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Genetic divergence studies on growth, yield and quality traits in pumpkin (*Cucurbita moschata* Duch. Ex. Poir.) Genotypes

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Abstract

Genetic divergence study was conducted on 33 pumpkin genotypes for thirty-seven characters at College of Horticulture, venkataramannagudem during 2022–2023. These genotypes were grouped into six clusters irrespective of geographic divergence, indicating no parallelism between geographic and genetic diversity. Cluster-I was the largest comprising 20 genotypes, followed by Cluster-II with 7 genotypes, Cluster-III with 3 genotypes. Clusters-IV, V and VI comprised one genotype each. As regards cluster means, Cluster-VI performed better in most of the biometric characters studied. Maximum inter-cluster distance was observed in Clusters-V and VI, followed by Clusters-IV and VI, and clusters-I and VI. Intra-cluster distance was highest in Cluster I.

Keywords: Pumpkin, genetic divergence, cluster

Introduction

The cucurbit family is one of the largest families in the plant kingdom including the largest number of edible species. There are 27 species in the genus Cucurbita, of which 5 are cultivated. These are *Cucurbita moschata*, *Cucurbita maxima*, *Cucurbita ficifolia*, *Cucurbita pepo* and *Cucurbita mixta*. Cucurbita moschata is commonly known as pumpkin and is a widely grown Cucurbita species and the fruit is highly prized for its long-storing ability and high nutritional value (Jahan *et al.* 2012) ^[5]. This species is compatible with *C. maxima*, *C. pepo* and *C. mixta* (Tindall, 1987) ^[24]. Pumpkins are widely used as vegetables in both the young and mature stages and ripe fruits can be stored for 2 to 4 months (Yawalkar, 1991) ^[25]. Fruits with yellow and orange flesh are rich in carotene (3,332 IU), a precursor to vitamin A with abundant vitamins B and C. This can help improve the nutritional status of the population, especially vulnerable groups in terms of vitamin A requirements (Satkar *et al.* 2013) ^[20]. In India, pumpkins are grown on an area of 106 million hectares, with a total production of 2,218 tons and a yield of about 20.92 tons/hectare (NHB database, 2021-22).

As for pumpkins, the main issue of consumer preference is the large fruit (4 to 5 kg), which is not very popular for a small family of 3 to 4 members. Furthermore, with the recent increase in the number of nuclear families in India, people prefer to buy only small and medium sized whole pumpkins rather than cut into pieces. Additionally, small fruits can be easily packed and transported without damage. Therefore, it is necessary to develop pumpkin varieties and hybrid pumpkin varieties with small to medium fruit sizes (2-3 kg). Several efforts have been made by the public and private sectors to develop high-yielding varieties and hybrids. However, the development of high-yield varieties and hybrids combined with medium-sized fruits with high beta-carotene content is still very limited. Pumpkins have received less attention in crop improvement compared to other cucurbits. Since ancient times, large amounts of genetic material have been available, but the conscious evaluation and exploitation of genetic material has only recently been realized. This is useful for the plant breeder when developing a commercially suitable variety for the market by identifying the constituent traits for which selection can be made based on improvement in yield and quality. Preliminary identification of early-maturing genotypes can be done based on characteristics such as days to appearance of first female flower, node number at which first female flower appeared, days to first fruit harvest.

Collection and evaluation of germplasm is a prerequisite for any breeding program aimed at selecting high-yielding genotypes with the desirable characteristics of high earliness, yield and

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quality. Therefore, characterization of the available pumpkin germplasm was carried out to identify potential earlymaturing, high-yielding cultivars of small and medium fruits as well as other cultivars. Quality parameters are improved. Furthermore, genetic advance can be used to predict selection efficiency. Mahalonobis (1936) ^[12] D² statistics provide a measure of the magnitude of divergence between two groups under comparison. D² genotype grouping will be useful in selecting suitable parental lines for heterosis breeding as well as deriving superior segregating lines from specified crosses.

