

## **Genetic estimates and diversity study in Sesame(*Sesamum indicum* L.)**

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**Abstract:** Genetic estimates and diversity of 33 genotypes sesame representing different eco-geographical regions were studied for number of morphological characters. Wide variation in plant habit (plant height and branching pattern), number of capsules per plant, number of seeds per capsule, mean seed weight, and yield per plant was recorded. The estimates of genotypic coefficient of variation (GCV) were less than its corresponding estimates of PCV for seed yield and yield component, exhibiting consistency in environmental interaction for character expression. Capsule per plant exhibited high heritability coupled with good genetic advance indicating that additive genetic effect was more prevalent in this trait. Correlation study at phenotypic level showed that seed yield per plant was significantly and positive associated with plant height, number of branches per plant, number of capsule per plant and capsule length. So selection for high no of capsule per plant, capsule length and no of branches per plant will lead towards high yield. The genotypes were further grouped through multivariate analysis based on 8 traits. The genotypes constituted three major clusters. Origin of genotypes did not play a significant role in constitution of clusters. Selection of parents based on parents belonging to different cluster would like to produce more desirable segregants.

**Keywords:** Cluster, Correlation, Genetic variability, Heritability and Sesame

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### **I. Introduction**

Sesame is one of the most ancient and important oilseed crops grown from ancient times. It was cultivated and domesticated on the Indian subcontinent during Harappan and Anatolian eras over 4,000 yrs ago. (Bedigian and Van der Mesen, 2003). Due to the stability of its healthy oil, easiness of extraction and resistance to drought, sesame was popular in the ancient world, Sesame is considered as a nutritious oilseed crop being rich source of protein (18–25 %), carbohydrate (13.5 %), minerals and polyunsaturated fatty acid (Bedigian et al. 1986). Sesame oil is favoured as a media of cooking by Indians and Africans. Presence of sesamol, a unique anti-oxidant and more poly-unsaturated fatty acid such as oleic acid (43 %), linoleic acid (35%), palmitic acid (11%) and stearic acid (7%), have made it to 'queen of oilseed crop' (Ashri 1989; Fukuda et al. 1986). Sesame ranks fifth for important edible oil crop in India after groundnut, rapeseed-mustard, sunflower and soybean. (www.agricoop.nic.in). India holds top position in the world in sesame-acreage (24%) and contribution in export (40%) (FAO Statistics Division, 2012) Raikwar and Srivastva, (2013). Productivity of sesame in India is low compared to other sesame producing countries. Improvement in productivity will definitely boost the oil crop market of India and other ancillary industries. One of the simple approach to improve production of any crop is to boost up productivity, Genetic up gradation of any crop depends primarily on utilization of existing genetic resource. A large germplasm resource is always favoured in plant breeding program as many desirable traits may obviously remain in the population which may exploit breeding program. Study on genetic diversity comparatively limited in sesame. Several genetic parameters, such as, phenotypic and genotypic coefficient of variation (PCV and GCV), heritability and genetic gain help to assess genetic diversity of experimental materials.

Multivariate analysis is an important biometric technique when different quantitative traits are usually pooled up together to reach towards a conclusive outcome of diversity. Multivariate analysis is often used in selection of parents for hybridization program in different crops like blackgram, (Dasgupta and Das, 1984,1991), horsegram ( Dasgupta et al 2005), mustard (Pandey et al 1984) and sesame (Tripathi et al 2013, Akbar et al 2011). The present study has been conducted to assess the diversity genotypes in sesame following different genetic estimate and multivariate analysis.

### **II. Material & Method**

33 sesame genotypes were collected from diverse eco-geographical regions of India and abroad. The genotypes were grown in summer season following Randomized Block Design (RBD) with 3 replications having 10cm spacing between plants and 40cm between rows at Agricultural Experimental Farm, University of Calcutta, Baruipur in 2015. The farm is situated at an elevation of 10 meter above sea level, at approximately 22°51' N latitude and 88° 24' E longitude. The N: P: K fertilizers were applied at the rate 50:25:25 as basal dose

during final soil preparation. Manganese sulphate at the rate of 5Kg per hectare was also given as basal. Normal culture practices were followed during cultivation and irrigations were applied whenever the soil becomes very dry. Statistical cluster analysis of 33 genotypes with respect to morphological characters was done by Average Linkage Between Groups (in other words UPGMA) Method. The divergence between accessions was evaluated using a Euclidean distance dissimilarity matrix. Euclidian distance between lines were calculated and UPGMA cluster analysis was performed with the help of IBM-SPSS software (Version: 16.0) for dendrogram formation

### III. Result & discussion

Eight morphological characters were studied and ANOVA (Analysis of Variance) revealed that genotypes were significantly different from each other indicating high diversity among the genotypes.

