_1d_1 + b_2d_2 + \dots + b_p d_p$$

Here,

The  $b_i$  value is to be estimated such that the ratio of variance between the populations to the variance within the populations is maximized. In terms of variances and covariance, the  $D^2$  value is obtained as follows:

$${}_pD^2 = W^{ij} (\bar{x}_i^1 - \bar{x}_i^2)(\bar{x}_j^1 - \bar{x}_j^2) \dots (1)$$

Where,

$W^{ij}$  is the inverse of estimated variance co-variance matrix.

## **EXPERIMENT II**

### **3.2.1 MATERIALS**

The genetic materials used in this experiment were six parents (IL<sub>4</sub>=P<sub>1</sub>, IL<sub>5</sub>=P<sub>2</sub>, IL<sub>18</sub>=P<sub>3</sub>, IL<sub>10</sub>=P<sub>4</sub>, IL<sub>23</sub>=P<sub>5</sub> and IL<sub>1</sub>=P<sub>6</sub>) and their F<sub>1</sub> which were P<sub>1</sub>×P<sub>2</sub>, P<sub>1</sub>×P<sub>3</sub>, P<sub>1</sub>×P<sub>4</sub>, P<sub>1</sub>×P<sub>5</sub>, P<sub>1</sub>×P<sub>6</sub>, P<sub>2</sub>×P<sub>3</sub>, P<sub>2</sub>×P<sub>4</sub>, P<sub>2</sub>×P<sub>5</sub>, P<sub>2</sub>×P<sub>6</sub>, P<sub>3</sub>×P<sub>4</sub>, P<sub>3</sub>×P<sub>5</sub>, P<sub>3</sub>×P<sub>6</sub>, P<sub>4</sub>×P<sub>5</sub>, P<sub>4</sub>×P<sub>6</sub> and P<sub>5</sub>×P<sub>6</sub>.

### **3.2.2 METHODS**

#### **Field Experiment**

The present investigation was carried out during 26 November 2009 and 28 November 2010 seasons at the experimental field, Department of Botany, University of Rajshahi, Bangladesh.

The genetic materials were crossed in half diallel fashion in 26 November 2009 and to obtain the F<sub>1</sub> seeds. The F<sub>1</sub> seeds of all the crosses with their parents were planted in the field in 28 November 2010 season for evaluation in a randomized block design with three replications. The experimental unit was single row of 3 m long. Inter-plant and inter-row distances were 10 and 30 cm, respectively. All other treatments were kept constant for the whole experiment.

#### **Collection of Data**

Five plants from each row were randomly selected for recording data on the studied characters such as days to tasseling, days to silking, days to maturity, plant height, ear height, cob length, cob diameter, number of rows/cob, number of kernels/row, number of kernels/cob and grain yield/plant.

#### **Techniques of Statistical Analysis of Data**

##### **(i) Diallel and Combining Ability Analysis**

The collected data were analyzed following the biometrical techniques of analyses according the Method 1 (Parents+F<sub>1</sub>s = half diallel) given by Griffings

(1956). In this study, six parents ( $n=6$ ) were involved in the diallel, producing 15  $F_1$ s [ $n(n-1)/2$ ] and without reciprocals i.e., there were 21 total entries, 15 crosses and 6 parents. Techniques of analyses of the data are described under the following sub-heads:

### Testing the significance differences

The data were first analyzed to test the significance of crossing differences. If the mean squares due to crossing are significant, there is need to proceed for further analyses. The total variability in the treatments was partitioned into components like variance due to replication, crosses (including parents), interaction ( $C \times R$ ) and within error.

The sums of squares were calculated as follows:

$$\text{Correction factor} = (\text{Grand total})^2 / r \times (n \times n) \times s$$

$$\text{Total}_{ss} = \text{Individual observation}^2 - \text{CF}$$

$$\text{Treatment}_{ss} = \sum X_i^2 / s - \text{CF}$$

$$\text{Replication}_{ss} = \frac{\sum X_j^2}{c \times s} - \text{CF}$$

$$\text{Cross (including parents)}_{ss} = \sum X_k^2 / rs - \text{CF}$$

$$C \times R_{ss} = \text{Total}_{ss} - \text{Treatment}_{ss} - \text{Replication}_{ss}$$

$$\text{Within error}_{ss} = \text{Total}_{ss} - \text{Treatment}_{ss}$$

Here,  $X_i$  = Replication total

$X_j$  = Treatment total

$X_k$  = Cross (including parents) total

$r$  = Number of replications

$n$  = Number of parents

$c$  = Number of crosses ( $n \times n$ )

$s$  = Number of plants or sibs



**Table 3.2.1.1. Preparation of ANOVA**

| Sources of variation               | df  | SS              | MS                                    | EMS   | VR <sub>1</sub>                  | VR <sub>2</sub>                  |
|------------------------------------|-----|-----------------|---------------------------------------|---|----------------------------------|----------------------------------|
| Treatment                          | 74  | SS <sub>1</sub> | SS <sub>1</sub> /df = MS <sub>1</sub> |   | MS <sub>1</sub> /MS <sub>5</sub> | MS <sub>1</sub> /MS <sub>4</sub> |
| Replication(R)                     | 2   | SS <sub>2</sub> | SS <sub>2</sub> /df = MS <sub>2</sub> | $\sigma^2w_{11} + \dots + cs\sigma^2R_{11}$           | MS <sub>2</sub> /MS <sub>5</sub> | MS <sub>2</sub> /MS <sub>4</sub> |
| Crosses (C)<br>(including parents) | 21  | SS <sub>3</sub> | SS <sub>3</sub> /df = MS <sub>3</sub> | $\sigma^2w_{11} + s\sigma^2I_{11} + rs\sigma^2C_{11}$ | MS <sub>3</sub> /MS <sub>5</sub> | MS <sub>3</sub> /MS <sub>4</sub> |
| Interaction<br>(I) (C×R)           | 48  | SS <sub>4</sub> | SS <sub>4</sub> /df = MS <sub>4</sub> | $\sigma^2w_{11} + s\sigma^2I_{11}$                    |                                  |                                  |
| Within error (W)                   | 300 | SS <sub>5</sub> | SS <sub>5</sub> /df = MS <sub>5</sub> | $\sigma^2w_{11}$                                      |                                  |                                  |
| Total                              | 374 |                 |                                       |   |                                  |                                  |

**Estimation of variance and covariance**

A number of first and second degree statistics (Mather, 1955) were calculated from the mean data. With the environmental expectation (E) included, the statistics of the above parameters may be shown as follows (Hayman, 1954 b):

$$\text{Parental mean} = \frac{\text{Sum of all the diagonal values}}{\text{Number of parents}}$$

$$V_0L_0 = \frac{1}{n-1} \left[ \sum \text{Diagonal values}^2 - \frac{(\sum \text{Diagonal values})^2}{\text{Number of diagonal values}} \right]$$

$$V_r = \frac{1}{n-1} \left[ \sum \text{Crosses involving a particular parent}^2 - \frac{(\sum \text{Crosses involving a particular parent})^2}{\text{Num of parents}} \right]$$

$$V_1L_1 = \frac{1}{n} \sum V_{ri}$$

$$W_r = \left[ \sum \text{Arrays} \times \text{no - recurring parents} - \frac{\sum \text{Arrays} \sum \text{Diagonal values}}{\text{Number of parents}} \right]$$

$$W_0L_0 = \frac{1}{n} \sum W_{ri}$$

$$V_0L_1 = \frac{1}{n-1} \left[ \sum \text{Array means}^2 - \frac{(\sum \text{Array means})^2}{\text{Number of arrays}} \right]$$

$$(ML_1 - ML_0)^2 = \left[ \frac{1}{n} \left\{ \frac{1}{n} \text{Grand total} - \text{Diagonal values} \right\} \right]^2$$

The above statistics may be defined as follows:

$$V_0L_0 = \text{Variance of parents.}$$

$V_r$  = Variance of each array.

$V_1L_1$  = Mean variance of the arrays.

$W_r$  = Covariance between parents and their offsprings.

$W_0L_0$  = Mean covariance between the parents and the arrays.

$V_0L_1$  = Variance of the mean arrays.

$(ML_1 - ML_0)^2$  = The difference between the mean of the parents and the mean of their  $n^2$  progenies.

The environmental variation (E) is calculated by using the following formula:

$$E = 1/r \left\{ \frac{\text{Interaction Error ss} + \text{Within Error ss} + \text{Rep. ss}}{\text{Interaction Error df} + \text{Within Error df} + \text{Rep. df}} \right\}$$

### Testing the validity of the hypothesis

The probable fulfillment of the hypothesis (Hayman, 1954 b) is tested by using the following formula:

$$t^2 = \frac{n-2}{4} \left[ \frac{\{\text{Var}(V_r) - \text{Var}(W_r)\}^2}{\text{Var } V_r \times \text{Var } W_r - \text{Cov}^2(V_r, W_r)} \right]$$

Which is an F with 4 and  $(n-2)$  degrees of freedom.

When,

$$\text{Var}(W_r) = \frac{1}{n-1} \left[ \{W_{ri}^2 - \frac{(\sum W_{ri})^2}{n}\} \right]$$

$$\text{Var}(V_r) = \frac{1}{n-1} \left[ \{V_{ri}^2 - \frac{(\sum V_{ri})^2}{n}\} \right]$$

$$\text{Cov}(V_r, W_r) = \frac{1}{n-1} \left[ \{V_r W_r - \frac{\sum V_r \sum W_r}{n}\} \right]$$

Here,

$\text{Var}(W_r)$  = Variance of  $W_r$

$\text{Var}(V_r)$  = Variance of  $V_r$

$\text{Cov}(V_r, W_r)$  = Covariance between  $V_r$  and  $W_r$

This is tested against the table value of “F” with 4 and  $(n - 2)$  degrees of freedom. Its significance indicates failure of the hypothesis. Another way of testing the hypothesis is through the regression coefficient, calculated by using the following formula:

$$b = \frac{\text{Cov}(V_r, W_r)}{\text{Var}(V_r)}$$

Where,

$$\text{Cov}(W_r, V_r) = \left[ \sum V_r W_r - \frac{\sum V_r \sum W_r}{n} \right] / (n - 1) \quad \text{and}$$

$$\text{Var}(V_r) = \left[ \sum V_r^2 - \frac{(\sum V_r)^2}{n} \right] / (n - 1)$$

Therefore,

$$b = \frac{\text{Cov}(W_r, V_r)}{\text{Var}(V_r)}$$

$$\text{Standard error (b)} = [(\text{Var } W_r - b \text{ Cov } W_r V_r) / \text{Var } V_r (n - 2)]^{1/2}$$

Now the significance of  $b$  from zero and unity can be tested as follows:

$$\begin{aligned} H_0 b &= 0 \\ &= (b - 0) / \text{SE}(b) \text{ and} \end{aligned}$$

$$\begin{aligned} H_0: b &= 1 \\ &= (1 - b) / \text{SE}(b) \end{aligned}$$

These values are tested against table value of “t” for  $(n - 2)$  degrees of freedom.

### Components of variation and their proportions

The expected values of the components of variation obtained by least square computations are as follows:

$$D = V_0 L_0 - E$$

$$F = 2V_0 L_0 - 4W_0 L_{01} - 2(n - 2) E/n$$

$$H_2 = 4V_1 L_1 - 4V_0 L_1 - 2E$$

$$h^2 = 4 (ML_1 - ML_0)^2 - 4 (n - 1) E/n^2$$

$$Fr = 2 (V_0L_0 - W_0L_{01} + V_1L_1 - W_r - V_r) - 2 (n - 2) E/n$$

The above components are genetic parameters:

D = Variation due to additive effect. F = The mean of “Fr” over the arrays.

H<sub>1</sub> = Component of variation due to the dominance effect of the genes.

h<sup>2</sup> = Dominance effect (as algebraic sum over all loci in heterozygous phase in all crosses).

Fr = The covariance of additive and dominance effects in a single array.

$$H_2 = H_1 [1 - (u - v)^2]$$

Where, U = Proportion of positive genes in the parents.

V = Proportion of negative genes in the parents.

$$\text{Thus, } H_2 = 4V_1L_1 - 4V_0L_1 - 2E$$

For testing the significance of each of these components, respective standard errors were calculated. Here the common multiplier or variance (S<sup>2</sup>) was calculated using the following formula:

$$S^2 = \frac{1}{2} [\text{Var} (W_r - V_r)]$$

And the specific multiplier was calculated with the following formula:

$$D = (n^5 + n^4)/n^5$$

$$F = (4n^5 + 20n^4 - 16n^3 + n^2)/n^5$$

$$H_1 = (n^5 + 41n^4 - 12n^3 + 4n^2)/n^5$$

$$H_2 = (36n^4)/n^5$$

$$h^2 = (16n^4 + 16n^2 - 32n + 16)/n^5$$

$$E = n^4/n^5$$

The standard errors for the different estimates were then calculated using the specific multiplier and common multiplier, which are as follows:

$$SE (D) = \{ \text{Specific multiplier} \times \text{Common multiplier} (s^2) \}^{1/2}$$

### Graphical analysis

Diallel analysis for the components of genetic variances and Vr-Wr graphs for all the characters studied were done according to Hayman (1954a, b).

A diallel table was prepared from the averages over all the three replicates and the following statistics were estimated. Vr = Variance of all progenies in each parental array (an array is a group of crosses involving a particular parents).

Wr = Covariance between parents and their offspring in each array. The validity of Hayman's hypothesis was tested for all the characters studied by the equation.

### Combining ability analysis

In the combining ability analysis the data are rearranged in Table 3.4. In this table, each value is the mean value in all the replications. The total variability in the population may, therefore, be partitioned into components like variance due to general combining ability (gca), specific combining ability (sca) and error. Using replicate mean the various sum of squares are obtained as follows:

$$\text{ss due to gca} = \frac{1}{2n} \left[ \sum (Y_{i.} + Y_{.j})^2 - \frac{2}{n} Y_{..}^2 \right]$$

$$\text{ss due to sca} = 1/2 \sum \sum Y_{ij}(Y_{ij} + Y_{ji}) - 1/2n \sum (Y_{.j} + Y_{i.})^2 + 1/n^2 Y^2$$

**Table 3.2.1.2. ANOVA for combining ability in method I**

| Source | df | SS              | MS              | EMS   |
|--------|----|-----------------|-----------------|---|
| gca    | 5  | SS <sub>1</sub> | MS <sub>g</sub> | $\sigma^2_e + \sigma^2_g \cdot 2(n-1) / n + 2n\sigma^2_g$ |
| sca    | 14 | SS <sub>2</sub> | MS <sub>s</sub> | $\sigma^2_e + 2(n^2 - n + 1) \sigma^2_s / n^2$            |
| Error  | 40 | SS <sub>3</sub> | MSe             | $\sigma^2_e$  |

Where,

gca = General combining ability.

sca = Specific combining ability.

$Y_{ij}$  = Mean of  $i \times j$  th cross

MSg = Mean square of gca effects.

MSs = Mean square of sca effects.

MSe = Mean square of error.

The mean of sum of squares due to error divided by the number of replications. Mean error variance,  $MS_g$  and  $MS_s$  have been calculated from the mean data, mean error variance is therefore, required for F-test.

$$\text{Thus } MS'(\text{error}) = \frac{MS(\text{error})}{\text{Number of replications}}$$

The general combining ability effects are defined as follows:

$$g_i = 1/2n(Y_{i.} + Y_{.j}) - 1/n^2Y_{..}$$

The specific combining ability effects are defined as follows:

$$S_{ij} = 1/2(Y_{ij} + Y_{ji}) - 1/2n(Y_{i.} + Y_{.i} + Y_{.j} + Y_{jj}) + 1/n^2Y_{..}$$

## (ii) Heterosis Study

A diallel cross of  $6 \times 6$  excluding reciprocals for obtaining  $F_1$  was conducted in the period from October, 2010 to March, 2011 was the materials of this investigation. Data on eleven quantitative traits were analyzed following the techniques given below. Techniques of the analysis of data:

### Estimation of mid-parent and better-parent

For estimation of heterosis in each parameter the mean values of the 15  $F_1$ s have been compared with better-parents (BP) for heterobeltosis and with mid-parent (MP) for heterosis over mid parent value.

Percent heterosis was calculated as

$$\text{Heterosis (MP)} = \frac{\bar{F}_1 - \text{MP}}{\text{MP}} \times 100$$

$$\text{Heterosis (BP)} = \frac{\bar{F}_1 - \text{BP}}{\text{BP}} \times 100$$

Standard error for each individual and overall heterosis was calculated. Significance tests for heterosis were done by using pooled error from the analysis of variance of  $F_1$  and parental populations.

$$\text{Mid-parent} = 1/2(P_1 + P_2)$$

$$\text{Variance of mid-parent} = 1/4(VP_1 + VP_2)$$

$$\text{Variance of } F_1 = VF_1$$

$$\text{Standard error of mean of MP and } F_1 = \sqrt{1/4VP_1 + 1/4VP_2 + VF_1}$$

Here, N = Total number of populations ( $P_1 + P_2 + F_1$ )

t = Estimated value of MP heterosis / Standard error of mean

$$\text{Standard of error of mean of BP} = \sqrt{VBP} / N$$

Here, N = Total number of populations ( $F_1$ )

t = Estimated value of BP heterosis / standard error of mean of BP

A general specification of heterosis must, therefore, be able to accommodate heterosis both in the positive ( $F_1 > P_1$ ) and in the negative ( $F_1 < P_2$ ) directions.

If heterosis is measured on a scale on which an additive–dominance model is adequate, then for positive heterosis, its expected magnitude is given by

Heterosis =  $\bar{F}_1 - P_1 = [h] - [d]$  and for heterosis to occur [h] must be positive and greater than [d]. For negative heterosis the comparable expectation is given

by Heterosis =  $\bar{F}_1 - P_2 = [h] - (-[d])$  and heterosis will occur only when [h] is

negative and greater than [d]. If the additive–dominance model is inadequate, its specification becomes complex.

### **Test of significance for heterosis**

The significance of heterosis was tested with the help of critical difference (CD) (Mian, 1985; Singh and Narayanan, 1993).

CD= SE of difference + “t” at 5% or 1% level of probability at respective degrees of freedom.

$$SE(H_1)=\sqrt{(3/2\times V_e/r)}$$

$$SE(H_1)=\sqrt{(2\times V_e/r)}$$

Where,

$V_e$ = error mean square of the ANOVA involving parents,  $F_1$ s and the commercial/ cultivated variety.



## RESULTS

### Experiment I

In the present study 25 maize inbred lines were evaluated for eleven quantitative characters *viz.*, days to tasseling, days to silking, days to maturity, plant height, ear height, cob length, cob diameter, no. of rows/cob, no. of kernels/row, no. of kernels/cob and grain(kernel)yield/plant. Collected data were analyzed in order to estimate mean with standard error, coefficient of variability, heritability, genetic advance, genetic advance as percentage of mean, correlation coefficient, path coefficient, selection index and genetic diversity with  $D^2$ - statistics. The results derived from these different statistical analyses are described under different heads.

#### 4.1.1 GENETIC VARIABILITY

##### Mean with Standard Error and Coefficient of Variability

Mean with standard error (SE) and coefficient of variability as percentage (CV %) for eleven characters for twenty five maize inbred lines was calculated from the pooled data and the results are shown in Tables 4.1.1- 4.1.6. The maximum days to tasseling (83.667) was recorded for IL-15, whereas the lowest days to tasseling (65.265) were noted for IL-4. On the other hand, CV% was highest in inbred line IL-10 and lowest in IL-5 for the days to tasseling (Table 4.1.1). The maximum days to silking (94.267) was exhibited by IL4, while the minimum value (84.667) was observed in IL-18. The highest CV% was recorded in IL-18 and the lowest in IL-7 (Table 4.1.1). The maximum days to maturity (129.733) was noted in IL-5 while the lowest (114.733) was in IL-10. The highest CV% was noted in IL-10 and the lowest CV% in IL-2 (Table 4.1.2). The highest plant height (111.529) was exhibited by IL-4, IL-5 and IL1-5 and the lowest was noted for IL-1 (96.459). The maximum CV% was recorded in IL-24 and the minimum in IL-3 and IL-13 (Table 4.1.2).

For ear height, the highest mean was recorded in IL-24, IL-21 (52.470) and the lowest mean was recorded in IL-1 and IL-2 (36.677). The highest CV% was noted in IL-14 and the lowest CV% was noted in IL-24 (Table 4.1.3).

The highest cob diameter was noted in IL-19, IL-17, IL-22, IL-14 (13.414) and the lowest was noted in IL-21 (9.933). The highest CV% was noted in IL-4 and IL-18 and the lowest in IL-12 (Table 4.1.3). For cob length, the maximum (13.181) was recorded in IL-24, IL-9 and the lowest (9.996) was recorded in the IL-2. The highest CV% was noted in the IL-15 and IL-21 and the lowest was noted in IL-3 (Table 4.1.4).

The maximum number of rows/cob (13.333) was noted in IL-23 while the lowest (10.733) was noted in IL-5. The highest CV% was noted in IL-2 and IL-22 and the lowest CV% in IL-7 (Table 4.1.4). For number of kernels/row, the maximum (15.333) was recorded in IL-4 and IL-13 and the minimum (12.600) was recorded in IL-22. The highest CV% was noted in IL-13 and the lowest was noted in IL-22 (Table 4.1.5). The maximum number of grains/cob was noted in IL-18 (72.133) and the lowest was noted in IL-8 (63.533). The highest CV% was noted in IL-10 and the lowest in IL-11 and IL-9 (Table 4.1.5). For grain yield (grain weight/plant), the maximum (84.904 g) was recorded in the inbred line IL-1 and the lowest (65.577 g) was recorded in IL-21.

The highest CV% was noted in IL-15 and the lowest was noted in IL-4 (Table 4.1.6). Twenty five inbred lines were evaluated to find out the extent of genetic variability for the respective characters (days to tasseling, days to silking, days to maturity, plant height, ear height, cob length, cob diameter, number of kernel rows/ear, number of kernels/row, number of grains/cob and grain yield/plant). Significant variations were existed among the inbred lines for the characters studied. Mean performances of 25 inbred lines are presented in Tables 4.1.1-4.1.6.

Duration of silking ranged from 84.667 to 94.266 days. The inbred line IL-4 and IL-5 took the longest time to silk and maturity which was statistically different from the other inbreds.

Inbreds IL-18, IL-10 and IL-23 took the shortest period for silking and maturity and were statistically different from the others. IL-4, IL-5 and IL-15 were the tallest inbred lines which were statistically identical with IL-3 and IL-19. The shortest height was produced by IL-1. Inbreds IL-21 and IL-24 showed the highest ear height where as IL-1 and IL-2 appeared as the shortest for the height. The inbred lines IL-19 and IL-24 showed the highest mean performance for ear length and appreciable ear diameter. Inbred line IL-23 had the highest number of kernel rows/cob which was statistically identical with IL-19 and IL-18.

The highest number of kernels/row was recorded in IL-6. Number of grains/cob ranged from 63.533 to 72.134. The grain yield/plant (weight/plant) ranged from 65.523 to 84.904 g. The inbred line IL-1 produced the highest grain yield which was statistically similar with IL-23, IL-18 and IL-14. Yield performance of IL-4 was poor. So, the best performance was observed in IL-4, IL-5, IL-18, IL-10, IL-23 and IL-1 inbred lines on the basis of tasseling, flowering and different important characters.

**Table 4.1.1. Mean with SE, CV % for days to tasseling and days to silking of 25 inbred lines of maize**

| Inbred lines | Days to tasseling |         | Days to silking |       |
|--------------|-------------------|---------|-----------------|-------|
|              | Mean±SE           | CV%     | Mean±SE         | CV%   |
| IL-1         | 78.600±1.530      | 7.5391  | 88.865±0.930    | 4.054 |
| IL-2         | 76.866±1.555      | 7.8346  | 90.867±1.009    | 4.300 |
| IL-3         | 70.265±1.675      | 9.2304  | 92.534±0.975    | 4.082 |
| IL-4         | 65.264±1.026      | 6.0866  | 94.265±0.796    | 3.269 |
| IL-5         | 75.867±0.956      | 4.8779  | 93.736±0.796    | 3.287 |
| IL-6         | 74.134±1.068      | 5.5820  | 89.667±1.330    | 5.744 |
| IL-7         | 71.933±1.329      | 7.1557  | 91.200±0.579    | 2.459 |
| IL-8         | 67.865±1.576      | 8.9951  | 91.800±0.611    | 2.578 |
| IL-9         | 72.933±1.148      | 6.0981  | 92.134±0.990    | 4.161 |
| IL-10        | 75.800±2.664      | 13.6119 | 92.535±0.668    | 2.796 |
| IL-11        | 75.000±1.082      | 5.5891  | 93.067±0.628    | 2.615 |
| IL-12        | 72.266±1.119      | 5.9968  | 92.736±0.918    | 3.834 |
| IL-13        | 77.200±1.096      | 5.5000  | 91.534±0.755    | 3.195 |
| IL-14        | 75.134±1.383      | 7.1297  | 91.538±0.682    | 2.887 |
| IL-15        | 83.667±2.184      | 10.1082 | 91.337±0.591    | 2.506 |
| IL-16        | 74.934±1.307      | 6.7572  | 89.400±0.872    | 3.777 |
| IL-17        | 78.465±1.764      | 8.7087  | 88.269±0.733    | 3.218 |
| IL-18        | 74.935±1.416      | 7.3176  | 84.663±1.701    | 7.779 |
| IL-19        | 79.266±1.465      | 7.1598  | 90.937±1.322    | 5.630 |
| IL-20        | 80.200±1.808      | 8.7307  | 92.800±1.176    | 4.908 |
| IL-21        | 76.066±2.207      | 11.2374 | 87.937±1.343    | 5.916 |
| IL-22        | 77.400±1.588      | 7.9464  | 87.338±0.860    | 3.814 |
| IL-23        | 78.600±2.210      | 10.8894 | 91.539±0.689    | 2.916 |
| IL-24        | 79.067±1.777      | 8.7032  | 90.737±1.569    | 6.697 |
| IL-25        | 79.865±1.496      | 7.2526  | 87.334±1.508    | 6.686 |
|              | LSD =6.822        |         | LSD= 6.551      |       |

**Table 4.1.2. Mean with SE, CV % for days to maturity and plant height (cm) of 25 inbred lines of maize**

| Inbred lines | Days to maturity |        | Plant height  |        |
|--------------|------------------|--------|---------------|--------|
|              | Mean±SE          | CV%    | Mean±SE       | CV%    |
| IL-1         | 124.665±0.659    | 2.0488 | 96.459±1.447  | 5.811  |
| IL-2         | 122.600±0.524    | 1.654  | 101.223±1.648 | 6.306  |
| IL-3         | 124.734±0.556    | 1.7265 | 108.927±1.332 | 4.737  |
| IL-4         | 123.738±0.714    | 2.235  | 111.529±1.961 | 6.811  |
| IL-5         | 129.780±0.796    | 2.376  | 111.123±2.101 | 7.323  |
| IL-6         | 127.812±1.135    | 3.438  | 106.630±1.542 | 5.600  |
| IL-7         | 125.600±0.809    | 2.496  | 106.300±1.943 | 7.081  |
| IL-8         | 125.134±0.970    | 3.005  | 103.153±2.412 | 9.056  |
| IL-9         | 123.150±1.009    | 3.173  | 104.307±1.806 | 6.706  |
| IL-10        | 114.735±3.202    | 10.808 | 104.793±2.297 | 8.488  |
| IL-11        | 124.732±1.333    | 4.137  | 103.079±2.026 | 7.611  |
| IL-12        | 123.401±1.644    | 5.159  | 101.869±1.888 | 7.179  |
| IL-13        | 123.800±1.314    | 4.109  | 99.257±1.228  | 4.791  |
| IL-14        | 120.335±1.804    | 5.805  | 100.503±2.222 | 8.564  |
| IL-15        | 123.370±0.950    | 2.981  | 111.007±1.860 | 6.488  |
| IL-16        | 124.665±0.766    | 2.380  | 104.975±2.107 | 7.774  |
| IL-17        | 122.136±1.664    | 5.278  | 105.394±2.238 | 8.223  |
| IL-18        | 125.930±0.848    | 2.607  | 104.053±1.846 | 6.871  |
| IL-19        | 122.920±1.136    | 3.578  | 106.065±2.141 | 7.817  |
| IL-20        | 124.825±1.294    | 4.014  | 98.586±1.855  | 7.287  |
| IL-21        | 124.461±1.059    | 3.296  | 106.351±1.677 | 6.108  |
| IL-22        | 118.412±1.447    | 4.732  | 104.735±1.976 | 7.307  |
| IL-23        | 117.465±1.064    | 3.508  | 97.860±2.896  | 11.459 |
| IL-24        | 119.861±2.225    | 7.189  | 100.483±3.434 | 13.235 |
| IL-25        | 121.012±0.946    | 3.028  | 101.740±1.999 | 7.611  |
|              | LSD = 7.359      |        | LSD=12.057    |        |