Material and Methods

Thirty-three genotypes (Table 1) of pumpkin having diverse origin were evaluated at the College of Horticulture, Venkataramannagudem, during the period November-April, 2022-23. Genotypes were evaluated using Augmented Block Design, with three blocks. Plants were grown at a spacing of 1.5 m \times 1.5 m adopting the package of practices as recommended. Observations were recorded on five randomly selected plants of each genotype in each block for thirty-seven

characters, viz., vine length (cm) at final harvest ; internodal length (cm) at 45 DAS, 90 DAS and final harvest; petiole length (cm) at 45 DAS, 90 DAS and final harvest; number of branches per plant at 45 DAS, 90 DAS and final harvest; number of leaves per plant at 45 DAS, 90 DAS and final harvest; days to appearance of first male flower; node number at which first male flower appeared; days to appearance of first female flower; node number at which first female flower appeared; number of male flowers per plant; number of female flowers per plant; sex ratio (male: female); days to first fruit harvest; number of fruits per vine; number of ridges per fruit; fruit length (cm); fruit diameter(cm); flesh thickness (cm); seed cavity diameter (cm); number of seeds per fruit; test weight (g); average fruit weight (kg); fruit yield per vine (kg); estimated fruit yield (t/ha); β-carotene (mg/ 100 g.f.w); reducing sugars (%): non-reducing sugars (%): total sugars (%) and total soluble solids (°brix). Genetic divergence was estimated using D² statistics of Mahalonobis (1928) ^[10] and the populations were grouped into clusters as per Rao (1952).

Table 1: List of genotypes of pumpkin and their source

S. No	Genotype	be Source			
1.	IC 284761	NBPGR Regional Station, Thrissur, Kerala			
2.	IC 333299	NBPGR Regional Station, Thrissur, Kerala			
3.	IC 395804	NBPGR Regional Station, Thrissur, Kerala			
4.	IC 599403	NBPGR Regional Station, Thrissur, Kerala			
5.	IC 599408	NBPGR Regional Station, Thrissur, Kerala			
6.	IC 599422	NBPGR Regional Station, Thrissur, Kerala			
7.	IC 599425	NBPGR Regional Station, Thrissur, Kerala			
8.	IC 599427	NBPGR Regional Station, Thrissur, Kerala			
9.	IC 599435	NBPGR Regional Station, Thrissur, Kerala			
10.	IC 599436	NBPGR Regional Station, Thrissur, Kerala			
11.	IC 599437	NBPGR Regional Station, Thrissur, Kerala			
12.	IC 613471	NBPGR Regional Station, Thrissur, Kerala			
13.	IC 618053	NBPGR Regional Station, Thrissur, Kerala			
14.	IC 618054	NBPGR Regional Station, Thrissur, Kerala			
15.	IC 618055	NBPGR Regional Station, Thrissur, Kerala			
16.	IC 618056	NBPGR Regional Station, Thrissur, Kerala			
17.	IC 618057	NBPGR Regional Station, Thrissur, Kerala			
18.	Coimbatore local	Coimbatore, Tamil Nadu			
19.	Pollachi local	Pollachi, Tamil Nadu			
20.	Mettupalyam local	Mettupalyam, Tamil Nadu			
21.	Balaram local	AnakapalleAndhra Pradesh			
22.	Jogumpeta local	Anakapalle, Andhra Pradesh			
23.	Kantaram local	Anakapalle, Andhra Pradesh			
24.	Komira local	Anakapalle, Andhra Pradesh			
25.	Jangareddygudem local	West Godavari, Andhra Pradesh			
26.	Martur local	Bapatla, Andhra Pradesh			
27.	Vemavaram local	Prakasam, Andhra Pradesh			
28.	Vittamrajupalli local	Guntur, Andhra Pradesh			
29.	Ummadivaram local	Guntur, Andhra Pradesh			
30.	Dechavaram local	Palnadu, Andhra Pradesh			
31.	Arka Suryamukhi (Check-1)	IIHR, Bangalore			
32.	Local Cultivar-1 (Check-2)	West Godavari, Andhra Pradesh			
33.	Local Cultivar-2 (Check-3)	West Godavari, Andhra Pradesh			

