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Genetic Diversity in Okra (Abelmoschus esculentus L.) Genotypes with Various Quantitative Traits

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Authors' contributions

This work was carried out in collaboration among all authors. All authors read and approved the final manuscript.

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ABSTRACT

An experiment was conducted at Horticulture Research Centre, Sardar Vallabhbhai Patel University of Agriculture & Technology, Meerut, India-250110. The analysis of variance data revealed substantial genetic differences for all morphological traits. According to Mahalanobis D² all the forty genotypes were found to have a significant amount of genetic diversity. The result indicated that the greatest mean value was found in cluster III for days to 50 per cent flowering, days to Ist flower initiation, days to Ist fruit set, days to Ist fruit picking, days to IInd fruit picking, number of primary branches, number of nodes / plant, duration of crop and also some characters was found highest mean value in cluster I like, plant height, fruit length, number of fruits / plant, length of internode, fruit yield / plant and fruit yield (q/ha.). Cluster II had the most genotypes recorded overall. (21 genotypes) followed by 11 genotypes in cluster I. The largest intra cluster distance was discovered in cluster III, whereas the largest inter cluster D² distance was measured between Cluster I and Cluster III. The genotypes of Cluster I and Cluster III are, according to the maximum inter-cluster D² distance. Among all the characters that were contributed, the length of internode had the highest contribution %.

Keywords: Cluster; genetic divergence; Okra; quantitative traits.

1. INTRODUCTION

Okra (Abelmoschus esculentus L.), is an annual plant, belongs to Malvaceae family. It is tropical and subtropical areas of the world; it is one of the most significant and major vegetable crops. A limited number of vegetables are readily available in the market during the summer, which satisfies customer demand for highly sought-after vegetables in the early summer. Because it contains a lot of iodine, the fruits of okra are most helpful in the treatment of goitre disease in human. In vegetable gardens, okra has the most exquisite blossoms that have great beauty value. According to [1], edible okra has a lot of nutrients, including protein, minerals, fat, fibre, calories, carbohydrate, maximum amount of water as well as good quantity of total phenolics, folate, and ascorbic acid.

Several experts have noted the medicinal and nutritional value of okra, including mineral components, protein, fibre, antioxidants, and vitamins [2]. Flavonoid and antioxidants such beta carotene, xanthein, and lutein are abundant in okra pods [3].

Okra is morphologically tall to medium in height plant and annual herbaceous nature. The leaves are alternate, broad, caudate, palmately with 3-7 lobed, hirsute, and serrate. The stem of okra is green or stained scarlet. Flowers are solitary, auxiliary, with a 2 cm long peduncle. The okra flower having 5 sepals and 5 petals. The petal colour of okra is yellow with a crimson spot. Okra flower has five to seven cm long staminal united column to the petal base with many stamens. Superior ovary with five to seven deep red stigmas.

Vegetables are mostly cultivated in India on an area of about 10.85 million hectares, total production about 200.4 million tonnes of produce year at a productivity of 18.55 tonnes per hectare [4]. Okra is grown on 530 thousand hectares, and annual production of 6465.59 metric tonnes [4]. Gujarat is the top state for okra production, producing 1019.42 thousand Tonnes from an area of 85.15 thousand hectares. West Bengal comes in second with production of about 882.59 thousand tonnes from an area of 78.64 thousand hectare [4].

The D² statistic, which calculates two levels of genetic divergence, namely; inter cluster and intra cluster levels, aids in the choosing genetically diverse parents for use in hybridization programmes [5]. Because hybrids between lines of different provenance typically demonstrate a stronger heterosis than those between closely related lines, genetic diversity exhibit plays a crucial role in breeding techniques [6]. Studying on okra can contribution to deeper understanding in plant biology and genetics.

