

Research Article

Assessment of genetic diversity among identified testers aids in development of hybrids in sesame (*Sesamum indicum*.L)

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Abstract

Diversity analyses on ninety two genotypes of sesame using ten morphological characters were studied. Results indicated that ten clusters were formed wherein maximum inter cluster distance was observed between cluster IX and X (239.87). Simultaneously crossing of these 90 genotypes with Co-1 and TMV-5 was carried out. The 180 hybrids generated using two lines (CO 1 and TMV 5) with these 90 genotypes, upon evaluation, indicated five elite hybrids namely TMV 5 X bardur local (B)(217.49 percent), TMV 5 X bardur local (W) (198.38 percent), CO 1 x RT 172 (163.47 percent), TMV 5 x Rama (161.04 percent) and TMV 5 x MT-10-23-3 (155.57 percent) significantly yielded than the standard check TMV 7 and their male parents were found to be from the cluster I which were moderately divergent (19.09) among them.

Key words

Sesamum, Diveristy and Heterosis

Introduction

Sesamum indicum L (2n= 26), an annual plant belonging to Pedaliaceae family is considered as the queen of the oilseeds originated from Africa. Kobayashi(1983). Its seed comprises 50 per cent oil with monounsaturated fatty acids especially oleic acid Gandhi(2009) that helps to keep the LDL lower. Sesame is rich in sesamol and sesaminol that eliminates harmful free radicals and the oil fraction shows a remarkable stability to oxidation. It is also a good source of B-complex such as niacin, folic acid, thiamine pyridoxine and riboflavin Borchani *et al.*(2010). India ranks second in production after China with a productivity of 413 kg/ha, cultivated in an area of 17.90 lakh hectares with production of 8.02 lakh tonnes NMOOP (2017). In India, West Bengal is the leading producer followed by Assam and Tamil Nadu. Despite the health benefits and highest area under cultivation, sesame ranks sixth in the world's production of oilseeds and twelfth for vegetable oil production Weiss(2001). This leaves a hope of improving sesame crop which can be attempted through heterosis breeding.

Information on genetic diversity and relationship among populations is important for plant breeding programs as it helps to select the right genetic material to be used Ganesh and Thangavelu, (1995). The analysis of genetic diversity using D^2 statistics enables one to discriminate between different cultivars according to the diversity present Mahalanobis (1936). Based on the magnitude of variation present in the population, individual character's contribution to the total spectrum of

variability differs. Number of capsules per plant and seed yield contributed mostly to divergence in many of the studies carried out Arriel *et al.*(2007). Sesame being a less explored crop when compared to the rest of the oilseeds need to be well studied and inculcating the production of hybrids may foster the productivity. In order to exploit the variability existing in the crop and produce superior hybrids this study was taken up by utilizing the diverse parents with good combiners of yield.

Materials and Methods

The experimental material consisted of 90 genotypes of sesame collected from different research stations all over India. These genotypes were utilized as male parents with two lines namely CO 1 and TMV 5 as female parents. Crossing programme was taken up in a Line x Tester fashion. All the 92 lines were first planted in two replications in Randomised Block Design and D^2 Mahalanobis (1936) analysis was done. Simultaneously, these 90 lines were crossed with CO1 and TMV 5 and their F_1 's were estimated for standard heterosis with TMV 7. Hybridisation was carried out by using the standard method of emasculation and dusting of pollens Ganesh and thangavelu (1995) in Line x Tester mating design. Ten characters namely days to first flowering, distance from base to capsule, number of branches, number of capsules in main stem, number of capsules in branches, capsule length, number of seeds per capsule, single plant yield, days to maturity and thousand seed weight were observed.

Results and Discussion

The morphological observations of 92 genotypes were analysed for their variability and diversity by Mahanobis (1936). The plant morphological characters observed showed considerable variation for all the traits (Table 5). The genotypes were grouped into ten clusters and their grouping depended on percentage contribution of characters wherein 59 per cent was contributed by number of capsules in main stem and 11.79 per cent by days to first flowering as shown in Table 2. The distribution pattern of genotypes indicated that genetic divergence was not based on their geographical differentiation as in accordance with Bandila *et al.* (2011).