**Table 4.1.3. Mean with SE, CV % for ear height (cm) and cob diameter (cm) of 25 inbred lines of maize**

| Inbred lines | Ear height   |        | Cob diameter |        |
|--------------|--------------|--------|--------------|--------|
|              | Mean±SE      | CV%    | Mean±SE      | CV%    |
| IL-1         | 39.160±1.789 | 17.69  | 10.553±0.339 | 12.451 |
| IL-2         | 39.677±1.346 | 13.142 | 11.682±0.358 | 11.879 |
| IL-3         | 45.058±2.205 | 18.949 | 10.839±0.351 | 12.552 |
| IL-4         | 41.913±1.744 | 16.119 | 11.884±0.429 | 13.983 |
| IL-5         | 43.260±0.963 | 8.62   | 12.216±0.403 | 12.778 |
| IL-6         | 47.546±2.012 | 16.387 | 10.536±0.47  | 17.278 |
| IL-7         | 48.817±1.415 | 11.227 | 11.391±0.507 | 17.224 |
| IL-8         | 50.281±1.173 | 9.034  | 12.283±0.39  | 12.304 |
| IL-9         | 48.330±1.352 | 10.838 | 10.863±0.48  | 17.099 |
| IL-10        | 44.965±2.228 | 19.194 | 12.299±0.468 | 14.726 |
| IL-11        | 51.768±1.516 | 11.342 | 10.637±0.37  | 13.454 |
| IL-12        | 49.764±2.177 | 16.942 | 12.562±0.289 | 8.905  |
| IL-13        | 48.578±2.457 | 19.593 | 12.748±0.367 | 11.154 |
| IL-14        | 50.643±2.858 | 21.857 | 13.080±0.238 | 7.047  |
| IL-15        | 47.139±1.554 | 12.771 | 10.043±0.307 | 11.845 |
| IL-16        | 46.287±1.689 | 14.134 | 12.977±0.343 | 10.229 |
| IL-17        | 50.021±2.074 | 16.057 | 13.153±0.383 | 11.276 |
| IL-18        | 49.160±2.323 | 18.302 | 12.284±0.435 | 13.726 |
| IL-19        | 48.915±2.164 | 17.131 | 13.415±0.327 | 9.455  |
| IL-20        | 49.043±2.129 | 16.811 | 11.696±0.349 | 11.563 |
| IL-21        | 52.559±1.944 | 14.33  | 9.935±0.35   | 13.628 |
| IL-22        | 49.930±1.322 | 10.258 | 13.217±0.409 | 11.988 |
| IL-23        | 50.761±2.091 | 15.951 | 12.754±0.452 | 13.719 |
| IL-24        | 52.470±1.048 | 7.738  | 12.339±0.384 | 12.058 |
| IL-25        | 46.089±2.308 | 19.396 | 11.537±0.332 | 11.134 |
| LSD = 7.359  |              |        | LSD=12.057   |        |

**Table 4.1.4. Mean with SE, CV % for cob length (cm) and for number of row/cob of 25 inbred lines of maize**

| Inbred lines | Cob length   |        | Number of rows/cob |        |
|--------------|--------------|--------|--------------------|--------|
|              | Mean±SE      | CV%    | Mean±SE            | CV%    |
| IL-1         | 10.901±0.318 | 11.315 | 12.134±0.435       | 13.885 |
| IL-2         | 9.996±0.230  | 8.908  | 11.868±0.542       | 17.696 |
| IL-3         | 12.012±0.237 | 7.627  | 11.867±0.413       | 13.463 |
| IL-4         | 11.551±0.373 | 12.522 | 11.334±0.303       | 10.369 |
| IL-5         | 12.055±0.361 | 11.561 | 10.738±0.463       | 16.691 |
| IL-6         | 12.303±0.395 | 12.423 | 12.736±0.463       | 14.069 |
| IL-7         | 10.829±0.392 | 14.004 | 12.467±0.274       | 8.503  |
| IL-8         | 12.226±0.415 | 13.154 | 11.734±0.33        | 10.908 |
| IL-9         | 13.095±0.330 | 9.748  | 11.400±0.412       | 13.988 |
| IL-10        | 12.926±0.526 | 15.754 | 11.866±0.477       | 15.56  |
| IL-11        | 11.911±0.391 | 12.712 | 11.938±0.316       | 10.247 |
| IL-12        | 11.960±0.335 | 10.851 | 12.135±0.435       | 13.885 |
| IL-13        | 10.959±0.36  | 12.722 | 11.936±0.316       | 10.247 |
| IL-14        | 11.581±0.418 | 13.984 | 11.464±0.291       | 9.815  |
| IL-15        | 11.225±0.479 | 16.533 | 11.931±0.396       | 12.853 |
| IL-16        | 10.898±0.402 | 14.284 | 12.601±0.375       | 11.54  |
| IL-17        | 10.246±0.389 | 14.688 | 12.000±0.39        | 12.599 |
| IL-18        | 11.005±0.451 | 15.862 | 13.002±0.352       | 10.483 |
| IL-19        | 12.433±0.466 | 14.518 | 13.200±0.49        | 14.374 |
| IL-20        | 11.616±0.372 | 12.412 | 12.401±0.349       | 10.905 |
| IL-21        | 10.692±0.449 | 16.265 | 11.266±0.396       | 13.613 |
| IL-22        | 12.389±0.400 | 12.496 | 11.934±0.53        | 17.193 |
| IL-23        | 11.598±0.460 | 15.356 | 13.266±0.473       | 13.801 |
| IL-24        | 13.181±0.310 | 9.11   | 12.334±0.287       | 9.022  |
| IL-25        | 12.225±0.441 | 13.946 | 12.067±0.371       | 11.914 |
| LSD= 2.5606  |              |        | LSD= 2.563         |        |

**Table 4.1.5. Mean with SE, CV % for number of kernes/row and number of grains/cob of 25 inbred lines of maize**

| Inbred lines | Number of kernels/row |             | Number of grains/cob |        |
|--------------|-----------------------|-------------|----------------------|--------|
|              | Mean±SE               | CV%         | Mean±SE              | CV%    |
| IL-1         | 14.600±0.375          | 9.959       | 68.732±2.409         | 13.576 |
| IL-2         | 13.534±0.376          | 10.768      | 66.001±1.93          | 11.324 |
| IL-3         | 14.00±0.436           | 12.074      | 69.136±1.956         | 10.96  |
| IL-4         | 15.334±0.433          | 10.932      | 66.864±1.561         | 9.042  |
| IL-5         | 14.067±0.452          | 12.449      | 66.067±1.089         | 6.383  |
| IL-6         | 14.735±0.502          | 13.198      | 69.000±2.287         | 12.835 |
| IL-7         | 12.867±0.363          | 10.939      | 67.810±1.831         | 10.462 |
| IL-8         | 13.000±0.458          | 13.637      | 63.531±1.104         | 6.727  |
| IL-9         | 12.739±0.384          | 11.674      | 66.400±0.94          | 5.483  |
| IL-10        | 13.265±0.441          | 12.889      | 69.412±2.635         | 14.703 |
| IL-11        | 13.067±0.419          | 12.43       | 65.665±0.871         | 5.137  |
| IL-12        | 12.938±0.431          | 12.894      | 66.732±1.62          | 9.4    |
| IL-13        | 15.267±0.539          | 13.665      | 68.131±2.394         | 13.609 |
| IL-14        | 12.930±0.431          | 12.894      | 66.610±1.253         | 7.285  |
| IL-15        | 12.739±0.248          | 7.548       | 67.665±1.861         | 10.652 |
| IL-16        | 13.600±0.335          | 9.547       | 67.869±1.226         | 6.997  |
| IL-17        | 13.336±0.361          | 10.48       | 65.801±1.662         | 9.785  |
| IL-18        | 13.612±0.349          | 9.943       | 72.132±1.486         | 7.979  |
| IL-19        | 13.067±0.267          | 7.904       | 65.268±0.897         | 5.322  |
| IL-20        | 13.400±0.363          | 10.478      | 69.332±2.37          | 13.238 |
| IL-21        | 13.335±0.454          | 13.195      | 68.269±1.127         | 6.396  |
| IL-22        | 12.612±0.214          | 6.572       | 66.264±1.274         | 7.447  |
| IL-23        | 14.339±0.494          | 13.359      | 64.869±1.287         | 7.683  |
| IL-24        | 14.412±0.466          | 12.533      | 67.605±2.441         | 13.987 |
| IL-25        | 14.131±0.456          | 12.505      | 71.412±2.428         | 13.168 |
| LSD= 2.583   |                       | LSD= 11.187 |                      |        |



**4.1.6. Mean with SE, CV % for grain yield/plant of 25 inbred lines of maize**

| Inbred lines | Mean±SE       | CV%    |
|--------------|---------------|--------|
| IL-1         | 84.904±2.9824 | 13.604 |
| IL-2         | 69.478±1.5444 | 8.6094 |
| IL-3         | 74.863±1.0745 | 5.5587 |
| IL-4         | 65.523±0.4783 | 2.8273 |
| IL-5         | 72.057±1.9259 | 10.351 |
| IL-6         | 71.326±2.2217 | 12.064 |
| IL-7         | 70.768±2.0575 | 11.26  |
| IL-8         | 65.679±1.9035 | 11.225 |
| IL-9         | 70.674±0.5981 | 3.2776 |
| IL-10        | 71.010±2.1374 | 11.658 |
| IL-11        | 78.892±1.8962 | 9.3088 |
| IL-12        | 71.116±0.6957 | 3.7887 |
| IL-13        | 72.955±1.7702 | 9.3975 |
| IL-14        | 82.376±1.466  | 6.8926 |
| IL-15        | 78.913±3.6075 | 17.705 |
| IL-16        | 75.559±2.1471 | 11.006 |
| IL-17        | 72.750±0.7011 | 3.7324 |
| IL-18        | 82.571±1.5644 | 7.3381 |
| IL-19        | 80.540±1.3992 | 6.7286 |
| IL-20        | 70.020±0.5706 | 3.1559 |
| IL-21        | 65.579±2.1321 | 12.592 |
| IL-22        | 79.535±1.7682 | 8.6101 |
| IL-23        | 83.998±2.3857 | 11.002 |
| IL-24        | 82.066±0.6435 | 3.0371 |
| IL-25        | 75.638±1.3539 | 6.9324 |
| LSD=11.01754 |               |        |

## Analysis of Variance

The analysis of variance for all the eleven characters was done separately and the results are shown in Tables 4.1.7– 4.1.9. For testing the effects of the main items and their interaction effects, a mixed model was followed.

Inbred line (L) item was highly significant at 5% and 1% level for all the characters. The interaction of line with replication (L×R) was also highly significant for most of the characters. Significant line item indicated that there were significant differences among the lines for these characters considered in this study.

**Table 4.1.7. Analysis of variance for days to tasseling, days to silking, days to maturity and plant height of 25 inbred lines of maize**

| Sources of variation | df  | Days to tasseling | Days to silking | Days to maturity | Plant height |
|----------------------|-----|-------------------|-----------------|------------------|--------------|
|                      |     | MS                | MS              | MS               | MS           |
| Total                | 374 | 21.331            | 19.546          | 34.610           | 74.704       |
| Treatment            | 74  | 41.372            | 37.519          | 97.611           | 170.041      |
| Line                 | 24  | 88.225**          | 79.734**        | 155.134**        | 248.028**    |
| Replication          | 2   | 64.904            | 42.674          | 98.696           | 305.064      |
| Line×Rep.            | 48  | 16.965**          | 16.196**        | 68.804**         | 125.422**    |
| Within error         | 300 | 16.388            | 15.113          | 19.070           | 51.187       |

**Table 4.1.8. Analysis of variance for ear height, cob diameter, cob length and number of rows/cob of 25 inbred lines of maize**

| Sources of variation | df  | Ear height | Cob diameter | Cob length | Number of rows/cob |
|----------------------|-----|------------|--------------|------------|--------------------|
|                      |     | MS         | MS           | MS         | MS                 |
| Total                | 374 | 69.119     | 3.138        | 2.999      | 2.621              |
| Treatment            | 74  | 136.633    | 7.330        | 3.263      | 3.871              |
| Line                 | 24  | 197.908**  | 16.265**     | 3.672*     | 5.463*             |
| Replication          | 2   | 750.804    | 0.021        | 9.739      | 7.784              |
| Line×Rep.            | 48  | 80.405**   | 3.167**      | 2.789**    | 2.911**            |
| Within error         | 300 | 52.466     | 2.105        | 2.934      | 2.313              |

**Table 4.1.9. Analysis of variance for number of kernels/row, and number of kernels/cob of 25 inbred lines of maize**

| Sources of variation | df  | Number of kernels/row | Number of kernels/cob | Grain yield/plant |
|----------------------|-----|-----------------------|-----------------------|-------------------|
|                      |     | MS                    | MS                    | MS                |
| Total                | 374 | 2.954                 | 519.562               | 47.596            |
| Treatment            | 74  | 5.407                 | 942.962               | 61.907            |
| Line                 | 24  | 9.395**               | 947.161**             | 68.778**          |
| Replication          | 2   | 36.642                | 9760.248              | 543.050           |
| Line×Rep.            | 48  | 2.112**               | 573.475**             | 43.425**          |
| Within error         | 300 | 2.349                 | 415.124               | 44.067            |

### Genetic Parameters

The estimates of phenotypic ( $\sigma^2_p$ ), genotypic ( $\sigma^2_g$ ) and error ( $\sigma^2_e$ ) components of variation were calculated separately for all the eleven agronomic characters of twenty five maize inbred lines and the results are presented in the Table 4.1.10.

#### (i) Phenotypic Variation ( $\sigma^2_p$ )

For all the characters phenotypic variation ( $\sigma^2_p$ ) was greater than those of  $\sigma^2_g$  and  $\sigma^2_e$  components of variation as expected. The phenotypic variation is the joint product of  $\sigma^2_g$  and  $\sigma^2_e$ . Table 4.1.10 shows that the greater portion of the total phenotypic variation appeared mostly due to error variation for all the characters. The highest value of  $\sigma^2_p$  was observed for number of grains /cob (1223.911) and the lowest was shown by cob length (5.418). The remaining characters followed with their lower to higher values were as days to silking, plant height, ear height, grain yield/plant, days to tasseling, days to maturity and cob diameter.

#### (ii) Genotypic Variation ( $\sigma^2_g$ )

The highest genotypic variation ( $\sigma^2_g$ ) was found for number of grains/cob with a value of 808.787, while the lowest genotypic variation was recorded for cob length with a value of 3.123 cm.

The other characters according to lower to higher values were as cob length, number of rows/cob, cob diameter, days to maturity, number of kernels/row, ear height, days to tasseling and grain yield/plant (Table 4.1.10.).

### **(iii) Error Variation ( $\sigma^2e$ )**

The highest error variation ( $\sigma^2e$ ) was recorded for number of grains/cob with a value of 415.124 and the lowest was noted for cob length with a value of 2.294 cm (Table 4.1.10.).

### **(iv) Coefficient of Variability**

The estimates of phenotypic (PCV), genotypic (GCV) and error (ECV) coefficient of variability for eleven characters were calculated and the results are presented in Table 4.1.10. In general, the phenotypic coefficient of variability (PCV) was greater than the genotypic and error coefficient of variability for all the characters. The estimates of the phenotypic coefficient of variability were the highest for number of grains/cob (788.765) and the lowest PCV was estimated for days to maturity (9.738). The PCVs cob diameter, days to silking and cob diameter were 32.0556, 28.962 and 24.636 (Table 4.1.10.). Genotypic co-efficient of variability was highest for number of grains/cob, while the lowest GCV was estimated for days to maturity.

Error coefficient of variability was high for number of grains/cob followed by grain yield/plant and ear height. However the lowest value of ECV was exhibited by days to maturity (Table 4.1.10.).

### **(v) Heritability ( $h^2b$ ), Genetic advance (GA) and Genetic advance in percentage of mean (GA %)**

For all the eleven grain yield and yield contributing characters, heritability in broad sense, genetic advance and genetic advance as percentage of mean were calculated separately and the results are shown in the Table 4.1.10.

**Heritability ( $h^2b$ )**

The highest heritability was observed for days to silking (93.406) and this was followed by days to maturity (88.638), days to tasseling (86.033), plant height (81.858), cob length (82.068), ear height (77.795), number of kernels/cob (77.183) and cob diameter (73.646) (Table 4.1.10.). The lowest value of heritability was 66.082 observed for number of grains/cob.

**Genetic advance (GA)**

The estimation of genetic advance shows the highest value for number of grains/cob (47.624) that was followed by days to silking (37.588), days to tasseling (29.206) and plant height (28.325 cm). The lowest genetic advance was 3.406 estimated for number of rows/cob (Table 4.1.10.).

**Genetic advance in percentage of mean (GA %)**

The estimation of genetic advance as percentage of mean reveals (Table 4.1.10.) the highest value 55.727 for days to silking and this was followed by number of kernels/row (40.189), days to tasseling (38.599), cob diameter (37.376 cm) and cob length (33.864 cm). The lowest value of GA% was 19.206 found for days to maturity.

**Table 4.1.10. Estimation of genetic parameters for grain yields and yield components in maize inbred**

| Characters | $\sigma^2_g$ | $\sigma^2_e$ | $\sigma^2_p$        | PCV%    | ECV%    | GCV%    | $h^2_b\%$ | GA     | GA(%)<br>of<br>mean |
|------------|--------------|--------------|---------------------|---------|---------|---------|-----------|--------|---------------------|
| DT         | 233.642      | 37.932       | 271.574             | 21.779  | 8.139   | 20.201  | 86.032    | 29.206 | 38.599              |
| DS         | 356.452      | 25.162       | 381.615             | 28.961  | 7.436   | 27.99   | 93.406    | 37.588 | 55.727              |
| DM         | 74.051       | 30.078       | 104.13              | 9.738   | 5.233   | 8.212   | 88.638    | 23.656 | 19.206              |
| PH         | 230.965      | 51.187       | 282.153             | 16.148  | 6.878   | 14.61   | 81.8582   | 28.325 | 27.231              |
| EH         | 180.447      | 52.918       | 233.366             | 32.055  | 15.105  | 28.273  | 77.795    | 24.491 | 51.371              |
| CL         | 3.123        | 2.294        | 5.418               | 20.03   | 8.482   | 18.146  | 82.068    | 4.763  | 33.864              |
| CD         | 7.137        | 2.554        | 9.691               | 24.636  | 12.647  | 21.142  | 73.6465   | 4.723  | 37.376              |
| NRC        | 4.692        | 2.313        | 7.006               | 21.94   | 11.659  | 17.956  | 72.152    | 3.406  | 26.059              |
| NKR        | 20.851       | 6.164        | 27.015              | 19.343  | 17.137  | 14.524  | 77.183    | 8.264  | 40.189              |
| NGC        | 808.787      | 415.124      | $\frac{1223.91}{1}$ | 788.765 | 267.532 | 521.233 | 66.082    | 47.624 | 30.691              |
| GYP        | 112.393      | 42.74        | 155.134             | 21.477  | 73.702  | 18.281  | 72.4492   | 18.588 | 32.054              |

#### 4.1.2 CORRELATION COEFFICIENT (r)

Correlation coefficients between grain yield and yield attributing characters and correlation coefficients between grain yield attributing characters of 25 maize inbred lines at the genotypic and phenotypic levels were calculated and the results are shown in Tables 4.1.12- 4.1.13.

##### **Genotypic Correlation Coefficient Between Grain Yield with its Components**

Genotypic correlation coefficient between grain yield and yield contributing characters is presented in Table 4.1.11. At the genotypic level, grain yield showed highly significant and positive correlation with days to maturity (0.5817\*), number of grains/cob (0.6534\*\*) and number of kernels/row

(0.5912\*\*). Cob length showed non significant but negative correlation with grain yield at the genotypic level.

### **Phenotypic Correlation Coefficient Between Grain Yield with its Components**

At the phenotypic level, the grain yield showed significant and positive correlation with days to maturity (0.627\*\*), cob diameter (0.655\*\*), number of kernels/row (0.3458\*\*) and number of grain/cob (0.9372\*\*) (Table 4.1.11).

**Table 4.1.11. Genotypic and phenotypic correlation coefficients between grain yield/plant and yield contributing characters in maize**

| Characters         | Correlation coefficient |            |
|--------------------|-------------------------|------------|
|                    | Genotypic               | Phenotypic |
| Days to maturity   | 0.5817*                 | 0.627**    |
| Plant height       | 0.208                   | 0.155      |
| Ear height         | 0.138                   | 0.141      |
| Cob length         | -0.023                  | -0.031     |
| Cob diameter       | 0.153                   | 0.655**    |
| No. of rows/cob    | 0.247                   | 0.220      |
| No. of kernels/row | 0.591*                  | 0.346*     |
| No. of grains/cob  | 0.653**                 | 0.937***   |

### **Genotypic Correlation Coefficient Between Different Pairs of Characters**

Genotypic correlation co-efficient among the different pairs of characters were estimated and the results are presented in Table 4.1.12. Among the different pairs of characters, days to maturity showed highly significant positive correlation with cob diameter, number of kernels/row and grain yield/plant.

On the other hand, genotypic correlation of days to maturity with plant height, ear height, cob length, number of rows/cob and number of grains /cob were non-significant.

Plant height exhibited highly significant and positive correlation with ear height and number of kernels/row. But this character showed non-significant correlation with cob length, cob diameter, number of rows/cob, number of grain/cob and grain yield, respectively. Number of kernels/row showed significant but positive correlation with days to maturity, plant height and cob diameter. On the other hand, this character was non-significant and negative correlation with rest of the characters.

Number of grain/cob showed significant positive correlation with number of row/cob, number of rows/cob and this character showed significant negative correlation with plant height. Genotypic correlation coefficient of days to maturity and number of kernels/row and number grains/cob with grain yield also was positive significant.

**Table 4.1.12. Genotypic correlation coefficient ( $r_g$ ) between different pairs of characters in maize**

| Characters | PH    | EH      | CL      | CD       | NRC      | NKR     | NGC      | GYP     |
|------------|-------|---------|---------|----------|----------|---------|----------|---------|
| DM         | 0.229 | 0.407   | -0.442* | 0.5412*  | 0.019    | 0.524*  | -0.196   | 0.582*  |
| PH         | 1.000 | 0.815** | -0.355  | 0.114    | 0.879**  | 0.549*  | -0.902** | 0.208   |
| EH         |       | 1.000   | 0.618** | -0.634** | -0.902** | 0.165   | -0.187   | 0.138   |
| CL         |       |         | 1.000   | -0.035   | 0.016    | -0.048  | 0.097    | -0.023  |
| CD         |       |         |         | 1.000    | 0.034    | 0.918** | 0.073    | 0.154   |
| NRC        |       |         |         |          | 1.000    | 0.089   | 0.718**  | 0.247   |
| NKR        |       |         |         |          |          | 1.000   | 0.512*   | 0.591*  |
| NGC        |       |         |         |          |          |         | 1.000    | 0.653** |



### **Phenotypic Correlation Coefficient Between Different Pairs of Characters**

Phenotypic correlation co-efficient among different pairs of characters are presented in table 4.1.13.

Among the different inter character associations, days to maturity showed significant positive correlation with cob length, number of rows/cob, number of grains/cob and grain yield.

The associations of days to maturity with other characters were non-significant. Association of plant height with cob length, cob diameter and number of grains/cob were highly significant.

The character, number of kernels/row showed significant positive correlation with cob length and number of rows/cob. Number of rows/cob exhibited non-significant correlation with plant height, ear height, and cob length and cob diameter.

Correlation coefficient of number of grains/cob with number of rows/cob and number of kernels/row were also non-significant. On the other hand, this trait showed non-significant and negative association with ear height and cob length.

The character, grain yield/plant exhibited non-significant and positive correlation with plant height, ear height, number of rows/cob and number of kernels/row.

Days to maturity, cob diameter and number of kernels/cob showed highly significant and positive correlation with grain yield/plant.

**Table 4.1.13. Phenotypic correlation coefficient ( $r_p$ ) between different pairs of characters in maize**

| Characters | PH    | EH    | CL     | CD      | NRC     | NKR      | NGC     | GYP     |
|------------|-------|-------|--------|---------|---------|----------|---------|---------|
| DM         | 0.198 | 0.063 | 0.564* | 0.1799  | 0.639** | 0.145    | 0.6142* | 0.627** |
| PH         | 1.000 | 0.233 | 0.581* | 0.907** | -0.001  | 0.1734   | -0.556* | 0.155   |
| EH         |       | 1.000 | 0.193  | 0.1386  | -0.059  | 0.108    | -0.194  | 0.142   |
| CL         |       |       | 1.000  | 0.0371  | -0.022  | -0.652** | -0.040  | -0.031  |
| CD         |       |       |        | 1.0000  | -0.005  | 0.451    | 0.872** | 0.655** |
| NRC        |       |       |        |         | 1.000   | 0.782**  | 0.429   | 0.219   |
| NKR        |       |       |        |         |         | 1.000    | 0.325   | 0.346   |
| NGC        |       |       |        |         |         |          | 1.000   | 0.937** |

### 4.1.3 PATH COEFFICIENT ANALYSIS

The correlation coefficients between grain yield and yield components were partitioned into direct and indirect effects through path coefficient analysis in order to find out more realistic picture of relationship. Path coefficient analysis was performed using the values of genotypic and phenotypic correlation and are presented in Tables 4.1.14 and 4.1.15, Figures 4.1.1 and 4.1.2.

The results of path coefficient analysis at genotypic and phenotypic levels are described below.

#### Path Coefficient at Genotypic Level

The results of path coefficient analysis at genotypic level are presented in Table 4.1.14. This table shows that the highest positive direct effect was contributed by number of rows/cob on grain yield and it was followed by number of kernels/row and cob length. Days to maturity, plant height, cob height, cob diameter, number of kernels/cob showed negative direct effect. Number of rows/cob had positive direct effect (0.1814) on grain yield.

However, number of kernels/row, days to maturity, number of grains/cob, number of rows/cob and cob diameter contributed to grain yield through large indirect effects of number of grains/cob (0.6723), number of kernels/row (0.3812), number of grains/cob (0.218) and number of kernels/row (0.1872), respectively. The total effect of cob length on grain yield was 0.4681.

Plant height expressed negative indirect effect on grain yield through number of grains/cob. The character days to maturity showed positive direct effect on grain yield (0.0396). This character showed positive indirect effect on grain yield through cob height, cob length, cob diameter, number of rows/plant and number of grains/cob. On the other hand, plant height had negative indirect effect on grain yield through days to maturity (-0.0829), cob length (-0.0614), cob diameter (-0.0177) and number of grains/cob (-0.1822). The total effect of this character on grain yield was 0.6463.

Ear height had direct negative effect on grain yield (-0.0128). Cob height also contributed to grain yield through positive indirect effect of days to maturity, plant height, cob length, number of kernels/row and number of grains/cob. The indirect effect of cob height through cob diameter and number of rows/cob were negative. The total effect of cob height was 2.3639.

Cob length had positive direct effect (0.0156) on grain yield. It had positive indirect effect through days to maturity, plant height, number of kernels/row and number of grains/cob. The indirect effect of cob length through ear height, cob diameter and number of rows/cob were negative. The total effect of cob length on grain yield was 0.0117.

The character cob diameter showed the negative direct effect (-0.0038) on grain yield. It contributed to grain yield greatly indirect effect through cob length (0.1066) followed by number of kernels/row (0.0753).

The indirect effect of cob diameter through days to maturity, cob length, number of rows/cob and number of kernels/row were positive.

But it showed negative indirect effect via plant height, cob height and number of grains/cob. The total effect of cob diameter was 0.0667.

Number rows/cob showed positive direct effect (0.1814) on grain yield. It showed positive indirect effect through days to maturity, plant height, cob length, cob diameter, and number of grains/cob. But this character exhibited the negative indirect influence on grain yield through cob height and number of kernels/row. The total effect of this trait was 0.3741.

Number of kernels/row had the highest positive direct effect (0.0307) on grain yield. Number of kernels/row had the highest positive indirect effect (0.3923) on grain yield through number of grains/cob followed by days to maturity (0.3812). The total effect of number of kernels/row on grain yield was 0.9364.

Number of grains/cob had the highest negative direct effect (-0.2704) on grain yield. This character had the highest positive indirect effect (0.6723) on grain yield through number of kernels/row followed by number of rows/cob (0.2156). The total effect of this trait was -5.9778.

Table 4.1.14 also shows that the highest total genotypic effect was observed for number of kernels/row (0.9364) and this was followed by number of kernels/cob (0.3741), cob diameter (0.0667). But number of grains/cob exhibited smaller effect of -5.9778.

The considerable amount of residual effect (0.8517) indicated that some other characters which have been included in this study had also effect on grain yield in this crop.

It may be concluded from the present study that number of rows/cob, days to maturity and number of kernels/row are the major components of grain yield in maize inbred lines and hence maximum stress should be given on these characters while selection is done for maximum grain yield.