Phenotypic coefficients of variation exhibited marginal higher values but maintained a close relation with genotypic coefficients of variation for all the traits. Highest coefficients of variation (phenotypic and genotypic) were exhibited by capsule per plant, seed yield/plant ( Table 2) Sumathi and Murlidharan (2010), Parameshwarappa *et al.*, (2009) and Sudhakar *et al.* (2007), and number of primary branches/plant similar finding was reported earlier Solanki and Gupta, (2003), Sudhakar *et al.* (2007), Saha *et al.* (2012), Gidey *et al.*, (2013), Iqbal and Dasgupta, (2015), in sesame. High GCV and PCV for capsule/plant, seed yield/plant and number of primary branches/plant suggest reasonably high variability in the studied materials and this would obviously help in upgrading the genotypes by simple selection. Heritability in broad sense was highest for 1000 seed weight followed by capsule per plant and seed yield/plant. Moderately heritability was observed for capsule length, plant height and days to 50% flowering on the contrary days to maturity exhibits very low heritability. Information on heritability of any traits aids to selection as traits with high heritability would like to exhibit similar character expression consistently over years. High heritability with high GCV of any trait is favourable combination for genetic up gradation of any trait as better estimates simultaneously in two biometrical parameter of any trait indicate that the genetic control of the trait is additive in nature( Pham *et al* 2011). The trait Capsule/plant followed by seed yield/plant showed such ideal combination in genetic estimates. High heritability with genetic advance of any trait point out that the trait is under additive genetic control. In self pollinated crop additive gene effect is more desirable than non-additive genetic effect. Capsule/plant showed high GA, high heritability and also GCV. So, capsule/plant exhibits innate potentiality for genetic improvement in sesame. Such combination was not observed in any other trait. Two other traits namely 1000 seed weight and seed yield/plant were characterized by high heritability with moderate genetic advance, moderate heritability with low genetic advance were found for plant height, branches per plant and days to flowering. The results confirm previous finding of Reddy *et al* (2001) , Iqbal and Dasgupta (2015).

Correlation study ( genotypic and phenotypic ) were showed that plant height, number of branches per plant, no of capsule per plant and capsule length were positively and significantly correlated with seed yield per plant ( Table 3). The results are in concurrence with the results of Uzun and Cagirgan (2001), and Sumathi *et al.* (2007) Subramanian and Subramanian (1990). Thus selection of any of these characters would lead to the improvement of seed yield/plant. On the contrary, 1000 seed weight showed negative relationship with seed yield *parsaeian et al.* (2010) .

A few inter- relationships showed significant and positive correlation namely plant height and number of branches/plant, number of capsule/plant and number of primary branches/plant, Plant height and number of capsule/plant, Capsule length and number of capsule/plant, number of capsules per plant and seed yield per plant. Negative interrelationship were found between days to 50% Flowering and 1000 seed weight, days to 50% flowering and capsule length. Days to maturity also had negative effect on capsule length and 1000 seed weight though the effect was not significant.

Combining all traits, selection for plant height would not only improve seed yield/plant but also will improve capsules/plant and branches/plant through correlated response. However in intensive cropping system the breeders are interested for early maturity though delayed maturity improves seed size.

Improvement of the trait capsule/plant would also improve plant height, branches/plant, capsule length and ultimately seed yield/plant. So, capsules/plant was the most desired trait for improvement of several traits and also seed yield/plant. It is interesting to note that capsules/plant had additive genetic control with high GCV, high GA and high heritability.

All 33 genotypes were further analyzed by SPSS for grouping. Three distinct clusters were found. (Figure 1 and Table 4). Cluster I comprised of 23 genotype where as cluster II consisted of 6 genotypes and cluster III consisted 4 genotypes. Four distinct sub-clusters were identified for cluster I. The sub- clusters IA, IB, IC and ID contained 8, 7, 2 and 6 genotypes respectively. Cluster IA has been further grouped into two distinct sub clusters IA-i and IA-ii comprising 4 genotypes and 2 genotypes each. The maximum Euclidean distance recorded between RT-346 and EC-303435(4) followed by EC-204704 and RT-346, IC-20477 and RT-346. Desirable segregates are expected if crossing is done between genotypes with high dissimilarities coefficient.