2. MATERIALS AND METHODS

The research materials, which included a total of 40 genotypes of okra populations Hisar Naveen, Varsha Uphar, Punjab Kranti, Arka Anamika, Arka Abhay, Pusa Sawani, Pusa A-4, Parbhani Kranti, Kashi Pragati, Mona, Kashi Kranti, U.S-8063, IC-090491, Y.V.S-9, IC-316/2,4,5, IIVR-II, VRO-3, VRO-4, VRO-5, VRO-6, 368-A, Hisar Unnat, IC-18530, EC-305642, EC-305643, EC-305644, EC-305645, EC-305639, IC-014026, EC-305638. EC-305637, EC-305635, EC-305768, IC-013356, IC-013664, EC-359637, IC-15027, EC-305640, IC-010265 and IC-14909 with 14 different characteristics, were gathered from a variety of locations in the okra-growing regions of HAU, Hisar, PAU, Ludhiana, IIHR, Bengaluru, Dr. PDKV, Akola, IARI, New Delhi, MPKV, Rahuri, IIVR, and Varanasi. Using a Randomized Block Design (RBD) with three replications, the experiment was carried out in the summer season of 2019 at the Horticulture Research Centre of the Sardar Vallabhbhai Patel University of Agriculture & Technology in Meerut. To determine the reliability of each variety's performance, the genotypes were analysed statistically for all of the morphological findings of a 60x30 cm row-to-row and plant-to-plant sowing. The experimental location is situated in SVP University of Agriculture & Technology, Meerut, which is about 70 km from the National Capital of Delhi on National Highway No. 58 (Delhi - Haridwar Highway). Geographically, the trial location in the North Western Plain Zone of Uttar Pradesh at an elevation of about 297 metres above MSL and at 29° 01 N latitude and 77° 45 E longitude.

Data were gathered on agronomic and morphological traits, namely days to Ist flower initiation, days to 50% flowering, plant height, number of primary branches, number of nodes / Plant, length of internode, fruit length, days to Ist fruit set, number of fruits / plants, days to Ist fruit picking, days to IInd fruit picking, duration of crop, fruit yield / plant and fruit yield. Following the analysis of variance, multivariate analysis (using the D² statistic) was performed on the data of 40 genotypes of okra [5]. By using crucial condensation, standardized uncorrelated variables were created from the original measurements. [7]. Tocher's method [7] was used to divide genotypes into several clusters, and [8] formula was used to discovered the parallel contribution of various traits to total divergence.

3. RESULTS AND DISCUSSION

The analysis of variance on the basis of data in Table 1 there is a highly significant variation for each character. The findings suggested that the genotypes under study exhibit adequate diversity. Okra has also seen this type of crop variability, according to [9]. According to D² analysis, there were three clusters formed. From the pattern of clustering, it might be extrapolated that sufficient divergence was present to enable the establishment of individual character. According to the clustering pattern, geographic diversity is not a reliable indicator of genetic diversity. As a result, Mahalanobis D² analysis of quantitative traits is an effective tool for assessing genetic divergence among materials chosen even from the same geographic region.

Table 2 shows the cluster mean value for the fourteen traits that were the focus of the inquiry. The results showed that Cluster III had the greatest mean value for days to 50% flowering (54.89) and Cluster II had the lowest mean value for days to 50% flowering (45.48). Days to Ist flower initiation showed that clusters III and II, respectively, had the greatest mean value (51.94) and the lowest mean value (42.47). The highest mean value for the character like days to Ist fruit set was in cluster III, while the lowest mean values were in clusters II (53.67) and (44.39), respectively. Cluster III had the greatest mean value for days till the first fruit picking, while cluster II had the lowest (58.30 and 49.32, respectively). Days second fruit picking showed that cluster III had the highest mean value and cluster II had the lowest mean value (63.04). (54.17). Plant height was showed highest mean value (93.64) and lowest mean value (71.17) found in cluster I and III. Fruit length was observed, and clusters I and III had the highest mean value (9.71) and the lowest mean value (8.83). For the number of fruits / plants, clusters I and III had the greatest mean values (11.84) and (9.60). Number of primary branches was showed highest mean value (2.92) and lowest mean value (2.03) found in cluster III and cluster II respectively. Number of nodes / Plant showed highest mean value (22.95) and lowest mean value (18.79) was found in cluster III and cluster II. Length of internode was recorded highest mean value in cluster I (4.88) and lowest cluster mean in cluster III (3.40) respectively. Fruit yield and its traits that influence fruit production, such as fruit yield / plant, had a mean value that was highest in cluster I (119.30) and lowest in cluster III. (98.47). Fruit yield highest mean value in cluster I (76.78) and lowest cluster III (65.45). In other character like duration of crop highest mean value in cluster III (101.55) and lowest in cluster II (109.88) [9], [10] and [11]. Breeding programmes may be using genotypes that performed good, meaning they have these traits in different clusters, to create high yielding okra cultivars.