**Table 4.1.14. Path coefficient analysis showing direct and indirect effects of yield components on grain yield/plant of maize at genotypic level**

| Characters   | Grain yield (kernel weight/plant) vs. |                |                |               |                |               |               |                | r <sub>g</sub> with grain yield |
|--------------|---------------------------------------|----------------|----------------|---------------|----------------|---------------|---------------|----------------|---------------------------------|
|              | DM                                    | PH             | EH             | CL            | CD             | NRC           | NKR           | NGC            |                                 |
| DM           | <u>0.0396</u>                         | -0.0829        | 0.0584         | 0.0017        | 0.0643         | 0.0076        | 0.3812        | 0.0544         | 0.5817*                         |
| PH           | -0.0203                               | <u>-0.0372</u> | 0.0340         | 0.0066        | -0.0130        | 0.1575        | 0.1946        | -0.5736        | 0.2076                          |
| EH           | 0.0214                                | 0.0510         | <u>-0.0128</u> | -0.0070       | -0.1120        | -0.0510       | 0.1918        | 0.1308         | 0.1383                          |
| CL           | -0.0038                               | -0.0614        | 0.0433         | <u>0.0156</u> | 0.1066         | 0.0750        | -0.1624       | -0.2229        | -0.0223                         |
| CD           | 0.0213                                | -0.0177        | -0.1011        | -0.0155       | <u>-0.0038</u> | 0.1029        | 0.1872        | -5.9840        | 0.1533                          |
| NRC          | 0.0016                                | 0.1323         | -0.0286        | -0.0068       | 0.0639         | <u>0.1814</u> | -0.2790       | 0.2156         | 0.2471                          |
| NKR          | -0.0254                               | 0.1059         | 0.0696         | 0.0095        | 0.0753         | -0.1808       | <u>0.0307</u> | 0.6723         | 0.5912*                         |
| NGC          | 0.0042                                | -0.1822        | 0.0277         | 0.0076        | -0.1146        | 0.0815        | 0.3923        | <u>-0.2704</u> | 0.653**                         |
| Total Effect | 0.0386                                | -0.0922        | 0.0905         | 0.0117        | 0.0667         | 0.3741        | 0.9364        | -5.9778        |                                 |

Residual effect = 0.8517, underlined values denote direct effect.

### Path Coefficient at Phenotypic Level

The results of path coefficient analysis at phenotypic level are presented in table 4.1.15. Number of kernels/cob had the highest positive direct effect (2.7688) on grain yield followed by number of kernels/row (0.6409), plant height (0.3518) and number of rows/cob (0.3213).

The highest direct negative effect towards grain yield was found for cob length (-0.0362).

The path analysis revealed that the most of the characters had positive direct effect on grain yield and cob length had negative direct effect on grain yield.

Days to maturity had positive direct effect of 0.0744 on grain weight/plant.

Days to maturity had indirect positive effect (0.222) via cob length followed by

number of rows/cob and number of kernels/row. The indirect effect via plant height, ear height, cob diameter and number of grains/cob was negative.

Plant height had direct positive effect on grain yield. Plant height exhibited the highest indirect positive effect of 0.2851 through number of grains/cob. The indirect effects of plant height on grain yield via days to maturity, ear height, cob length, number of rows/cob and number of kernels/row were negative. The total effect was 0.3042.

The character ear height showed positive direct effect (0.1603) on grain yield. On the other hand, it had indirect positive effect through number of grains/cob. The indirect effects of ear height through most of the characters were negative. The total effect was 0.104.

Cob length indicated the negative direct effect (-0.0362) on grain yield. Cob length showed highest indirect positive effect (0.0062) on grain yield via number of grains/cob. The indirect effects through days to maturity, cob diameter, number of rows/cob, number of kernels/row were negative.

Cob diameter showed the positive direct effect on grain yield. This character showed the highest positive indirect effect (0.0091) through plant height.

However, cob diameter showed negative indirect effects (-0.0130, -0.0056, -0.0085, -0.0004, and -0.0415) through days to maturity, ear height, cob length, number of rows/cob, number of kernels/row but the value was least and negligible.

Number of rows/cob had positive direct effect of 0.03213 on grain yield. The indirect effects of this character on grain yield via plant height, ear height, cob diameter and number of grains/cob were negative. The total effect was 0.013.

Number of kernels/row showed positive direct effect (0.6409) on grain yield. On the other hand, it had indirect positive effect through days to maturity, cob length, and number of rows/cob. The indirect effects of number of kernels/row through most of the characters were negative. The total effect was -0.1732.

The character ear height showed positive direct effect (0.1603) on grain yield. On the other hand, it had indirect positive effect through number of grains/cob. The indirect effects of ear height through most of the characters were negative. The total effect was 0.104.

Number of grains/cob had highest positive direct effect of 2.7688 on grain yield. The indirect effects of this character on grain yield via days to maturity, cob length, number of rows/cob and number of kernels/row were negative. The total effect was 2.2317 (the highest value).

The highest direct positive effect towards grain yield was found for number of grains/cob (2.7688) followed by number of kernels/row (0.6409) and plant height (0.3518). The lowest direct negative effect  $-0.0362$  towards grain yield was found for cob length. The residual effect at phenotypic level was 0.9451.

The path coefficient analysis at phenotypic level in the present study reveals that number of grains/cob had highest total effect (2.2317) on grain yield, which was followed by plant height, ear height and days to maturity with the values of 0.3042, 0.104 and 0.0525, respectively.

**Table 4.1.15. Path coefficient analysis showing direct and indirect effects of yield components on grain yield/plant of maize at phenotypic level**

| Characters   | Grain yield (grain weight/plant) vs. |               |               |                |               |               |               |               | $r_p$ with grain yield |
|--------------|--------------------------------------|---------------|---------------|----------------|---------------|---------------|---------------|---------------|------------------------|
|              | DM                                   | PH            | EH            | CL             | CD            | NRC           | NKR           | NGC           |                        |
| DM           | <u>0.0744</u>                        | -0.0520       | -0.0110       | -0.0110        | -0.0130       | 0.0068        | 0.0231        | -0.1768       | 0.0627                 |
| PH           | -0.0210                              | <u>0.3518</u> | -0.0095       | 0.0067         | 0.0091        | -0.1558       | -0.3311       | 1.7255        | 0.1549                 |
| EH           | -0.0049                              | -0.0105       | <u>0.1603</u> | 0.0073         | -0.0053       | -0.0059       | -0.0538       | 0.2272        | 0.1407                 |
| CL           | 0.0222                               | -0.0336       | -0.0329       | <u>-0.0362</u> | -0.0085       | 0.0145        | 0.0709        | -0.1855       | -0.0311                |
| CD           | -0.0125                              | 0.0215        | -0.0113       | -0.0085        | <u>0.0859</u> | -0.0004       | -0.2192       | 0.1386        | 0.0655                 |
| NRC          | 0.0019                               | -0.1098       | -0.0038       | -0.0020        | -0.0004       | <u>0.3213</u> | 0.1342        | -1.0784       | 0.2197                 |
| NKR          | 0.0042                               | -0.1483       | -0.0219       | -0.0064        | -0.0415       | 0.0853        | <u>0.6409</u> | -1.1877       | 0.3458                 |
| NGC          | -0.0118                              | 0.2851        | 0.0341        | 0.0062         | 0.0097        | -0.2528       | -0.4382       | <u>2.7688</u> | 0.9372                 |
| Total Effect | 0.0525                               | 0.3042        | 0.104         | -0.0439        | 0.036         | 0.013         | -0.1732       | 2.2317        |                        |

Residual effect = 0.9451 Underlined values denote the direct effect.



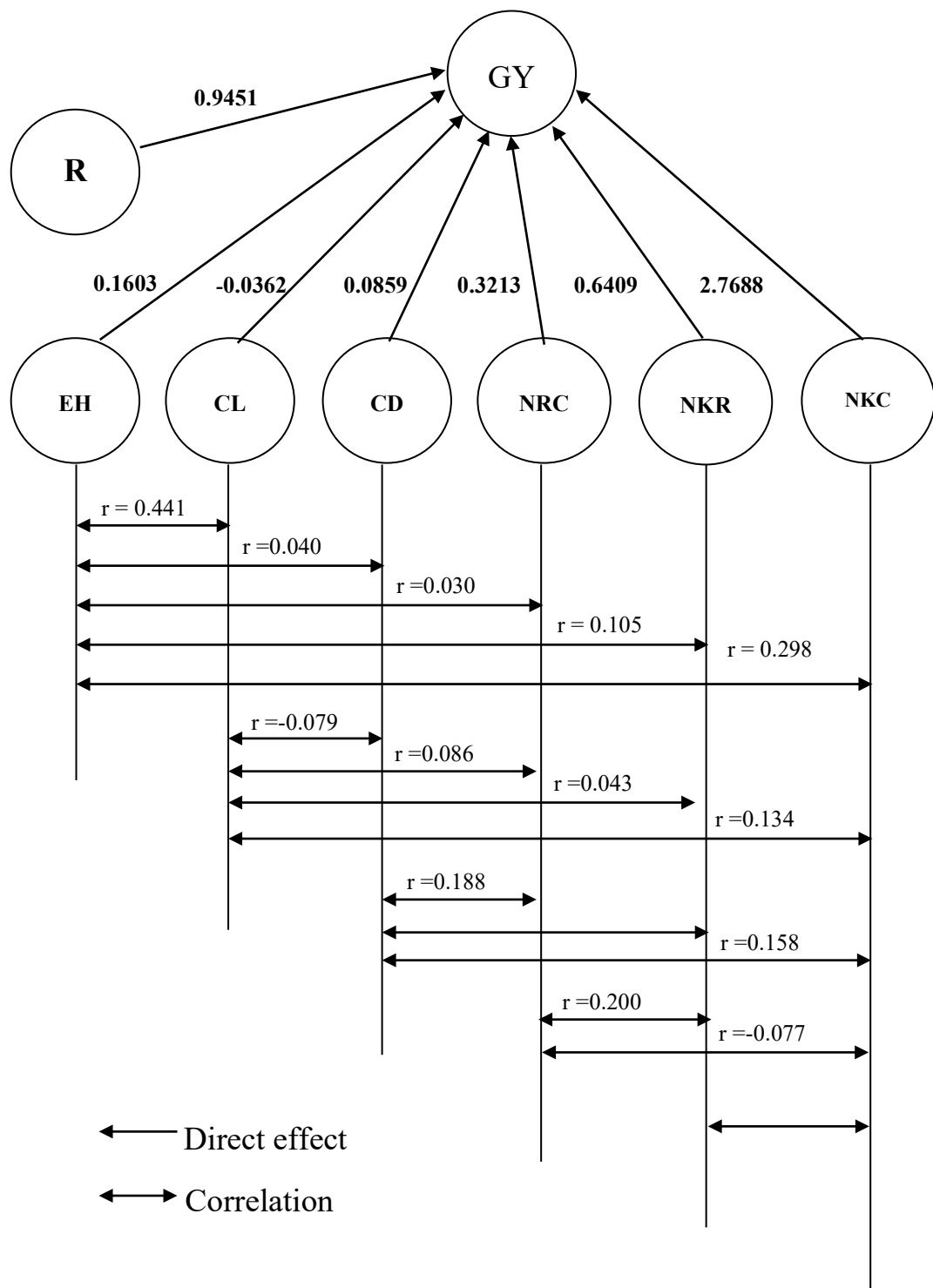


Figure 4.1.1. Path diagram of different grain yield contributing characters on yield at phenotypic level

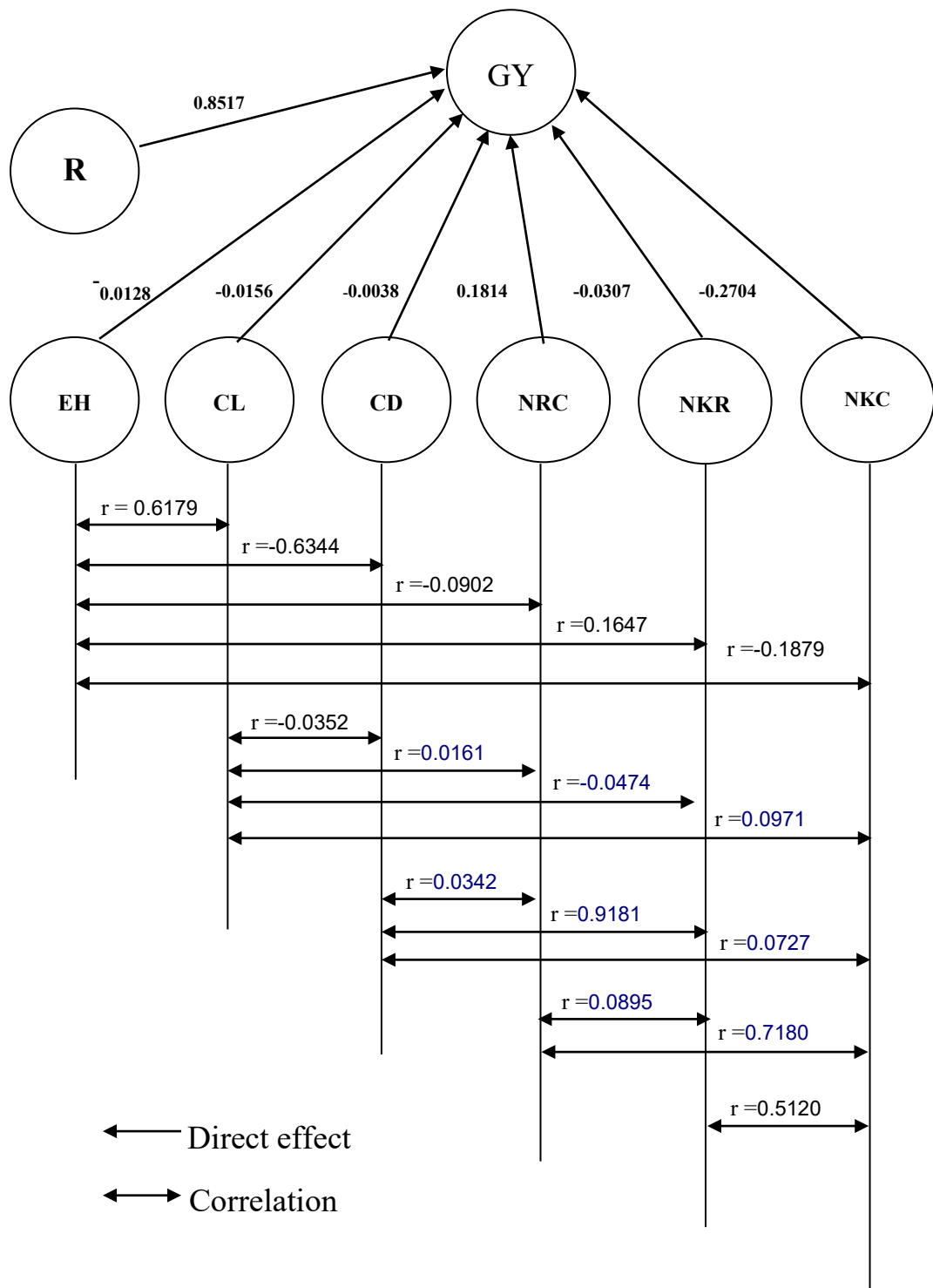


Figure 4.1.2. Path diagram of different grain yield contributing characters on yield at genotypic level

#### 4.1.4 SELECTION INDEX

Selection indices for grain yield were constructed to identify the character or character association over straight selection, which may be useful during selection breeding programme for higher yield. In constructing the selection indices, all the nine quantitative characters *viz*, DM, PH, EH, CL, CD, NRC, NKR, NKC and GYP were included of which GYP was dependent character.

The results obtained for different indices, containing GYP and its components with expected gain in percentage over straight selection are shown in Table 4.1.16. This table showed that the maximum genetic (expected) gain of 192.534% was exhibited when cob height, number of rows/cob, number of kernels/row and grain yield/plant were included in the discriminant function. This value was followed by 158.199% GA which was obtained when cob length, cob diameter, number of row/cob, number of kernels/row and number of grains/cob were included in the discriminant function. In the present investigation, discriminant function analyses have been done considering individual character separately and are shown in Table 4.1.16.

Table 4.1.16 exhibited that individual character expect days to maturity, plant height, cob height and cob diameter showed positive expected gain and among them number of kernels/row exhibited highest genetic gain, followed by number of grain/cob, number of rows/cob and number of kernels/row. Grain yield/plant in combination with two or more characters gave the highest positive expected gain but in the remaining cases it showed negative genetic gain. Number of rows/cob in combination with number of kernels/row (NRC+NKR) gave the highest positive gain of 186.457% followed by 175.429 % ( NKR+GYP), 153.913 % ( DM+NKR), 122.758 % (DM+CL), 105.593 % (NKR+NGC) and 83.458 (DM+NGC).

In the discriminant function analyses when selection index included three characters, the maximum genetic gain was recorded as 168.975% for

CL+NRC+NKR followed by 159.806% for CL+CD+NKR, 147.618% for DM+CL+NKR.

In the same way, when four characters were included in the discriminant function, GYP in combination with CD, NRC and NKR gave the highest GA% of 162.534% and next were 142.482% and 139.817% and 137.338 by DM+CL+CD+NKR+NGC, DM+PH+CL+NRC+GYP and DM+CL+NRC+NKR, respectively. Similarly, when five characters were included in the discriminant function, GYP in combination with DM, PH, CL and NRC exhibited the highest genetic gain of 139.817%, which was the highest value among all the selection indices followed by 135.467%(CL+CD+NRC+NKR+GYP), 132.804% (DM+PH+CL+NKR+GYP), 131.706 % (DM+PH+CH+CL+GYP) and 125.705 % (DM+CL+NKR+NGP+GYP).

In case of discriminant function when six characters were included, GYP combination with DM, CL, CD, NRC and NRK exhibited the highest genetic gain of 129.328%, followed by 128.498% (EH+CL+CD+NRC+NKR+GYP) and 125.359% (DM+PH+CL+CD+NKR+GYP).

Table 4.1.16 showed the highest value of GA% of 125.0154% when seven characters, such as DM+PH+CL+CD+NRC+NKR+GYP were in combination.

This combination was followed by PH+CL+CD+NRC+NKR+NGC+GYP, DM+PH+EH+CL+NRC+NKR+GYP and DM+PH+EH+CD+NRC+NGC+GYP with the values of 124.113%, 122.015% and 119.584%, respectively.

Similarly, when eight characters were included in the selection index, GYP in combination with DM+PH+EH+CD+NRC+NKR+NGC showed the highest genetic gain of 121.582% and the next were 119.459% and 119.263% for (DM+PH+EH+CL+NRC+NKR+NGC) and (DM+PH+EH+CL+CD+NRC+NKR).

When all the nine quantitative characters were considered the expected genetic gain was found to be 158.723% and this was the second highest in the present discriminant function analysis.

**Table 4.1.16. Expected gain in percentage of grain yield over selection from the use of various selection indices in maize inbred lines**

| Selection index | Expected gain | Selection index | Expected gain | Selection index | Expected gain |
|-----------------|---------------|-----------------|---------------|-----------------|---------------|
| X1              | -129.809      | X5+X6           | 3.4302        | X1+X7+X8        | 107.6842      |
| X2              | -52.9325      | X5+X7           | -173.994      | X1+X7+X9        | 52.7325       |
| X3              | -61.6707      | X5+X8           | 74.989        | X1+X8+X9        | 68.486        |
| X4              | 31.29928      | X5+X9           | 4.3224        | X2+X3+X4        | 70.442        |
| X5              | -41.1064      | X6+X7           | 186.457       | X2+X3+X5        | 69.356        |
| X6              | 51.52128      | X6+X8           | -75.516       | X2+X3+X6        | -69.478       |
| X7              | 241.5108      | X6+X9           | 8.0630        | X2+X3+X7        | 81.4340       |
| X8              | 76.337        | X7+X8           | 105.593       | X2+X3+X8        | 80.2636       |
| X9              | 25.820        | X7+X9           | 175.429       | X2+X3+X9        | 60.0067       |
| X1+X2           | 70.963        | X8+X9           | -60.790       | X2+X4+X5        | -58.044       |
| X1+X3           | -79.148       | X1+X2+X3        | 76.8540       | X2+X4+X6        | -58.092       |
| X1+X4           | 122.758       | X1+X2+X4        | 72.3973       | X2+X4+X7        | -83.033       |
| X1+X5           | 74.8090       | X1+X2+X5        | 70.3667       | X2+X4+X8        | 77.6917       |
| X1+X6           | -82.3619      | X1+X2+X6        | -70.742       | X2+X4+X9        | 45.6736       |
| X1+X7           | 153.9128      | X1+X2+X7        | -91.083       | X2+X5+X6        | 56.6788       |
| X1+X8           | 83.4584       | X1+X2+X8        | 81.4317       | X2+X5+X7        | 73.5281       |
| X1+X9           | -31.539       | X1+X2+X9        | 57.0823       | X2+X5+X8        | 76.837        |
| X2+X3           | 69.1001       | X1+X3+X4        | -80.002       | X2+X5+X9        | 46.290        |
| X2+X4           | -56.6846      | X1+X3+X5        | -76.0016      | X2+X6+X7        | -78.972       |
| X2+X5           | 54.8994       | X1+X3+X6        | 76.9749       | X2+X6+X8        | 94.8053       |
| X2+X6           | 54.859        | X1+X3+X7        | 117.563       | X2+X6+X9        | 45.6466       |
| X2+X7           | 82.504        | X1+X3+X8        | -83.969       | X2+X7+X8        | 84.6101       |
| X2+X8           | -77.015       | X1+X3+X9        | -57.171       | X2+X7+X9        | -56.053       |
| X2+X9           | 42.6151       | X1+X4+X5        | 76.3463       | X2+X8+X9        | -69.128       |
| X3+X4           | 64.8754       | X1+X4+X6        | 82.2569       | X3+X4+X5        | -63.482       |
| X3+X5           | -60.545       | X1+X4+X7        | 147.618       | X3+X4+X6        | -64.194       |
| X3+X6           | -61.246       | X1+X4+X8        | 83.8938       | X3+X4+X7        | 119.235       |
| X3+X7           | -121.665      | X1+X4+X9        | 36.5366       | X3+X4+X8        | 80.2595       |
| X3+X8           | 79.6717       | X1+X5+X6        | 68.1518       | X3+X4+X9        | 44.7202       |
| X3+X9           | -41.325       | X1+X5+X7        | 137.917       | X3+X5+X6        | 61.0801       |
| X4+X5           | 7.4460        | X1+X5+X8        | 81.7069       | X3+X5+X7        | 106.245       |
| X4+X6           | 8.8190        | X1+X5+X9        | -37.5137      | X3+X5+X8        | -79.036       |
| X4+X7           | -204.633      | X1+X6+X7        | -141.782      | X3+X5+X9        | -45.323       |
| X4+X8           | -77.3202      | X1+X6+X8        | -82.2772      | X3+X6+X7        | 112.098       |
| X4+X9           | 9.8926        | X1+X6+X9        | -36.4947      | X3+X6+X8        | 79.3153       |

X1=DM, X2=PH, X3=EH, X4= CL, X5= CD,X6= NRC, X7= NKR, X8= NKC and X9= GYP

**Table 4.1.16. (Continued)**

| Selection index | Expected gain | Selection index | Expected gain | Selection index | Expected gain |
|-----------------|---------------|-----------------|---------------|-----------------|---------------|
| X3+X6+X9        | -44.633       | X1+X2+X5+X6     | -70.451       | X1+X5+X6+X7     | 128.593       |
| X3+X7+X8        | 89.049        | X1+X2+X5+X7     | 85.6111       | X1+X5+X6+X8     | -80.976       |
| X3+X7+X9        | 56.429        | X1+X2+X5+X8     | 81.0226       | X1+X5+X6+X9     | -41.450       |
| X3+X8+X9        | 69.531        | X1+X2+X5+X9     | -58.889       | X1+X5+X7+X8     | 4.91097       |
| X4+X5+X6        | 20.389        | X1+X2+X6+X7     | 86.9667       | X1+X5+X7+X9     | -55.180       |
| X4+X5+X7        | 159.805       | X1+X2+X6+X8     | 81.2140       | X1+X5+X8+X9     | -69.225       |
| X4+X5+X8        | -75.995       | X1+X2+X6+X9     | -58.603       | X1+X6+X7+X8     | 98.6625       |
| X4+X5+X9        | -6.4526       | X1+X2+X7+X8     | 87.7271       | X1+X6+X7+X9     | -54.857       |
| X4+X6+X7        | 168.975       | X1+X2+X7+X9     | -66.010       | X1+X6+X8+X9     | -69.146       |
| X4+X6+X8        | 76.5041       | X1+X2+X8+X9     | 73.6565       | X1+X7+X8+X9     | -74.563       |
| X4+X6+X9        | -3.583        | X1+X3+X4+X5     | -77.058       | X2+X3+X4+X5     | -70.595       |
| X4+X7+X8        | 104.233       | X1+X3+X4+X6     | 77.9722       | X2+X3+X4+X6     | -70.729       |
| X4+X7+X9        | -31.310       | X1+X3+X4+X7     | -116.167      | X2+X3+X4+X7     | 81.991        |
| X4+X8+X9        | -62.261       | X1+X3+X4+X8     | 84.3197       | X2+X3+X4+X8     | -80.704       |
| X5+X6+X7        | 147.216       | X1+X3+X4+X9     | -59.004       | X2+X3+X4+X9     | -61.415       |
| X5+X6+X8        | 74.6180       | X1+X3+X5+X6     | 74.9819       | X2+X3+X5+X6     | -69.797       |
| X5+X6+X9        | 7.50199       | X1+X3+X5+X7     | 109.622       | X2+X3+X5+X7     | 80.0006       |
| X5+X7+X8        | 90.2890       | X1+X3+X5+X8     | -83.145       | X2+X3+X5+X8     | -80.111       |
| X5+X7+X9        | 32.4701       | X1+X3+X5+X9     | -58.997       | X2+X3+X5+X9     | -61.480       |
| X5+X8+X9        | -62.214       | X1+X3+X6+X7     | 112.212       | X2+X3+X6+X7     | 80.4996       |
| X6+X7+X8        | 92.3308       | X1+X3+X6+X8     | -83.446       | X2+X3+X6+X8     | 80.2267       |
| X6+X7+X9        | -31.167       | X1+X3+X6+X9     | -58.722       | X2+X3+X6+X9     | -61.246       |
| X6+X8+X9        | 62.0051       | X1+X3+X7+X8     | 92.0462       | X2+X3+X7+X8     | 85.290        |
| X7+X8+X9        | 68.8779       | X1+X3+X7+X9     | -66.960       | X2+X3+X7+X9     | -67.322       |
| X1+X2+X3+X4     | -77.600       | X1+X3+X8+X9     | 74.1928       | X2+X3+X8+X9     | -74.223       |
| X1+X2+X3+X5     | 76.4847       | X1+X4+X5+X6     | -70.610       | X2+X4+X5+X6     | -59.426       |
| X1+X2+X3+X6     | 76.7074       | X1+X4+X5+X7     | 133.944       | X2+X4+X5+X7     | -78.596       |
| X1+X2+X3+X7     | 86.2987       | X1+X4+X5+X8     | -82.220       | X2+X4+X5+X8     | 77.4897       |
| X1+X2+X3+X8     | 83.2653       | X1+X4+X5+X9     | -41.540       | X2+X4+X5+X9     | -48.899       |
| X1+X2+X3+X9     | -67.350       | X1+X4+X6+X7     | 137.338       | X2+X4+X6+X7     | 79.8349       |
| X1+X2+X4+X5     | 71.7317       | X1+X4+X6+X8     | -82.766       | X2+X4+X6+X8     | -77.647       |
| X1+X2+X4+X6     | 72.1065       | X1+X4+X6+X9     | -40.697       | X2+X4+X6+X9     | -48.341       |
| X1+X2+X4+X7     | 90.8431       | X1+X4+X7+X8     | 106.930       | X2+X4+X7+X8     | 84.9401       |
| X1+X2+X4+X8     | 81.8660       | X1+X4+X7+X9     | -55.460       | X2+X4+X7+X9     | -58.021       |
| X1+X2+X4+X9     | -58.813       | X1+X4+X8+X9     | 69.4410       | X2+X4+X8+X9     | 69.9424       |

