

Divergent gene pools in rice improvement

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Summary. The objective of the present study was to investigate the genetic architecture of yield in diverse populations of rice. Multivariate analysis by Mahalanobis's D^2 statistic and canonical (vector) analysis revealed that panicle weight, days to maturity, plant height and seed size were the important forces contributing towards divergence. One hundred rice strains were grouped into nine clusters with the help of D^2 and canonical analysis. The grouping pattern of the varieties were quite at random indicating that the geographical and genetic diversity were not related.

Key words: Rice – Genetic divergence – Gene pools

Introduction

Genetic uniformity has set the stage for crop destructive insect and disease epidemics in the past (Nat. Acad. Sci 1972). The diffusion of narrow genetic bases, such as 'Dee-Gee-Woo-Gen' and 'I-Geo-Tse', as the dwarfing genes, have increased the vulnerability of present day high yielding rice varieties. Thus, scientists engaged in rice improvement have an interest in the genetic diversity of improved rice varieties. This calls for strong and autonomous breeding programmes using genetically divergent gene pools and for identification of the alternative sources of dwarfism (Vairavan et al. 1973, Hargrove 1979; Hargrove et al. 1979; Murty 1979; Singh et al. 1979).

In our endeavour to diversify the genomes of rice we have collected local germplasm from U.P. (India) and parts of Nepal which appear to be phenotypically diverse. However, genetic diversity may not always be

synonymous with geographical and phenotypical diversity (Murty 1979). Thus, measurements such as distance, canonical and cluster analysis based on genetic criteria quantifying diversity have been carried in an effort to locate divergent gene pools for recombination breeding.

Materials and methods

The experimental material comprised of 100 rice strains which included: (i) 74 indigenous collections having about 10 dwarfs and semi dwarfs, (ii) improved plant types from IRRI, CRR1, and AICRIP and (iii) commercially grown cultivars endemic to the area. The material was grown in a replicated trial at the Institute of Agricultural Sciences, B.H.U., Varanasi during 1979. Random and competitive five plants were tagged in each replication during emergence in order to record biometrical observations. Characters of fitness and physiological efficiency were scored at the appropriate developmental stages of plant growth. These included: (1) days to emergence, (2) days to maturity, (3) effective tillers per plant, (4) height of the plant at maturity, (5) panicle length, (6) panicle weight, (7) numbers of grains per panicle, (8) length of grain, (9) breadth of grain, (10) L/B ratio, (11) test weight and (12) seed yield per plant.

A 12×12 dispersion matrix was used for the simultaneous test of significance of difference in the mean values of 12 variables based on Wilk's criterion (Rao 1952). Mahalanobis's D^2 statistic was used for assessing the genetic divergence between the populations. Simple criteria, suggested by Tocher (Rao 1952), for determining the group constellations was used. The canonical analysis was carried out by the method described by Arunachalam (1967). All these statistics have been established to be sensitive tools for the determination of genetic affinities.

Results

The populations differed significantly with regard to the characters studied individually (Table 1) and had a marked divergence when subjected to Wilk's criterion

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Table 1. Analysis of variance and estimates of parameters of variation in rice (*Oryza sativa* L.)

Sources of variation	df	Mean squares											
		X ₁	X ₂	X ₃	X ₄	X ₅	X ₆	X ₇	X ₈	X ₉	X ₁₀	X ₁₁	X ₁₂
Replication	1	–	11.00	0.52	216.90	–	12.00	807.50	0.0037	–	0.058	2.02	11.25
Treatment	99	444.60*	390.19*	11.57*	1724.07*	11.78*	2.29*	4891.57*	0.0223*	0.002257*	0.37*	42.45*	35.25*
Error	99	5.33	5.59	2.91	55.61	1.24	0.18	422.85	0.0009	0.000115	0.21	0.52	13.02
Grand mean		99.53	133.37	8.88	115.94	23.33	3.53	153.28	0.71	0.27	2.60	20.26	16.43
SE (d)		±2.30	±2.36	±1.71	±7.46	±1.11	±0.43	±20.56	±0.03	±0.01	±0.14	±0.71	±3.61
C.V. (g)		14.97	10.40	23.42	24.91	9.84	29.05	30.84	14.54	11.92	16.09	22.59	20.29
C.V. (p)		15.15	10.55	30.29	25.73	10.94	31.53	33.63	15.15	12.54	17.02	22.86	29.91
C.V. (e)		2.32	1.77	19.20	6.43	4.78	12.25	13.41	4.25	3.90	5.56	3.54	21.96
h ² (Broad sense)		0.98	0.97	0.60	0.94	0.81	0.85	0.85	0.92	0.90	0.89	0.97	0.46
G.A.		30.34	28.16	3.32	57.61	4.25	1.95	89.29	0.20	0.06	0.82	9.32	4.46
Genetic advance over discriminant function coefficient = 6.11													
Expected gain (%) = 543.74													