An investigation of the cluster composition revealed that each of the two clusters consisted of varieties belonging to different origin, i.e. from different states of India and also from different countries. This indicates that genetic divergence of genotypes is independent of geographic origin. Similar findings were also reported by Banerjee and Kole (2009), Kandamoorthy and Govindarasu (2005), Banumathy *et al.* (2010) and Saha *et al.* (2012).

#### IV. Conclusion

Sesame genotypes showed considerable genetic variability and divergence. It was found that additive genetic effect was more prevalent for capsule per plant. Correlation study at phenotypic level showed that seed yield per plant was significantly and positively associated with plant height, number of branches per plant, number of capsules per plant and capsule length, so selection for high number of capsules per plant, capsule length and number of branches per plant will lead towards high yield. The cluster analysis helped in grouping the genotypes into different clusters having specific characteristic traits which may be helpful in selecting parents for future breeding programs. Crossing between the genotypes RT-346, EC-303435(4), EC-204704 and IC-20477 would most likely express considerable amount of heterosis in F1 generation and also provide a wide spectrum of recombinants in segregating generations. Grouping of genotypes based on multivariate analysis was independent of origin of cultivars. The conventional assumption that selecting genotypes of different geographical origin will maximize the diversity available to a breeding project does not follow in sesame.

#### Acknowledgement

We acknowledge the university grants commission for providing the financial support to carry out the study under UGC Major Project on sesame titled Genetic diversity among genotypes and molecular linkage map construction in sesame (*Sesamum indicum L.*) UGC Reference No. F. 42-721/2013(SR)

**Table 1 : List of Genotypes**

Serial No.	Name of genotypes	Origin	Serial No.	Name of genotypes	Origin
1	Savitri	Malda	18	TKG-352	Tikamgarh
2	Rama	Malda	19	TKG-22	Tikamgarh
3	Saheb	West Bengal	20	IC-20477	India
4	Osc-207	Odisha	21	IC-26230	India
5	Cums-20	West Bengal	22	IC-141464	India
6	DSS-09	Karnatak	23	EC-182832(26)	Bulgaria
7	B-76	South 24 parganas	24	EC-334988(3)	Bulgaria
8	V-13	West Bengal	25	EC-303435(4)	Bulgaria
9	V-15	Hoogly	26	EC-161492-A	Bulgaria
10	CST-2001	West Bengal	27	EC-164966(50)	USA
11	Haveri	West Bengal	28	EC-204704	USA
12	Utawadia	West Bengal	29	EC-41923(B)	USA
13	Cums-11	West Bengal	30	EC-303433	USA
14	Cums-9	West Bengal	31	EC-303442(32)	Bulgaria
15	VRI-1	Tamil Nadu	32	EC-100043-A	Bulgaria
16	RT-346	Rajasthan	33	EC-100049(3)	Bulgaria
17	GT-Black	Gujarat			

**Table.2. Estimates of variability, Heritability and Genetic Advance in sesame**

	50% Flowering	Days to Maturity	1000 Seed Weight	Plant Height	Branches per Plant	Capsule Length	Capsule per Plant	Seed yield Per Plant
<b>GCV</b>	5.76	1.97	26.11	14.64	29.31	13.15	62.56	36.37
<b>PCV</b>	7.92	4.34	26.30	20.73	42.21	16.34	67.70	45.32
<b>H%</b>	52.88	20.61	98.57	49.9	48.21	64.79	85.39	64.41
<b>GA</b>	2.69	0.74	1.27	13.57	1.63	0.38	72.09	2.14
<b>Mean</b>	42.98	88.899	2.479(gm)	90.111(cm)	5.596	2.176(cm)	65.576	4.383(gm)
<b>Range</b>	36.33-48.00	83.00-94.67	1.64-3.13	70.33-121.67	2.67-8.00	1.57-3.07	29.00-193.00	4.04-8.22

**Table- 3:** Phenotypic Correlation Matrix

	50% flowering	Days to maturity	1000 Seed weight	Plant Height	Branches per Plant	Capsule Length	Capsule per Plant	Seed yield Per Plant
50% flowering	1.000							
Days to maturity	0.285	1.000						
1000 Seed weight	-0.120	0.049	1.000					
Plant Height	0.353*	0.240	-0.056	1.000				
Branches per Plant	0.149	-0.050	0.058	0.151	1.000			
Capsule Length	0.234	-0.138	-0.153	-0.065	0.044	1.000		
Capsule per Plant	0.232	0.052	-0.026	0.080	0.646**	0.088	1.000	
Seed yield Per Plant	0.613**	0.074	-0.070	0.461**	0.435**	0.414**	0.536**	1.000