**Table 4.1.16. (Continued)**

| Selection index | Expected gain | Selection index | Expected gain | Selection index | Expected gain |
|-----------------|---------------|-----------------|---------------|-----------------|---------------|
| X2+X5+X6+X7     | 76.0454       | X4+X5+X8+X9     | -63.540       | X1+X2+X5+X6+X7  | 80.889        |
| X2+X5+X6+X8     | 76.889        | X4+X6+X7+X8     | 92.0625       | X1+X2+X5+X6+X8  | 139.76        |
| X2+X5+X6+X9     | -48.844       | X4+X6+X7+X9     | -36.629       | X1+X2+X5+X6+X9  | 86.643        |
| X2+X5+X7+X8     | 83.6136       | X4+X6+X8+X9     | -63.357       | X1+X2+X5+X7+X8  | 133.068       |
| X2+X5+X7+X9     | -58.017       | X4+X7+X8+X9     | -69.891       | X1+X2+X5+X7+X9  | 125.919       |
| X2+X5+X8+X9     | -69.840       | X5+X6+X7+X8     | 87.428        | X1+X2+X5+X8+X9  | 86.996        |
| X2+X6+X7+X8     | 83.9615       | X5+X6+X7+X9     | 192.534       | X1+X2+X6+X7+X8  | 133.166       |
| X2+X6+X7+X9     | -57.722       | X5+X6+X8+X9     | 136.693       | X1+X2+X6+X7+X9  | 125.958       |
| X2+X6+X8+X9     | -69.747       | X5+X7+X8+X9     | 130.451       | X1+X2+X6+X8+X9  | 122.390       |
| X2+X7+X8+X9     | 73.9894       | X6+X7+X8+X9     | 130.501       | X1+X2+X7+X8+X9  | 76.0556       |
| X3+X4+X5+X6     | -63.716       | X1+X2+X3+X4+X5  | 77.2097       | X1+X3+X4+X5+X6  | 108.462       |
| X3+X4+X5+X7     | -103.379      | X1+X2+X3+X4+X6  | 77.431        | X1+X3+X4+X5+X7  | 83.5154       |
| X3+X4+X5+X8     | 79.6256       | X1+X2+X3+X4+X7  | 86.561        | X1+X3+X4+X5+X8  | 139.375       |
| X3+X4+X5+X9     | -48.184       | X1+X2+X3+X4+X8  | 83.578        | X1+X3+X4+X5+X9  | 111.039       |
| X3+X4+X6+X7     | 110.132       | X1+X2+X3+X4+X9  | 131.706       | X1+X3+X4+X6+X7  | 83.808        |
| X3+X4+X6+X8     | -79.898       | X1+X2+X3+X5+X6  | 76.459        | X1+X3+X4+X6+X8  | 139.612       |
| X3+X4+X6+X9     | -47.598       | X1+X2+X3+X5+X7  | 84.678        | X1+X3+X4+X6+X9  | 92.031        |
| X3+X4+X7+X8     | 89.1700       | X1+X2+X3+X5+X8  | 82.991        | X1+X3+X4+X7+X8  | 131.833       |
| X3+X4+X7+X9     | -58.512       | X1+X2+X3+X5+X9  | 131.809       | X1+X3+X4+X7+X9  | -125.209      |
| X3+X4+X8+X9     | 70.3641       | X1+X2+X3+X6+X7  | 85.190        | X1+X3+X4+X8+X9  | 95.869        |
| X3+X5+X6+X7     | 87.7470       | X1+X2+X3+X6+X8  | 83.124        | X1+X3+X5+X6+X7  | -82.781       |
| X3+X5+X6+X8     | -78.837       | X1+X2+X3+X6+X9  | 131.922       | X1+X3+X5+X6+X8  | 139.632       |
| X3+X5+X6+X9     | -48.083       | X1+X2+X3+X7+X8  | 87.585        | X1+X3+X5+X6+X9  | 89.903        |
| X3+X5+X7+X8     | 87.0188       | X1+X2+X3+X7+X9  | 127.098       | X1+X3+X5+X7+X8  | 132.240       |
| X3+X5+X7+X9     | -58.367       | X1+X2+X3+X8+X9  | 122.652       | X1+X3+X5+X7+X9  | 125.430       |
| X3+X5+X8+X9     | -70.208       | X1+X2+X4+X5+X6  | 71.736        | X1+X3+X5+X8+X9  | -90.511       |
| X3+X6+X7+X8     | 87.6214       | X1+X2+X4+X5+X7  | 85.912        | X1+X3+X6+X7+X8  | 132.299       |
| X3+X6+X7+X9     | -58.098       | X1+X2+X4+X5+X8  | 81.455        | X1+X3+X6+X7+X9  | 125.453       |
| X3+X6+X8+X9     | -70.129       | X1+X2+X4+X5+X9  | 139.566       | X1+X3+X6+X8+X9  | 121.611       |
| X3+X7+X8+X9     | 74.713        | X1+X2+X4+X6+X7  | 87.163        | X1+X3+X7+X8+X9  | 125.705       |
| X4+X5+X6+X7     | 139.289       | X1+X2+X4+X6+X8  | 81.643        | X1+X4+X5+X6+X7  | -81.509       |
| X4+X5+X6+X8     | -75.597       | X1+X2+X4+X6+X9  | 139.817       | X1+X4+X5+X6+X8  | 155.091       |
| X4+X5+X6+X9     | -15.860       | X1+X2+X4+X7+X8  | 87.915        | X1+X4+X5+X6+X9  | -94.524       |
| X4+X5+X7+X8     | 90.241        | X1+X2+X4+X7+X9  | 132.804       | X1+X4+X5+X7+X8  | 142.481       |
| X4+X5+X7+X9     | -37.660       | X1+X2+X4+X8+X9  | 125.750       | X1+X4+X5+X7+X9  | -129.891      |

Table 4.1.16. (Continued)

| Selection index | Expected gain | Selection index   | Expected gain | Selection index   | Expected gain |
|-----------------|---------------|-------------------|---------------|-------------------|---------------|
| X1+X4+X5+X8+X9  | -97.224       | X2+X4+X5+X8+X9    | -84.3102      | X1+X2+X3+X4+X6+X8 | 131.037       |
| X1+X4+X6+X7+X8  | 142.736       | X2+X4+X6+X7+X8    | -140.498      | X1+X2+X3+X4+X6+X9 | -87.764       |
| X1+X4+X6+X7+X9  | 129.958       | X2+X4+X6+X7+X9    | -129.484      | X1+X2+X3+X4+X7+X8 | 126.399       |
| X1+X4+X6+X8+X9  | 124.759       | X2+X4+X6+X8+X9    | -125.393      | X1+X2+X3+X4+X7+X9 | 122.238       |
| X1+X4+X7+X8+X9  | 91.493        | X2+X4+X7+X8+X9    | -83.163       | X1+X2+X3+X4+X8+X9 | -83.937       |
| X1+X5+X6+X7+X8  | 142.996       | X2+X5+X6+X7+X8    | 140.516       | X1+X2+X3+X5+X6+X7 | -82.897       |
| X1+X5+X6+X7+X9  | 130.159       | X2+X5+X6+X7+X9    | -129.58       | X1+X2+X3+X5+X6+X8 | 131.135       |
| X1+X5+X6+X8+X9  | 125.175       | X2+X5+X6+X8+X9    | -125.628      | X1+X2+X3+X5+X6+X9 | -86.961       |
| X1+X5+X7+X8+X9  | 125.150       | X2+X5+X7+X8+X9    | 125.650       | X1+X2+X3+X5+X7+X8 | 126.668       |
| X1+X6+X7+X8+X9  | -70.953       | X2+X6+X7+X8+X9    | -87.727       | X1+X2+X3+X5+X7+X9 | 122.403       |
| X2+X3+X4+X5+X6  | 80.609        | X3+X4+X5+X6+X7    | -79.414       | X1+X2+X3+X5+X8+X9 | -87.180       |
| X2+X3+X4+X5+X7  | 80.541        | X3+X4+X5+X6+X8    | 149.386       | X1+X2+X3+X6+X7+X8 | 126.692       |
| X2+X3+X4+X5+X8  | 137.244       | X3+X4+X5+X6+X9    | -87.242       | X1+X2+X3+X6+X7+X9 | 122.415       |
| X2+X3+X4+X5+X9  | -81.088       | X3+X4+X5+X7+X8    | -139.803      | X1+X2+X3+X6+X8+X9 | 119.683       |
| X2+X3+X4+X6+X7  | -80.657       | X3+X4+X5+X7+X9    | 129.015       | X1+X2+X3+X7+X8+X9 | -84.064       |
| X2+X3+X4+X6+X8  | 137.454       | X3+X4+X5+X8+X9    | -87.817       | X1+X2+X4+X5+X6+X7 | -81.314       |
| X2+X3+X4+X6+X9  | 85.554        | X3+X4+X6+X7+X8    | 140.026       | X1+X2+X4+X5+X6+X8 | 138.343       |
| X2+X3+X4+X7+X8  | 131.664       | X3+X4+X6+X7+X9    | -129.085      | X1+X2+X4+X5+X6+X9 | -86.868       |
| X2+X3+X4+X7+X9  | 125.238       | X3+X4+X6+X8+X9    | -124.669      | X1+X2+X4+X5+X7+X8 | 131.982       |
| X2+X3+X4+X8+X9  | -79.440       | X3+X4+X7+X8+X9    | -86.074       | X1+X2+X4+X5+X7+X9 | 125.358       |
| X2+X3+X5+X6+X7  | -80.116       | X3+X5+X6+X7+X8    | 140.168       | X1+X2+X4+X5+X8+X9 | -87.210       |
| X2+X3+X5+X6+X8  | 137.403       | X3+X5+X6+X7+X9    | -124.977      | X1+X2+X4+X6+X7+X8 | 132.064       |
| X2+X3+X5+X6+X9  | -84.725       | X3+X5+X7+X8+X9    | -124.976      | X1+X2+X4+X6+X7+X9 | 125.391       |
| X2+X3+X5+X7+X8  | 131.844       | X3+X6+X7+X8+X9    | -87.618       | X1+X2+X4+X6+X8+X9 | 121.933       |
| X2+X3+X5+X7+X9  | 125.370       | X4+X5+X6+X7+X8    | -158.199      | X1+X2+X4+X7+X8+X9 | -86.108       |
| X2+X3+X5+X8+X9  | -84.938       | X4+X5+X6+X7+X9    | 135.467       | X1+X2+X5+X6+X7+X8 | 132.306       |
| X2+X3+X6+X7+X8  | -131.944      | X4+X5+X6+X8+X9    | -129.515      | X1+X2+X5+X6+X7+X9 | 125.551       |
| X2+X3+X6+X7+X9  | 125.412       | X4+X5+X7+X8+X9    | -129.552      | X1+X2+X5+X6+X8+X9 | 122.204       |
| X2+X3+X6+X8+X9  | -122.275      | X4+X6+X7+X8+X9    | 129.865       | X1+X2+X5+X7+X8+X9 | 122.190       |
| X2+X3+X7+X8+X9  | -77.084       | X5+X6+X7+X8+X9    | -77.158       | X1+X2+X6+X7+X8+X9 | -94.904       |
| X2+X4+X5+X6+X7  | -77.517       | X1+X2+X3+X4+X5+X6 | -85.012       | X1+X3+X4+X5+X6+X7 | -83.155       |
| X2+X4+X5+X6+X8  | 148.824       | X1+X2+X3+X4+X5+X7 | -83.302       | X1+X3+X4+X5+X6+X8 | 138.144       |
| X2+X4+X5+X6+X9  | -83.973       | X1+X2+X3+X4+X5+X8 | 130.937       | X1+X3+X4+X5+X6+X9 | -90.002       |
| X2+X4+X5+X7+X8  | 140.244       | X1+X2+X3+X4+X5+X9 | -85.503       | X1+X3+X4+X5+X7+X8 | 131.136       |
| X2+X4+X5+X7+X9  | -129.400      | X1+X2+X3+X4+X6+X7 | -83.434       | X1+X3+X4+X5+X7+X9 | 124.865       |



Table 4.1.16. (Continued)

| Selection index   | Expected gain | Selection index      | Expected gain | Selection index            | Expected gain |
|-------------------|---------------|----------------------|---------------|----------------------------|---------------|
| X1+X3+X4+X5+X8+X9 | -90.582       | X2+X4+X6+X7+X8+X9    | 125.281       | X1+X3+X5+X6+X7+X8+X9       | -75.723       |
| X1+X3+X4+X6+X7+X8 | 131.178       | X2+X5+X6+X7+X8+X9    | -86.336       | X1+X4+X5+X6+X7+X8+X9       | -84.739       |
| X1+X3+X4+X6+X7+X9 | 124.882       | X3+X4+X5+X6+X7+X8    | 138.507       | X2+X3+X4+X5+X6+X7+X8       | 130.285       |
| X1+X3+X4+X6+X8+X9 | 121.161       | X3+X4+X5+X6+X7+X9    | 128.498       | X2+X3+X4+X5+X6+X7+X9       | -121.657      |
| X1+X3+X4+X7+X8+X9 | -88.889       | X3+X4+X5+X6+X8+X9    | -124.393      | X2+X3+X4+X5+X7+X8+X9       | -121.653      |
| X1+X3+X5+X6+X7+X8 | 131.546       | X3+X4+X5+X7+X8+X9    | -124.387      | X2+X3+X4+X6+X7+X8+X9       | -121.854      |
| X1+X3+X5+X6+X7+X9 | 125.090       | X3+X4+X6+X7+X8+X9    | -124.672      | X2+X3+X5+X6+X7+X8+X9       | -75.2753      |
| X1+X3+X5+X6+X8+X9 | 121.498       | X3+X5+X6+X7+X8+X9    | 128.986       | X2+X4+X5+X6+X7+X8+X9       | 124.113       |
| X1+X3+X5+X7+X8+X9 | 121.463       | X4+X5+X6+X7+X8+X9    | -84.295       | X3+X4+X5+X6+X7+X8+X9       | -86.844       |
| X1+X3+X6+X7+X8+X9 | -91.459       | X1+X2+X3+X4+X5+X6+X7 | -83.204       | X1+X2+X3+X4+X5+X6+X7+X8    | 125.653       |
| X1+X4+X5+X6+X7+X8 | 140.908       | X1+X2+X3+X4+X5+X6+X8 | 130.314       | X1+X2+X3+X4+X5+X6+X7+X9    | 119.263       |
| X1+X4+X5+X6+X7+X9 | 129.328       | X1+X2+X3+X4+X5+X6+X9 | -87.154       | X1+X2+X3+X4+X5+X7+X8+X9    | 119.240       |
| X1+X4+X5+X6+X8+X9 | 124.535       | X1+X2+X3+X4+X5+X7+X8 | 126.012       | X1+X2+X3+X4+X6+X7+X8+X9    | 119.459       |
| X1+X4+X5+X7+X8+X9 | 124.505       | X1+X2+X3+X4+X5+X7+X9 | -122.007      | X1+X2+X3+X5+X6+X7+X8+X9    | 121.581       |
| X1+X4+X6+X7+X8+X9 | 124.887       | X1+X2+X3+X4+X5+X8+X9 | -87.368       | X1+X2+X4+X5+X6+X7+X8+X9    | -120.925      |
| X1+X5+X6+X7+X8+X9 | -80.054       | X1+X2+X3+X4+X6+X7+X8 | 126.029       | X1+X3+X4+X5+X6+X7+X8+X9    | -121.461      |
| X2+X3+X4+X5+X6+X7 | -80.535       | X1+X2+X3+X4+X6+X7+X9 | 122.015       | X2+X3+X4+X5+X6+X7+X8+X9    | -119.148      |
| X2+X3+X4+X5+X6+X8 | 136.221       | X1+X2+X3+X4+X6+X8+X9 | -119.352      | X1+X2+X3+X4+X5+X6+X7+X8+X9 | 158.723       |
| X2+X3+X4+X5+X6+X9 | -84.999       | X1+X2+X3+X4+X7+X8+X9 | -86.646       |                            |               |
| X2+X3+X4+X5+X7+X8 | 130.909       | X1+X2+X3+X5+X6+X7+X8 | 126.277       |                            |               |
| X2+X3+X4+X5+X7+X9 | 124.861       | X1+X2+X3+X5+X6+X7+X9 | -122.172      |                            |               |
| X2+X3+X4+X5+X8+X9 | -85.208       | X1+X2+X3+X5+X6+X8+X9 | 119.583       |                            |               |
| X2+X3+X4+X6+X7+X8 | 130.994       | X1+X2+X3+X5+X7+X8+X9 | 119.562       |                            |               |
| X2+X3+X4+X6+X7+X9 | 124.897       | X1+X2+X3+X6+X7+X8+X9 | -86.348       |                            |               |
| X2+X3+X4+X6+X8+X9 | 121.850       | X1+X2+X4+X5+X6+X7+X8 | -68.705       |                            |               |
| X2+X3+X4+X7+X8+X9 | -84.463       | X1+X2+X4+X5+X6+X7+X9 | 125.015       |                            |               |
| X2+X3+X5+X6+X7+X8 | 131.163       | X1+X2+X4+X5+X6+X8+X9 | -121.767      |                            |               |
| X2+X3+X5+X6+X7+X9 | 125.023       | X1+X2+X4+X5+X7+X8+X9 | -121.750      |                            |               |
| X2+X3+X5+X6+X8+X9 | 122.064       | X1+X2+X4+X6+X7+X8+X9 | -122.003      |                            |               |
| X2+X3+X5+X7+X8+X9 | 122.063       | X1+X2+X5+X6+X7+X8+X9 | -93.413       |                            |               |
| X2+X3+X6+X7+X8+X9 | -83.530       | X1+X3+X4+X5+X6+X7+X8 | 130.517       |                            |               |
| X2+X4+X5+X6+X7+X8 | 138.933       | X1+X3+X4+X5+X6+X7+X9 | -75.450       |                            |               |
| X2+X4+X5+X6+X7+X9 | -128.866      | X1+X3+X4+X5+X6+X8+X9 | -78.932       |                            |               |
| X2+X4+X5+X6+X8+X9 | -125.04       | X1+X3+X4+X5+X7+X8+X9 | -78.971       |                            |               |
| X2+X4+X5+X7+X8+X9 | -125.067      | X1+X3+X4+X6+X7+X8+X9 | -78.657       |                            |               |

#### 4.1.5 GENETIC DIVERGENCE ANALYSIS

In order to find out the extent of genetic diversity among the 25 inbred lines cluster, variate and vector analyses were performed for different characters. The results of these analyses are described below.

##### Cluster Analysis

By application of non-hierarchical clustering using co-variance matrix, 25 lines were grouped into five different clusters. Compositions of different clusters with their corresponding inbred lines in each cluster are presented in Table 4.1.17. It is revealed from Table 4.1.17 that clusters I, II and IV had the maximum number of lines and cluster III had the minimum number.

Cluster III had three inbred lines *viz.*, IL-2, IL-4 and IL-22. Cluster V consisted of four inbred lines *viz.*, IL-11, IL-17, IL-21 and IL-22. Cluster I had six inbred lines which were IL-1, IL-3, IL-5, IL-7, IL-8 and IL-12. Cluster II also comprised six inbred lines, which were IL-6, IL-9, IL-17, IL-18, IL-21 and IL-25. Six inbred lines *viz.*, IL-10, IL-13, IL-14, IL-15, IL-16 and IL-20 were found in cluster IV.

##### Canonical Variate Analysis

Canonical variate analysis was done to calculate intra-cluster Mahalanobis's values. The intra and inter-cluster distance ( $D^2$ ) values are presented in Table 4.1.18. Inbred lines grouped in the intra-cluster are expected to be genetically more similar to each other while inbred lines grouped in inter clusters as genetically more divergent. Intra-group distances appeared much smaller than inter-group, suggesting a lower genetic diversity among the lines of the same group than those from different groups. The intra-cluster divergence among the inbred lines under different clusters varied from 0.252 to 1.642. The highest intra-cluster distance was observed among the lines in cluster III; whereas, it was minimum in cluster V. The intra-cluster distance of clusters I, II and IV was 0.492, 0.683 and 0.871, respectively. Inter-cluster distances ranged from 3.766 to 19.279.

The maximum inter-cluster distance was observed between clusters I and II (19.279) indicating that the inbred lines grouped in these clusters were highly divergent from each other. On the other hand, the distance between clusters I and V was 18.844 and between II and III, I and IV and III and V were approximately same values. The lowest inter-cluster distance was observed between the clusters II and IV (3.766).

### **Cluster Means**

The genetic differences between clusters were reflected in their cluster means. Mean values for different clusters are presented in Table 4.1.19. The highest mean values for days to tasseling and silking, days to maturity, ear length and grain yield/plant were observed in the cluster V. On the other hand, in cluster V the highest values were recorded for days to maturity, grain yield/plant and days to tasseling. Number of kernels/ear, plant height, days to maturity, grain yield/plant and days to silking showed the highest cluster means in clusters IV, V, II, and I (Table 4.1.19).

### **Contribution of Characters towards Divergence of the inbred lines**

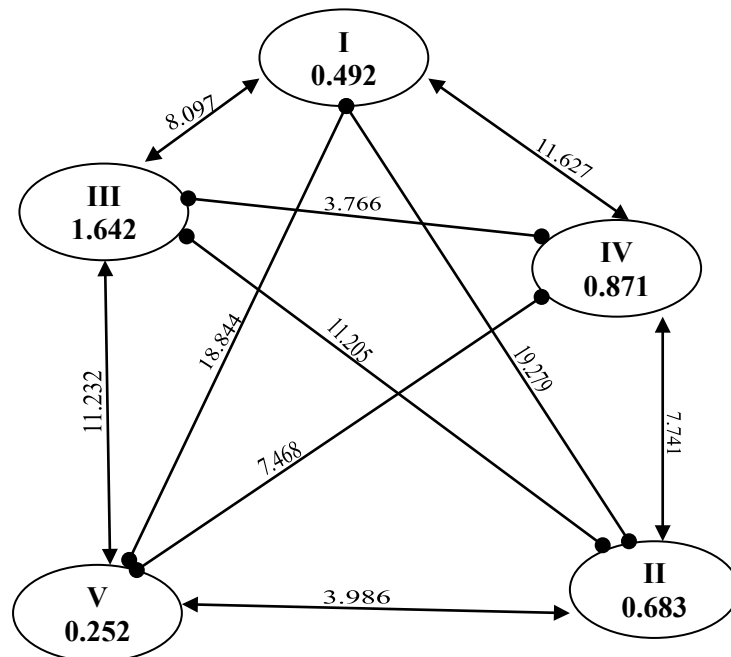
Contribution of characters towards divergence is presented in Table 4.1.20. Results showed that, Vector I obtained from principal component analysis expressed that the important characters responsible for genetic divergence in the major axis of differentiation were days to silking, ear height, cob diameter and number of kernels/ear. In vector II, which is the second axis of differentiation, the responsible characters were ear diameter and number of kernel rows/ear, played their major role on genetic divergence. Days to tasseling, plant height, cob length, cob diameter, number of kernels/ear and kernel yield showed positive values in respect to both the vectors were the major important traits responsible for genetic divergence in the major axis of differentiation. So, the greater divergence in the present materials due to these six characters will offer a good scope for improvement of yield through selection of parents.

**Table 4.1.17. Distribution of 25 maize inbred lines in five different clusters**

| Group/Cluster no. | No. of inbred lines | Inbred lines in different clusters |
|-------------------|---------------------|------------------------------------|
| I                 | 6                   | IL1, IL3, IL5, IL7, IL8, IL12      |
| II                | 6                   | IL6, IL9, IL17, IL18, IL21, IL25   |
| III               | 3                   | IL2, IL4, IL22                     |
| IV                | 6                   | IL10, IL13, IL14, IL15, IL16, IL20 |
| V                 | 4                   | IL11, IL17, IL21, IL22             |
| Total=25          |                     |                                    |

**Table 4.1.18. Inter and intra-cluster (bold) distance ( $D^2$ ) for 25 maize inbred lines obtained by canonical variate analysis**

| Clusters | I            | II           | III          | IV           | V            |
|----------|--------------|--------------|--------------|--------------|--------------|
| I        | <b>0.492</b> |              |              |              |              |
| II       | 19.279       | <b>0.683</b> |              |              |              |
| III      | 8.097        | 11.205       | <b>1.642</b> |              |              |
| IV       | 11.627       | 7.741        | 3.766        | <b>0.871</b> |              |
| V        | 18.844       | 3.986        | 11.232       | 7.468        | <b>0.252</b> |

**Figure 4.1.3. Diagram showing intra and inter-cluster distance of 30 maize inbred lines**

**Table 4.1.19. Cluster means for 11 characters of 25 maize inbred lines**

| Characters             | Clusters |        |        |        |        |
|------------------------|----------|--------|--------|--------|--------|
|                        | I        | II     | III    | IV     | V      |
| Days to tasseling      | 64.75    | 65.55  | 64.67  | 65.41  | 65.88  |
| Days to silking        | 66.73    | 65.80  | 64.89  | 66.21  | 66.18  |
| Days to maturity       | 102.40   | 105.55 | 101.89 | 102.31 | 103.51 |
| Plant height           | 114.40   | 119.15 | 105.85 | 109.55 | 102.25 |
| Ear height             | 45.39    | 49.91  | 43.84  | 47.96  | 49.75  |
| Cob length             | 12.08    | 13.50  | 10.93  | 13.70  | 11.76  |
| Cob diameter           | 11.20    | 10.85  | 12.10  | 11.50  | 11.25  |
| No. of kernel rows/cob | 13.75    | 12.85  | 12.15  | 11.90  | 12.25  |
| No. of kernels/row     | 23.55    | 21.85  | 23.31  | 20.50  | 19.90  |
| No. of kernels/cob     | 261.56   | 193.25 | 230.00 | 275.70 | 198.65 |
| Grain yield/plant(GYP) | 73.23    | 70.08  | 71.51  | 75.14  | 80.62  |

**Table 4.1.20. Relative contributions of 11 characters to the total divergence in maize**

| Characters             | Vector I | Vector II |
|------------------------|----------|-----------|
| Days to tasseling      | 0.1511   | 0.1426    |
| Days to silking        | 0.4569   | -0.8246   |
| Days to maturity       | -0.3794  | -0.2930   |
| Plant height           | 0.0888   | 0.0997    |
| Ear height             | 0.4263   | -0.0549   |
| Cob length             | 0.4260   | 0.6849    |
| Cob diameter           | 0.8342   | 2.6272    |
| No. of kernel rows/cob | -0.0093  | 0.5307    |
| No. of kernels/row     | -0.9731  | -1.1659   |
| No. of kernels/cob     | 0.2166   | 0.0030    |
| Grain yield/plant(GYP) | 0.1445   | 0.1480    |

## EXPERIMENT II (6×6 DIALLEL CROSS)

### 4.2.1 DIALLEL ANALYSIS

#### Mean Performance of F<sub>1</sub> Hybrids Involving 25 Inbred Lines

Statistically significant variation was observed among all the crosses. From Table 4.2.1, it is observed that a number F<sub>1</sub> yielded higher than selfing. The maximum yield was obtained from the F<sub>1</sub>s P<sub>1</sub>×P<sub>5</sub>, P<sub>5</sub>×P<sub>6</sub> and P<sub>3</sub>×P<sub>5</sub> followed by P<sub>4</sub>×P<sub>6</sub>. The minimum yield was recorded from P<sub>2</sub>×P<sub>5</sub> followed by P<sub>2</sub>×P<sub>4</sub>. Among the F<sub>1</sub>s, P<sub>1</sub>×P<sub>2</sub> followed by P<sub>1</sub>×P<sub>3</sub> and P<sub>1</sub>×P<sub>5</sub> showed identical and took maximum days to tasseling. The shorter time to tasseling was found for P<sub>3</sub>×P<sub>4</sub> followed by P<sub>2</sub>×P<sub>6</sub> and P<sub>2</sub>×P<sub>3</sub>. Regarding days to silking a wide range of variation (92.50-100.25) was observed for the trait. The F<sub>1</sub>s P<sub>1</sub>×P<sub>2</sub>, P<sub>1</sub>×P<sub>5</sub> and P<sub>1</sub>×P<sub>3</sub> took identical and took the maximum time. The early maturing F<sub>1</sub> was P<sub>3</sub>×P<sub>5</sub>, which was statistically identical with P<sub>2</sub>×P<sub>5</sub> and P<sub>3</sub>×P<sub>4</sub> followed by P<sub>2</sub>×P<sub>4</sub>. A range from 121.62-160.10cm was observed for plant height. Among the F<sub>1</sub>s produced significantly highest plant height followed by P<sub>2</sub>×P<sub>3</sub>, P<sub>4</sub>×P<sub>6</sub> and P<sub>3</sub>×P<sub>4</sub>. The shortest plant height was recorded from P<sub>3</sub>×P<sub>5</sub>. In case of ear height a range of 54.92-75.76 cm was found among the F<sub>1</sub>s. P<sub>2</sub>×P<sub>3</sub> produced significantly highest ear height followed by P<sub>2</sub>×P<sub>4</sub>. The lowest ear height was found for P<sub>1</sub>×P<sub>5</sub>. The highest cob length was recorded in P<sub>2</sub>×P<sub>6</sub> followed by P<sub>2</sub>×P<sub>5</sub>, P<sub>1</sub>×P<sub>4</sub> and P<sub>1</sub>×P<sub>3</sub>. P<sub>1</sub>×P<sub>6</sub> produced the shortest cob length. The maximum cob diameter was found in P<sub>4</sub>×P<sub>5</sub>, P<sub>1</sub>×P<sub>3</sub>, P<sub>2</sub>×P<sub>3</sub> and P<sub>1</sub>×P<sub>5</sub> and minimum was produced by P<sub>3</sub>×P<sub>5</sub>. Regarding number of rows/cob, hybrids P<sub>1</sub>×P<sub>2</sub>, P<sub>1</sub>×P<sub>3</sub> and P<sub>2</sub>×P<sub>3</sub> produced maximum number which was significant. P<sub>2</sub>×P<sub>5</sub> gave the minimum number. P<sub>1</sub>×P<sub>3</sub> produced significantly the highest number of kernels/row and minimum number was in P<sub>2</sub>×P<sub>4</sub>. The highest number of grains/cob was obtained from P<sub>2</sub>×P<sub>3</sub> followed by P<sub>1</sub>×P<sub>2</sub>, P<sub>1</sub>×P<sub>3</sub> and P<sub>5</sub>×P<sub>6</sub>. Now based on yield and desired quantitative characters F<sub>1</sub> hybrids P<sub>1</sub>×P<sub>5</sub>, P<sub>5</sub>×P<sub>6</sub>, P<sub>4</sub>×P<sub>5</sub>, P<sub>3</sub>×P<sub>5</sub> and P<sub>4</sub>×P<sub>6</sub> are considered as better performer.

### Testing the Validity of the Hypothesis

The validity of the postulated hypothesis for diallel was tested by  $t^2$  and the obtained values were 0.076, 0.52, 0.0028, 0.002, 0.224 and 0.0212, respectively for characters days to tasselling, days to silking, plant height, number of rows/cob, ear height and cob length. The values suggested the probable fulfilment of the postulated hypothesis.