According to Ghadri *et al.* (1984), increasing the parental distance implies a great number of contrasting alleles at the desired loci. Maximum genetic recombination is expected from the hybridization of the parents selected from distant groups. In the 10 clusters, the maximum inter cluster distance was observed between cluster IX and X (239.87) followed by cluster VII and IX (226.91) as reported in Table 3.

Hence, genotypes from clusters IX (China and LT 8), X (CO 1) and VII (JCS 2454) were highly diverse genotypes and crossing between them may yield hybrids.

Regarding the intra cluster values, Cluster IX had the maximum intra cluster (45.91) distance as compared to cluster I and III (Table 3). Although only two genotypes were found to be in the cluster, these genotypes China and LT 8 (cluster IX) varied in number of capsules in main stem and flowering which was found to contribute more towards divergence. The cluster means for 10 characters spatially varied among genotypes and clusters, the maximum mean values were observed in cluster VII with high values for number of capsules in main stem and branches, number of branches, number of seeds per capsule, single plant yield and thousand seed weight (Table 4).

Cluster VII had a single genotype JCS 2454 and this genotype was found to be completely deviant from other clusters indicating its potential use as a parent in hybridization programmes for realizing higher single plant yield (20.80) and early duration (99) on an average. Though 77 genotypes clustered in single cluster, their mean performance compared to the other clusters were poor indicating the grouping of poor performers under a single cluster. The cluster VII (JCS 2454), IV (JCS 2498) and V (CST-2008-2) had higher cluster means for single plant yield as well as thousand seed weight and hence could be used in improvement of yield parameters. The cluster IX comprised China which

is a monostem variety with highest capsule length and number of seeds per capsule coupled with higher number of capsules, could be used as a parent in hybridization for breeding monostem varieties with moderate yield. Although Variety Co 1 in cluster X showed highest mean for duration, number of capsules in branches, capsule length it had moderate values for single plant yield indicating that the shattering losses while harvesting as a constraint.

This study revealed high variability for ten characters studied with maximum genotypic and phenotypic variations for number of capsules in branches followed by distance from base to first capsule and number of seeds per capsule as mentioned in Table 6. As these characters recorded high GCA coupled with GA, selection for these characters would be effective as they are controlled by additive gene action Burton & Devane (1953). These similar results were also found in the findings of Tripathi *et al.* (2013) and Parimala *et al.* (2013).

Kiranmayi *et al.* (2014) stated that the difference between GCV and PCV indicates the level of environmental influence whereas this experiment showed narrow differences for all 10 traits. Therefore, high genetic variability present for these characters in the given population can be ardently exploited for breeding sesame with more number of capsules in branches along with more seeds per capsule.

GCV coupled with high heritability would be more valuable in selection programmes Burton & Devane (1953). Number of capsules in branches, distance from the base to first capsule, number of capsules in main stem, seed yield per plant and days to first flowering exhibited high heritability and GCV indicating the predominance of additive gene action and selection for these characters is effective to improve seed yield by practicing simple selection. Genetic advance has considerable importance as it indicates the magnitude of the expected genetic gain from one cycle of selection Burton & Devane (1953). In the present investigation, all the traits except days to maturity, capsule length and number of seeds per capsule recorded high genetic advance as percent of mean indicating the scope of selection for all these characters.

Among the 180 crosses obtained, five of them exhibited heterosis against the standard check TMV 7 (Table 6). The female parent of four of these hybrids is TMV 5 inferring the potential of TMV 5 as a general good combiner for seed yield which was also acclaimed by Vidhyavathi *et al.* (2005).

The elite hybrids possessed TMV 5 as their female and male parents from the clusters I, II and X. Although their genetic distance were not found to be wider, it exhibited heterosis for yield which might be due to combination of favourable alleles from the parents.

Among the five hybrids, TMV 5 x Bardur local (B) exhibited higher percentage of heterosis and overall performance of the successful hybrids as given in the table 7. These hybrids can be test verified further for stability of their performance over environments in future.