Results and Discussion

Analysis of variance indicated that mean sum of squares in pumpkin genotypes were not highly significant for all characters except number of male flowers per plant and fruit yield per vine (kg). Having computed D^2 values for all possible pairs, the thirty-three genotypes were classified into six groups of gene constellations. These indicated a large genetic diversity (Table 2) and (Fig. 1). Maximum number of genotypes (20) grouped under Cluster-I, followed by Clusters-II and III, with 7 and 3 genotypes each. Clusters-IV, V and VI comprised one genotype each (mono-genotypic clusters).

Table 2: Distribution of pumpkin genotypes into different clusters

Cluster No.	Number of genotypes	Name of the genotypes		
Cluster I 20 618054, IC 618056, IC 618057, Coimbatore Local, Pollachi Local, Mettupalyam Local, Kan		IC 284761, IC 333299, IC 599403, IC 599425, IC 599427, IC 599435, IC599436, IC 599437, IC 613471, IC 618054, IC 618056, IC 618057, Coimbatore Local, Pollachi Local, Mettupalyam Local, Kantaram Local, Vemavaram Local, Vittamrajupalli Local, Ummadivaram Local, Dechavaram Local		
Cluster II	7	IC 395804, IC 599422, Komira Local, Jangareddygudem Local, Arka Suryamukhi, Local Cultivar-1, Local Cultivar-2		
Cluster III	3	IC 618055, Balaram Local, Jogumpeta Local		
Cluster IV	1	IC 618053		
Cluster V	1	IC 599408		
Cluster VI	1	Martur Local		

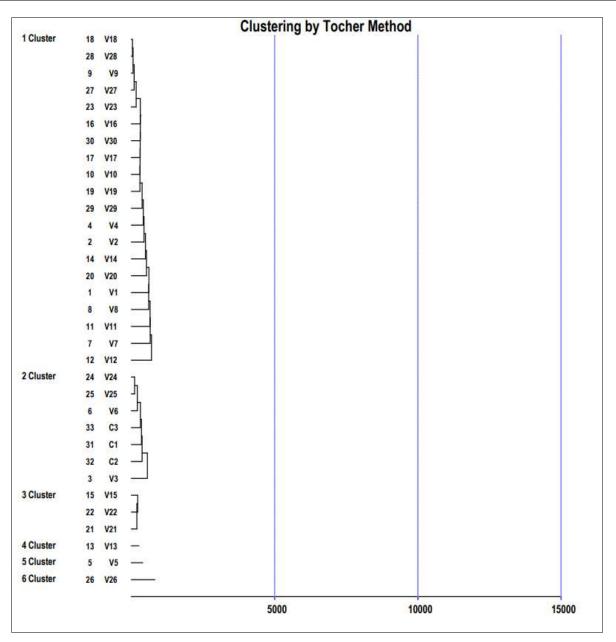


Fig 1: Dendrogram showing clustering pattern for divergence in pumpkin genotypes

Average inter and intra-cluster distances in pumpkin genotypes

Intra and inter-cluster distances are an index of genetic diversity among clusters as shown in (Table 3) and (Fig. 2). Inter-cluster distances were greater than intra-cluster distances, revealing a considerable amount of genetic diversity among the genotypes studied. Intra-cluster distance was highest in Cluster-I (1441.37), followed by Clusters-II

and III (1278.25 and 725.98, respectively). Highest intercluster distance was observed in Clusters-V and VI (19260.38), followed by Clusters-IV and VI (13564.75) and Clusters-I and VI (12735.94). Genetic distance (D²) between Cluster-VI was larger than in Clusters- I, IV and V. Minimum inter cluster distance was observed between Clusters-I and V (2135.77) indicating close relationship among genotypes. Data clearly indicated that the genotypes did not cluster