### Genetic Components of Variation and Their Proportions

Estimates of genetic components of variation and their proportion are presented in Tables 4.2.2-4.2.12. The sign of component F indicates the relative frequencies of dominance and recessive alleles in the parents. In the present study value of F for all the characters except cob diameter were positive and greater than zero, which expressed that dominant alleles were more frequent than recessive alleles. The estimate of additive genetic variance (D) was significant for days to silking, days to maturity, ear height and number of kernel rows/ear indicating the importance of additive gene effect in their inheritance. The component  $H_1$  which measures the dominance variation was highly significant for all the characters which indicate the dominance gene effect in the inheritance of these characters. The component  $H_2$  was highly significant for all the characters indicating the dominance with asymmetry of positive and negative gene effect in controlling these characters. Thus highly significant values of the components D,  $H_1$  and  $H_2$  indicated the importance of both additive and dominant gene effect for the characters under study. The magnitude of  $H_1$  was greater than D indicating the predominance of dominance effect over additive effect for expression of the characters. The value of  $h^2$  which measures the dominant effect over all loci was non-significant for all the characters. The magnitude of E (environmental variance) for each character was much lower than the respective value of D and  $H_1$ .

This indicated that the environment had lesser effect on the characters than the additive and dominant effects. The average degree of dominance  $(H_1/D)^{1/2}$  was more than unity for all the characters suggesting the importance of over dominance for these traits.

The proportion (ratio) of  $H_2/4H_1$  provides an estimate of the average frequency of positive and negative alleles in the parents. The values of this ratio were smaller than 0.25 for all the characters which indicated that positive and negative alleles were not distributed in equal proportion in the parents. However, the values of this ratio were near 0.25 for cob diameter which indicated that positive and negative alleles were distributed nearly in equal proportion for the trait. The ratio of  $[4DH_1]^{1/2} + F/[4DH_1]^{1/2} - F$  estimates the relative proportion of dominant and recessive alleles in the parents. In the present investigation the values of this ratio were greater than unity in all the characters except cob diameter, suggesting excess of dominant alleles and minority of recessive alleles i.e., asymmetrical distribution of dominant and recessive alleles in the parents for the characters. Another ratio  $h^2/H_2$  provides an estimate the number of groups of genes that control a character and exhibit degree of dominance to some extent. If the dominance effect of the genes differs in size or sign, the ratio would be underestimated. Also it does not provide any information about genes exhibiting little or no dominance. The values of the ratio were greater than one indicating many groups of genes were responsible for their genetical control.

### **Graphical Analysis (Wr-Vr graph)**

Wr-Vr graphs, the two directional depictions were made based on the parent-offspring co-variance ( $W_r$ ) and parental variance ( $V_r$ ) is presented in Figures 4.2.1 to 4.2.11. By plotting the paired of values of  $V_r$  and  $W_r$  the position of arrays was obtained in the  $W_r/V_r$  graphs. The position of the dominant homozygote on the regression line will be near to the origin as compared to that of the recessive homozygote.



Thus, the positions of the parents on the line will indicate the direction of dominance. In the absence of dominance, both the parental points will cluster together and will thus make a single point.

The analysis of variance due to diallel progenies indicated significant differences among themselves (Table 4.2.13), which warrants for further analysis. Hayman's graphic approach to diallel analysis is based on monogenic additive model.

Hence testing the adequacy of the model is important to detect the presence or absence of epistasis. Different assumptions underlying the diallel cross analysis were tested by  $t^2$  (test of homogeneity of  $W_r$ - $V_r$  variances). The  $t^2$  values for all the characters studied were insignificant indicated the validity of the hypothesis i.e., the basic assumption (including the absence of epistasis) made for the simple additive-dominance model was satisfied for all the traits studied.

**Days to tasseling:** The observed regression line intercepted the  $W_r$  axis above the point of origin showing partial dominance gene action in the expression of the character. The scattered distribution of array points indicated the diversity existed among the parents. It is clear from the graph that the array points representing their respective parental number are clustered into three major groups along the regression line.

**Days to silking:** The observed regression line passed through the  $W_r$  axis above the point of origin showing over dominance. Wide distribution of array points in the  $V_r$ - $W_r$  graph showed genetic diversity among the parents. It is clear from the graph that the array points representing their respective parental number are clustered into two major groups along the regression line.

**Days to maturity:** The observed regression line intercepted the  $W_r$  axis above the point of origin showing partial dominance gene action in the expression of the character. The striking discontinuity between array points indicated that there was a wide genetic diversity between and within the parental groups.

The parent  $P_2$  had maximum concentration of dominant genes due to their proximity to the points of origin whereas  $P_3$  far away from the origin carried maximum number of recessive genes.

**Plant height:**  $W_r$  -  $V_r$  graph for plant height is presented in Figure 4.2.4. The graph showed that the regression line passed just through the point of origin which indicated the presence of an average complete dominance over all arrays

**Ear height:** Regression line intercepted Y axis above the point of origin indicating partial dominance. The striking discontinuity among the array points representing their respective parental number indicated that there was a wide genetic diversity between and within the parents. Parent  $P_6$  had maximum concentration of dominant genes due to its proximity to the points of origin whereas the parent  $P_3$  being away from the origin carried maximum number of recessive genes. The other parents scattered along the regression line of the graph contained equal frequencies of dominant and recessive alleles.

**Cob length:** The observed regression line intercepted the  $W_r$  axis above the point of origin showing partial dominance gene action in the expression of the character. The striking discontinuity between the array points indicated that there was a wide genetic diversity between and within the parental groups.

**Cob diameter:** For cob diameter, the observed regression line passed through the  $W_r$  axis above the point of origin showing partial dominance. Wide distribution of array points in the  $W_r$ -  $V_r$  graph showed genetic diversity among the parents. It is clear from the graph that the array points representing their respective parental number are clustered into three major groups along the regression line. In the first group parent  $P_2$  and four parents had maximum concentration of dominant genes due to their proximity to the points of origin.

**Number of rows/cob:** For number of rows per cob, the observed regression line passed through the  $W_r$  axis above the point of origin showing partial dominance. Wide distribution of array points in the  $W_r$ - $V_r$  graph showed genetic diversity among the parents. Parent  $P_6$  had maximum concentration of

dominant genes due to its position nearest to the point of origin and parent P<sub>3</sub> fall furthest away from the origin carried maximum number of recessive genes containing the character. All other parents lying scattered along the regression line of the graph contained equal frequencies of dominant and recessive alleles.

**Number of kernels/ row:** Partial dominance effect of the parents was observed due to the interception of regression line above the point of origin of Wr axis. The scattered distribution of array points representing their respective parental number indicated the wide diversity among the parents.

All the parents except P<sub>4</sub> and P<sub>5</sub> formed one group where the parent P<sub>3</sub> had maximum concentration of dominant genes due to its proximity to the point of origin. The other parents occupied at the middle of the graph contained both dominant and recessive alleles.

**Number of grains/cob:** The graph showed that the regression line passed above the point of origin which indicated the presence of over dominance over all arrays. In the graph, the parent P<sub>5</sub> was nearest to the origin contained the maximum concentration of dominant genes and the parent P<sub>3</sub> being away from the origin carried maximum number of recessive genes. Other parents remained scattered along the regression line of the graph contained equal frequencies of dominant and recessive alleles.

**Grain yield:** The observed regression line intercepted the Wr axis above the point of origin showing partial dominance gene action in the expression of the character. The scattered distribution of array points indicated the diversity existed among the parents. It is clear from the graph that the array points representing their respective parental number are clustered into three major groups along the regression line. The parents P<sub>2</sub>, P<sub>3</sub> and P<sub>4</sub> nearer to the origin formed one group whereas the parent P<sub>5</sub> had maximum concentration of dominant genes due to their proximity to the points of origin. The parents P<sub>1</sub> and P<sub>6</sub> which were away from the origin formed second group where P<sub>2</sub> being away from the origin carried maximum number of recessive genes.

The other parents formed the third group occupied at the middle of the graph contained equal frequencies of dominant and recessive alleles.

**Table 4.2.1. Mean performances of 21 F<sub>1</sub> hybrids in 6×6 diallel crosses in maize**

| Crosses                | DT     | DS      | DM      | PH      | EH      | CL     | CD     | NRC    | NKR    | NGC     | GYP     |
|------------------------|--------|---------|---------|---------|---------|--------|--------|--------|--------|---------|---------|
| P1×P1                  | 99.250 | 100.250 | 132.400 | 135.500 | 65.200  | 12.500 | 13.500 | 13.500 | 26.500 | 357.750 | 83.134  |
| P1×P2                  | 97.600 | 100.200 | 147.000 | 136.800 | 55.930  | 13.300 | 14.600 | 14.700 | 28.900 | 423.530 | 108.617 |
| P1×P3                  | 97.444 | 98.889  | 149.472 | 135.167 | 59.011  | 13.849 | 15.072 | 14.389 | 31.000 | 446.056 | 96.962  |
| P1×P4                  | 95.556 | 98.611  | 142.111 | 129.408 | 61.994  | 13.982 | 14.342 | 13.778 | 28.167 | 388.074 | 103.853 |
| P1×P5                  | 96.956 | 99.944  | 142.472 | 129.797 | 54.938  | 13.574 | 14.588 | 13.722 | 27.361 | 375.455 | 128.245 |
| P1×P6                  | 94.750 | 97.611  | 151.278 | 127.154 | 59.468  | 12.683 | 14.567 | 12.722 | 24.583 | 312.755 | 101.034 |
| P2×P2                  | 90.500 | 92.500  | 153.200 | 140.500 | 65.100  | 12.500 | 15.200 | 13.500 | 26.700 | 360.450 | 103.681 |
| P2×P3                  | 93.611 | 97.972  | 156.722 | 142.138 | 75.758  | 13.538 | 14.633 | 13.611 | 27.583 | 375.440 | 99.595  |
| P2×P4                  | 94.944 | 97.528  | 150.417 | 127.522 | 72.014  | 13.276 | 13.356 | 13.222 | 24.139 | 319.170 | 95.011  |
| P2×P5                  | 94.222 | 96.944  | 150.833 | 138.586 | 63.653  | 13.997 | 14.447 | 12.472 | 28.556 | 356.151 | 89.551  |
| P2×P6                  | 93.472 | 95.917  | 148.528 | 132.322 | 61.972  | 14.553 | 14.142 | 13.000 | 27.889 | 362.556 | 100.109 |
| P3×P3                  | 90.500 | 93.800  | 152.500 | 130.500 | 59.400  | 15.200 | 13.900 | 14.200 | 30.100 | 427.420 | 121.970 |
| P3×P4                  | 93.139 | 94.944  | 152.528 | 138.253 | 71.900  | 13.847 | 13.742 | 13.028 | 29.611 | 385.767 | 104.007 |
| P3×P5                  | 95.444 | 97.778  | 143.806 | 121.619 | 71.900  | 13.296 | 13.568 | 12.889 | 28.833 | 371.630 | 120.664 |
| P3×P6                  | 95.139 | 97.889  | 146.722 | 135.206 | 69.213  | 13.585 | 13.253 | 12.778 | 29.722 | 379.784 | 104.089 |
| P4×P4                  | 92.600 | 99.500  | 143.500 | 136.800 | 71.900  | 13.500 | 15.600 | 12.500 | 26.780 | 334.750 | 117.826 |
| P4×P5                  | 94.444 | 96.694  | 144.750 | 137.161 | 60.102  | 13.033 | 16.113 | 13.250 | 24.556 | 325.361 | 126.973 |
| P4×P6                  | 93.972 | 95.667  | 148.889 | 138.394 | 71.089  | 13.186 | 14.512 | 12.639 | 29.889 | 377.762 | 116.321 |
| P5×P5                  | 95.200 | 95.700  | 150.500 | 145.400 | 102.500 | 13.200 | 12.800 | 13.000 | 32.400 | 421.200 | 122.914 |
| P5×P6                  | 94.528 | 96.694  | 148.111 | 160.100 | 81.883  | 13.186 | 13.392 | 13.222 | 31.194 | 412.460 | 127.150 |
| P6×P6                  | 94.100 | 95.050  | 150.700 | 142.900 | 85.100  | 14.000 | 14.500 | 13.400 | 25.900 | 347.060 | 115.469 |
| CV%                    | 2.214  | 2.156   | 3.528   | 5.866   | 16.354  | 4.762  | 5.769  | 4.555  | 8.0548 | 9.7949  | 12.001  |
| LSD 5%                 | 3.052  | 2.971   | 3.723   | 9.833   | 18.778  | 3.148  | 2.425  | 1.153  | 1.364  | 95.841  | 27.132  |
| SE of mean             | 0.233  | 0.226   | 0.284   | 0.751   | 1.434   | 0.241  | 0.185  | 0.088  | 0.529  | 7.321   | 2.073   |
| Level of significance. | *      | *       | *       | **      | **      | *      | *      | *      | *      | **      | **      |

**Table 4.2.2. Components of variation and their proportions for days to tasseling**

| Notation | Components of Variation | Proportional Values                         |                  |
|----------|-------------------------|---|------------------|
|          | Estimated Values        | Proportion                                  | Estimated Values |
| D        | 3.8697±1.824            | $\sqrt{H1/D}$                               | 1.548            |
| F        | 6.519±4.457             | H2/4H1                                      | 0.149            |
| H1       | 9.274±4.6318            | $[(4DH_1)^{1/2} + F] / [(4DH_1)^{1/2} - F]$ | 3.386            |
| H2       | 5.5496±4.1377           | h2/H2                                       | -0.2349          |
| h2       | -1.3037±2.784           | V1L1/WoLo1                                  | 4.236            |
| E        | 2.9607±0.689            | h <sup>2</sup> NS                           | 0.11008          |

**Table 4.2.3. Components of variation and their proportions for days to silking**

| Notation | Components of Variation | Proportional Values                         |                  |
|----------|-------------------------|---|------------------|
|          | Estimated Values        | Proportion                                  | Estimated Values |
| D        | 3.121±2.290             | $\sqrt{H1/D}$                               | 2.068            |
| F        | 6.445±5.595             | H2/4H1                                      | 0.184            |
| H1       | 13.356±5.814            | $[(4DH_1)^{1/2} + F] / [(4DH_1)^{1/2} - F]$ | 2.993            |
| H2       | 9.850±5.193             | h2/H2                                       | 0.1077           |
| h2       | 1.061±3.495             | V1L1/WoLo1                                  | 9.824            |
| E        | 2.850±0.865             | h <sup>2</sup> NS                           | 0.0166           |

**Table 4.2.4. Components of variation and their proportions for days to maturity**

| Notation | Components of Variation | Proportional Values                         |                  |
|----------|-------------------------|---|------------------|
|          | Estimated Values        | Proportion                                  | Estimated Values |
| D        | 1.0168±0.814            | $\sqrt{H1/D}$                               | 4.559            |
| F        | 0.396±1.990             | H2/4H1                                      | 0.166            |
| H1       | 3.509±2.068             | $[(4DH_1)^{1/2} + F] / [(4DH_1)^{1/2} - F]$ | 1.694            |
| H2       | 2.3414±1.847            | h2/H2                                       | -0.235           |
| h2       | -0.551±1.243            | V1L1/WoLo1                                  | -73.065          |
| E        | 0.993±0.308             | h <sup>2</sup> NS                           | 0.160            |

**Table 4.2.5. Components of variation and their proportions for plant height**

| Notation | Components of Variation |  | Proportional Values                         |                  |
|----------|-------------------------|--|---|------------------|
|          | Estimated Values        |  | Proportion                                  | Estimated Values |
| D        | 67.2918±10.879          |  | (H1/D)1/2                                   | 0.848            |
| F        | 68.4061±25.577          |  | H2/4H1                                      | 0.1805           |
| H1       | 48.4904±27.618          |  | $[(4DH_1)^{1/2} + F] / [(4DH_1)^{1/2} - F]$ | 3.984            |
| H2       | 35.0134±24.671          |  | h2/H2                                       | 4.0846           |
| h2       | 143.017±16.605          |  | V1L1/WoLo1                                  | 1.1516           |
| E        | 18.4208±4.112           |  | h <sup>2</sup> NS                           | 0.1853           |

**Table 4.2.6. Components of variation and their proportions for ear height**

| Notation | Components of Variation |  | Proportional Values                         |                  |
|----------|-------------------------|--|---|------------------|
|          | Estimated Values        |  | Proportion                                  | Estimated Values |
| D        | 7.3689±8.463            |  | (H1/D)1/2                                   | 2.606            |
| F        | 12.991±20.677           |  | H2/4H1                                      | 0.2095           |
| H1       | 50.049±21.486           |  | $[(4DH_1)^{1/2} + F] / [(4DH_1)^{1/2} - F]$ | 2.0222           |
| H2       | 41.945±19.194           |  | h2/H2                                       | 0.3466           |
| h2       | 14.539±12.918           |  | V1L1/WoLo1                                  | 11.239           |
| E        | 4.8068±3.199            |  | h <sup>2</sup> NS                           | 0.0750           |

**Table 4.2.7. Components of variation and their proportions for cob length**

| Notation | Components of Variation |  | Proportional Values                         |                  |
|----------|-------------------------|--|---|------------------|
|          | Estimated Values        |  | Proportion                                  | Estimated Values |
| D        | 0.6944±0.4515           |  | $\sqrt{H1/D}$                               | 1.6187           |
| F        | 1.4062±1.1031           |  | H2/4H1                                      | 0.1463           |
| H1       | 1.8197±1.146            |  | $[(4DH_1)^{1/2} + F] / [(4DH_1)^{1/2} - F]$ | 4.3398           |
| H2       | 1.0651±1.023            |  | h2/H2                                       | 0.1803           |
| h2       | 0.1920±0.689            |  | V1L1/WoLo1                                  | 5.8734           |
| E        | 0.7643±0.1706           |  | h <sup>2</sup> NS                           | 0.0203           |

**Table 4.2.8. Components of variation and their proportions for cob diameter**

| Notation | Components of Variation | Proportional Values                         |                  |
|----------|-------------------------|---|------------------|
|          | Estimated Values        | Proportion                                  | Estimated Values |
| D        | 1.10970±0.399           | $\sqrt{H1/D}$                               | 1.522            |
| F        | 2.1049±0.977            | H2/4H1                                      | 0.1387           |
| H1       | 2.5713±1.015            | $[(4DH_1)^{1/2} + F] / [(4DH_1)^{1/2} - F]$ | 4.3060           |
| H2       | 1.4268±0.9070           | h2/H2                                       | -0.3579          |
| h2       | -0.5107±0.610           | V1L1/WoLo1                                  | 5.1063           |
| E        | 0.9260±0.1511           | h <sup>2</sup> N                            | 0.0549           |

**Table 4.2.9. Components of variation and their proportions for number of rows/cob**

| Notation | Components of Variation | Proportional Values                         |                  |
|----------|-------------------------|---|------------------|
|          | Estimated Values        | Proportion                                  | Estimated Values |
| D        | 1.5357±0.607            | $\sqrt{H1/D}$                               | 1.6218           |
| F        | 2.1809±1.485            | H2/4H1                                      | 0.2068           |
| H1       | 4.0396±1.543            | $[(4DH_1)^{1/2} + F] / [(4DH_1)^{1/2} - F]$ | 2.5575           |
| H2       | 3.3417±1.378            | h2/H2                                       | -0.241           |
| h2       | -0.807±0.927            | V1L1/WoLo1                                  | 3.648            |
| E        | 1.464±0.229             | h <sup>2</sup> NS                           | 0.0113           |

**Table 4.2.10. Components of variation and their proportions for number of kernels/row**

| Notation | Components of Variation | Proportional Values                         |                  |
|----------|-------------------------|---|------------------|
|          | Estimated Values        | Proportion                                  | Estimated Values |
| D        | -0.5696±1.795           | $\sqrt{H1/D}$                               | 6.2486           |
| F        | 1.8679±4.386            | H2/4H1                                      | 0.2171           |
| H1       | 22.2435±4.558           | $[(4DH_1)^{1/2} + F] / [(4DH_1)^{1/2} - F]$ | 1.7113           |
| H2       | 19.3221±4.072           | h2/H2                                       | 6.8059           |
| h2       | 131.5054±2.740          | V1L1/WoLo1                                  | 27.125           |
| E        | 2.988±0.402             | h <sup>2</sup> NS                           | 0.265            |

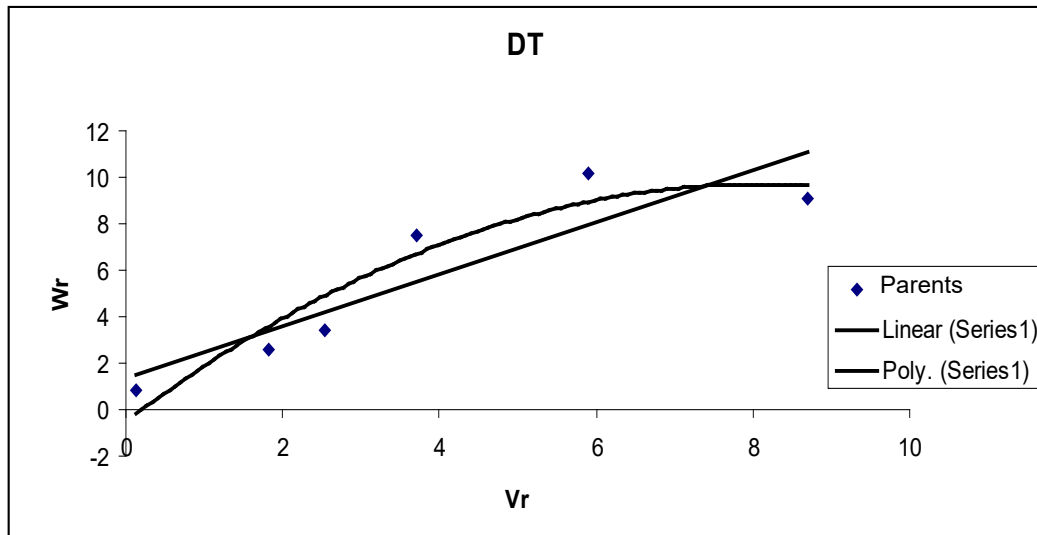
**Table 4.2.11. Components of variation and their proportions for number of grains/cob**

| Notation | Components of Variation | Proportional Values                         |                  |
|----------|-------------------------|---|------------------|
|          | Estimated Values        | Proportion                                  | Estimated Values |
| D        | 77.847±29.919           | $\sqrt{H1/D}$                               | 1.4743           |
| F        | 37.384±73.0943          | H2/4H1                                      | 0.2243           |
| H1       | 169.2121±75.954         | $[(4DH_1)^{1/2} + F] / [(4DH_1)^{1/2} - F]$ | 1.3890           |
| H2       | 151.8614±67.8519        | h2/H2                                       | 1.1937           |
| h2       | 181.288±45.668          | V1L1/WoLo1                                  | 1.8398           |
| E        | 7.2261633±11.308        | h <sup>2</sup> NS                           | 0.3901           |

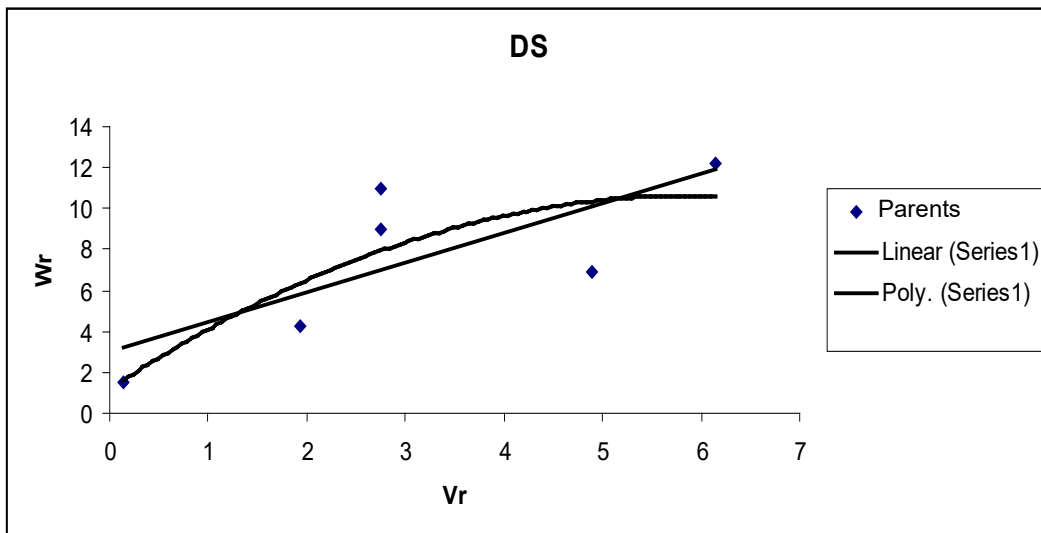
**Table 4.2.12. Components of variation and their proportions for grain yield/plant**

| Notation | Components of Variation | Proportional Values                         |                  |
|----------|-------------------------|---|------------------|
|          | Estimated Values        | Proportion                                  | Estimated Values |
| D        | 16.3356±10.529          | $\sqrt{H1/D}$                               | 1.9605           |
| F        | 1.5754±25.724           | H2/4H1                                      | 0.2319           |
| H1       | 62.793±26.730           | $[(4DH_1)^{1/2} + F] / [(4DH_1)^{1/2} - F]$ | 1.051            |
| H2       | 58.269±23.879           | h2/H2                                       | 1.024            |
| h2       | 63.071±16.072           | V1L1/WoLo1                                  | 2.533            |
| E        | 9.4061±3.979            | h <sup>2</sup> NS                           | 0.318            |

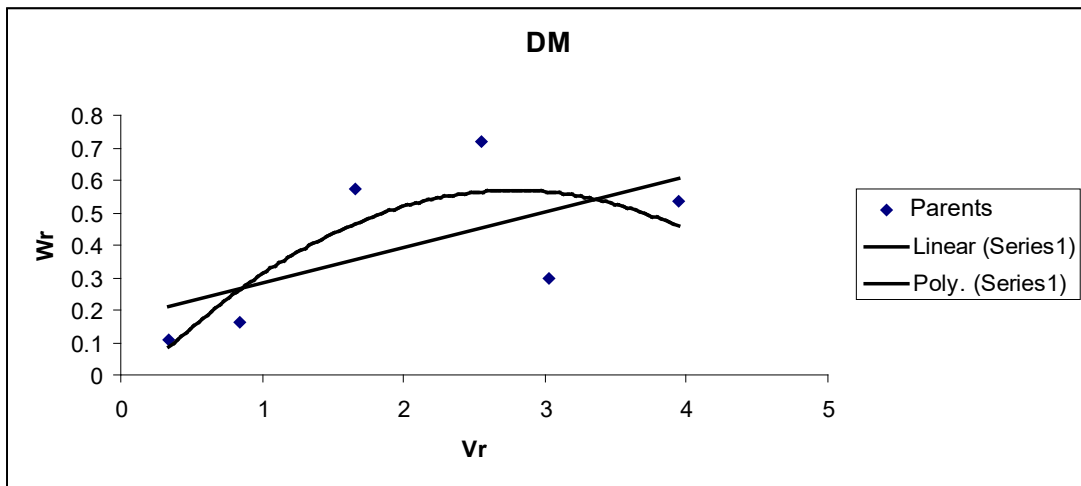




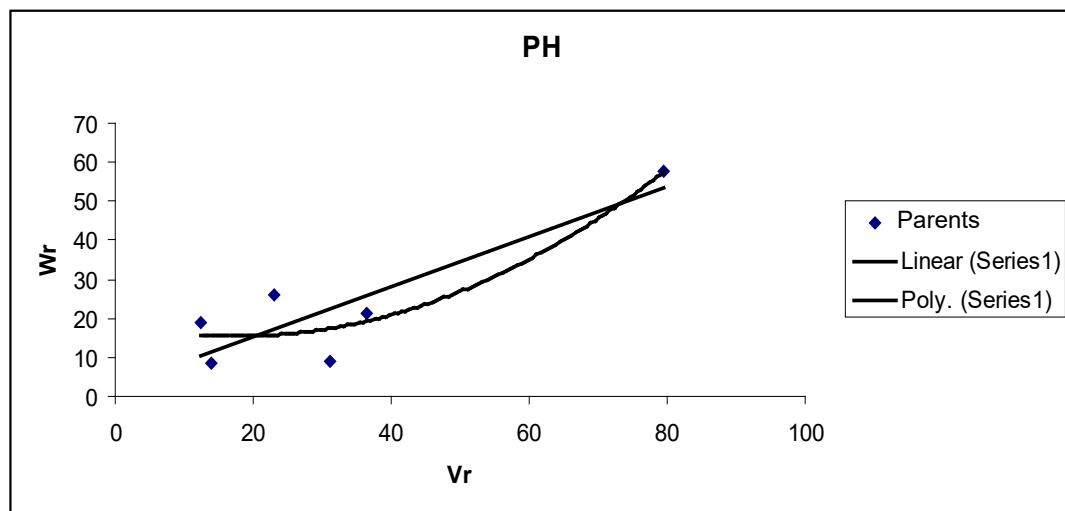
**Graph 4. 2.1. Vr–Wr graph for days to tasseling in 6x6 diallel cross in maize**



