



Genetic divergence in mulberry (*Morus* spp.)

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Abstract

Genetic divergence among 98 mulberry (*Morus* spp.) genotypes (63 exotic, 35 indigenous) of different eco-geographic origin was assessed using Mahalanobis D^2 statistics. The total genotypes were grouped into seven clusters. Maximum number of genotypes were grouped in cluster III (19), IV (19), II (16), IV (16), VII (13) and V (12), respectively. Cluster I had only 3 exotic genotypes. All the clusters having both the exotic and indigenous genotypes except cluster I. The genotypes falling in cluster III had the maximum divergence followed by cluster I and II. The maximum and minimum divergence were revealed between cluster I and VI and between cluster V and VI, respectively. The cluster I and VI showed higher and lower mean values for most of the characters. So, mulberry crop improvement programme may be tried with the genotypes of divergent clusters for better heterotic effects.

Key words : Mulberry, exotic genotypes, indigenous genotypes, genetic divergence

Introduction

Mulberry is the sole food of silkworm (*Bombyx mori* L.). The plants are mainly grown for its foliage and breeding of mulberry are directed towards higher foliage production. To meet this objective, genetically diverse parents are required which could produce high heterotic effects with more variability in segregating generations. Till date, mulberry breeding has mainly confined to include indigenous genotypes to evolve improved variety. The available exotic genotypes collected from different eco-geographic origin has not been properly exploited to create variability for more production. The genetic diversity among mulberry genotypes have been reported by several authors [1,2,3]. But the information was based on limited parameters and genotypes. So, the present study was carried out to assess the genetic divergence among the genotypes using Mahalanobis D^2 statistics and to select the suitable genotypes for further utilization in breeding programme.

Materials and methods

The study was carried out at CSR&TI, Berhampore, West Bengal, during 1993-1995. The mulberry genotypes were maintained as high bush, spacing 1.5 m \times 1.5m with once annual pruning. Recommended agronomic practices were followed to maintain the plantation. Each genotype represented by a row of 5 plants. Growth parameters were recorded after 90 days of pruning of 5 plants and the plant attained 5 years when data was recorded. Leaf moisture and moisture retention capacity in harvested leaves were calculated following the standard procedure.

The observation on 11 parameters i.e., Moisture %, moisture retention capacity %, laminar index %, nodal distance, 100 leaves dry wt., lenticel/sq.cm. no. of buds sprouted, days to sprout, biomass wt., growth rate at 90 days and no. of twigs/plant were recorded from all 5 plants. The statistical analysis was carried out using Mahalanobis D^2 method [4] and the genotypes were grouped into different clusters following Tocher's method as described by Rao [5].

Results and discussion

The mulberry genotypes representing different eco-geographic origins are presented in Table 1. Based on divergence and magnitude of D^2 values, 98 genotypes were grouped into 7 clusters (Table 2). The distribution of different genotypes revealed that cluster III and IV having maximum number of genotypes. Cluster II and VI, both having 16 genotypes each. The clusters VI and V had 13 and 12 genotypes, respectively. The cluster I had only 3 exotic genotypes. The genotypes had distributed randomly in different clusters irrespective of geographic origin. Further, the grouping of genotypes did not show any relationship between genetic divergence and geographic diversity. The same observations were also reported by several authors (1, 2,3,7). The genetic drift and selection in different environments could cause greater diversity than geographic distance [6].

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Table 4. Clusterwise mean values of 11 characters in mulberry

Characters	Clusters							Contribution towards divergence %
	I	II	III	IV	V	VI	VII	
MC (%)	76.44	72.20	71.56	71.31	72.72	70.35	72.90	0.08
MRC (%)	95.89	93.94	91.82	93.33	93.72	90.77	93.37	0.04
LI (%)	86.10	84.91	83.29	85.88	81.61	81.33	83.77	0.44
NC (cm)	3.30	3.47	3.93	3.38	3.45	3.95	3.74	0.00
100 LDW (gm)	134.66	75.12	30.87	51.02	51.19	28.26	82.27	41.41
LT (sq. cm)	7.62	6.54	6.04	6.32	6.04	5.24	6.85	0.00
NBD (no.)	102.22	79.40	114.65	57.99	39.03	33.50	36.90	41.43
DSP	9.56	10.02	8.54	9.81	9.67	8.48	10.57	0.17
BIO (wt.)	17.00	14.54	16.86	9.45	9.62	5.73	7.37	1.58
GR (cm)	0.78	1.02	1.27	0.91	1.11	1.07	0.66	0.00
TWG (no.)	102.33	76.33	99.39	56.98	39.22	30.46	33.87	14.83

MC = Moisture content, MRC = Moisture retention capacity, LI = Laminar index, ND = Nodal distance, 100 LDW = 100 leaves dry wt., LT = Lenticel/sq. cm, NBS = No. of buds sprouted, DS = Days to sprout, BIO = Biomass wt., GR = Growth rate at 90 days, TWG = Twig no./ plant.

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