# Genetic diversity analysis for qualitative and quantitative traits of Indian mustard (*Brassica juncea* L. Czern & Coss) under timely sowing condition in District Amroha

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#### **Abstract**

All the 25 genotypes were grouped into 6 clusters based on  $D^2$  analysis. The cluster-I with 9 strains had maximum genotypes among all the clusters followed by cluster-III, II, IV, V and VI. The inter cluster distance was recorded highest between cluster-III and cluster-IV (90.88). The minimum inter cluster distance was observed between cluster-I and IV (15.38) indicating their close relationship.

**Key words:** Clusters, Genetics divergence (D<sup>2</sup>) and Indian mustard

#### Introduction

Botanically, the genus *Brassica* comprises six species (B. nigra, B. oleracea, B. campestris, B. carinata, B. juncea, and B. napus). Among them first three species are elementary and diploid with 2n=16, 18 and 20 chromosomes and other three are tetraploids with chromosomes numbers 2n=34, 36 and 38. All these crops are grown under wide range of agroclimatic conditions. Indian mustard [Brassica juncea (L.) Czern & Coss], which is cultivated under the genus Brassica is cultivated all over India and it is throughout the world belongs to family Cruciferae (Brassicaceae). It has 38 to 42 % oil and 24% protein. The availability of genetic variation is advantageous for crop improvement. Such type of variability brought about by a group of genes which have a small individual effect, can be studied through quantitative measurements. The genetic facts are inferred from observations on phenotypes. Since phenotype is determined by the joint effect of genotype and environment, non-genetic

<sup>1</sup>Professor & Head, Department of Genetics & Plant Breeding, Country Head, AICRP on Mustard & Linseed, Section of Oilseeds, CSAUA&T, Kanpur-208002, U.P., India. parts exert large influence on genetic variability.

The literature available on these aspects in Indian mustard is relevant to the materials and environments of respective studies and cannot be generalized. Therefore, study on the above aspects on the available germplasm under the prevailing environment, where it is to be exploited is essential for successful utilization of germplasm resources for the development of superior varieties. In eastern Uttar Pradesh, a large acreage of mustard is under saline-alkaline soil condition. However, for such situation, screening of genotypes will help in identification of suitable genotypes. The exploitable variability is, therefore, required to be judged through various genetic parameters like heritability, genetic advance and others. Such a study appears to be extremely necessary for planning genetic improvement in Indian mustard. It is generally assumed by the plant breeders that cultivars originating from widely separated parts of the world are more likely to be genetically different. Therefore, the more diverse the parents, the more chance of increased spectrum of variability. On the basis such cultivars are included in hybridization programme in

the hope that their presumed genetic diversity would provide a greater likelihood of promising genetic rearrangements.

#### **Materials and Methods**

The present experiment was carried out during rabi, 2022-23 using 25 germplasms namely; DRMRIJ-31, Basanti, LAHAR, Pusa Bahar, NRH-101, NRC-DR-2, Mutant Varuna, RH-749, NRCHB-101, Pusa Bold, RH-406, Vardan, Pusa Krishma, Ashirvadh, Nav Gold, Pusa Barani, Pusa Jai Kisan, Kranti, Vaibhav, PM-26, Urvashi, Maya, Agarani, NDR-8501 and RLM-198 of Indian mustard made available and collected from the Section of Oilseed, Department of Genetics and Plant Breeding of Chandra Shekhar Azad University of Agriculture and Technology Nawabgani, Kanpur. The experiment was laid out in Randomized Block Design (RBD) with three replications. These lines were grown in single row plot of 5-meter length. The spacing between row to row and plant to plant was 45 cm and 15 cm, respectively maintained by thinning in the Technology Park/ Crop Cafeteria of Krishi Vigyan Kendra, Gajraula, Amroha, SVPUA&T, Meerut. Recommended agronomic practices were adopted to raise a good crop. Five competitive plants from each plot were randomly selected for recording observations for all the quantitative characters except days to flowering and days to maturity which were recorded on the plot basis. The data were recorded for thirteen characters namely; days to 50% flowering, days to maturity, plant height (cm), number of primary branches per plant, number of secondary branches per plant, length of main raceme (cm), number of siliquae per plant, number of seeds per siliqua, 1000-seed weight (g), harvest index (%), biological yield per plant (g), oil content (%) and seed yield per plant (g). Oil content was estimated using NMR method. D² analysis is done as per P.C. Mahalanobis (1928).

## **Results and Discussion**

25 strains/varieties of Indian mustard were grouped into 6 clusters under normal sown condition (Table 1). The genotypes from one source of origin clustered with the genotypes of other source of origin. This indicated that there was no parallelism between geographical distribution and genetic diversity. Anand and Rawat (1984), and Verma and Sachan (2000), also found the similar trend. The grouping of genotypes from same geographical origin into different clusters may be due to the different

Table 1: Distribution of 25 genotypes of Indian Mustard in different clusters.