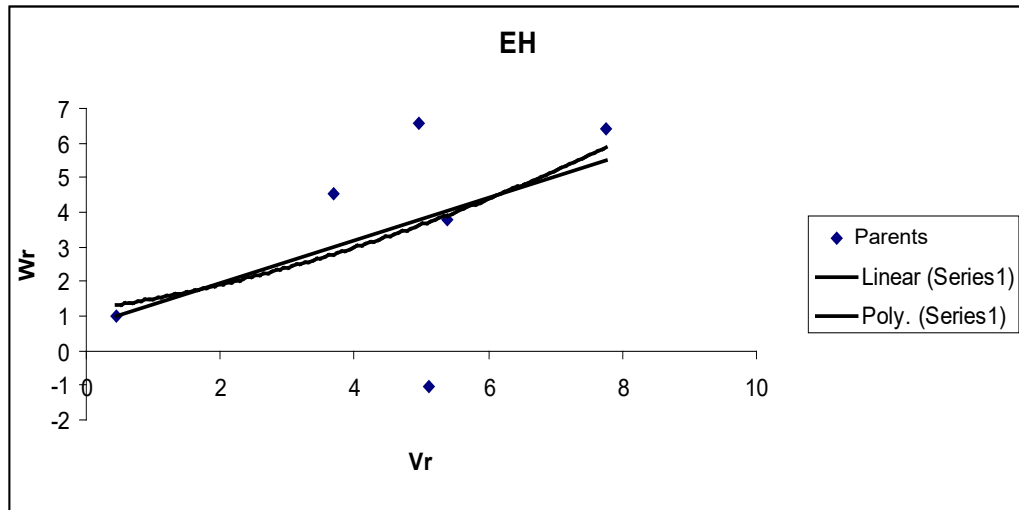
**Graph 4. 2. 2. Vr–Wr graph for days to silking in 6x6 diallel cross in maize**



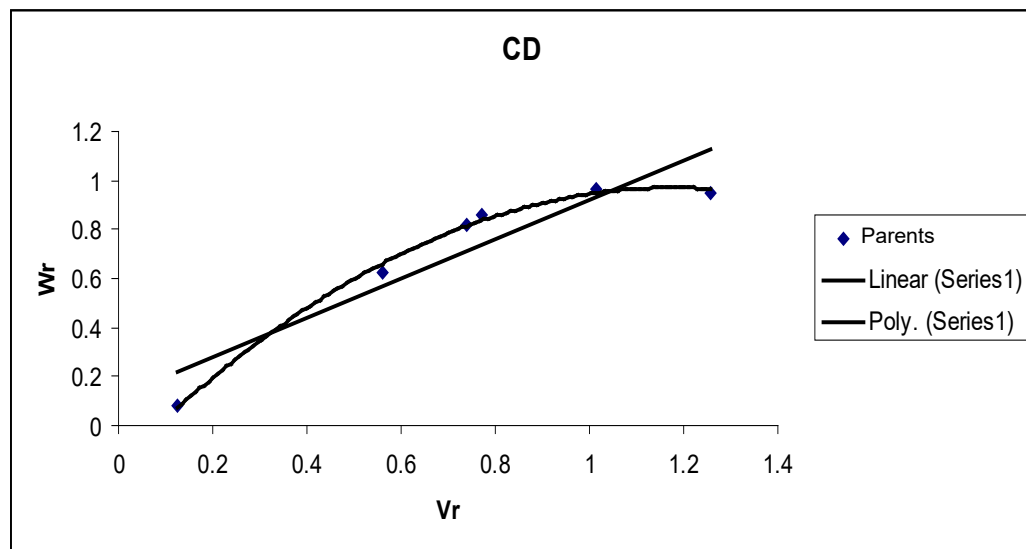
Graph 4. 2. 3. Vr–Wr graph for days to maturity in 6x6 diallel cross in maize



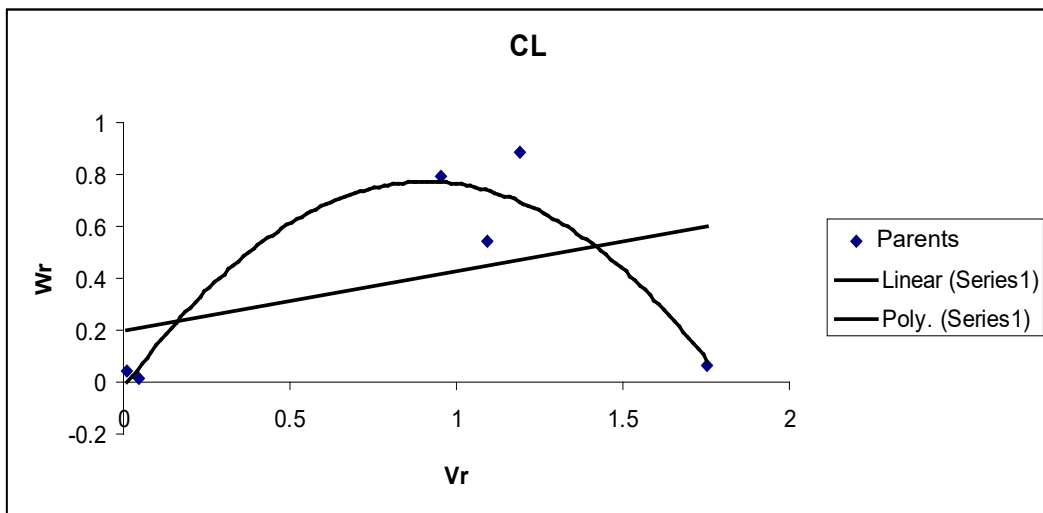
Graph 4. 2. 4. Vr–Wr graph for plant height in 6x6 diallel cross in maize



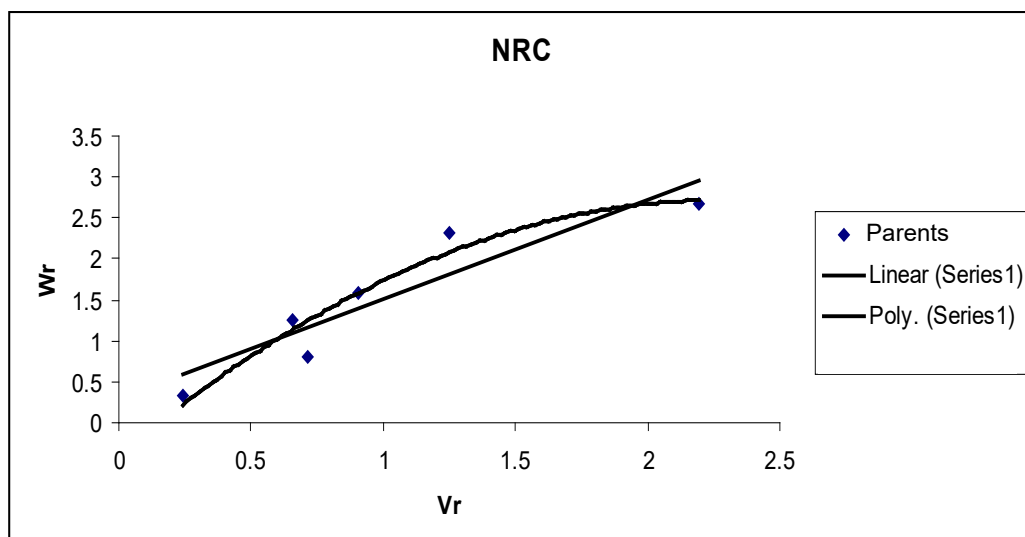
Graph 4. 2. 5. Vr–Wr graph for ear height in 6x6 diallel cross in maize



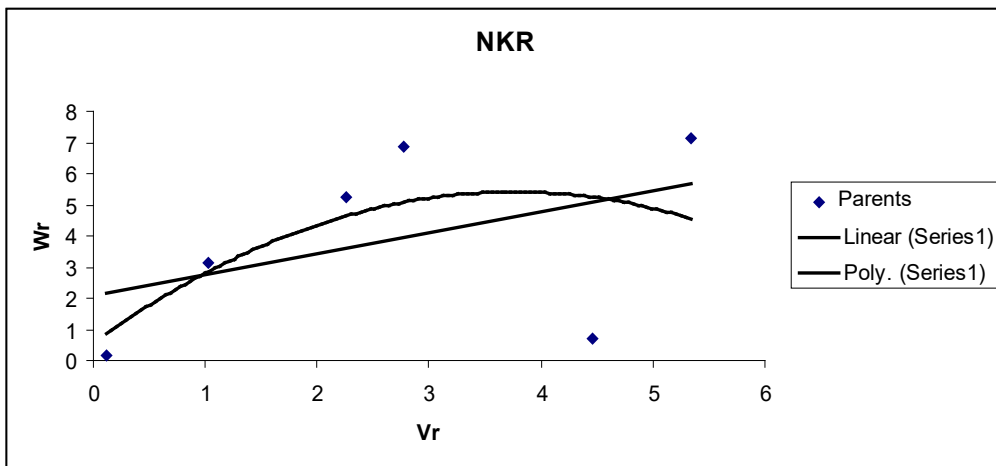
Graph 4. 2. 6. Vr–Wr graph for cob diameter in 6x6 diallel cross in maize



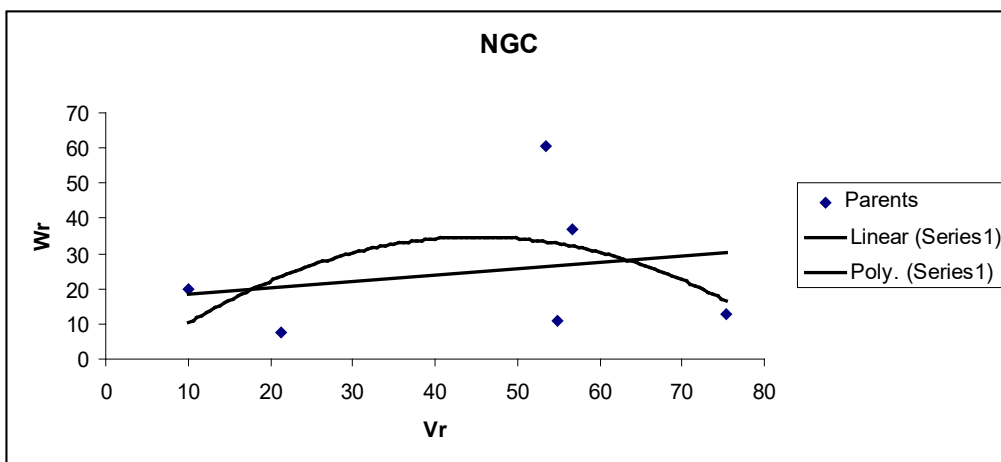
Graph 4. 2.7. Vr–Wr graph for cob length in 6x6 diallel cross in maize



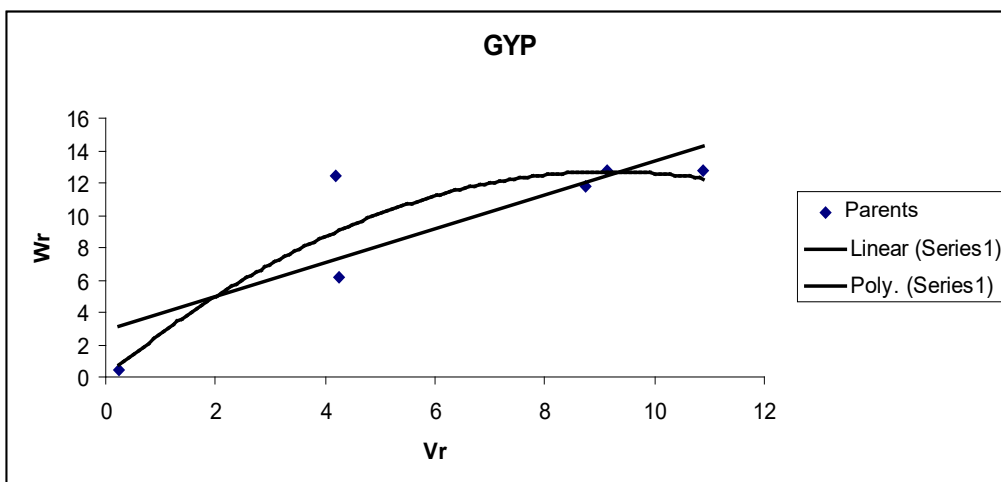
Graph 4. 2. 8. Vr–Wr graph for number of kernels/cob in 6x6 diallel cross in maize



Graph 4. 2. 9. Vr–Wr graph for number of kernels/row in 6x6 diallel cross in maize



Graph 4. 2.10. Vr–Wr graph for number of grains/cob in 6x6 diallel cross in maize



Graph 4. 2.11. Vr–Wr graph for grain yield/plant in 6x6 diallel cross in maize

## 4.2.2 COMBINING ABILITY FOR GRAIN YIELD AND YIELD COMPONENTS

### Variance Analysis

The analysis of variance revealed highly significant difference for all the characters, indicating the existence of wider genetic variability among the lines. Both general combining ability (GCA) and specific combining ability (SCA) variances were found to be highly significant (Table 4.2.13) for all the characters studied which indicated that both additive and non-additive gene action played predominant role for expression of these characters. Relative importance of GCA and SCA was calculated, closer the ratio is to unity greater is the predictability based on GCA alone.

In the present study GCA/SCA variance was observed to be unity for cob length, cob diameter and number of rows/cob indicating equal importance of both additive and non additive gene effects.

The SCA component of variance was higher than GCA component of variance for the characters like plant height, cob diameter and number of rows/cob indicating the predominance of non-additive or dominant gene action.

High magnitude of SCA components were predominant indicating the dominance and epistatic interaction for these characters. Average GCA: SCA ratio for four characters was above unity.

### General Combining Ability

The GCA effect ( $g_i$ ) represents the additive nature of gene action. The nature (direction or sign) and magnitude of  $g_i$  both are considered. Besides, *per se* performance of the parent is also considered together with  $g_i$  since the former offers authenticity to  $g_i$  as a guide to select the parent. GCA and SCA variances with each parent play a significant role in the choice of the parent. A parent with higher positive significant GCA effect is considered as a good general combiner. A parent showing high GCA and SCA variances is a better parent for creating high yielding specific combination.

The magnitude and direction of the significant effect for six parents provides meaningful comparison and would give clue to the future breeding program.

General combining ability (GCA) effects for grain yield and yield components are presented in Table 4.2.14. In this study, days to maturity, cob height and number of kernels/cob of the inbred lines with significant and negative effects were considered as good general combiners, while grain yield and other yield components of inbred with significant and positive effects were considered as good general combiners.

Highly significant and positive GCA effect of the parents P<sub>4</sub>, P<sub>5</sub> and P<sub>6</sub> were observed for grain yield (Table 4.2.14.). P<sub>1</sub> and P<sub>2</sub> parents showed significant and negative GCA effect. Inbred parent P<sub>3</sub> showed non-significant GCA effect for this character.

For days to tasseling the P<sub>2</sub>, P<sub>3</sub>, P<sub>4</sub> and P<sub>6</sub> parents showed non-significant negative GCA. Estimates of GCA were positively non-significant for P<sub>1</sub> and P<sub>5</sub>. Inbred parents P<sub>2</sub>, P<sub>3</sub>, P<sub>4</sub> and P<sub>6</sub> showed non-significant negative GCA effect for days to silking but the parents P<sub>1</sub> and P<sub>5</sub> showed positive non-significant GCA (Table 4.2.14).

The GCA effect of P<sub>1</sub>, P<sub>2</sub> and P<sub>3</sub> was significantly negative for days to maturity. Parents P<sub>4</sub>, P<sub>5</sub> and P<sub>6</sub> were found to show non-significant negative GCA effect for the character (Table 4.2.14). For plant height the GCA effects of the parents, P<sub>1</sub>, P<sub>2</sub> and P<sub>3</sub> showed non-significant positive whereas negative non-significant GCA effect was observed in P<sub>4</sub>, P<sub>5</sub> and P<sub>6</sub> (Table 4.2.14). The inbred parents P<sub>1</sub>, P<sub>5</sub> and P<sub>6</sub> showed positive significant GCA for cob height, other parents showed non-significant negative GCA effect for the character (Table 4.2.14).

The parent P<sub>3</sub> showed non-significant positive GCA for cob length. Non-significant negative GCA was found for other parents (Table 4.2.14). For cob diameter it was observed that the parents P<sub>3</sub>, P<sub>5</sub> and P<sub>6</sub> showed non significant

negative GCA. The remaining three parents showed non-significant positive GCA for cob diameter (Table 4.2.14).

The inbreds P<sub>1</sub>, P<sub>2</sub> and P<sub>3</sub> showed non-significant positive GCA for number of rows/cob and also P<sub>4</sub>, P<sub>5</sub> and P<sub>6</sub> showed non-significant negative GCA for this trait (Table 4.7.2). For number of kernels/row it was found that parents P<sub>3</sub>, P<sub>5</sub> and P<sub>6</sub> showed non-significant positive GCA. Other parents like P<sub>1</sub>, P<sub>2</sub> and P<sub>4</sub> showed non-significant negative GCA for the trait (Table 4.2.14).

The parents P<sub>1</sub>, P<sub>2</sub>, P<sub>3</sub>, and P<sub>5</sub> showed significant positive GCA for number of grains/cob. Significant negative GCA was found for P<sub>4</sub> and P<sub>6</sub> (Table 4.2.14).

Inbred parents P<sub>1</sub>, P<sub>2</sub>, P<sub>4</sub>, P<sub>5</sub> and P<sub>6</sub> showed significant positive GCA effect for grain yield but the parents P<sub>3</sub> showed positive non-significant GCA (Table 4.2.14).

From the study, it is concluded that the parents P<sub>1</sub>, P<sub>2</sub>, P<sub>4</sub> and P<sub>5</sub> are good general combiner and might be used for hybrid variety development program.

### **Specific Combining Ability**

The SCA effects signify the role of non-additive gene action in the expression of the characters. It denotes the highly specific combining ability leading to the highest performance of some specific cross combinations. For this reason it relates to a particular cross. The estimates of SCA effects are presented in Tables 4.2.14 - 4.2.25.

SCA effects for grain yield was observed among 15 crosses, eight crosses like P<sub>1</sub>×P<sub>2</sub>, P<sub>1</sub>×P<sub>5</sub>, P<sub>2</sub>×P<sub>5</sub>, P<sub>3</sub>×P<sub>5</sub>, P<sub>4</sub>×P<sub>5</sub>, P<sub>2</sub>×P<sub>6</sub>, P<sub>4</sub>×P<sub>6</sub> and P<sub>5</sub>×P<sub>6</sub> showed highly significant and positive SCA. Five other crosses, P<sub>1</sub>×P<sub>3</sub>, P<sub>2</sub>×P<sub>4</sub>, P<sub>3</sub>×P<sub>4</sub>, P<sub>1</sub>×P<sub>6</sub> and P<sub>3</sub>×P<sub>6</sub> showed significant negative SCA effect for the character.

For days to tasseling seven crosses P<sub>1</sub>×P<sub>2</sub>, P<sub>2</sub>×P<sub>4</sub>, P<sub>1</sub>×P<sub>5</sub>, P<sub>2</sub>×P<sub>5</sub>, P<sub>3</sub>×P<sub>5</sub>, P<sub>1</sub>×P<sub>6</sub> and P<sub>3</sub>×P<sub>6</sub> showed significant negative SCA which is desirable. Regarding days to silking three crosses, P<sub>3</sub>×P<sub>4</sub>, P<sub>1</sub>×P<sub>5</sub> and P<sub>3</sub>×P<sub>6</sub> showed desired significant negative SCA, P<sub>2</sub>×P<sub>5</sub>, P<sub>3</sub>×P<sub>5</sub>, P<sub>1</sub>×P<sub>6</sub> and P<sub>3</sub>×P<sub>6</sub> showed significant negative SCA, five crosses P<sub>3</sub>×P<sub>4</sub>, P<sub>1</sub>×P<sub>5</sub>, P<sub>4</sub>×P<sub>5</sub>, P<sub>3</sub>×P<sub>6</sub> and P<sub>5</sub>×P<sub>6</sub>.



For days to maturity, five crosses  $P_1 \times P_2$ ,  $P_1 \times P_3$ ,  $P_2 \times P_3$ ,  $P_3 \times P_4$  and  $P_1 \times P_6$  were positive significant and the rest three crosses  $P_3 \times P_5$ ,  $P_2 \times P_6$  and  $P_3 \times P_6$  showed significant negative SCA effect for the character (Table 4.2.17.).

Regarding plant height seven crosses,  $P_1 \times P_4$ ,  $P_1 \times P_5$ ,  $P_1 \times P_6$ ,  $P_2 \times P_4$ ,  $P_2 \times P_5$ ,  $P_2 \times P_6$  and  $P_3 \times P_5$  and  $P_3 \times P_6$  showed desired significant negative SCA, six crosses  $P_1 \times P_2$ ,  $P_1 \times P_3$ ,  $P_2 \times P_3$ ,  $P_3 \times P_4$ ,  $P_4 \times P_6$  and  $P_5 \times P_6$  showed positive significant for this trait.(Table 4.2.18). Regarding ear height  $P_1 \times P_2$ ,  $P_1 \times P_5$ ,  $P_2 \times P_5$ ,  $P_3 \times P_5$ ,  $P_4 \times P_5$ ,  $P_1 \times P_6$  and  $P_2 \times P_4$  showed significant negative SCA. The rest six crosses  $P_1 \times P_3$ ,  $P_2 \times P_3$ ,  $P_1 \times P_4$ ,  $P_2 \times P_4$ ,  $P_4 \times P_6$  and  $P_5 \times P_6$  exhibited significant positive SCA for the trait (Table 4.2.19). Regarding cob length three crosses like  $P_2 \times P_4$ ,  $P_1 \times P_5$  and  $P_2 \times P_6$  showed significant positive SCA, four crosses had significant negative SCA (Table 4.2.20). For cob diameter two crosses  $P_1 \times P_3$  and  $P_3 \times P_5$  exhibited positive significant SCA and the two crosses  $P_3 \times P_4$  and  $P_1 \times P_5$  exhibited significant negative SCA. Other crosses showed non-significant SCA for the trait (Table 4.2.21). For number of rows/cob three crosses  $P_1 \times P_2$ ,  $P_1 \times P_4$  and  $P_1 \times P_5$  showed significant positive SCA. On the other hand, negative significant SCA was observed for four crosses  $P_2 \times P_3$ ,  $P_3 \times P_5$ ,  $P_1 \times P_6$  and  $P_1 \times P_6$  for the trait (Table 4.2.22).

For number of kernels/ row five crosses  $P_1 \times P_2$ ,  $P_1 \times P_5$ ,  $P_4 \times P_5$ ,  $P_4 \times P_6$  and  $P_5 \times P_6$  showed significant positive SCA whereas three crosses  $P_2 \times P_4$ ,  $P_2 \times P_5$  and  $P_3 \times P_6$  showed significant negative SCA and the other seven crosses exhibited non significant SCA for the trait (Table 4.2.23). Regarding number of grains/cob, six crosses  $P_1 \times P_2$ ,  $P_1 \times P_4$ ,  $P_1 \times P_5$ ,  $P_4 \times P_5$  and  $P_5 \times P_6$  showed positive significant SCA whereas nine crosses showed significant negative SCA (Table 4.2.24). For grain yield/plant, twelve crosses showed significant SCA whereas six crosses  $P_1 \times P_2$ ,  $P_1 \times P_5$ ,  $P_2 \times P_5$ ,  $P_4 \times P_5$ ,  $P_2 \times P_6$ ,  $P_4 \times P_6$  and  $P_5 \times P_6$  were positive and the rest six were negative for the character (Table 4.2.25).

The proportion of GCA/SCA was more than unity for all the studied characters, suggesting that additive gene effects were more important than the non-additive ones in the expression of these characters.

**Table 4.2.13. Analysis of variance for combining ability analysis for grain yield and its components in 6×6 diallel cross in maize**

| Characters | Source of variation | df | Sum of squares | Mean squares | F.value | GCA/SGA |
|------------|---------------------|----|----------------|--------------|---------|---------|
| DT         | GCA                 | 5  | 75.048         | 15.009       | 1.532   | 3.159   |
|            | SCA                 | 15 | 71.9284        | 4.795        |         |         |
|            | Crosses             | 20 | 146.976        | 7.348        |         |         |
|            | Error               | 40 | 136.938        | 4.795        |         |         |
| DS         | GCA                 | 5  | 52.365         | 10.473       | 1.831   | 2.365   |
|            | SCA                 | 15 | 66.406         | 4.427        |         |         |
|            | Crosses             | 20 | 118.771        | 5.938        |         |         |
|            | Error               | 40 | 129.709        | 3.242        |         |         |
| DM         | GCA                 | 5  | 927.511        | 185.502      | 15.279  | 4.414   |
|            | SCA                 | 15 | 630.427        | 42.028       |         |         |
|            | Crosses             | 20 | 1557.938       | 77.896       |         |         |
|            | Error               | 40 | 203.957        | 5.09892      |         |         |
| PH         | GCA                 | 5  | 730.553        | 146.11069    | 5.295   | 0.7025  |
|            | SCA                 | 15 | 3034.03        | 202.2687     |         |         |
|            | Crosses             | 20 | 3764.584       | 188.229      |         |         |
|            | Error               | 40 | 1421.831       | 35.545       |         |         |
| EH         | GCA                 | 5  | 3533.856       | 706.771      | 2.812   | 2.8225  |
|            | SCA                 | 15 | 3756.084       | 250.405      |         |         |
|            | Crosses             | 20 | 7289.941       | 364.497      |         |         |
|            | Error               | 40 | 5185.474       | 129.636      |         |         |
| CL         | GCA                 | 5  | 58.668         | 11.733       | 2.178   | 1.7577  |
|            | SCA                 | 15 | 100.127        | 6.675        |         |         |
|            | Crosses             | 20 | 158.796        | 7.939        |         |         |
|            | Error               | 40 | 145.775        | 3.644        |         |         |
| CD         | GCA                 | 5  | 20.124         | 4.024        | 2.174   | 0.8165  |
|            | SCA                 | 15 | 73.927         | 4.928        |         |         |
|            | Crosses             | 20 | 94.052         | 4.702        |         |         |
|            | Error               | 40 | 86.51          | 2.162        |         |         |
| NRC        | GCA                 | 5  | 2.167          | 0.433        | 1.112   | 0.745   |
|            | SCA                 | 15 | 8.718          | 0.581        |         |         |
|            | Crosses             | 20 | 10.886         | 0.544        |         |         |
|            | Error               | 40 | 19.564         | 0.489        |         |         |
| NKR        | GCA                 | 5  | 470.348        | 94.069       | 1.843   | 7.764   |
|            | SCA                 | 15 | 181.74         | 12.116       |         |         |
|            | Crosses             | 20 | 652.088        | 32.604       |         |         |
|            | Error               | 40 | 707.584        | 17.689       |         |         |
| NGC        | GCA                 | 5  | 64582.993      | 12916.598    | 1.749   | 3.6169  |
|            | SCA                 | 15 | 53567.158      | 3571.143     |         |         |
|            | Crosses             | 20 | 118150.15      | 5907.507     |         |         |
|            | Error               | 40 | 135068.29      | 3376.707     |         |         |
| GYP        | GCA                 | 5  | 5175.866       | 1035.173     | 1.749   | 3.6167  |
|            | SCA                 | 15 | 4293.246       | 286.216      |         |         |
|            | Crosses             | 20 | 9469.112       | 473.455      |         |         |
|            | Error               | 40 | 10825.162      | 270.629      |         |         |

**Table 4.2.14. Estimation of GCA effects of the parents for different characters in maize**

| Parents        | DT     | DS     | DM       | PH     | CH       | CL     | CD     | NRC    | NKR    | NGC      | GYP      |
|----------------|--------|--------|----------|--------|----------|--------|--------|--------|--------|----------|----------|
| P <sub>1</sub> | 1.422  | 1.265  | -4.448** | 0.388  | -8.390** | -0.045 | 0.051  | 0.388  | -0.237 | 18.422** | -8.301** |
| P <sub>2</sub> | -0.743 | -0.134 | 2.659**  | 0.132  | -0.208   | -0.018 | 0.336  | 0.132  | -0.846 | 7.419**  | -7.018** |
| P <sub>3</sub> | -0.428 | -0.267 | 2.975**  | 0.173  | -1.030   | 0.248  | -0.271 | 0.173  | 1.045  | 19.055** | 0.358    |
| P <sub>4</sub> | -0.547 | -0.487 | -1.402   | -0.198 | -0.891   | -0.199 | 0.320  | -0.198 | -1.062 | -15.88** | 3.178**  |
| P <sub>5</sub> | 0.452  | 0.114  | -0.576   | -0.252 | 7.130**  | -0.059 | -0.373 | -0.252 | 0.789  | 2.703**  | 8.877**  |
| P <sub>6</sub> | -0.155 | -0.491 | 0.793    | -0.242 | 3.390**  | 0.074  | -0.063 | -0.242 | 0.311  | -2.880** | 2.906**  |

**Table 4.2.15. Specific combining ability (SCA) effect for days to tasseling in 6×6 diallel cross in maize**

| Parent         | P <sub>2</sub> | P <sub>3</sub> | P <sub>4</sub> | P <sub>5</sub> | P <sub>6</sub> |
|----------------|----------------|----------------|----------------|----------------|----------------|
| P <sub>1</sub> | 2.476**        | 1.6009         | -0.203         | 0.221          | -1.201         |
| P <sub>2</sub> |                | 0.0967         | 1.549          | 0.0538         | -0.3127        |
| P <sub>3</sub> |                |                | -0.569         | 0.9588         | 1.0355         |
| P <sub>4</sub> |                |                |                | 0.0780         | -0.0086        |
| P <sub>5</sub> |                |                |                |                | -0.230         |

