

Original Research Article

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Study of Genetic Divergence in Soyabean (*Glycine max.* (L.) Merrill) Germplasm

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ABSTRACT

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Genetic divergence of 40 soybean genotypes was studied for eleven quantitative traits using Mahalanobis's D² statistics during *Kharif*-2019 in Randomized Block Design with three replications. Analysis of Variance showed significant differences for all characters under study at 1% and 5% Level of Significance. High estimates of GCV and PCV were observed for number of pods per plant followed by seed yield per plant. High heritability coupled with high genetic advance was recorded for plant height followed by test weight. Based on the relative magnitude of D² values, the genotypes were grouped into nine clusters. Cluster III constituted maximum number of genotypes with 19 genotypes followed by cluster I. Maximum intra-cluster distance was found in cluster III followed by cluster II. Maximum inter-cluster distance (D²) was observed between cluster III and V. Test weight contributed maximum towards genetic divergence. Based on inter cluster distances and *Per se* performance, the genotypes TNAU 20051, MAUS 128 and KB 17 with test weight traits should be given top priority during selection of the genetically divergent parents in future breeding program.

Introduction

Soybean (*Glycine max* (L.) Merrill) popularly known as “Miracle crop” is a member of family Leguminosae and is widely used as oilseed crop. Soybean is a self-pollinated crop having chromosome number of $2n = 40$. Soybean is a very good source for both protein and oil as it has protein content of 40 to 42% which is highest among food crops and 18 to 22% oil content comprising 85% unsaturated fatty acids and free from

cholesterol, therefore it is highly desirable in the human diet (Antalina *et al.*, 1999). Global area under soybean cultivation is around 124.90 million hectares and 350.0million ton production where USA, Brazil, Argentina, China and India accounts more than 85% acreages and 88.5% of production among all soybean growing countries (FAO, 2017-18). India ranks 4th in terms of soybean area but 5th in terms of production after USA, Brazil, Argentina and China, however productivity of soybean in India is very low (1.04 t/ha) as

compared to average global productivity (2.9 t/ha). During 2017-18 the production of soybean in India was 10.98 million tons from 10.47 million-hectare area with productivity of 1.04 t/ha (Directorate of Economics & Statistics, DAC&FW). The major soybean growing states in India are Madhya Pradesh, Maharashtra and Rajasthan which accounts around 93% in terms of both acreages and production among all soybean growing states. Area under soybean in Uttar Pradesh (UP) is currently very low and so is the case of productivity. Looking to very low productivity of soybean in India, it's a need to improve its productivity by both genetical and agronomical means where genetical means of improving productivity of any crops play a major role.

Genetically diverse parent is a pre-requisite to improve the chances of selecting better segregants for various characters. When such parents are utilized in hybridization program, they are likely to produce high heterotic effect and wide spectrum of variability (Barhet *et al.*, 2014). The assessment of genetic diversity is important not only for crop improvement but also for efficient management and conservation of germplasm resources. Keeping this in view, the present investigation was carried out to assess the nature and magnitude of genetic diversity in soybean germplasm.

Materials and Methods

The experiment was carried out in the Field Experimentation Centre, Department of Genetics and Plant Breeding, Naini Agricultural Institute, Sam Higginbottom University of Agriculture, Technology and Sciences, Prayagraj, U.P, India. The experimental materials for the present study consisted of 40 genotypes of Soybean including 1 local check was procured from Indian Institute of Soybean Research (IISR),

Indore. The seeds of all soybean germplasm were planted in experimental field in a Randomized Block Design (RBD) trial with three replications. All necessary precautions were taken to maintain uniform plant population in each treatment and replications. All the recommended package of practices was followed along with necessary prophylactic plant protection measures to raise a good crop. Observations were recorded at different stage of plant growth and at harvest. The data was subjected to statistical analysis. The variability was estimated as per procedure for analysis of variance suggested by Fischer (1936), Phenotypic Coefficient of Variance (PCV) and Genotypic Coefficient of Variance GCV were calculated using formula given by Burton (1952), heritability in broad sense (h^2) as suggested by Burton and De Vane (1953), genetic advance i.e., the expected genetic gain were calculated by using the procedure given by Johnson *et al.*, (1955), and genetic divergence was calculated by using D^2 Statistics given by Mahalonobis (1936).

Results and Discussion

Analysis of variance showed significant differences among 40 soybean genotypes for all the characters under study. Similar variations in soybean have been also reported by Karad *et al.*, in 2005. The Phenotypic Coefficient of Variation were higher in magnitude than the Genotypic Coefficient of Variation for all the characters and the difference between PCV and GCV was low for most of the characters, this indicates less influence of environment in the expression of these characters in soybean germplasm. Similar information was also reported in soybean by Chandrawat *et al.*, in 2017. The magnitude of GCV and PCV was highest for no. of pods per plant (35.11&35.58). The heritability estimates were found to be high (more than 70%) for all the characters except

numbers of seeds per pod (65.00). High heritability coupled with high genetic advance as per cent mean (>20) in the present genotypes were recorded for plant height (98.60% and 49.61%) indicating

predominance of additive gene effects and the possibilities of effective selection for the improvement of these characters (Fig. 1 and 2).