* Significant at 1% level of probability

X₁ = Days to floweringX₂ = Days to maturityX₃ = Effective tillers per plantX₄ = Height of the plant at maturityX₅ = Panicle lengthX₆ = Panicle weightX₇ = Grains per panicleX₈ = Length of grainX₉ = Breadth of grainX₁₀ = L/B ratioX₁₁ = Seed indexX₁₂ = Seed yield per plant**Table 2.** Distribution of 100 strains of rice in different clusters

Cluster	Strains included	N	Mode of origin
I	Muturi (7), Nagina-32 (58), IET-2683 (89), Bala (93), Jakkoku × Serapheehil-214-55-102 (97) and CRM-13-3241 (100)	6	Nepal, U.P., CRRI, IRRI and A.P.
II	Parwanipur (48), IR 4412-207-2-3 (78), IR 2307-217-2-3 (79), Dhaneshwar (88) and Jaya (91)	5	Nepal, U.P., CRRI and IRRI
III	Kachani (3), Ramachur (8), Saro (35), Sonkharcha (43), N ₂₂ (50), Ch ₄ (51), Duddhee (53), Silhat (54), Japanee (57), Kabiraj (65), IR 3941-25-1 (81), Rasi (84), IR 36 (86), IR 30 (87), Ratna (90), Sherammi × Serocephchehil-215-54-1A (98) and Sherammi × Eyoinchich-219-61-8 (99)	17	U.P., CRRI and IRRI
IV	Samsafiroj (2), Bachee (26), Parhane (38), Baburam (49), Ramjiyawan (68) and Badshaw Pasand (69)	6	U.P.
V	MPR-7 (6), Singar (20), T 39 (22), Sadhuya (24), Lezura (27), Bhanslote (47), Safari (56), Ghoghar (66) and Bansee (74)	9	M.P., U.P. and Nepal
VI	Mahishra (1), Sathari (9), Anandi-II (11), Mansara-I (17), Dedaie (23), Zelona (25), Sarju-49 (28), Mansora II (30), Bakail (31), Badali (32), Sudas (33), Kalakant (42), Amawadhawad (44), Zhanghir (45), Madrasi (52), Motichur (59), Chanaw (62), Palamu (67), Rajal (70), Ahraura (71), Oorairoot (73), IR 4417-179-1-5-2 (75), IR 5853-118-5 (76), IR 2863-38-1-2 (77), IR-2863-35-3-3 (80), IR 2070-414-3-9 (83), IET 5656 (85), IR 34 (95) and IR 46 (96)	29	U.P., A.P. and IRRI
VII	Amzhos (18), Zhingee (60) and IR 3880-10 (82)	3	U.P. and IRRI
VIII	Lawangchur (4), Anandi-I (5), Sathari (10), Bajar Bhang (14), Bhajarbong (15), Latera (16), Basmati (19), Gauriya I (21), Kapoorsal (29), Nazir (34), Tulsi Ram (37), Ram Bhoj (39), Barhany (40), Morangee (46), Udraj (72), Mahsuri (92) and IR 42 (94)	17	U.P. and Nepal
IX	Bahraini (12), Adamchini (13), Gauriya II (36), Kanakzeera (41), Brahma (55), Zeera (61), Sonabhusi (63) and Manbhog (64)	8	U.P.