**Table 4.2.16. Specific combining ability effect for days to silking in 6×6 diallel cross in maize**

| Parent         | P <sub>2</sub> | P <sub>3</sub> | P <sub>4</sub> | P <sub>5</sub> | P <sub>6</sub> |
|----------------|----------------|----------------|----------------|----------------|----------------|
| P <sub>1</sub> | 4.757**        | 0.6708         | 0.614          | -2.343**       | -0.384         |
| P <sub>2</sub> |                | 2.153**        | 0.927          | -0.254         | -0.678         |
| P <sub>3</sub> |                |                | -3.519**       | 0.708          | -3.427**       |
| P <sub>4</sub> |                |                |                | -3.154**       | -0.154         |
| P <sub>5</sub> |                |                |                |                | -2.146**       |

**Table 4.2.17. Specific combining ability effect for days to maturity in 6×6 diallel cross in maize**

| Parent         | P <sub>2</sub> | P <sub>3</sub> | P <sub>4</sub> | P <sub>5</sub> | P <sub>6</sub> |
|----------------|----------------|----------------|----------------|----------------|----------------|
| P <sub>1</sub> | 2.244**        | 3.178**        | 0.192          | -0.270         | 7.163**        |
| P <sub>2</sub> |                | 3.319**        | 1.391          | 0.981          | -2.694**       |
| P <sub>3</sub> |                |                | 3.185**        | -6.364**       | -4.813**       |
| P <sub>4</sub> |                |                |                | -1.039         | 1.730          |
| P <sub>5</sub> |                |                |                |                | 0.191          |

**Table 4.2.18. Specific combining ability effect for plant height in 6×6 diallel cross in maize**

| Parent         | P <sub>2</sub> | P <sub>3</sub> | P <sub>4</sub> | P <sub>5</sub> | P <sub>6</sub> |
|----------------|----------------|----------------|----------------|----------------|----------------|
| P <sub>1</sub> | 2.130**        | 4.121**        | -2.574**       | -7.139**       | -7.866**       |
| P <sub>2</sub> |                | 7.677**        | -7.871**       | -3.763**       | -6.110**       |
| P <sub>3</sub> |                |                | 5.929**        | -15.662**      | -0.155         |
| P <sub>4</sub> |                |                |                | -1.058         | 2.095**        |
| P <sub>5</sub> |                |                |                |                | 18.843**       |

**Table 4.2.19. Specific combining ability effect for ear height in 6×6 diallel crosses in maize**

| Parent         | P <sub>2</sub> | P <sub>3</sub> | P <sub>4</sub> | P <sub>5</sub> | P <sub>6</sub> |
|----------------|----------------|----------------|----------------|----------------|----------------|
| P <sub>1</sub> | -7.487**       | 6.7617**       | 7.332**        | -9.197*8       | -5.590**       |
| P <sub>2</sub> |                | 5.367**        | 6.191**        | -3.174*8       | -17.434**      |
| P <sub>3</sub> |                |                | 1.986          | -10.609**      | 1.394          |
| P <sub>4</sub> |                |                |                | -10.489**      | 2.534**        |
| P <sub>5</sub> |                |                |                |                | 10.205**       |

**Table 4.2.20. Specific combining ability effect for cob length in 6×6 diallel crosses in maize**

| Parent         | P <sub>2</sub> | P <sub>3</sub> | P <sub>4</sub> | P <sub>5</sub> | P <sub>6</sub> |
|----------------|----------------|----------------|----------------|----------------|----------------|
| P <sub>1</sub> | -0.065         | 0.171          | 0.752          | 2.202**        | -0.822         |
| P <sub>2</sub> |                | -3.168**       | 4.022**        | 0.602          | 3.021**        |
| P <sub>3</sub> |                |                | 0.325          | -3.367**       | -2.215**       |
| P <sub>4</sub> |                |                |                | -4.183**       | -0.1613        |
| P <sub>5</sub> |                |                |                |                | -0.304         |

**Table 4.2.21. Specific combining ability effect for cob diameter in 6×6 diallel crosses in maize**

| Parent         | P <sub>2</sub> | P <sub>3</sub> | P <sub>4</sub> | P <sub>5</sub> | P <sub>6</sub> |
|----------------|----------------|----------------|----------------|----------------|----------------|
| P <sub>1</sub> | -0.102         | 2.202**        | -0.248         | -3.147**       | 0.741          |
| P <sub>2</sub> |                | 1.173          | 0.296          | 0.630          | -0.103         |
| P <sub>3</sub> |                |                | -2.949**       | 4.361**        | 0.623          |
| P <sub>4</sub> |                |                |                | 0.250          | 0.373          |
| P <sub>5</sub> |                |                |                |                | -0.516         |

**Table 4.2.22. Specific combining ability effect for number of rows/cob in 6×6 diallel cross in maize**

| Parent         | P <sub>2</sub> | P <sub>3</sub> | P <sub>4</sub> | P <sub>5</sub> | P <sub>6</sub> |
|----------------|----------------|----------------|----------------|----------------|----------------|
| P <sub>1</sub> | 7.603**        | 0.253          | 2.248**        | 3.398**        | -4.790**       |
| P <sub>2</sub> |                | -2.179**       | -0.457         | -0.617         | 0.093          |
| P <sub>3</sub> |                |                | -0.480         | -5.041**       | -5.247**       |
| P <sub>4</sub> |                |                |                | 0.231          | 0.208          |
| P <sub>5</sub> |                |                |                |                | 0.451          |

**Table 4.2.23. Specific combining ability effect for number of kernels/row in 6×6 diallel cross in maize**

| Parent         | P <sub>2</sub> | P <sub>3</sub> | P <sub>4</sub> | P <sub>5</sub> | P <sub>6</sub> |
|----------------|----------------|----------------|----------------|----------------|----------------|
| P <sub>1</sub> | 3.034**        | -1.253         | -0.324         | 4.1236**       | 1.300          |
| P <sub>2</sub> |                | -0.058         | -3.579**       | -4.298**       | -1.121         |
| P <sub>3</sub> |                |                | -1.101         | -1.635         | -2.392**       |
| P <sub>4</sub> |                |                |                | 3.126**        | 4.253**        |
| P <sub>5</sub> |                |                |                |                | 2.801**        |

**Table 4.2.24. Specific combining ability effect for number of grains/cob in 6×6 diallel cross in maize**

| Parent         | P <sub>2</sub> | P <sub>3</sub> | P <sub>4</sub> | P <sub>5</sub> | P <sub>6</sub> |
|----------------|----------------|----------------|----------------|----------------|----------------|
| P <sub>1</sub> | 56.096**       | -11.127**      | 3.251**        | 16.295**       | -5.745**       |
| P <sub>2</sub> |                | -6.360**       | -32.514**      | -71.930**      | -13.545**      |
| P <sub>3</sub> |                |                | -26.795**      | -23.411**      | -25.542**      |
| P <sub>4</sub> |                |                |                | 24.238**       | 7.699**        |
| P <sub>5</sub> |                |                |                |                | 25.821**       |

**Table 4.2.25. Specific combining ability effect for grain yield/plant in 6×6 diallel cross in maize**

| Parent         | P <sub>2</sub> | P <sub>3</sub> | P <sub>4</sub> | P <sub>5</sub> | P <sub>6</sub> |
|----------------|----------------|----------------|----------------|----------------|----------------|
| P <sub>1</sub> | 15.880**       | -3.150**       | 0.920          | 4.613**        | -1.626         |
| P <sub>2</sub> |                | -1.800         | -9.205**       | 20.363**       | 3.834**        |
| P <sub>3</sub> |                |                | -7.585**       | -6.627**       | -7.231**       |
| P <sub>4</sub> |                |                |                | 6.861**        | 2.181**        |
| P <sub>5</sub> |                |                |                |                | 7.309**        |

### 4.2.3 HETEROSIS STUDY

#### Estimation of Mid-parent and Better-parent

The estimation of percent heterosis observed in  $F_1$  generation over mid-parent and better-parent for different characters are presented in Tables 4.2.26- 4.2.31. Heterosis over mid-parent for different crosses was recorded non-significant in some crosses viz.,  $P_2 \times P_6$ ,  $P_3 \times P_4$ ,  $P_4 \times P_5$  and  $P_4 \times P_6$  for days to tasseling. The highest percent of heterosis over mid-parent was recorded to be 4.165 in  $P_1 \times P_2$  for this character (Table 4.2.26). Out of the cross combinations, eleven crosses showed significant heterosis over better-parent for days to tasseling. Most of the crosses showed highly significant positive heterosis both over mid-parent and better-parent for days to silking (Table 4.2.26.).

Most of the crosses showed significant positive heterosis over mid-parent and better-parent for days to maturity. The highest heterosis with 10.185 was recorded in  $P_1 \times P_6$  over mid-parent and that of 10.663 was recorded in  $P_3 \times P_5$  over better-parent.

For plant height nine crosses showed negative but significant heterosis and rest of them showed positive significant heterosis over mid-parent. All the crosses except only one cross  $P_3 \times P_6$  showed positive significant heterosis over better-parent for this character.

Heterosis over mid-parent for different crosses was recorded significant but negative heterosis in some crosses viz.,  $P_1 \times P_2$ ,  $P_1 \times P_5$ ,  $P_1 \times P_6$ ,  $P_2 \times P_5$ ,  $P_2 \times P_6$ ,  $P_3 \times P_5$ ,  $P_4 \times P_5$ ,  $P_4 \times P_6$  and  $P_5 \times P_6$  for ear height. The highest percent of heterosis over mid-parent was recorded to be 9.060 in  $P_2 \times P_3$  for this character (Table 4.2.28). All the cross combinations except  $P_2 \times P_4$  showed significant heterosis over better-parent and the highest percent positive heterosis 45.863 was recorded in  $P_1 \times P_5$ . Regarding mid-parent and better-parent heterosis, most of the crosses for cob length exhibited non-significant results. From Table 4.2.28, it is observed

that three crosses showed significant positive heterosis over mid parent and better parent but rest of the crosses showed significant negative heterosis.

Two crosses showed non-significant negative heterosis in cob diameter. Out of fifteen cross combinations, five  $F_1$  showed positive significant heterosis over mid-parent for number of rows/cob. Rest of them was found to be non-significant. Four non-significant values of -2.506, -3.531, -2.473 and -1.874 were estimated in  $P_1 \times P_5$ ,  $P_1 \times P_6$ ,  $P_2 \times P_4$  and  $P_3 \times P_5$  over mid-parent heterosis for number of kernels/row.

The highest significant positive heterosis percent over mid-parent of 3.043 was recorded in  $P_4 \times P_6$ . Different crosses for this character exhibited non-significant to significant, negative to positive heterosis over better-parent. The highest percent better-parent heterosis of 7.210 was recorded in  $P_4 \times P_5$ . All the  $F_1$ s showed significant negative heterosis over mid-parent though six positive significant estimations were found in  $P_1 \times P_2$ ,  $P_1 \times P_3$ ,  $P_1 \times P_4$ ,  $P_3 \times P_4$ ,  $P_4 \times P_6$  and  $P_5 \times P_6$ , respectively for number of grains/cob (Table 4.2.30). The highest significant positive heterosis percent over mid-parent of 59.096 was recorded in  $P_1 \times P_2$  while all non-significant heterosis over better-parent were estimated for the above character. From Table 4.2.31, it is evident that most of the crosses over mid parent and better parent showed significant negative but rest of them showed both significantly positive and negative heterotic effect.

A wide range of variation from 1.732 to 15.209% over mid parent and -40.363 to 4.936% over better parent was observed. The cross  $P_1 \times P_2$  showed the highest and significantly positive heterosis over mid parent (15.209%) and better parent (4.936%).



**Table 4.2.26. Heterotic effect in F<sub>1</sub> generation over mid parent and better parent for days to tasseling and days to silking**

| Crosses                        | Days to tasseling |         |        | Days to silking |         |         |
|--------------------------------|-------------------|---------|--------|-----------------|---------|---------|
|                                | Mean              | MP      | BP     | Mean            | MP      | BP      |
| P <sub>1</sub> ×P <sub>2</sub> | 97.840            | 4.165** | 1.756* | 100.110         | 3.485** | 2.360** |
| P <sub>1</sub> ×P <sub>3</sub> | 97.280            | 3.105** | 1.196* | 98.890          | 2.282** | 1.140*  |
| P <sub>1</sub> ×P <sub>4</sub> | 95.356            | 0.732*  | 0.726* | 98.613          | 1.438*  | 0.863*  |
| P <sub>1</sub> ×P <sub>5</sub> | 96.556            | 0.765*  | 0.473  | 99.943          | 2.718** | 2.193** |
| P <sub>1</sub> ×P <sub>6</sub> | 94.750            | -0.658* | 1.333* | 97.610          | 0.526*  | 0.140   |
| P <sub>2</sub> ×P <sub>3</sub> | 93.610            | 1.843*  | 1.343* | 97.973          | 2.490** | 2.473** |
| P <sub>2</sub> ×P <sub>4</sub> | 94.943            | 2.726** | 1.776* | 97.526          | 1.476*  | 0.926*  |
| P <sub>2</sub> ×P <sub>5</sub> | 94.223            | 0.840*  | 1.276* | 96.946          | 0.846*  | 0.246   |
| P <sub>2</sub> ×P <sub>6</sub> | 93.473            | 0.473   | 1.260* | 95.916          | -0.041  | 0.500*  |
| P <sub>3</sub> ×P <sub>4</sub> | 93.140            | 0.423   | 0.026  | 94.946          | -1.086  | 1.653*  |
| P <sub>3</sub> ×P <sub>5</sub> | 95.443            | 1.560*  | 0.056  | 97.77           | 1.693*  | 1.076*  |
| P <sub>3</sub> ×P <sub>6</sub> | 95.136            | 1.636*  | 0.403  | 97.890          | 1.948*  | 1.473*  |
| P <sub>4</sub> ×P <sub>5</sub> | 94.443            | 0.110   | 1.056* | 96.693          | 0.043   | 0.006   |
| P <sub>4</sub> ×P <sub>6</sub> | 93.973            | 0.023   | 0.760* | 95.666          | -0.841* | 0.933*  |
| P <sub>5</sub> ×P <sub>6</sub> | 94.526            | -0.590* | 0.973* | 96.696          | 0.138   | 0.003   |

**Table 4.2.27. Heterotic effect in F<sub>1</sub> generation over mid parent and better parent for days to maturity and plant height (cm)**

| Crosses                        | Days to maturity |          |          | Plant height |           |          |
|--------------------------------|------------------|----------|----------|--------------|-----------|----------|
|                                | Mean             | MP       | BP       | Mean         | MP        | BP       |
| P <sub>1</sub> ×P <sub>2</sub> | 146.223          | 3.681**  | 5.243**  | 136.246      | -2.186**  | 4.253**  |
| P <sub>1</sub> ×P <sub>3</sub> | 149.473          | 5.431**  | 4.993**  | 135.166      | 1.766*    | 1.200*   |
| P <sub>1</sub> ×P <sub>4</sub> | 142.110          | 4.185**  | 0.123    | 129.406      | -6.276**  | 6.960**  |
| P <sub>1</sub> ×P <sub>5</sub> | 144.473          | 0.731    | 7.393**  | 129.800      | -11.667** | 16.775** |
| P <sub>1</sub> ×P <sub>6</sub> | 151.276          | 10.185** | 2.710**  | 127.153      | -8.996**  | 9.213**  |
| P <sub>2</sub> ×P <sub>3</sub> | 156.723          | 3.756**  | 2.256**  | 142.134      | 6.670**   | 1.635**  |
| P <sub>2</sub> ×P <sub>4</sub> | 152.416          | 3.566**  | 1.050*   | 127.523      | -10.226** | 12.973** |
| P <sub>2</sub> ×P <sub>5</sub> | 150.833          | 0.166    | 0.633    | 138.590      | -4.943**  | 7.976**  |
| P <sub>2</sub> ×P <sub>6</sub> | 148.526          | 3.490**  | 2.940**  | 132.323      | -5.893**  | 8.176**  |
| P <sub>3</sub> ×P <sub>4</sub> | 153.526          | 4.176**  | 1.940*   | 138.253      | 5.536**   | 3.253**  |
| P <sub>3</sub> ×P <sub>5</sub> | 143.803          | -8.363** | 10.663** | 121.620      | -16.880** | 24.946** |
| P <sub>3</sub> ×P <sub>6</sub> | 146.723          | -4.793** | 7.743**  | 135.206      | 2.023**   | 0.726    |
| P <sub>4</sub> ×P <sub>5</sub> | 144.750          | -1.301   | 5.116**  | 137.160      | -3.623**  | 9.406    |
| P <sub>4</sub> ×P <sub>6</sub> | 149.890          | 3.490**  | 0.323    | 138.393      | 2.926**   | 2.460**  |
| P <sub>5</sub> ×P <sub>6</sub> | 148.176          | -1.040** | 1.690**  | 160.100      | 18.850**  | 13.532** |

**Table 4.2.28. Heterotic effect in F<sub>1</sub> generation over mid and better parent for ear height (cm) and cob length (cm)**

| Crosses                        | Ear height |           |          | Cob length |         |         |
|--------------------------------|------------|-----------|----------|------------|---------|---------|
|                                | Mean       | MP        | BP       | Mean       | MP      | BP      |
| P <sub>1</sub> ×P <sub>2</sub> | 56.160     | -9.873**  | 16.740** | 13.587     | -0.497  | -1.247* |
| P <sub>1</sub> ×P <sub>3</sub> | 59.013     | -0.820    | 1.486**  | 14.536     | 0.687   | -0.297  |
| P <sub>1</sub> ×P <sub>4</sub> | 61.993     | -0.790    | 4.406**  | 16.700     | 3.600** | 1.867*  |
| P <sub>1</sub> ×P <sub>5</sub> | 54.936     | -25.046** | 45.863** | 16.263     | 2.555*  | 1.430*  |
| P <sub>1</sub> ×P <sub>6</sub> | 59.466     | -10.250** | 20.800** | 11.883     | -2.100* | -2.950* |
| P <sub>2</sub> ×P <sub>3</sub> | 75.760     | 9.060**   | 2.860**  | 11.667     | -1.433* | -1.667* |
| P <sub>2</sub> ×P <sub>4</sub> | 72.016     | 2.366**   | 0.883    | 10.950     | -1.400* | -2.383* |
| P <sub>2</sub> ×P <sub>5</sub> | 63.653     | -23.196** | 37.146** | 13.250     | 0.233   | -0.083  |
| P <sub>2</sub> ×P <sub>6</sub> | 61.970     | -14.613** | 18.296** | 13.000     | 2.343*  | -0.333  |
| P <sub>3</sub> ×P <sub>4</sub> | 71.900     | 8.450**   | 5.50**   | 14.460     | -1.375* | 1.593*  |
| P <sub>3</sub> ×P <sub>5</sub> | 71.90      | -8.750**  | 28.90**  | 13.00      | -1.625* | -1.767* |
| P <sub>3</sub> ×P <sub>6</sub> | 69.213     | -1.170    | 11.053** | 12.200     | -0.800* | -0.933* |
| P <sub>4</sub> ×P <sub>5</sub> | 60.103     | -23.496** | 40.696** | 11.600     | -1.375* | -1.983* |
| P <sub>4</sub> ×P <sub>6</sub> | 71.086     | -2.246*   | 9.180**  | 12.700     | 0.450   | -0.433  |
| P <sub>5</sub> ×P <sub>6</sub> | 81.883     | -8.650**  | 18.916** | 13.830     | -0.075  | -0.350  |

**Table 4.2.29. Heterotic effect in F<sub>1</sub> generation over mid and better parent for cob diameter (cm) and number of rows/cob**

| Crosses                        | Cob diameter |         |         | Number of rows/cob |         |        |
|--------------------------------|--------------|---------|---------|--------------------|---------|--------|
|                                | Mean         | MP      | BP      | Mean               | MP      | BP     |
| P <sub>1</sub> ×P <sub>2</sub> | 13.956       | 0.734*  | 0.617*  | 14.666             | 1.400*  | 1.333* |
| P <sub>1</sub> ×P <sub>3</sub> | 15.073       | 3.167*  | 2.234*  | 14.386             | 0.970*  | 0.753* |
| P <sub>1</sub> ×P <sub>4</sub> | 14.343       | -0.457  | -1.707* | 13.776             | 0.910*  | 0.576* |
| P <sub>1</sub> ×P <sub>5</sub> | 15.586       | -0.641* | -0.950* | 13.723             | 0.873*  | 0.523* |
| P <sub>1</sub> ×P <sub>6</sub> | 14.566       | 1.383*  | 1.283*  | 12.712             | -0.443  | 0.476  |
| P <sub>2</sub> ×P <sub>3</sub> | 12.633       | 2.250*  | 1.200*  | 13.610             | 0.126   | 0.023  |
| P <sub>2</sub> ×P <sub>4</sub> | 13.354       | 0.250   | -0.933* | 13.220             | 0.286   | 0.113  |
| P <sub>2</sub> ×P <sub>5</sub> | 14.450       | 1.248*  | 0.823*  | 12.470             | -0.446  | 0.863* |
| P <sub>2</sub> ×P <sub>6</sub> | 15.143       | 0.650*  | 0.633*  | 13.001             | -0.235  | 0.331  |
| P <sub>3</sub> ×P <sub>4</sub> | 13.743       | -2.916* | -5.100* | 13.026             | -0.056  | 0.606* |
| P <sub>3</sub> ×P <sub>5</sub> | 12.570       | 2.108*  | 1.483*  | 12.886             | -0.180  | 0.746* |
| P <sub>3</sub> ×P <sub>6</sub> | 13.253       | 1.507*  | 0.473   | 12.776             | -0.606* | 0.856* |
| P <sub>4</sub> ×P <sub>5</sub> | 15.056       | -0.175  | -1.733* | 13.250             | 0.733*  | 0.716* |
| P <sub>4</sub> ×P <sub>6</sub> | 14.513       | 0.083   | -1.066* | 12.640             | -0.193  | 0.493  |
| P <sub>5</sub> ×P <sub>6</sub> | 12.393       | -0.0917 | -0.500* | 13.223             | 0.406   | 0.0901 |

**Table 4.2.30. Heterotic effect in F<sub>1</sub> generation over mid and better parent for number of kernels/row and number of grains/cob**

| Crosses                        | Number of kernels/row |          |          | Number of grains/cob |           |            |
|--------------------------------|-----------------------|----------|----------|----------------------|-----------|------------|
|                                | Mean                  | MP       | BP       | Mean                 | MP        | BP         |
| P <sub>1</sub> ×P <sub>2</sub> | 28.780                | 1.296*   | 0.813*   | 383.670              | 53.725**  | 17.436**   |
| P <sub>1</sub> ×P <sub>3</sub> | 31.000                | 2.183**  | 1.333*   | 342.503              | -19.745** | -88.336**  |
| P <sub>1</sub> ×P <sub>4</sub> | 28.166                | 1.070    | 0.200    | 366.843              | 11.913**  | -49.360**  |
| P <sub>1</sub> ×P <sub>5</sub> | 27.360                | -2.506** | 4.406**  | 400.020              | 23.741**  | 58.880**   |
| P <sub>1</sub> ×P <sub>6</sub> | 24.583                | -3.531** | -3.683** | 356.886              | 6.118**   | -50.993**  |
| P <sub>2</sub> ×P <sub>3</sub> | 27.583                | -0.750*  | 2.083**  | 351.803              | -46.733** | -79.037**  |
| P <sub>2</sub> ×P <sub>4</sub> | 24.140                | -2.473** | 2.860**  | 335.610              | -55.608** | -80.593**  |
| P <sub>2</sub> ×P <sub>5</sub> | 28.556                | -0.826*  | 3.210**  | 316.326              | -96.240** | -142.573** |
| P <sub>2</sub> ×P <sub>6</sub> | 27.890                | 0.256    | 0.376    | 353.620              | -33.437** | -54.260**  |
| P <sub>3</sub> ×P <sub>4</sub> | 29.610                | 1.663*   | 0.056    | 367.387              | -56.135** | -63.453**  |
| P <sub>3</sub> ×P <sub>5</sub> | 28.831                | -1.874*  | -2.912** | 390.903              | -53.967** | -67.997**  |
| P <sub>3</sub> ×P <sub>6</sub> | 29.723                | 0.756*   | 0.059    | 367.680              | -51.680** | -63.160**  |
| P <sub>4</sub> ×P <sub>5</sub> | 24.556                | -4.440** | 7.21**   | 448.883              | 10.962**  | -10.387**  |
| P <sub>4</sub> ×P <sub>6</sub> | 30.290                | 3.043**  | 2.023**  | 410.883              | -1.158    | -5.320*    |
| P <sub>5</sub> ×P <sub>6</sub> | 31.193                | 1.176*   | 0.573    | 449.137              | 15.747**  | -9.763**   |

**Table 4.2.31. Heterotic effect in F<sub>1</sub> generation over mid and better parent for grain yield/plant (g)**

| Crosses                        | Grain yield/plant |           |           |
|--------------------------------|-------------------|-----------|-----------|
|                                | Mean              | MP        | BP        |
| P <sub>1</sub> ×P <sub>2</sub> | 108.617           | 15.209**  | 4.936**   |
| P <sub>1</sub> ×P <sub>3</sub> | 96.962            | -5.590**  | -25.008** |
| P <sub>1</sub> ×P <sub>4</sub> | 103.853           | 3.372**   | -13.973** |
| P <sub>1</sub> ×P <sub>5</sub> | 113.245           | 6.721**   | -16.669** |
| P <sub>1</sub> ×P <sub>6</sub> | 101.034           | 1.732*    | -14.435** |
| P <sub>2</sub> ×P <sub>3</sub> | 99.595            | -13.23**  | -22.375** |
| P <sub>2</sub> ×P <sub>4</sub> | 95.011            | -15.742** | -15.742** |
| P <sub>2</sub> ×P <sub>5</sub> | 89.551            | -27.246** | -40.363** |
| P <sub>2</sub> ×P <sub>6</sub> | 100.109           | -9.466**  | -15.360** |
| P <sub>3</sub> ×P <sub>4</sub> | -15.360           | -15.891** | -17.963** |
| P <sub>3</sub> ×P <sub>5</sub> | 110.664           | -15.278** | -19.250** |
| P <sub>3</sub> ×P <sub>6</sub> | 104.089           | -14.630** | -17.881** |
| P <sub>4</sub> ×P <sub>5</sub> | 126.973           | 3.103**   | -2.941**  |
| P <sub>4</sub> ×P <sub>6</sub> | 116.321           | -0.327    | -1.505*   |
| P <sub>5</sub> ×P <sub>6</sub> | 127.150           | 4.457**   | -2.764**  |

## DISCUSSION

### GENETIC VARIABILITY

The magnitude of variability present in a crop species is of utmost importance as it provides the basis for effective selection. In the present study, genetic variability among twenty five inbred lines of maize was assessed for grain yield and important yield components. The results (Table 4.1.10) revealed that GCV was less than its corresponding estimates of PCV for all the traits which indicated significant role of environment in the expression of these traits. The high values of genotypic coefficient of variation for cob height, days to silking, number of kernels/cob and grain yield indicated high degree of genetic variability for these characters. Similar results have also been obtained by Alam (2009). On the contrary, Satyanarayana and Saikumar (1995) recorded low genotypic coefficient of variation for grain yield. In crop improvement, only the genetic component of variation is important since only it is transmitted to the next generation. Although high heritability estimates have been found to be helpful in making selection of superior lines.

On the basis of phenotypic performance, Wannows *et al.* (2010) suggested that heritability estimates along with genetic gain were more useful in predicting the effect for selecting the best individual. Among the quantitative characters studied, the degree of heritability was found to be different.

Heritability estimate in broad sense was highest for days to silking followed by days to maturity, cob length and days to tasseling. Results showed that high heritability estimates were detected for days to silking, cob height, number of kernels/row, days to tasseling and yield, emphasizing that the additive genetic variation was the major component of genetic variation in the inheritance of these traits and the effectiveness of selection in the early segregating generations of the studied hybrids for improving these traits. Higher genetic advance for number of grains/cob, days to silking, days to tasseling, plant height and grain yield depicts additive gene effects.

High heritability estimates for plant height, cob height, number of rows/cob and number of kernels/row were also reported by Yasien (2000) and Abd El-Sattar (2003). Similar results have been reported in maize by Wannows *et al.* (2010) and Alam (2009).

High genotypic coefficient of variation and high to moderate heritability coupled with high genetic advance as percentage of mean for cob height, number of grains/cob, days to silking, cob diameter and grain yield indicating that these characters might be transmitted to the progenies. Therefore, selection of parents based on the superior phenotypes for these characters may be effective in hybridization program to develop high performing maize hybrids.

#### **CORRELATION COEFFICIENT**

The correlation coefficient between grain yield/plant and its component characters and between the various components themselves were estimated at the genotypic and phenotypic levels. The study reveals that in most of the cases, the values of  $r_g$  were higher than the corresponding  $r_p$  indicating less pronounced environmental effect. Lower  $r_p$  than  $r_g$  indicates that both environmental and genotypic correlations in those cases act in the same direction and finally maximize their expression at phenotypic level.

In this study, the traits studied were positively correlated with grain yield (Tables 4.1.12 and 4.1.13). The highest significant positive correlation with grain yield was shown by number of grains/cob followed by number of kernels/row and days to maturity. Similar results have been reported in maize by Swarnalatha and Mohammad (2001), Mohan *et al.* (2002), Abd EL-Aty and Katta (2002), Mohammadia *et al.* (2003), Ahmed (2004), Sadek *et al.* (2006) and Aydin *et al.* (2007).