Table.1 Area, production and Productivity of soybean during 2017-2018

Particulars	Area (Mha)	Production (Mt)	Productivity (t/ha)
World	124.90	348.71	2.79
India	10.47	10.98	1.04

Source: FAO Statistics, 2017-18

Table.2 List of soybean genotypes used in the present investigation

S. No.	Name of Genotype	S. No.	Name of Genotype	S. No.	Name of Genotype	S. No.	Name of Genotype
1.	NRC 37	11.	EC 15961	21.	UPSM 1087	31.	SL 432
2.	EC 251541	12.	JSM 245	22.	SL 752	32.	LEE 75
3.	UPSM 55	13.	SQL 37	23.	NRC 801	33.	TG 5404E
4.	PK 258	14.	JSM 195	24.	TNAU 20051	34.	MACS 1259
5.	RKS 54	15.	PK 701	25.	PS 1467	35.	PK 1038
6.	UGM 75	16.	EC 251682	26.	PS 1336	36.	RKS 48
7.	JSM 232	17.	JS 9631	27.	PS 1475	37.	TNAU 2024
8.	EC 381884	18.	KB 17	28.	JSM 222	38.	MAUS 128
9.	NRC 2006M	19.	UPSL 152	29.	SL 744	39.	G 91
10.	UPSM 57	20.	UPSM 77	30.	MACS 171	40.	JS 2029 (Check)

Table.3 Analysis of Variance for different quantitative characters in soybean genotypes

Characters	Mean sum of squares		
	Replication(d. f.= 2)	Treatments (d. f.= 39)	Error (d. f.= 78)
Days to 50% flowering	0.433	44.294**	1.322
Plant height	4.662	809.366**	3.935
No. of primary branches	0.012	2.652**	0.030
No. of nodes per plant	0.026	11.122**	0.095
No of clusters per plant	0.156	2.916**	0.117
No. of pods per plant	2.561	906.717**	8.116
No. of seeds per pod	0.009	0.115**	0.017
Biological yield	6.939	90.952**	3.953
Harvest index	7.252	69.992**	5.253
Test weight	0.231	15.858**	0.09
Seed yield per plant	0.02	6.173**	0.062

*Indicates significant at 5 % level of significance

** Indicates significant at 1% level of significance

Table 4 Estimation of genetic variability parameters for different quantitative characters

S.No.	Characters	GCV	PCV	H ² (%)	GA (5%)	GA as % of mean (5%)
1	Days to 50% flowering	8.22	8.59	91.50	7.46	16.19
2	Plant height (cm)	24.26	24.44	98.60	33.51	49.61
3	No. of primary branches	26.49	26.94	96.70	1.89	53.64
4	No. of nodes per plant	16.07	16.27	97.50	3.90	32.68
5	No. of clusters per plant	15.15	16.08	88.80	1.88	29.41
6	No. of pods per plant	35.11	35.58	97.40	35.18	71.36
7	No. of seeds per pod	7.50	9.31	65.00	0.30	12.45
8	Biological yield	22.67	24.16	88.00	10.41	43.09
9	Harvest Index	21.35	23.81	80.40	8.58	39.45
10	Test weight	22.59	22.78	98.30	4.68	46.14
11	Seed yield per plant	28.13	28.56	97.00	2.90	57.09

Table.5 Distribution of genotypes into different clusters

Cluster	Number of genotypes	Genotypes
I	11	NRC 37, PK 258, EC 15961, PK 701, SL 752, MACS 171, SL 432, LEE 75, PK 1038, RKS 48, G 91
II	4	UPSM 57, UPSM 77, UPSM 1087, UPSL 152
III	19	UPSM 55, RKS 54, UGM 75, JSM 232, EC 381884, NRC 2006M, JSM 245, JSM 195, JS 9631, KB 17, NRC 801, PS 1467, PS 1336, PS 1475, JSM 222, SL 744, TG 5404E, MACS 1259, TNAU 2024
IV	1	SQL 37
V	1	MAUS 128
VI	1	EC 251682
VII	1	TNAU 20051
VIII	1	EC 251541
IX	1	JS 2029

Table.6 Cluster mean value for 11 characters in soybean genotypes

Cluster	Days to 50% flowering	Plant height (cm)	No. of Primary Branches	No. of Nodes per plant	No. of clusters per plant	No. of Pods per plant	No. of seeds per pod	Biological yield per plant (g)	Harvest Index (%)	Test weight (g)	Seed yield per plant (g)
I	46.36	82.43	3.92	12.22	6.19	43.73	2.41	22.86	22.02	11.26	4.94
II	52.42	78.98	4.25	14.32	7.50	74.97	2.48	33.06	21.47	8.80	7.03
III	44.70	54.32	2.91	10.71	5.93	41.08	2.42	20.31	21.67	9.62	4.34
IV	41.00	57.17	4.00	9.27	6.13	40.47	2.33	21.30	29.98	14.30	6.33
V	50.00	90.80	5.00	13.87	8.40	90.80	2.33	31.02	21.75	12.17	6.72
VI	46.33	52.97	2.87	14.00	7.40	59.00	2.07	31.02	23.29	12.16	7.18
VII	46.33	74.17	5.00	12.00	8.47	95.93	2.47	31.37	28.95	11.05	9.07
VIII	49.00	95.87	4.00	16.00	6.67	39.27	2.00	31.30	14.74	6.41	4.59
IX	41.00	76.20	4.93	16.93	7.20	84.93	2.47	34.83	11.67	8.04	4.04