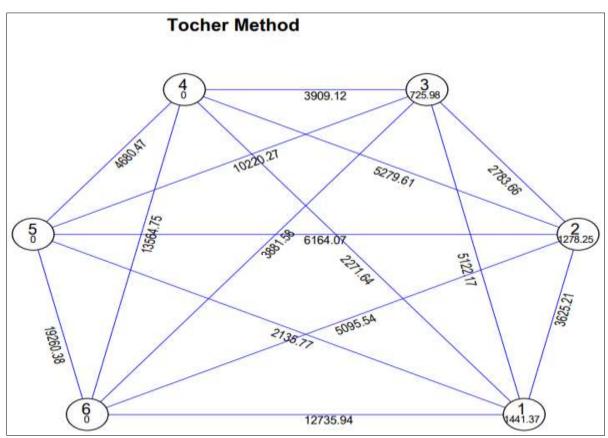
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according to their geographical distribution. In general, the pattern of distribution of genotypes from various regions into different clusters was seen to be random. Similar observations were also reported by Lovely (2001)^[9] in ash gourd, Kale *et al.* (2002)^[6] and Lakshmi *et al.* (2003)^[8] in pumpkin, Kandasamy (2004)^[7] in melon, Maharana *et al.* (2006)^[13] in ivy gourd, and by Devmore *et al.* (2007)^[2] and Dey *et al.* (2007)^[3] in bitter gourd. One possible reason may be that it is very difficult to establish the actual place of origin of a genotype. Free and frequent exchange of genetic material among breeders in the country makes it very difficult to maintain the real identity of a genotype. Absence of relationship between genetic diversity and geographical

distance indicates that forces other than geographical origin (such as exchange of genetic stock, genetic drift, natural mutation, spontaneous variation or natural and artificial selection) may be responsible for the genetic diversity. Another possibility may be that estimates of diversity based on characters used in the present investigation may not be sufficient to account for variability caused by some other traits of physiological / biochemical nature (which could be important in depicting the total genetic diversity in a population). Therefore, selection of genotypes for hybridization should be based on genetic diversity other than geographic divergence.

Table 3: Average inter and intra–cluster (diagonal) distance D² values in pumpkin genotypes

	Cluster I	Cluster II	Cluster III	Cluster IV	Cluster V	Cluster VI
Cluster I	1441.37	3625.21	5122.17	2271.64	2135.77	12735.94
Cluster II		1278.25	2783.66	5279.61	6164.07	5095.54
Cluster III			725.98	3909.12	10220.27	3881.58
Cluster IV				0.00	4680.47	13564.75
Cluster V					0.00	19260.38
Cluster VI						0.00



Mahalonobis Euclidean Distance (Not to the Scale)

Fig 2: Cluster diagram showing average intra and inter-cluster D² values among pumpkin genotypes

Cluster means of thirty-seven characters in pumpkin genotypes

The cluster means for each of thirty-seven characters are presented in (Table 4). Cluster I included the genotypes with highest sex ratio (male: female) (2.52) and with lowest internodal length at final harvest (21.34 cm), petiole length at 45 DAS (14.27 cm), number of female flowers per plant (3.27) and number of fruits per vine (2.11). Cluster II included the genotypes with lowest days to appearance of first

female flower (57.52), number of male flowers per plant (7.09), test weight (10.26 g), estimated fruit yield (16.17 t/ha) and reducing sugars (3.28%). Cluster III included the genotypes with highest number of leaves per plant at 90 DAS (247.93) and number of leaves per plant at final harvest (317.93) and with lowest number of branches per plant at final harvest (11.67), number of female flowers per plant (3.27) and non-reducing sugars (2.39%).Cluster IV included the genotype with highest petiole length at final harvest