Table 1. ANOVA for 14 characters in Okra	[Abelmoschus esculentus (L.) Moench]
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Source of	DF	DFI	DF	PH (cm)	NPB	NN/P	LI	FL	DFS	NF/P	DFP	DSFP	DC	FY/P	FY
variation							(cm)	(cm)						(gm)	(q/ha)
Replication	2	1.25	3.48	0.08	0.03	0.67	0.04	0.24	0.49	0.29	3.71	0.38	7.78	17.69	3.09
Treatment	39	54.73**	52.77**	428.93**	1.73**	33.75**	1.81**	2.25**	52.96**	6.29**	49.82**	50.88**	56.28**	502.64**	152.76**
Error	78	3.77	3.80	4.28	0.06	1.10	0.12	0.66	3.86	0.17	5.00	5.22	17.25	14.01	10.41

*, ** significant at 5% and 1% level, respectively. DFI (Days to Ist flower initiation, DF (days to 50 % flowering), PH (plant height), NPB (number of primary branches), NN/P (number of nodes/P), LI (length of internode), FL (fruit length), DFS (days to Ist fruit set), NF/P (number of fruits/plant), DFP (days to Ist fruit picking), DSFP (days to IInd fruit picking), DC (duration of crop), FY/P (fruit yield/plant) and FY/h (fruit yield/plant)

Table 2. Cluster means

Characters		DFI	DF	PH	NPB	NN/P	LI	FL (cm)	DFS	NF/P	DFP	DSFP	DC	FY/P	FY
				(cm)			(cm)							(gm)	(q/ha)
Cluster I	Mean	43.05	46.19	93.64	2.71	20.03	4.88	9.71	45.09	11.84	50.10	55.02	98.20	119.30	76.78
	± SE	2.18	2.08	8.24	0.43	0.99	0.50	0.76	2.18	1.65	2.31	2.71	2.61	13.21	7.34
Cluster II	Mean	42.47	45.48	75.06	2.03	18.79	4.08	8.87	44.39	9.64	49.32	54.17	94.98	99.96	66.06
	± SE	1.87	1.77	8.24	0.31	2.27	0.45	0.81	1.93	0.77	1.78	1.95	3.70	7.27	4.03
Cluster III	Mean	51.94	54.89	71.17	2.92	22.95	3.40	8.83	53.67	9.60	58.30	63.04	101.55	98.47	65.45
	± SE	2.46	2.35	7.27	1.35	5.67	0.97	0.80	2.40	0.73	2.42	2.26	4.22	9.34	5.00

The 40 genotypes in total were divided by Tocher's methods, which are shown in Table 3 & Fig. 1, into three distinct genetic clusters based on genetic affinity or diversity. Cluster I comprised of 11 genotypes viz, Varsa Uphar, Punjab Kranti, Arka Anamika, Arka Abhay, Parbhani Kranti, Kashi Pragati, VRO-3, VRO-5, EC-305635, IC-14909. Cluster II 368-A. comprised of 21 genotypes viz, Hisar Naveen, Pusa Sawani, Pusa A-4, Mona, Kashi Kranti, Y.V.S-9, IIVR-II, VRO-4, VRO-6, Hisar Unnat, IC-18530, EC-305644, EC-305645, EC-305639, IC-014026, EC-305638, EC-305637, IC-013356, IC-013664, IC-15027, IC-010265. Cluster III comprised of 8 genotypes viz, U.S-8063, IC-090491, IC-316/2,4,5, EC-305642, EC-305643,

EC-305768. EC-359637. EC-305640. The parallels between genetic and geographic variety were evident. The viewpoint. which indicates a large deal of variability among the clusters, has been hold up by the study of [9], [10] and [11]. This shows how drastically different the genetic architecture of the lines in one cluster is from the genetic architecture of the lines in the other cluster. Based on the minimal cluster distance found between clusters I and III, in order to create the desired recombinants, a hybridization application may use the genotypes in this cluster as parents. The varieties that will act as parents in a hybridization are chosen using the inter cluster distance criterion in a D² analysis.