References

- Arriel.N.H.C, A.OrlandoDi-Mauro, E.F.Arriel, S.Helena, Unêda-Trevisoli, M.M.Costa, I.M.Bárbaro, and F.R.S.Muniz.2007.Genetic divergence in sesame based on morphological and agronomic traits,Crop Breed.Appl .*Biotechnol.*,**7**: 253-261.
- Bandila.S, A.Ghanta, S.Natarajan, Sivaramakrishnan and Subramaniam .2011. Determination of Genetic Variation in Indian Sesame (*Sesamum indicum* L.) Genotypes for Agro-Morphological Traits, *J.Agric Res.***7** (2):88-99.
- Borchani.S, Besbes, Ch. Blecker, and H. Attia.2010.Chemical characteristics and oxidative stability of sesame seed, sesame paste and olive oils *J. Agr. Sci. Tech.*,**12**: 585-596.
- Burton, W.G. and E.H. Devane. 1953. Estimating heritability in tall fescue (*Festuca arundinacea*) from replicated clonal material. *Agron. J.*, **45**: 478-481.
- Gandhi A.P. 2009. Simplified process for the production of sesame seed (*Sesamum indicum* L.) butter and its nutritional profile. *Asian journal of Food and Agro Industry.*,**2**: 24-27
- Ganesh S. K. and S. Thangavelu 1995. Genetic divergence in sesame (*Sesamum indicum* L.). *Mad. Agric. J.*, **82**: 263–265.
- Ghadri.H,P. J.Shea,G.A.Wicks and L.C Harderli, Atrazine dissipation conventional-till and no-till sorghum. *J Environ.Qual.***13**:549-552
- Kiranmayi.L.S, V.Roja, N.Sivaraj and S.Sivaramakrishnan.2014.Morphological and molecular diversity analysis in Sesame (*Sesamum indicum* L.), *Agrotechnol.***2**:4.
- Kobayashi, T. 1983. Type diversity and evolution in Sesamum (*Sesamum indicum* L.), In: Eds: Zerihun, *Proc.IV. SABRAO Congress*, Malaysia p: 361–366.
- Mahalanobis, P.C.1936. On the generalized distance in statistics. In: *Proc. Natl. Inst. Sci.*,**2**: 49-55.
- Parimala K, I.Swarnalatha Devi, V.Bharathi, B.Raghu, K.Srikrishnalatha and A.Vishnuvardhan Reddy.2013. Heterosis for yield and its component traits in sesame (*Sesamum indicum* L.), *International journal of Applied biology and Pharmaceutical technology.*, **4**(4):65-68.
- Tripathi.A, R.Bisen P.Ravindra. Ahirwal, S.Paroha, R.Sahu and A. R. G. Ranganatha.2013. Study on genetic divergence in sesame (*Sesamum indicum* L.) germplasm based on morphological and quality traits, *The Biosacan.*, **8**(4): 1387-1391.
- Vidhyavathi.R, N. Manivanan, and V. Murlidharan. 2005. Line x Tester analysis in sesame (*Sesamum indicum* L.).*Indian J.Agric.,sci.*,**39** (3):225-228.
- Weiss. EA. 2000. *Oil Seed Crops*. 2nd Edition Blackwell Longman Group Ltd. USA.