(21.75 cm) and number of leaves per plant at 45 DAS (94.20) and with lowest internodal length at 45 DAS (9.90), number of branches per plant at 45 DAS (4.20), number of branches per plant at 90 DAS (7.40), sex ratio (male: female) (2.24), fruit length (14.27 cm), fruit diameter (16.89 cm), number of seeds per fruit (258.80), average fruit weight (1.97 kg) and total soluble solids (3.87 °brix). Cluster V included the genotype with lowest vine length at final harvest (449.13cm), internodal length at 90 DAS (15.40 cm), petiole length at 90 DAS (18.62 cm), petiole length at final harvest (20.86 cm), number of leaves per plant at 90 DAS (214.60), number of leaves per plant at final harvest (284.60), days to first fruit harvest (88.40), number of ridges per fruit (14.00), flesh thickness (2.55 cm), seed cavity diameter (9.92 cm), fruit yield per vine (7.23 kg), beta-carotene (2.30) and total sugars (5.85%). Cluster VI included the genotype with highest vine length at final harvest (569.66 cm), internodal length at 45 DAS (11.98 cm), internodal length at 90 DAS (19.42 cm), internodal length at final harvest (23.11 cm), petiole length at 45 DAS (16.61 cm), petiole length at 90 DAS (20.41 cm), number of branches per plant at 45 DAS (5.80),number of branches per plant at 45 DAS (5.80),number of branches per plant at 90 DAS (9.60),number of branches per plant at final harvest (14.60), number of female flowers (4.40), number of fruits per vine (3.40), number of ridges per fruit (19.40), fruit length (23.05 cm), fruit diameter (21.38 cm), flesh thickness (3.51 cm), seed cavity diameter (12.26 cm), number of seeds per fruit (338.20), test weight (12.40 g), average fruit weight (2.90 kg), fruit yield per vine (10.77 kg), estimated fruit yield (23.96 t/ha), beta-carotene (3.15), reducing sugars (4.16%), non-reducing sugars (2.78%), total sugars (6.93%) and total soluble solids (5.39 °brix).

Table 4: Cluster means for growth, yield and quality traits in pumpkin genotypes

S. No	Character	Cluster I	Cluster II	Cluster III	Cluster IV	Cluster V	Cluster VI
5.110		Frowth chai				Cluster V	
1	Vine length at final harvest (cm)	479.18	518.74	533.55	491.36	449.13	569.66
2	Internodal length at 45 DAS (cm)	10.38	10.92	10.87	9.90	11.21	11.98
3	Internodal length at 45 DAS (cm)		17.65	18.59	17.53	15.40	19.42
4	Internodal length at final harvest (cm)	16.84 21.34	21.92	22.24	22.39	22.38	23.11
5	Petiole length at 45 DAS (cm)	14.27	14.32	15.05	14.46	15.54	16.61
6	Petiole length at 90 DAS (cm)	18.64	19.23	19.49	18.69	18.62	20.41
7	Petiole length at final harvest (cm)	21.10	21.00	21.27	21.75	20.86	21.52
8	Number of branches per plant at 45 DAS	4.63	4.39	4.77	4.20	5.10	5.80
9	Number of branches per plant at 90 DAS	8.60	8.54	8.27	7.40	9.00	9.60
10	Number of branches per plant at final harvest	12.57	12.54	11.67	13.00	13.00	14.60
11	Number of leaves per plant at 45 DAS	75.48	73.70	87.47	94.20	77.60	68.20
12	Number of leaves per plant at 90 DAS	230.25	222.07	247.93	245.00	214.60	235.60
13	Number of leaves per plant at final harvest	300.22	288.57	317.93	317.00	284.60	305.60
15	Floral characters						
14	Days to appearance of first male flower	54.10	56.21	54.80	51.60	51.00	56.40
15	Node number at which first male flower appeared	13.17	13.59	12.20	17.40	13.00	15.80
16	Days to appearance of first female flower	57.72	57.52	61.33	69.60	58.00	65.00
17	Node number at which first female flower appeared	24.14	24.07	25.60	22.60	25.20	21.40
18	Number of male flowers per plant		7.09	8.07	7.60	8.50	10.80
19	Number of female flowers per plant	8.12 3.27	3.30	3.27	3.40	3.60	4.40
20	Sex ratio (male: female)	2.52	2.38	2.50	2.24	2.36	2.45
		Fruit chara	oters	•		-	
21	Days to first fruit harvest	93.44	94.37	96.13	93.20	88.40	92.80
22	Number of fruits per vine	2.11	2.38	2.27	2.60	2.20	3.40
23	Number of ridges per fruit	15.77	16.75	15.53	16.00	14.00	19.40
24	Fruit length (cm)	14.90	14.97	16.18	14.27	16.96	23.05
25	Fruit diameter (cm)	17.62	18.22	18.93	16.89	19.07	21.38
26	Flesh thickness (cm)	2.65	2.63	2.68	2.60	2.55	3.51
27	Seed cavity diameter (cm)	10.54	10.90	10.26	11.01	9.92	12.26
28	Number of seeds per fruit	279.10	303.79	299.93	258.80	279.40	338.20
29	Test weight (g)	10.60	10.26	10.60	10.31	10.71	12.40
30	Average fruit weight (kg)		2.28	2.36	1.97	2.32	2.90
31	Fruit yield per vine (kg/vine)	7.98	8.08	8.41	7.90	7.23	10.77
32	Estimated fruit yield (t/ha)	17.76	16.17	18.76	19.50	16.58	23.96
Quality characters							
33	β -carotene (mg/100g.f. w)	2.32	2.52	2.74	2.49	2.30	3.15
34	Reducing sugars (%)	3.41	3.28	3.68	3.35	3.32	4.16
35	Non-reducing sugars (%)	2.53	2.63	2.39	2.63	2.52	2.78
36	Total sugars (%)	5.93	5.91	6.07	5.98	5.85	6.93
37	Total soluble solids (°Brix)	4.27	4.41	4.69	3.87	4.12	5.39