Table.7 Intra and inter cluster distance in soybean

Clusters	I	II	III	IV	V	VI	VII	VIII	IX
I	80.15	258.69	259.64	175.21	306.51	193.56	430.95	270.5	343.62
II		105.45	470.2	490.07	192.86	202.38	201.17	225.49	202.78
III			124.68	243.84	748.16	207.8	744.29	431.26	489.74
IV				0	493.55	221.3	507.92	659.09	636.67
V					0	402.44	101.63	503.18	346.61
VI						0	363.74	320.51	336.75
VII							0	650.06	438.75
VIII								0	182.95
IX									0

Fig.1 Clustering by Tocher's method

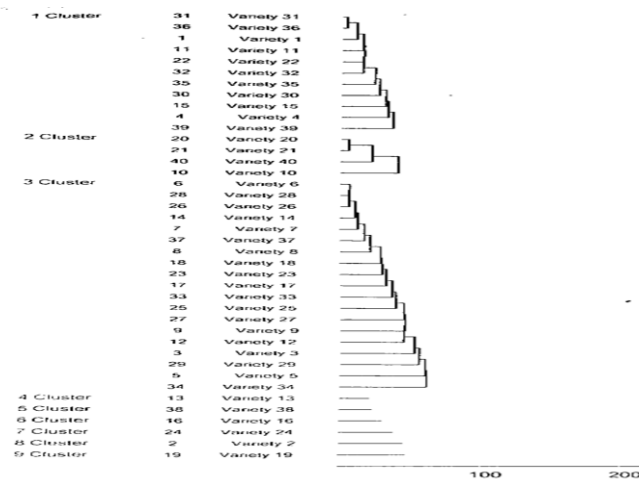
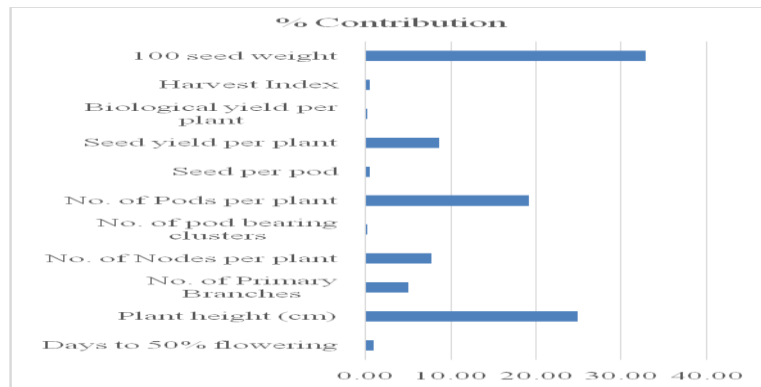


Fig.2 Contribution of different characters towards divergence



On the basis of Mahalanobis D₂ statistics, the 40 soybean genotypes were grouped into 9 clusters. The maximum number of genotypes

was included in cluster III comprising of 19 genotypes followed by cluster I with 11 genotypes, cluster II with 4 genotypes and

clusters IV, V, VI, VII, VIII and IX with one genotype each indicating a high degree of heterogeneity among the genotypes. The maximum intra-cluster distance was registered for, cluster III (124.68), followed by cluster II (105.45) and cluster I (80.15).

The minimum intra cluster distance was observed in cluster IV, VII, VIII, IX (0.00). The inter-cluster distance (D2) was found maximum between cluster III and V (748.16) followed by the distance between cluster III and VII (744.29). Therefore, the genotypes present in these clusters should be given priority during selection of parents for further hybridization programmes.

It is concluded that Analysis of Variance showed significant differences for all characters under study at 1% and 5% Level of Significance. High estimates of GCV and PCV were observed for number of pods per plant followed by seed yield per plant. High heritability coupled with high genetic advance was recorded for plant height followed by test weight. Based on the relative magnitude of D2 values, the genotypes were grouped into nine clusters. Cluster III constituted maximum number of genotypes with 19 genotypes followed by cluster I. Maximum intra-cluster distance was found in cluster III followed by cluster II. Maximum inter-cluster distance (D2) was observed between cluster III and V. Test weight contributed maximum towards genetic divergence. Based on inter cluster distances and *Per se* performance, the genotypes TNAU 20051, MAUS 128 and KB 17 with test weight traits should be given top priority during selection of the genetically divergent parents in future breeding program.

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