Cob length showed significant and positive genotypic correlation with cob height; on the other hand, it was significantly and negatively correlated with days to maturity. This result agreed with those mentioned by Soliman *et al.* (1999), Yasien (2000), Mohammadia *et al.* (2003) and Sadek *et al.* (2006).

Cob diameter had significant and positive correlations with days to maturity while it showed significant and negative correlations with cob height. Number of rows/cob showed significant and negative correlations with cob height. Such results are in harmony with those obtained by Salami *et al.* (2007), Yasien (2000), Amin *et al.* (2003) and Mohammadia *et al.* (2003).

As yield (grain weight/plant) is the ultimate goal, the positive association of these characters will help for selecting best genotype. Similar results have also been reported by Mohan *et al.* (2002), Alam (2009) and Rafiq *et al.* (2010). In general, the existence of positive associations in the present study among the grain yield and cob length, number of kernels/row and number of grains/cob suggests that an increment of production may be achieved upon improving either one or more of these traits.

### **PATH COEFFICIENT**

The analysis of path coefficient has (Tables 4.1.14 and 4.1.15) been made to identify the important yield contributors by estimating the direct effects of the contributing characters to yield and separating the direct from the indirect effects through other related characters by partitioning the correlation coefficient and finding out the relative importance of different characters as selection criteria. The estimates of direct and indirect effects of the eight yield related characters *viz.* days to maturity, plant height, ear height, cob length, cob diameter, number of rows/cob, number of kernels/row, number of grains/cob on grain yield are presented in Table 4.1.14.

The highest direct effect on grain yield was exhibited by number of rows/cob followed by days to maturity, number of kernels/row and cob length. Number

of grains/cob had highest indirect effect on grain yield through number of kernels/row followed by days to maturity through number of kernels/row and number of rows/cob through number of grains/cob. These results are agreed with some researchers such as Swarnalatha and Mohammad (2001), Mohsan *et al.* (2002), Viola *et al.* (2003 and Alam (2009).

At phenotypic level, highly significant undesirable direct effect on grain yield was found for number of grains/cob which is in agreement with the results of Alvi *et al.* (2003), Sofi and Rather (2007) and Alam (2009); but contrary to the results of path analysis which in their research found Akbar *et al.* (2008) and Najeeb *et al.* (2009).

Path coefficient values based on phenotypic correlation revealed that most of traits except cob length had direct positive effect towards grain yield also having positive correlation with grain yield. Therefore, proper attention should be given to the above characters for the improvement of grain yield. These results are in agreement with the results of many authors (Alvi *et al.*, 2003; Sumathi *et al.*, 2005; Sofi and Rather, 2007; Najeeb *et al.*, 2009 and Alam, 2009).

In the present study, path coefficient analysis suggests that during selection more emphasis should be given on cob length, number of rows/cob, number of kernels/row and number of grains/cob. Since these characters, had high correlation and high direct effect on grain yield. Nemati *et al.* (2009) reported that ear weight had direct effect on grain yield.

## SELECTION INDEX

Different selection indices were formulated using different combinations of grain yield and yield contributing characters and their expected genetic gain were estimated (Table 4.1.16). It was observed that among all the selection indices, the index based on cob height + number of rows/cob + number of

kernels/row + grain yield/plant had the maximum genetic gain. Among the others, the indices based on cob length+ number of rows/cob + number of kernels/row over straight selection for grain yield alone.

Therefore, improvement of grain yield through these selection indices is suggested. There are scarcity papers of selection index in maize. More or less similar results were also reported by Paul *et al.* (1978) in mustard. Similar opinions were also reported by Singh *et al.* (1999) in maize and Mondal (2003) in potato.

Bergele *et al.* (2002) suggested that the number of spikes per plant, grains per spike and harvest index must be given preference in selection along with optimum plant height and days to flowering to select the superior wheat genotypes. Similar opinions were also reported by Ferdous *et al.* (2010) in wheat.

Shiv *et al.* (2008) suggested that number of tillers per plant, numbers of spikelets per ear, number of grains per ear, grain weight per ear, 100-grains weight and biological yield could form effective selection indices for selection of high yielding genotypes of wheat.

## GENETIC DIVERGENCE

Clustering pattern of inbred lines showed considerable genetic diversity among themselves by occupying five different clusters. Cluster analysis revealed that the 25 maize inbred lines could be grouped into 5 different clusters of which maximum number of inbreeds (each containing 6 inbreeds) was included in cluster I, II and IV. Cluster III had only three lines and that was the lowest. Similar opinions were also reported by Singh *et al.* (2005) in maize and Alam (2009) in maize inbred lines.

The maximum inter-cluster divergence (Table 4.1.17) was observed between the clusters I and II and it was minimum inter-cluster divergence between clusters III and IV. The maximum intra-cluster distance was observed in cluster III and minimum in cluster V. The crosses involving parents from most divergent clusters are expected to manifest maximum heterosis and generate wide variability in genetic architecture.

Intra-cluster distance was much lower than the inter-cluster one, suggesting, heterogeneous and homogeneous nature between and within groups, respectively. This was further supported by an appreciable variation observed for cluster means (Table 4.1.18). Similar results were reported by Singh *et al.* (2005), Liu YuAi *et al.* (2006) in maize. Another study was carried out by Chen FaBo *et al.* (2007) who reported that 186 maize genotypes could be classified into ten clusters.

Mean values of days to maturity, plant height, and ear height and cob length were highest in cluster II and cob diameter and number of kernels/row in III and grain yield in cluster V.

A wide range of variations for several characters among the multi genotypic cluster was observed. However, the difference was clear for plant height, cob length, cob diameter, number of kernels/row and grain yield, which contributed largely to the total divergence. Similar results have also been reported by Singh *et al.* (2005) and Chen FaBo *et al.* (2007). Hence, for the improvement of different characters *viz.* cob length, cob diameter, number of kernels/row, grain number and grain yield under the present study, inbred lines should be selected from clusters II, III and V.

The principal component analysis revealed that in major vector I the important characters responsible for genetic divergence in the major axis of differentiation were days to tasseling, days to silking, plant height, ear height and cob length, cob diameter, number of kernels/cob and grain yield/plant

(Table 4.1.19). In vector II, which was the second axis of differentiation, the characters like days to tasseling, plant height, cob diameter, number of rows/cob, number of kernels/cob and grain yield were important.

The role of plant height, days to tasseling and silking, cob length and diameter, number of kernels/cob and grain yield/plant for both the vectors was positive across two axis which is the indication of the important components of genetic divergence in this material. Similar results have been reported in maize by Singh *et al.* (2005), Chen FaBo *et al.* (2007) and Azad *et al.* (2012).

Clustering  $D^2$ -statistics is useful in this matter. The inbred lines grouped together are less divergent than the ones which fall into different clusters. In selecting lines from the already chosen groups, other important characteristics like disease resistance, earliness, quality or even performance of particular character should also be considered.

#### **DIALLEL AND COMBINING ABILITY**

The combining ability of inbred lines is the ultimate factor determining future usefulness of the lines for hybrid development. In the present study, the combining ability of the inbred was assessed for grain yield and important yield contributing characters in a six inbreds diallel cross.

The results of this study showed that the expression of all these characters studied was found to be controlled by both additive and non-additive genes. Griffing analysis of variances for diallel crosses showed significant GCA and SCA mean squares for most of the traits (Table 4.2.14) which is an indication of the importance of both additive and non-additive genetic effects. This result is in agreement with the results of Ahmed and Salem (2003), Glover *et al.* (2005), Uddin *et al.* (2008), Alam (2009) and Afshar and Bahram (2012). Parents P<sub>5</sub> and P<sub>4</sub> exhibited significant positive GCA effects for number of grains/cob and grain yield, but parent P<sub>2</sub> showed negative GCA effect. Therefore, parent P<sub>5</sub> could be a donor parent for yield and earliness in

hybridization programme. The parent P<sub>1</sub> showed significant negative GCA effect for days to maturity and cob height.

Gichuru *et al.* (2011) evaluated different maize inbred lines for days to silking and introduced VHCY with the highest significantly negative (-3.194) GCA effect as the best combiners for earliness.

Roy *et al.* (1998), Debnath *et al.* (1988), Zelleke (2000), Choudhary *et al.* (2000), Desai and Singh (2001), Dubey *et al.* (2001), Hussain *et al.* (2003) and Uddin *et al.* (2008) found two and one inbred lines of maize, respectively, as well as a good general combiner of earliness in two separate experiments. Thus, the inbred lines which exhibited good general combining ability for at least one character can be used for development of early maturity and high grain yield. So, these three parents could be used extensively in hybrid breeding program with a view to increasing the yield level. Similar results have been reported in maize by earlier workers (Beck *et al.*, 1990; Das and Islam, 1993; Odongo and Bockholt, 1995; Spaner *et al.*, 1996; Preciado *et al.*, 1997; Dahlan *et al.*, 1997; San-vicente *et al.*, 1998; Zhao, 1999; Lemos *et al.*, 1999; Desai and Singh, 2001; Uddin *et al.*, 2008; Alam *et al.*, 2008 and Alam, 2009). However, additive genetic effects were preponderance for the expression of days to silking and days to maturity, plant height, ear length and ear girth, number of kernels/ear and 1000-kernel weight.

More importance of additive gene action for these attributes has been reported by Crossa *et al.* (1990), Mahajan and Khehra (1991), Pal and Prodhan (1994), Das and Islam (1994), Altinbas (1995), Odongo and Bockhot (1995), Tulu and Ramachandrappa (1998), Roy *et al.* (1998), Paul and Debnath (1999) and Zelleke (2000).

On the contrary, predominance of non-additive type of gene action for these characters has also been reported earlier (Alika, 1994; El-Hosary *et al.*, 1994; Dehghanpour *et al.*, 1996; Singh and Singh, 1998 and Suneetha *et al.*, 2000). The results of present study elucidated that for grain yield, number of rows/cob,

and cob diameter, non-additive gene action was more important in controlling their expression. Similar gene effect for these characters has been reported by El-Hosary *et al.* (1994), Pal and Prodhan (1994), Mostafa *et al.* (1996), Kim and Ajla (1996), Dehghanpour *et al.* (1996), Joshi *et al.* (1998), Roy *et al.* (1998), San-Vicente *et al.* (1998), Singh and Singh (1998), Ramech *et al.* (2000), Zelleke (2000) and Alam (2009).

However, the results differed from the findings of earlier reports (Das and Islam, 1994; Ferrao *et al.*, 1994; Szatmari, 1996; Tulu and Ramachandrappa, 1998; Mathur *et al.*, 1998 and Choukan, 1999) where additive gene effects were of major contributor for these characters.

For days to maturity and silking, additive gene action was important which is in conformity with the works of Pal and Prodhan (1994) and Zelleke (2000) as they showed a greater influence of additive component of gene action in the expression of maturity. In general, the crosses showing significant specific combining ability (SCA) effect for different characters also possess high mean performance.

In this study, P<sub>1</sub>xP<sub>2</sub>, P<sub>2</sub>xP<sub>5</sub>, P<sub>4</sub>xP<sub>5</sub> and P<sub>5</sub>xP<sub>6</sub> showed positively significant cross combinations were the superior specific combiner for grain yield and other yield components, indicating that most probably the inbred lines involved in producing each one of these crosses belongs to the different heterotic pattern.

It was found that in most of the crosses having the highest mean performance for different characters showed the best or at least good specific combiner in respect of these characters.

Thus, the present results indicate a relationship between mean performance of crosses and SCA effects. However, a few crosses appeared to have high mean value but non-significant SCA effects and vice-versa. Significant positive SCA effect was observed in P<sub>1</sub>xP<sub>2</sub>, P<sub>1</sub>xP<sub>5</sub>, P<sub>4</sub>xP<sub>5</sub>, P<sub>4</sub>xP<sub>6</sub> and P<sub>5</sub>xP<sub>6</sub> for number of kernels/row. The crosses P<sub>1</sub>xP<sub>2</sub>, P<sub>1</sub>xP<sub>4</sub>, P<sub>1</sub>xP<sub>5</sub>, P<sub>4</sub>xP<sub>5</sub>, P<sub>4</sub>xP<sub>6</sub> and P<sub>5</sub>xP<sub>6</sub> showed significantly positive SCA effects for number of grains/cob. Significant

positive SCA represents dominance and epistatic component of variation. This indicated that high *per se* value of cross may not necessarily indicate their potentiality in crosses.

Uddin *et al.* (2006), Uddin (2008), Alam *et al.* (2008) and Alam (2009) reported that any combination among the parents may produce hybrid vigour over the parents which might be due to dominant, over dominant or epistatic gene action. Choudhary *et al.* (2000) observed that SCA effects of the crosses were closely associated with their performance for days to silking and days to maturity. Earlier reporters also identified superior combinations in maize by estimating specific combining ability for different traits (Zelleke, 2000; Choudhary *et al.*, 2000; Desai and Singh, 2001 and Dubey *et al.*, 2001).

From the present study, the inbred parents P<sub>1</sub>, P<sub>2</sub>, P<sub>4</sub>, P<sub>5</sub>, and P<sub>6</sub> can be selected for development of high yielding hybrids for their good general combining ability of grain yield and other yield components. Based on mean performance and SCA effects, the crosses P<sub>1</sub>xP<sub>2</sub>, P<sub>2</sub>xP<sub>3</sub>, P<sub>4</sub>xP and P<sub>5</sub>xP<sub>6</sub> were found to be superior. These hybrids can be selected as single cross hybrids for verifying their performance over environments. The importance of GCA compared to SCA for grain yield agrees with previous findings of Kim and Ajala (1996), Ogunbodede *et al.* (2000), Alam *et al.* (2008) and Alam (2009).

In the present study value of F (relative frequencies of dominance and recessive alleles in the parents) for all the characters was positive and greater than zero. It indicates that dominant alleles were more frequent than recessive alleles.

The component D was significant for days to tassel, days to silking, plant height, ear height, number of kernel rows/cob, number of grains/cob and grain yield indicating the importance of additive gene effect in their inheritance.

The component H<sub>1</sub> was highly significant for all the characters which indicate the dominance gene effect in the inheritance of these characters.



The component  $H_2$  was highly significant for all the characters indicating the dominance with asymmetry of positive and negative gene effect in controlling these characters.

Thus highly significant values of the components D,  $H_1$  and  $H_2$  indicated the importance of both additive and dominant gene effect for the characters under study.

The value of  $h^2$  which measures the dominant effect over all loci was non-significant with non-significant environmental variance E for most of the characters.

The environmental component (E) for cob length, cob diameter, days to maturity, number of kernels /cob, number of kernels/row showed lower values indicating less influence of environment and might be less than additive and dominant effects.

Alam (2009) reported that kernels weight was controlled by environmental factors up to 60% in maize. This contradicts with the findings of Debnath and Sarker (1989).

The average degree of dominance  $(H_1/D)^{1/2}$  was more than unity for all the characters suggesting the importance of over dominance. Over dominance as well as predominant role non-additive genetic variance in the inheritance of grain yield was reported by Gardner and Lonnquist (1961), Genova (1984), Shahi and Singh (1985) and Genov (1987). Over dominance and predominant genetic variance was observed by Debnath and Sarker (1989) for grain yield, kernel rows per ear and 1000-kernel weight; by Nawar *et al.* (1980) for kernel rows per ear, and by Gamble (1962b) for kernel weight. Gardner and Lonnquist (1959), Johnson (1973), Shahi and Singh (1985), Lin and Chen (1986) and Debnath and Sarker (1989) observed partial dominance for ear length and number of kernels per row. Debnath and Sarker (1989) also showed complete dominance for ear diameter.

The values of  $H_2/4H_1$  were smaller than 0.25 for all the characters which indicated that positive and negative alleles were not distributed in equal proportion in the parents. Another values of  $[4DH_1]^{1/2} + F/[4DH_1]^{1/2} - F$  was greater than unity for all the characters except grain yield, suggesting asymmetrical distribution of dominant and recessive alleles in the parents for the characters.

Another ratio  $h^2/H_2$ , carried values greater than one indicating many groups of genes were responsible for their genetical control. For grain yield, the presence of an excess of dominant alleles than recessive alleles was observed in the parents.

Almost all the characters were observed to be under polygenic control. However, maximum number of genes or gene groups controlled grain yield. The highest number of genes or gene groups for grain yield was also reported by Debnath and Sarker (1989) and Alam (2009).

The low narrow-sense heritability values indicated that genotypic variance was governed by non-additive gene action. Similar results have been reported in maize by Alam (2009) and Azad *et al.* (2012).

$V_r$ - $W_r$  graphs indicate that it is difficult to attain simultaneous improvement for all the characters, as there is involvement of partial dominance and over dominance. The combined improvement of such characters should be based upon exploitation of both fixable and non-fixable components of genetic variance.

Utilization and exploitation of yield and yield components of maize require a clear understanding of their genetic architecture. A great deal of genetic architecture in maize has been studied which revealed that the nature of inheritance regarding gene action varied from material to material.

Over dominance as well as importance of non-additive gene action in the inheritance of grain yield was observed by Gamble (1962a), Darrah and Hallauer (1972), Genov (1987) and Debnath and Sarker (1989, 1990b). On the

other hand, Lonquist and Castro (1967) and Murthy (1978) reported partial dominance as well as predominant role of additive genetic variance of this trait. Epistasis was also observed in grain yield by Gamble (1962a) and Darrah and Hallauer (1972).

Result of the present study revealed that sufficient genetic diversity existed among the parents, which is essential for the improvement of a crop. Partial dominance or over dominance with non-allelic interaction was observed in most of the characters. For grain yield, over dominance as well as non-allelic interaction in the inheritance was evident.

It has been noticed that expression of dominant and recessive alleles in the parents was influenced by environment as the same parent showed different positions on Vr-Wr graphs.

## **HETEROSIS**

Heterosis is directly proportional to the existence of non-additive (dominance and epistasis) genetic variance in a population. The existence of significant amount of dominance variance is a prerequisite for exploitation of heterosis. Heterosis is also associated with wide adaptability of parents.

The cross having high heterosis coupled with high SCA can be utilized for commercial exploitation of heterosis. For commercial usefulness, economic or useful heterosis also compared for hybrid selection.

Pal and Prodhan (1994) suggested that selection on the basis of specific combining ability for the most useful heterotic crosses and thereafter development of single or double cross hybrid would be more effective in achieving genetic amelioration of maize for grain yield and oil content.

For days to tasseling,  $P_1 \times P_2$ ,  $P_1 \times P_3$ ,  $P_1 \times P_4$ ,  $P_1 \times P_5$ ,  $P_2 \times P_3$ ,  $P_2 \times P_4$  and  $P_2 \times P_5$  over mid parent and better parent showed significant positive heterotic effect.  $P_1 \times P_6$  and  $P_5 \times P_6$  crosses over mid parent and better parent showed significant negative heterosis.

For silking, ten crosses over mid parent and eleven crosses over better parent showed significant positive heterosis whereas the crosses  $P_2 \times P_6$  and  $P_3 \times P_4$  showed significant negative heterosis. For maturity, three crosses over mid parent showed negative and useful heterosis whereas twelve crosses over better parent showed significant positive heterosis in these crosses,  $P_3 \times P_6$  showed highest and earliest performance in maturity stage. Negative heterosis for earliness in maize was reported by Vasal *et al.* (1992b), Alam *et al.* (2008) and Alam (2009).

Five crosses for plant height and three crosses for ear height showed significant positive heterosis over mid parent and over better parent. Rest of the crosses in these traits had significant negative heterosis. Cob length and cob diameter are important yield contributing characters of maize inbred lines.

Most of the crosses showed significant negative and useful heterosis over mid parent and over better parent for cob length and diameter. Paul *et al.* (1995) reported significant negative heterosis over high parent in most of the crosses for cob length.

For number of kernels/row,  $P_1 \times P_2$ ,  $P_1 \times P_3$  and  $P_4 \times P_6$  over mid parent and better parent showed significant positive heterotic effect.  $P_1 \times P_6$  and  $P_3 \times P_5$  crosses over mid parent and better parent showed significant negative and useful heterosis. Highest significant positive heterosis over mid parent was found by the cross  $P_4 \times P_6$  and better parent in  $P_4 \times P_5$ .

Most of the crosses over mid parent and better parent showed significant heterosis for number of grains per cob. The crosses  $P_1 \times P_2$  and  $P_1 \times P_5$  respectively showed the highest and significantly positive heterosis over mid parent and over better parent in this trait.

Most of the crosses over mid parent and better parent showed significant negative and useful heterosis for grain yield. Maryam and Jones (1985), Vasal *et al.* (1992b) and Alam (2009) reported that yield is associated with delayed maturity and increased plant height which support the present investigation.

Among the characters studied for grain (kernel) yield and other yield contributing characters, most of the crosses are considered to be the most excellent, as they showed significant negative and useful heterosis.

$P_2 \times P_5$  showed the highest negative heterosis percent (-27.25 and -40.37) for grain weight while 15.30% positive heterosis over mid-parent was recorded for grain yield in  $P_1 \times P_2$ .

The results revealed that there is enough heterosis for all the characters studied in maize. It is well established that the incidence and magnitude of heterosis have positive association with the presence and magnitude of non-allelic interaction (Singh and Narayanan, 1993). It also indicates the possibility of increasing kernel yield by exploiting heterosis.

The presence of high heterosis indicates genetic diversity among the parents. Similar results have been reported in maize by Alam *et al.* (2008), Uddin *et al.* (2008) and Alam (2009). More importance of additive gene action for days to silking and days to maturity, plant height, ear length and girth, number of kernels/ear and 1000-kernel weight has been reported by Paul and Debnath (1999) and Zelleke (2000).

## SUMMARY AND CONCLUSION

In the present investigation, 25 different maize inbred lines were evaluated for eleven quantitative characters *viz.*, days to tasseling, days to silking, days to maturity, plant height, ear height, cob length, cob diameter, number of rows/ear, number of kernels/row, number of kernels/cob and grain yield/plant were selected. Statistical analyses such as mean, partition of components of variation, coefficient of variability, heritability, genetic advance, genetic advance as percentage of mean, correlation coefficient, selection index, genetic divergence through  $D^2$ - statistics, diallel analysis, combining ability and heterosis study were carried out.

An analysis of variance reveals that the item inbred line was highly significant indicating that 25 maize inbred lines were genetically different from each other. The estimates of different components of variation and coefficient of variability, such as phenotypic, genotypic and error were more or less high for days to silking, days to tasseling, number of grains/cob, number of kernels/row and yield, which indicates the wide scope of improvement of these characters through selection. The highest of  $h^2b$  was recorded for days to silking, cob diameter and the lowest in number of grains/cob. Genetic advance was highest in plant height and lowest in number of rows/cob. The highest value of GA% was found for cob diameter and the lowest in days to silking.

High values of genotypic coefficient of variation and high heritability estimates coupled with high genetic advance for days to silk, plant height, cob diameter, number of kernels/cob and number of kernels/row suggesting that phenotypic selection of parental inbred lines for hybridization program based on these characters would be effective. Phenotypic coefficient of variation for all the characters was higher than the genotypic coefficient of variation suggesting the more environmental effect on phenotypic complexes.

Grain yield was positively and significantly associated at the genotypic as well as the phenotypic levels with days to maturity, number of kernels/row and number of grains/cob. So, selection on the basis of these characters should get preference for breeding programme.

Path coefficient analysis using genotypic correlation revealed that days to maturity, cob length, number of rows/cob, and number of kernels/row had direct positive influence on grain yield. At the phenotypic level, days to maturity, plant height, ear height, cob length, number of rows/cob, number of kernels/row and grains number showed direct positive effect on grain yield and also positively correlated with grain yield. These results suggest that during selection more emphasis should be given on days to maturity, number of rows/cob and number of kernels/row.

The high expected genetic gains were more frequent through the different sets of data, when more character combinations were studied in the function. Large values for expected gains were obtained when all the four characters were included in a combination. Among the indices, the combinations, cob diameter + number of rows/cob + number of kernels/row + grain yield gave high values for expected gains over all sets of data.

Cluster analysis showed that maize inbred lines could be grouped into five different clusters. The highest inter-cluster distance was observed between I and II and the lowest inter-cluster distance was observed between clusters III and IV. Cluster III had maximum intra-cluster distance. However, the differences were clearer for days to silking, cob diameter, number of kernels/row, number of kernels/cob and grain yield/plant which had contributed largely to the total genetic divergence. Cluster I had the highest mean values for number of kernels/ear, plant height, days to maturity and grain weight/cob; cluster III had highest mean values for number of kernels/cob, plant height and days to maturity. Cluster IV had the highest mean value for days to silking and

days to maturity, cob length and number of kernels/cob. Therefore, for the improvement of maize, inbred lines should be selected from III and IV.

Combining ability analysis was carried out in a half diallel cross among six inbred lines for grain yield and its contributing characters. The expression of all these characters studied was found to be controlled by both additive and non-additive genes. However, additive genetic effects were preponderance for the expression of days to silking and days to maturity, and ear height, cob length and cob diameter, number of kernels/cob and grain weight.

The results of the present investigation elucidated that for grain yield, number of rows/cob, and ear height non-additive gene action was more important in controlling their expression. Thus, the present results indicated that the genetic improvement for the characters having additive genetic effects would be possible through the exploitation of such gene effects and for the characters, the predominance of non-additive genetic variance offered the scope for exploitation of heterosis utilizing such gene action.

Among the inbred lines, P<sub>1</sub>, P<sub>2</sub>, P<sub>5</sub> and P<sub>6</sub> were found to have good general combiners for grain yield and other yield contributing characters. These inbred lines can be used for the development of high yielding hybrids in maize.

Mean performance of heterotic crosses and SCA effects for different characters showed a good association. Based on *per se* performance and SCA effects, superior cross combinations P<sub>1</sub>xP<sub>2</sub>, P<sub>1</sub>xP<sub>3</sub>, P<sub>1</sub>xP<sub>5</sub>, P<sub>2</sub>xP<sub>3</sub>, P<sub>2</sub>xP<sub>4</sub>, P<sub>4</sub>xP<sub>5</sub> and P<sub>5</sub>xP<sub>6</sub> can be selected as promising single cross hybrids. Among the parents, these crosses P<sub>1</sub>xP<sub>2</sub>, P<sub>2</sub>xP<sub>5</sub>, P<sub>4</sub>xP<sub>5</sub> and P<sub>5</sub>xP<sub>6</sub> were considered to be the most excellent for exploitation of heterosis. After verifying the results of these promising hybrids over environments, good performers can be selected for commercial utilization.



Gene action of parents showed highly significant values of the components D,  $H_1$  and  $H_2$  indicated the importance of both additive and dominant gene effects for the characters under study.

The value of  $h^2$  which measures the dominant effect over all loci was significant with significant environmental variance E for all the characters which suggested the importance of dominance effect. As an indicator of the relative frequency of dominant and recessive alleles in the parents, the F value was found to be positive but significant for most of the characters, which means either that no alleles exhibit dominance or else that the dominant and recessive alleles are distributed equally among the parents.

In this study, the latter alternative may apply since the variances for  $H_1$  and  $H_2$  were significantly different from zero. It may thus be concluded that the dominant and recessive alleles of the related genes are distributed equally among the parents. Since the mean dominance effect of the heterozygote locus ( $h^2$ ) was significant, high heterotic effect values would be expected for most of the traits among the crosses. The parameters E, an estimate of the genotypic environmental variation and D, the additive genetic variance, were not different from zero.

The parameter D, which may also include a portion of the additive x additive epistatic variances as well as additive genetic variance itself, was significant for grain yield. Dominance variance ( $H_1$ ) and corrected dominance variance ( $H_2$ ) were significantly different from zero. It may thus be concluded that grain yield is under the dominance gene effect.

Graphical analysis revealed over dominance gene action for most of the characters, whereas partial dominance gene action was recorded for cob length and cob diameter and number of grains/cob. Partial dominance was indicated for the inheritance of grain yield since the regression line of the  $W_r/V_r$  graph cut the

W<sub>r</sub> axis above the origin. With regard to grain yield, the parents P<sub>3</sub> and P<sub>6</sub> had more dominant genes whereas P<sub>5</sub>, P<sub>4</sub>, P<sub>2</sub> and P<sub>6</sub> carried more recessive genes.

The following inferences are made on the basis of overall performance of the maize inbred lines.

- » Additive gene action with partial dominance was found for days to silking and maturity, and cob length and diameter, number of kernels/cob and grain weight under the situations.
- » Over- dominance type of gene action was observed for days to tasseling and plant height.
- » Additive gene action for grain yield, number of rows/cob and cob height changed to over-dominance.
- » The best combinations on the basis of mean performance were P<sub>1</sub>xP<sub>2</sub>, P<sub>2</sub>xP<sub>5</sub>, P<sub>4</sub>xP<sub>5</sub> and P<sub>5</sub>xP<sub>6</sub>.
- » Parents P<sub>1</sub>, P<sub>2</sub>, P<sub>5</sub> and P<sub>6</sub> were found to be best parents. These parents may be exploited in future breeding programs.

IL18 (P<sub>5</sub>) had the highest number of kernels/cob and grain yield/plant compared with the other parents and it is considered the best combiner to improve the kernels and grain weight in plants, because it showed the highest positive and significant GCA.

Better performing four crosses (P<sub>1</sub>xP<sub>4</sub>, P<sub>1</sub>xP<sub>5</sub>, P<sub>4</sub>xP<sub>5</sub> and P<sub>5</sub>xP<sub>6</sub>) can be utilized for developing high yielding hybrid varieties as well as for exploiting hybrid vigor. These crosses also need to be evaluated through multiplications.

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