Relative contribution of different characters towards divergence

The percent contribution towards genetic divergence by all the thirty-seven characters is furnished in the (Table 5) and (Fig. 3). The trait reducing sugars (%) had shown the highest contribution towards divergence by ranking first with a contribution of 21.03% followed by vine length (cm) at final harvest with 18.21%. The characters *viz.*, number of seeds per fruit, sex ratio (male: female), fruit yield per vine (kg), days to appearance of first female flower, number of leaves per plant at 45 DAS, average fruit weight (kg), number of leaves per plant at final harvest, number of leaves per plant at 90 DAS, beta-carotene, fruit length (cm), test weight (g), number of fruits per vine, total soluble solids (°brix), node number at which first female flower appeared, days to first fruit harvest, estimated fruit yield (t/ha), flesh thickness (cm), number of male flowers per plant, number of female flowers per plant and non-reducing sugars (%) contributed 13.72, 8.72, 6.76, 6.54, 5.49, 3.57, 3.03, 2.46, 2.44, 2.25, 1.54, 1.19, 0.75, 0.64, 0.38, 0.38, 0.26, 0.25, 0.25, 0.13 per cent respectively, but however internodal length at 45 DAS (cm), internodal length at 90 DAS (cm), internodal length at final harvest (cm), petiole length at 45 DAS (cm), petiole length at 45 DAS (cm), number of branches per plant at 45 DAS, number of branches per plant at 90 DAS, number of branches per plant at 90 DAS, number of first male flower, node number at which first male flower appeared, number of ridges per fruit, fruit diameter (cm), seed cavity diameter (cm) and total sugars (%) did not contribute anything to the diversity.