Table 1. List of parents and their place of collection

S.NO	PARENTS	PLACE	S.NO	PARENTS	PLACE
1.	AT 178	AMRELI	48.	Jinbakkia	VRIDHACHALAM
2.	AT 222	AMRELI	49.	OMT-30	VRIDHACHALAM
3.	AT 229	AMRELI	50.	ORM-17	VRIDHACHALAM
4.	AT 231	AMRELI	51.	KMR-77-1	VRIDHACHALAM
5.	AT 238	AMRELI	52.	NIC 8984	VRIDHACHALAM
6.	China	AMRELI	53.	Nirmala	VRIDHACHALAM
7.	Wild dhanuka	AMRELI	54.	N 32	VRIDHACHALAM
8.	JR 22	AMRELI	55.	VS-07-023	VRIDHACHALAM
9.	RSS 106	AMRELI	56.	GPC-12-06	JAGTIAL
10.	RSE-3	AMRELI	57.	JCS 1020	JAGTIAL
11.	Shelna 5	AMRELI	58.	JCS 1942	JAGTIAL
12.	U-76-10	AMRELI	59.	JCS 1935	JAGTIAL
13.	IS 196	AMRELI	60.	JCS 2454	JAGTIAL
14.	B-90-1	AMRELI	61.	JCS 2498	JAGTIAL
15.	Vardi local	JALGAON	62.	Madhavi	JAGTIAL
16.	Amalgaon sel	JALGAON	63.	JCS 1673	JAGTIAL
18.	JL-SG-06-6	JALGAON	64.	JCS 1860	JAGTIAL
19.	JLT 501	JALGAON	65.	JCS 2698	JAGTIAL
20.	JLS-510-3	JALGAON	66.	Chandhana	JAGTIAL
21.	Gopi	JALGAON	67.	Haryana til-2	JABALPUR
22.	NIC 7913	JALGAON	68.	PT 1	JABALPUR
23.	KMR 18	JALGAON	69.	Brijeshwari	JABALPUR
24.	M-2	JALGAON	70.	T 78	JABALPUR
25.	JLS-110-12	JALGAON	71.	TC-289	JABALPUR
26.	IC 413187	JALGAON	72.	Amrit	JABALPUR
27.	IC 204484	DHARWAD	73.	Usha	JABALPUR
28.	E8	DHARWAD	74.	Kalaika	JABALPUR
29.	TKG 377	DHARWAD	75.	TKG-306	JABALPUR
30.	DS 21	DHARWAD	76.	JT-7	JABALPUR
31.	DS 11	DHARWAD	77.	Thilathara	JABALPUR
32.	DS 7	DHARWAD	78.	Rama	JABALPUR
33.	RT 311	MANDOR	79.	Savitri	JABALPUR
34.	RT 321	MANDOR	80.	Bardur local(W)	DHARWAD
35.	RT 194	MANDOR	81.	Bardur local(B)	DHARWAD
36.	RT 201	MANDOR	82.	MT-10-23-3	DHARWAD
37.	LT 8	MANDOR	83.	PKVNT 11	DHARWAD
38.	SI 3160	MANDOR	84.	Shekar	DHARWAD
39.	CST-2001-1	MANDOR	85.	DS 3	DHARWAD
40.	CST-2008-2	MANDOR	86.	RT 172	MANDOR
41.	SP 44	MANDOR	87.	RT 180	MANDOR
42.	HT 9713	MANDOR	88.	RT 202	MANDOR
43.	PAIYUR 1	VRIDHACHALAM	89.	RT 209	MANDOR
44.	TNAU 64	VRIDHACHALAM	90.	RT 215	MANDOR
45.	Veppur local	VRIDHACHALAM	91.	TMV 5	VRIDHACHALAM
46.	Thumkur local	VRIDHACHALAM	92.	CO 1	COIMBATORE
47.	Cardeberga	VRIDHACHALAM			

Table 2. Percentage contribution of characters

S.No	Character	Contribution (%)
1	Days to first flowering	11.79
2	Days to maturity	9.01
3	Distance from the base to first capsule	0.15
4	No. of primary branches	5.19
5	No. of capsules in main stem	59.10
6	No. of capsules in branches	0.12
7	Capsule length	0.02
8	No. of seeds per capsule	0.02
9	Seed yield /plant	11.49
10	Thousand seed weight	3.10

Table 3. Inter and intra cluster D² values and cluster distances

	I	II	III	IV	V	VI	VII	VIII	IX	X
I	19.09	25.63	37.24	25.98	58.01	36.88	76.26	41.39	160.70	88.80
II		0	23.34	35.13	42.07	34.41	61.35	39.61	180.35	68.59
III			21.96	45.20	31.39	30.91	50.49	50.08	190.93	64.39
IV				0	61.79	43.23	82.05	41.19	154.31	93.50
V					0	36.31	27.24	62.40	210.81	49.35
VI						0	51.69	46.07	178.84	75.24
VII							0	82.58	226.91	54.83
VIII								0	167.03	74.85
IX									45.91	239.87
X										0