Clusters Strains/variety	No.
1. NRC-DR-2, Nav Gold, Kranti, Vaibhav, RH-30, Selection 2016/10,	
Selection ns/4, Pusa Jai Kisan, DRMRIJ-31	9
2. Pusa Bold, Ashirvadh, Pusa Bahar, Vardan, Mutant Varuna, Basanti	6
3. Urvashi, NDR-8501, Agarni, Maya, Pusa Barani, RLM-198, KR-5610	7
4. B-85	1
5. LAHAR	1
6. NRH-101	1

Table 2: The average intra and inter cluster value of different clusters in Indian mustard (Brassica juncea)

Clusters	1 cluster	2 cluster	3 cluster	4 cluster	5 cluster	6 cluster
Cluster 1	11.887	26.954	52.704	15.383	31.070	34.592
Cluster 2		18.368	48.633	39.952	27.233	30.768
Cluster 3			35.054	57.755	58.158	90.883
Cluster 4				0.000	29.720	58.423
Cluster 5					0.000	66.710
Cluster 6						0.000

Table 3: Cluster mean for 13 characters in Indian mustard

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	Oil Seed yield/ Content plant (%) (g)	12.000 11.444 11.857 11.000 12.000		Oil Seed yield/ Content plant (%) (g)	11.57		Oil Seed yield/ Content plant (%) (g)	26	
	Oil Se Content (%)	39.287 38.678 38.971 40.840 37.950 38.753		Oil Se Content (%)	38.07 40.26		Oil Se Content (%)	13	
	Harvest index (%)	23.228 22.097 22.785 21.055 22.953 23.510		Harvest index (%)	22.31 23.50		Harvest index (%)	5 1.67	
	Biological Yield/ plant (g)	51.630 51.833 51.905 52.333 52.333		Biological Yield/ plant (g)	51.33 53.33		Biological Yield/ plant (g)	3	
	1000seed Weight (g)	3.388 3.600 3.703 2.960 2.327 4.453		No. of No. of 1000seed siliquae/ seeds/ Weight plant siliqua (g)	3.13		1000seed Weight (g)	40	
	l \	13.148 12.889 13.048 15.000 13.333 14.333		No. of / seeds/ siliqua	12.66	ıstard	No. of seeds/ siliqua	25	
	No. of No. of siliquae/ seeds/m) plant siliqua	325.556 311.722 313.143 307.667 329.000 300.333		No. of No. of siliquae/ seeds/ m) plant siliqua	295.33 329.41	Indian mu		2 0.67	
	Length of main traceme (c	56.843 45.871 57.668 60.700 44.973	rd	Length of main t raceme (c)	44.97 59.50	naracters in	Length of main t raceme (c	84	
	Plant No. of No. of Length No. of No. of height primary secondary of main siliquae/ seeds (cm) branch/plant branch/plant raceme (cm) plant siliqua	18.667 18.556 15.429 21.333 20.000	ters in musta	No. of Length No. of secondary of main siliqua branch/plant raceme (cm) plant	10.00	nce for 13 cl	No. of No. of Length No. of primary secondary of main siliquaranch/plant branch/plant raceme (cm) plant	6	
	No. of primary anch/plant	7.963 7.667 7.476 9.333 9.333 8.000	rent charact	Plant No. of height primary (cm) branch/plant	7.41	rds diverge	No. of primary anch/plant	1 0.33	
	Plant height (cm) br	174.424 170.838 155.975 166.477 175.143 182.163	n for diffe	Plant height (cm) br	145.51 176.88	cter to wo	Plant height (cm) br	3	
	Days to maturity	131.148 128.778 122.190 133.333 127.333	luster mea	Days to maturity	117.41	each chara	Days to maturity	3	
	Days 50% flowering	80.444 75.500 67.048 78.000 75.333 84.667	Table 4: Range among cluster mean for different characters in mustard	Days 50% flowering	65.25 83.50	Table 5: Contribution of each character to words divergence for 13 characters in Indian mustard	Days 50% Days to flowering maturity	1 strime 89 %)	
	Clusters	Cluster1 Cluster2 Cluster3 Cluster4 Cluster5 Cluster5	Table 4: Ra	Range	Lower	Table 5: Cc	Range	No.of time appearing 1stime 89 Per-cent (%)	

genetic backgrounds and wide divergence in features. Different genetic background is perhaps due to the free exchange of materials among different regions of country for breeding purpose; genetic drift and selection in different environments could be the other important factors contributing to the divergence. Singh and Gupta (1984), also reported similar reasons for genetic diversity.

In present investigation, on the basis of magnitude of D<sup>2</sup> values, 25 genotypes of Indian mustard were grouped into 6 clusters. The distribution of genotypes in both the environments was different. Maximum genotypes (8) were present in cluster-I. The perusal of Table 2 revealed that the maximum inter cluster distance was observed between cluster-III and cluster-IV (90.88) indicated wide diversity between these groups. Hybridization among the genotypes separated by high inter cluster distance will result in most heterotic crosses. The estimates of genetic divergence for most of the characters under study are in accordance with earlier reports. Amar Singh *et al.* (2005), Binesh Goyal *et al.* (2012), Ratnesh Pandey *et al.* (2013).

The maximum intra cluster distance was observed for cluster-III (35.05) followed by cluster-II and cluster-I. The maximum intra cluster value indicated maximum divergence among various genotypes within the cluster. A comparison of cluster mean for thirteen characters under study revealed considerable genetic differences between the clusters regarding one or more characters. The maximum character contribution towards divergence was observed for days to 50% flowering (29.67%) (Table 5). Similar findings were also reported by Shalini *et al.* (2000), Patel and Patel (2006), Singh *et al.* (2007), Doddabhimappa *et al.* (2010) and Binesh Goyal *et al.* (2012)

# Conclusion

The present investigations on the basis of studies made on genetic divergence it was suggested that cross between the genotypes of clusters-III and IV may give better results during hybridization programme. The maximum contribution towards divergence for days to 50% flowering (29.66%). It indicates that the germplasms are contributed to cluster-III namely; Urvashi, NDR-8501, Agarni, Maya, Pusa Barani, RLM-198, Pusa Krishma and

for clusters-IV namely; RH-406 can be utilized for future breeding programme.

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