Short Communication



Genetic diversity in germplasm of mulberry (*Morus* spp.) on root proliferation

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Exploitation of heterosis is considered as an outstanding success of plant breeding. The magnitude of heterosis gained by the hybrids largely depends on the genetic divergence between the parents. This information facilitates the breeders in selection of parents for crossing and development of hybrids. The genetic diversity of the parents is not always related with the geographical diversity [1]. Assessment of genetic divergence for parents' selection is necessary for which statistical procedures, such as, D² statistics and nonhierarchical Euclidean cluster analysis [2, 3] is considered. Development of superior varieties with higher survival is one of the important factors in mulberry breeding. In sericulture, mulberry (Morus spp.) plays the key role due to the monophagous habit of the silkworm (Bombyx mori L.) where production and guality of silk largely depend on the quality of mulberry. Vegetative nature of mulberry propagation allows perpetuation of the characters once it is introduced through crossing. In vegetative propagated plants, rooting ability is the most important criteria, since a well developed root system determines the utilization of nutrients from the soil for growth and development [4]. As a large number of indigenous and exotic mulberry germplasm accessions are available, study of genetic divergence of the parents are necessary to identify genetically diverse genotypes with higher adaptability for selection and utilization in breeding programme.

Fifty-six indigenous and 29 exotic mulberry accessions collected from 12 different countries. maintained in the germplasm bank (GPB) at Central Sericultural Research and Training Institute, Berhampore West Bengal, India (24º6'N and 88º15'E) were studied. The experiment was conducted during 2000-2001 in the polythene tube condition with three replications. Stem cuttings of each accession were planted in the polythene tube and nurtured. On 90th day of cutting plantation, plants were gently removed from the soil and root proliferations viz., number of roots, root length, fresh and dry weight of roots, root hair zone and root volume were recorded. Data were subjected to multivariate analysis utilizing Mahalanobis D² statistic [2] and the accessions were grouped into different clusters following Tocher's method of cluster formation [5].

The analysis of variance showed significant difference among the accessions irrespective of their country of origin. Using Tocher's procedure, the accessions were grouped into 6 clusters. Among these, cluster I was the largest and consists 66 accessions followed by cluster II with 10 accessions and cluster III with 6 accessions. Clusters IV, V and VI were unique, since each had only one accession (Table 1). The random distribution of the accessions was evident from cluster I having maximum number with diverse distribution. The grouping pattern did not show any relationship between genetic divergence and geographic diversity [6]. The inclusion of accessions of different origin together in the same cluster indicated the ancestral relationship among the accessions. Average inter-cluster divergence among the accessions showed highest value (77.26) between the clusters IV and VI followed by clusters II and V (72.70), clusters I and V (66.85), clusters V and VI (66.18) and clusters III and IV (65.71) whereas the least value (29.42) was observed between the clusters I and III (Table 2).

The data on intra- and inter-cluster distances are given in Table 3. The D² values among 85 accessions revealed that cluster III showed minimum intracluster distance (3.40), which indicate that the accessions within the cluster were similar. While cluster II showed maximum D^2 value (3.74) followed by cluster I (3.70). Maximum inter-cluster distance (8.79) was observed between cluster IV and VI and minimum between cluster I and III. High values of D² indicate genetic divergence among the accessions and least value showed close affinity among the accessions of the clusters. However in all the cases, inter-cluster distances were greater than the intra-cluster distance indicating the degree of genetic diversity present among the accessions of different clusters. The average cluster means of six characters revealed that only one accession placed in cluster V was highest in fresh weight of root (12.21 g), root volume (39.67 ml) and root hair zone (5.61 cm) but moderate in length of root (38.67 cm) and number (14.22). The accession Matigara white of cluster IV was highest in root length (60.44 cm). The average number of roots (22) was highest in the accessions of cluster II.

Table 1. Distribution of 85 mulberry germplasm accessions in different clusters based on D² values

Clusters	Accessions	Origin/country	Number of accessions
ł	Indigenous Kajli, Ber-A, Ber-20, <i>M. indica</i> (x), CSRS-2, Assambola, Kolitha-7, Kolitha-8, Kolitha-9, Dudhia red, Dudhia white, Bishnupur-4, Bishnupur-9, Tollygunge, Bush malda-A, Bush malda-B, Matigata black, Tista valley, Kurseong, Sujanpur, Sultanpur, KPG-1, KPG-2, Kanva-2, MS-1, MS-5, <i>M. indica</i> (HP), Sujanpur-5, RFS-175, OPH-1, Ace-105, Nagaland local, Kaliakotai, Golaghat, Jatinuni, Punjab local, MS-6, Mysore local, Almora local, Kakpillai, FGDTR-9, MS-8, Jodhpur, S 1635	India	45
	Exotic China black-B, China white Kokuso-13 Philippine Burma-8, Mandalaya, Thailand (unlobed), Monla-1, Monlai, Molai Bogura-4, Shrim-2, Shrim-5, Shrim-8 M. australis Calabresa, Fernandodias <i>M. multicaulis</i> <i>M. rotandiloba</i> , Multicaulis <i>M. nigra</i>	China Japan Philippine Burma Bangladesh Australia Paraguay Russia France Indonesia	21
	Indigenous M. indica black, MR-1, Surat	India	3
	Italian mulberry Okinowaso Australia Cyprus Muroso Thailand (lobed) <i>M. cathyana</i>	Italy Japan Australia Greece Paraguay Burma Indonesia	7
111	Ber-B Ber-6 Kolitha-3 Black cherry MS-7 MS-9	India	6
IV	Matigara white	India	1
V	VI	India	1
VI	Bogura-1	Bangladesh	1

Table 2. Average intra and inter-cluster values of D² among different clusters in mulberry

Table 3. Cluster mean of different root characters

	anterer	n cluster	s in muli	berry		
Cluster	1			IV	V	VI
i	13.69					
	(3.70)					
11	42.21	14.00				
	(6.50)	(3.74)				
111	29.43	43.26	11.59			
	(5.43)	(6.58)	(3.40)			
IV	37.76	50.75	65.71	0.00		
	(6.15)	(7.12)	(8.11)	(0.00)		
V	66.85	72.70	47.55	60.26	0.00	
	(8.18)	(8.53)	(6.90)	(7.76)	(0.00)	
VI	47.35	58.34	31.96	77.26	66.18	0.00
	(6.88)	(7.64)	(5.65)	(8.79)	(8.14)	(0.00)

Antiqued values are intra-cluster distances; D² values are in parenthesis

It can be concluded that while selecting the accessions as parents in breeding for obtaining potential hybrids, the parents of distant clusters with higher genetic divergence in root characters may be used to create high adaptability.

References

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Cluster	Number of roots	Length of root (cm)	Fresh wt. of root (g)	Root volume (mł)	Root hair zone (cm)	Dry wt. of root (g)
1	12.47	26.72	3.46	16.24	3.96	3.17
11	22.48	27.47	3.97	15.45	4.32	2.38
111	7.5	19.31	4.51	13.22	3.21	1.97
IV	16.78	60.44	3.87	24.33	3.49	25.77
V	14.22	38.67	12.21	39.67	5.61	3.09
VI	11.89	27.89	5.93	13.11	4.83	2.32

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