Table 4. Cluster means for different characters of 90 Sesame lines

Cluster	DFE	DB	NPB	NMCP	NBRCP	CPL	NSCP	DM	SPYLD	TSDWT
I	37.20	41.38	4.25	28.62	77.14	2.44	59.07	100.11	13.88	3.49
II	34.50	45.79	4.00	25.25	86.39	2.40	58.00	100.50	13.12	2.68
III	44.08	41.47	4.67	30.29	92.75	2.44	60.08	106.42	14.96	4.03
IV	40.00	46.92	3.50	28.61	73.11	2.55	60.00	114.00	23.13	3.20
V	47.00	44.79	4.50	41.94	104.29	2.81	71.50	119.00	20.47	3.11
VI	47.00	42.62	5.00	50.03	89.80	2.65	68.00	105.50	15.37	3.36
VII	34.50	34.71	5.25	53.53	115.22	2.80	76.00	99.00	20.80	4.20
VIII	45.00	82.22	4.75	32.28	79.39	2.85	69.00	112.00	11.53	3.55
IX	36.00	42.51	0.00	42.77	0.00	2.85	64.00	100.00	14.18	2.94
X	41.50	83.69	5.00	30.49	116.60	2.90	70.00	118.00	13.41	4.18

*DFE : DAYS TO FIRST FLOWERING

DB : DISTANCE FROM THE BASE TO FIRST CAPSULE (cm)

NPBR : NUMBER OF PRIMARY BRANCHES

NMCP : NUMBER OF CAPSULES IN MAIN STEM

NBRCP : NUMBER OF CAPSULES IN BRANCHES

CPL : CAPSULE LENGTH (cm)

NSCP : NUMBER OF SEEDS PER CAPSULE

SPYLD: SEED YIELD PER PLANT (g)

TSDWT: THOUSAND SEED WEIGHT (g)

Table 5. ANOVA for the mean performance of 90 genotypes

Source of Variation	SS	Df	MS	F	P-value	F crit
Genotypes	7706.856	89	86.59388	2.145296	3.68E-08	1.27805
Traits	928480.8	9	103164.5	2555.821	0	1.891552
Error		32332		801		40.36454
Total			968519.6		899	

Table 6. Co-efficient of Variation, Heritability (broad sense) and Genetic advance in 92 sesame accessions

S.No	Characters	PCV%	GCV%	ECV%	H ² (%)	GA	GA%
1.	Days to first flowering	29.73	28.67	1.05	96.44	10.83	28.54
2.	Days to maturity	51.63	39.48	12.14	76.48	11.32	11.17
3.	Distance from the base to first capsule (cm)	57.13	55.92	1.20	97.88	15.24	35.92
3.	No. of primary branches	0.65	0.54	0.10	83.92	1.39	33.17
4.	No. of capsules in main stem	30.40	29.70	0.69	97.71	11.09	37.32
5.	No. of capsules in branches	214.31	214.03	0.28	99.87	30.11	38.68
6.	Capsule length (cm)	0.13	0.07	0.05	58.41	0.43	17.70
7.	No. of seeds per capsule	86.74	45.96	40.77	52.99	10.16	16.97
9.	Seed yield /plant (g)	7.01	6.77	0.24	96.57	5.26	37.12
10.	Thousand-seed weight (g)	0.41	0.36	0.04	88.91	1.18	33.65

PCV (%) - Phenotypic coefficient of variation

GCV (%) - Genotypic coefficient of variation

ECV (%) - Environmental coefficient of variation

H² - Heritability

GA - Genetic advance

GA (%) - Genetic advance as percent of mean

Table 7. Overall performance of superior five heterotic crosses for yield and other component characters

HYBRIDS	DFP	DB	NPB	NMCP	NBRCP	CPL	NSCP	TSWD	DM
TMV5 x Bardur local (B)	2.47	29.64*	0.00	22.72*	33.77**	16**	28.57**	11.89**	2.40**
TMV5 x Bardur local (W)	2.47	69.94**	16.67**	7.62	21.05*	0.00	14.29**	-15.47*	0.48
CO1 x RT172	8.64* *	36.32*	4.17	49.77*	-52.93*	4.00*	21.43**	12.97**	19.71**
TMV5 x Rama	2.47	17.11	50.00**	19.15*	27.81	0.00*	7.14	-15.14**	10.58
TMV5 x MT 10-23-3	12.35**	2.59**	29.17*	60.42*	74.30*	-8.00	-3.57	5.41	1.44