Clusters	Number of genotypes	Genotypes Name
Cluster I	11	Varsa Uphar, Punjab Kranti, Arka Anamika, Arka Abhay, Parbhani Kranti, Kashi Pragati, VRO-3, VRO-5, 368-A, EC- 305635, IC-14909
Cluster II	21	Hisar Naveen, Pusa sawani, Pusa A-4, Mona, Kashi Kranti, Y.V.S-9, IIVR-II, VRO-4, VRO-6, Hisar Unnat, IC-18530, EC- 305644, EC-305645, EC-305639, IC-014026, EC-305638, EC-305637, IC-013356, IC-013664, IC-15027, IC-010265.
Cluster III	8	U.S-8063, IC-090491, IC-316/2,4,5, EC-305642, EC- 305643, EC-305768, EC-359637, EC-305640.

Table 4. Inter and intra cluster distances

Clusters	Cluster I	Cluster II	Cluster III	
Cluster I	2.59			
Cluster II	3.59	2.17		
Cluster III	6.14	5.52	3.32	

Table 5. Contribution of various characters towards total genetic divergence

S. No	Characters	Contribution %
1.	DFI	5.83
2.	DF	7.69
3.	PH (cm)	4.29
4.	NPB	7.24
5.	NN/P	7.20
6.	LI (cm)	8.72
7.	FL (cm)	6.45
8.	DFS	7.25
9.	NF/P	7.69
10.	DFP	7.09
11.	DSFP	8.14
12.	DC	6.25
13.	FY/P (gm)	8.01
14.	FY (q/ha)	8.15







Fig. 1. Clustering pattern of 40 genotypes of okra on the basis of genetic divergence

Fig. 2. Contribution of Various Characters Towards Total Genetic Divergence

The average D² values between and within clusters are shown in Table 4. Cluster I and Cluster III had the largest inter cluster D² distance (6.14), whereas Cluster I and Cluster II had the smallest inter cluster D² distance (3.59). Cluster III (3.32) had the greatest intra-cluster distance, followed by cluster I. (2.59). Cluster II recorded the shortest distance within a cluster (2.17). The genotypes of Cluster I and Cluster III are not closely related, based on the maximum inter-cluster D² distance, although the genotypes of Cluster I and Cluster II are, as determined by the minimum inter-cluster D² distance. As can be seen from the modest variance of D² values, the genotypes of the cluster do not significantly differ in terms of their relative genetic distance. In the past, reported similar outcomes [9,10] and [11].

Crosses suggesting parent belonging to most divergence clusters would be expected to manifest highest heterosis and also wide heterogeneity of genomic architecture. In order to create high heterotic populations, segregates, and/or to be used for the production of hybrid in okra, crossings between the genetically varied genotypes of clusters I and III may be based on supremacy for any characteristic or genetic diversity.

Contributions of several characters towards overall genetic divergence are reported in Table 5 & Fig. 2. The genetic divergence was influenced by measuring the D² value for each character individually and summing the rank for each trait across all entries and alterable. Among all the characters that were supplied, the length of the internode had the highest contribution rate (8.72%), followed by fruit yield (8.15%), days to IInd fruit picking (8.14 %), fruit yield / plant (8.01 %), days to 50 per cent flowering (7.69 %), number of fruits / plant (7.69 %), days to Ist fruits set (7.25 %), number of primary branches (7.24 %), number of nodes / plant (7.20 %), days to Ist fruit picking (7.09 %), fruit length (6.45 %), duration of crop (6.25 %), days to Ist flower initiation (5.83 %) and lowest contribution of plant height (4.29 %). The maximum divergence was directly brought on by [9] and [10]. Thus, choosing variable parents based on these characteristics that indicated to producing highquality hybrids as well as segregants in okra, according to [11].

4. CONCLUSION

It is suggested that genetic diversity rather than geographic location should be considered when choosing the parent for a hybridization. Crosses between the genetically distinct genotypes of clusters I and III were suggested by a divergence analysis. From a breeding perspective, the genotype may be identified and involved in crossing to produce desirable segregates in okra based on genetic diversity and superiority with regard to any trait.

DISCLAIMER (ARTIFICIAL INTELLIGENCE)

Author(s) hereby declare that NO generative AI technologies such as Large Language Models (ChatGPT, COPILOT, etc) and text-to-image generators have been used during writing or editing of this manuscript.

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COMPETING INTERESTS

Authors have declared that no competing interests exist.

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