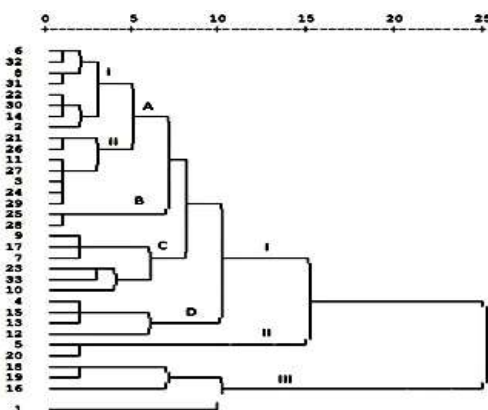
\* and \*\*: significant at 5% and 1% level respectively

**Table 4:** Cluster composition of 33 genotypes

CLUSTER	SUB CLUSTER	NUMBERS OF GENOTYPES	ACCESSION NUMBER
CLUSTER-I	A	i	8 DSS-09, EC-100043-A, V-13, EC-303442(32), IC-141464, EC-303433, Cums-9, Rama
		ii	7 IC-26230, EC-161492-A, Haveri, EC-164966(50), Saheb, EC-334988(3), EC-41923(B),
	B	2 EC-303435(4), EC-204704	
	C	6 V-15, GT-Black, B-76, CST-2001, EC-182832(26), EC-100049(3)	
	D	4 Osc-207, Cums-11, VR-I, Utawadia	
CLUSTER- II		2	Cums-20, IC-20477
CLUSTER- III		4	Savitri, TKG-352, TKG-22, RT-346

**Table 5:** Euclidean distance between 33 genotypes

	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33				
1	0	104	973	524	455	758	519	627	574	514	167	757	412	301	63	527	428	342	247	554	1015	717	675	55	1521	563	122	1034	1071	1017	874	752	732				
2		0	243	155	327	195	421	162	219	200	147	475	237	52	124	244	105	1712	1445	573	2105	125	571	241	246	242	105	129	272	107	210	212	212				
3			0	424	719	312	424	252	201	525	111	711	415	157	151	412	1116	1221	1757	1757	1757	1757	1757	1757	1757	1757	1757	1757	1757	1757	1757	1757	1757				
4				0	475	164	348	312	412	166	214	224	162	229	157	164	414	167	164	177	419	242	219	419	157	425	167	167	169	169	169	169	169	169			
5					0	515	517	311	351	219	115	174	415	115	115	115	115	115	115	115	115	115	115	115	115	115	115	115	115	115	115	115	115	115			
6						0	342	116	112	219	219	412	147	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112			
7							0	244	154	419	419	119	219	147	147	147	147	147	147	147	147	147	147	147	147	147	147	147	147	147	147	147	147	147			
8								0	571	311	115	174	115	115	115	115	115	115	115	115	115	115	115	115	115	115	115	115	115	115	115	115	115	115	115		
9									0	342	219	115	174	115	115	115	115	115	115	115	115	115	115	115	115	115	115	115	115	115	115	115	115	115	115		
10										0	424	217	51	167	214	164	164	164	164	164	164	164	164	164	164	164	164	164	164	164	164	164	164	164	164	164	
11											0	167	424	114	117	117	117	117	117	117	117	117	117	117	117	117	117	117	117	117	117	117	117	117	117		
12												0	212	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	
13													0	167	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	
14														0	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	
15															0	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	
16																0	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	
17																	0	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	
18																		0	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	
19																			0	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	
20																				0	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	
21																					0	112	112	112	112	112	112	112	112	112	112	112	112	112	112	112	
22																						0	112	112	112	112	112	112	112	112	112	112	112	112	112	112	
23																							0	112	112	112	112	112	112	112	112	112	112	112	112	112	112
24																								0	112	112	112	112	112	112	112	112	112	112	112	112	112
25																									0	112	112	112	112	112	112	112	112	112	112	112	112
26																										0	112	112	112	112	112	112	112	112	112	112	112
27																											0	112	112	112	112	112	112	112	112	112	112
28																												0	112	112	112	112	112	112	112	112	112
29																													0	112	112	112	112	112	112	112	112
30																														0	112	112	112	112	112	112	112
31																															0	112	112	112	112	112	112
32																																0	112	112	112	112	112
33																																	0	112	112	112	112



**Fig: 1** Dendrogram showing the distribution of 33 genotypes in clusters

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