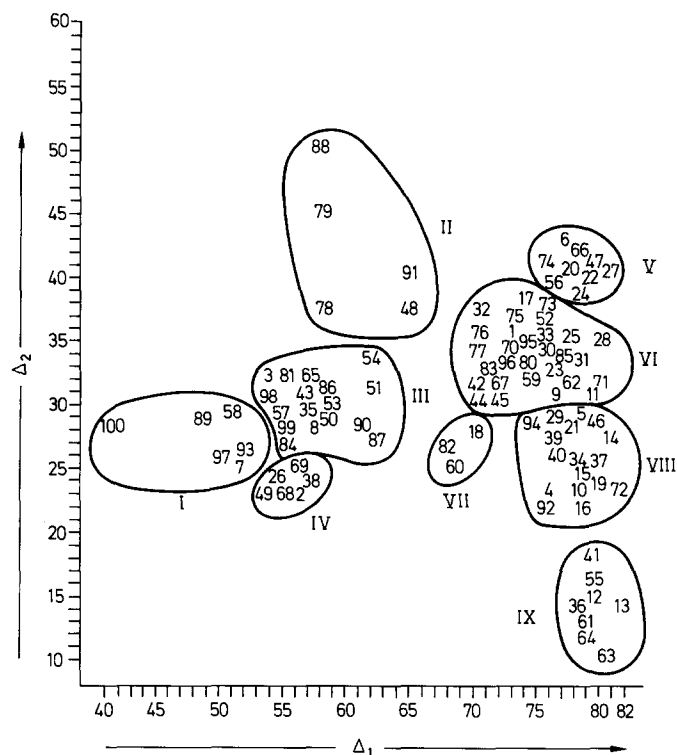


Fig. 2. Group constellations of 100 populations of rice (*Oryza sativa* L.) in Δ_1 and Δ_2 chart

Discussion

The sensitiveness and the utility of Mahalanobis's D^2 technique in identifying genetically diverse parents for obtaining success in recombination breeding by arriving at the gene constellations of divergent origins have been emphasized by several workers (Dhawan and Singh 1961; Matzinger et al. 1962; Chandrasekhariah et al. 1969; Sethi et al. 1978).

Earlier workers assumed that geographical diversity reflected genetic diversity. However, Moll et al. 1962, Timothy 1963, Murty and Arunachalam 1966, Arunachalam and Ram 1967 and Narsinghani et al. 1978 could not find any direct relationship between the two. The clustering pattern in the present case also supports the above view as the genotypes from different origins cluster together in 6 out of 9 cases.

A crossing of genotypes belonging to the same cluster would not be expected to yield desirable segregates. Consequently, a crossing programme involving genetically diverse parents belonging to different topologically distinct clusters would provide an opportunity for bringing together gene constellations of divergent origins as isolation in time and space results in the locking up the genes in different constellations.

The statistical distance (D) given in Table 3 represents the index of the genetic diversity among the clusters. In the present study the maximum distance of

$D = 34.95$ exists between cluster I and IX. The second largest distance of $D = 34.28$ appears between cluster II and IX. The 36 possible combinations of 9 clusters may be arranged in a descending order of magnitude of distance. The mean statistical distance ($D = 21.75$) may be considered as a guideline and crosses belonging to different clusters showing an inter-cluster distance of 21.75 or more should be attempted. However, other practical considerations e.g. disease reaction, lodging index and maturity etc., should also be taken into account when choosing between genotypes from a cluster.

The relative contribution of each character to the total divergence shows that yield per se had a low contribution (7.4%). Therefore, the parental material, chosen on the basis of a single complex character, i.e. yield, may not necessarily throw transgressive segregates for yield potential. However, relatively simply inherited characters, such as panicle weight, contributed maximally towards total divergence (52.9%). Therefore, it is possible that a crossing programme involving parental material selected on the basis of total divergence for the character studied might lead to an overall improvement in yield through considerable improvement in panicle weight, which is an important component of rice plant.

The idea put forth for choosing the parents on the basis of total divergence is based on the assumption

that genetic drift and selection in different environments could cause greater diversity than geographical distance.

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