S. No	Source	Contribution (%)	Number of times ranked first
		n characters	
1	Vine length at final harvest (cm)	18.21	136
2	Internodal length at 45 DAS (cm)	0.00	0
3	Internodal length at 90 DAS (cm)	0.00	0
4	Internodal length at final harvest (cm)	0.00	0
5	Petiole length at 45 DAS (cm)	0.00	0
6	Petiole length at 90 DAS (cm)	0.00	0
7	Petiole length at final harvest (cm)	0.00	0
8	Number of branches per plant at 45 DAS	0.00	0
9	Number of branches per plant at 90 DAS	0.00	0
10	Number of branches per plant at final harvest	0.00	0
11	Number of leaves per plant at 45 DAS	5.49	29
12	Number of leaves per plant at 90 DAS	2.46	13
13	Number of leaves per plant at final harvest	3.03	16
	Floral	characters	
14	Days to appearance of first male flower	0.00	0
15	Node number at which first male flower appeared	0.00	0
16	Days to appearance of first female flower	6.54	51
17	Node number at which first female flower appeared	0.64	5
18	Number of male flowers per plant	0.25	1
19	Number of female flowers per plant	0.25	1
20	Sex ratio (male: female)	8.72	68
	Fruit	characters	
21	Days to first fruit harvest	0.38	3
22	Number of fruits per vine	1.19	19
23	Number of ridges per fruit	0.00	0
24	Fruit length (cm)	2.25	36
25	Fruit diameter (cm)	0.00	0
26	Flesh thickness (cm)	0.26	2
27	Seed cavity diameter (cm)	0.00	0
28	Number of seeds per fruit	13.72	107
29	Test weight (g)	1.54	12
30	Average fruit weight (kg)	3.57	57
31	Fruit yield per vine (kg/vine)	6.76	104
32	Estimated fruit yield (t/ha)	0.38	2
		v characters	
33	β-carotene (mg/100g.f. w)	2.44	19
34	Reducing sugars (%)	21.03	164
35	Non-reducing sugars (%)	0.13	1
36	Total sugars (%)	0.00	0
37	Total soluble solids (°Brix)	0.75	12

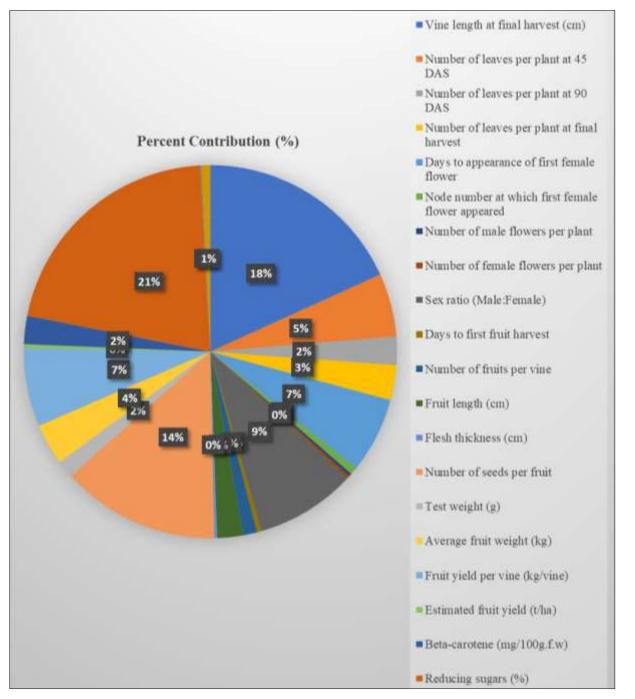


Fig 3: Relative contribution of characters towards genetic divergence in pumpkin

Based on these results, Mahalonobis D^2 was found to be a useful tool in grouping genotypes phenotypically and geographically. Findings revealed that in pumpkin, there is a vast scope for developing new varieties with greater yield potential and to better other attributes of economic importance, using this elite germplasm. In crop improvement programmes, intercrossing among genotypes with outstanding mean performance for these characters would prove to be effective. For recovering improved progenies for growth, floral, fruit and quality characters, crosses can be attempted between the genotypes belonging to clusters I, IV and V with cluster VI as revealed by divergence studies.

Conclusion

Multivariate analysis considering 37 morphological traits following Mahalanobis D^2 statistic revealed good diversity

among 33 genotypes of pumpkin, which were grouped into six distinct clusters. The genotypes of diverse clusters I and VI, clusters IV and VI and clusters V and VI could be used in hybridization programme either to produce highly heterotic F_{1S} or to generate wide range of transgressive segregants in population to develop high yielding varieties of pumpkin. The characters reducing sugars (%), vine length (cm) at final harvest and number of seeds per fruit contributed maximum towards divergence among the pumpkin genotypes.

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