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Analysis of genetic divergence in basil (Ocimum spp.)

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Abstract

Analysis of 21 basil (*Ocimum* spp.) genotypes grown at Raipur (Chhattisgarh) revealed that a maximum of five genotypes were accommodated in clusters I and II followed by four genotypes in clusters III and V, and three genotypes in cluster IV. The intra-cluster distance between members of cluster IV was maximum followed by clusters I, II, III and V, suggesting that the genotypes in cluster IV were relatively more diverse.

Keywords: basil, cluster analysis, genetic divergence, Ocimum spp.

Basil (*Ocimum* spp.) has long been recognized as a diverse and rich source of cytotaxonomical, genetical and chemical variability in different geographical races (Pushpangadan *et al.* 1979). However, not much effort has been made for its genetic improvement. Hence, the present study was undertaken to classify and understand the nature and magnitude of genetic diversity present in 21 indigenous basil genotypes available at Indira Gandhi Agricultural University, Raipur (Chhattisgarh).

The material for the present investigation included 21 genotypes/accessions of *Ocimum* spp. collected from different sources. These were grown in a complete randomized block design with three replications at Raipur, during the wet season of 2004. Appropriate management and cultural practices were undertaken. Each genotype was grown in plots of 3 rows of 2 m length at spacing of 45 cm between row and 30 cm between plants in a row. A uniform rate of fertilizer dose of NPK @ 60, 40, 40 kg ha⁻¹ with farmyard manure @ 10 t ha⁻¹ was applied. Observations were recorded in five randomly selected plants of each accession. The data were subjected to the Mahalanobis D² analysis (1936) to measure the genetic divergence.

The genotypes were grouped into a number of clusters. The criteria was that any two genotypes on an average showing lower D² value were grouped in the same cluster, while those showing high D² values were grouped into different clusters. Based on the relative magnitude of D² values, the 21 genotypes grouped into five were clusters (Table 1). A maximum of five genotypes were accommodated in clusters I and II followed by four genotypes each in clusters III and V, and three genotypes in cluster IV. The intracluster distance between members of cluster IV was maximum followed by clusters I, II, III and V, suggesting that genotypes in cluster IV were relatively more diverse. The minimum intra-cluster distance exhibited by cluster V indicated limited genetic diversity among the constituents genotypes (Table 2).

Cluster	No. of genotypes included	Genotype	Place of origin/adaptation
Ι	5	OCL-1 OCL-3, OCL-10 OCL-11, OCL-12	Neemuch (Madhya Pradesh) Akola, Pune (Maharashtra) Raipur (Chhattishgarh)
П	5	<i>O. sanctum</i> -1 <i>O. kilimandscharicum</i> -1 RRL-Og-14 <i>O. kilimandscharicum</i> -2 <i>O. sanctum</i> -2	RRL-Jammu (Jammu & Kashmir) RRL-Jammu (Jammu & Kashmir) RRL-Jammu (Jammu & Kashmir) RRL-Jammu (Jammu & Kashmir) Mandi (Himachal Pradesh)
Ш	4	OCL-4 CIM-Shayam, O. basilicum, RRL-OC-12	Rajgarh (Madhya Pradesh) CIMAP (Uattar Pradesh) RRL-Jammu (Jammu & Kashmir)
IV	3	OCL-5 OCL-6 OCL-8	Raipur (Chhattishgarh) Palampur (Himachal Pradesh) Rewa (Madhya Pradesh)
V	4	OCL-2 OCL-9 OCL- 7 RRL-OC-11	Raipur (Chhattishgarh) Rajnandgaon (Chhattishgarh) Jaipur (Rajasthan) RRL-Jammu (Jammu & Kashmir)

Table 1. Clustering pattern of 21 genotypes of basil

Table 2.	Mean intra- (diagonal and bold) and	
	inter-cluster D ² values of 21 geno-	
	types of basil	

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Cluster	Ι	П	III	IV	V
Ι	2.492				
Π	3.859	2.396			
III	3.582	3.247	2.203		
IV	5.461	4.249	4.021	2.779	
V	3.238	4.350	3.601	4.386	2.046

The relative divergence of each cluster from the other cluster (inter-cluster distance) indicated greater divergence between cluster I and IV (D²=5.461) followed by cluster IV and V (D²=4.386), cluster II and V (D²=4.350) and cluster II and IV (D²=4.249). The selection of diversed genotypes from these clusters would produce a broad spectrum of variability for oil yield, which may enable further selection and improvement of yield components. The hybrids developed from the selected genotypes within the limits of compatibility of these clusters, may produce high magnitude of heterosis or desirable trangressive segregants, which would be rewarding in a breeding programme. Due

emphasis in breeding programme should be given to genotypes OCL-3 and OCL-11 of cluster I; OCL-5 and OCL-8 of cluster IV; RRL-OC-11 and OCL-7 of cluster V. Crosses among parents having genetic divergence are likely to yield desirable combinations therefore, a crossing programme should be initiated between genotypes belonging to different clusters. Particular genotypes may be selected from selected clusters for consideration as parent in crossing schemes. There was a wide range of variation in cluster mean values for most of the traits except number of inflorescence plant⁻¹ (Table 3). The maximum and minimum mean value for characters were, days to 50% flowering: 51.44 (IV) and 84.80 (II); days to maturity: 124.54 (V) and 178.67 (II); plant height: 66.32 (I) and 97.18 (IV); number of primary branches plant⁻¹: 6.11 (IV) and 13.42 (V); number of secondary branches plant⁻¹: 3.20 (II) and 9.58 (III); length of inflorescences: 8.39 (IV) and 16.46 (I); plant spread: 64.52 (II) and 80.74 (V); dry plant biomass: 155.23 (I) and 293.11 (IV); fresh plant biomass: 328.17 (I) and 814.67 (IV); acid value of oil: 1.85 (III) and 5.16 (I); ester value of oil: 17.11 (IV) and 46.32 (V) and

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essential oil per cent: 0.38 (III) and 0.61 (II), respectively. This indicated that while planning hybridization programmes, genotypes like OCL-3 and OCL-11 from cluster I (for characters such as dwarfness, length of inflorescence and acid value of oil); OCL-5 and OCL-8 of cluster IV (for tallness, earliness, dry and fresh plant biomass); RRl-OC-11 and OCL-7 of cluster V (for primary branches plant-1 and ester value of oil) and genotypes O. kilimandscharicum-1 and O. kilimandscharicum-2 of cluster II (for oil %, number of inflorescence plant⁻¹, least number secondary branches plant⁻¹, least plant spread); are expected to give promising and desirable recombinants in segregating generations as they possess desirable features as seen from their cluster means.

References

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Table 3. Cluster means for various quantitative characters of basil

1	int		9	1	8	ß	2
Oil	conte	%)	0.46	0.61	0.38	0.55	0.55
Acid Ester	value	of oil	5.16 29.32	18.81	1.85 20.41	17.11	46.32
	value value content	of oil		4.10 18.81	1.85	4.87 17.11	1.93
Fresh	plant	cences (cm) biomass biomass of oil of oil (%) (cm) (g) (g)	69.20 155.23 328.17	607.70	749.53	814.67	635.38 1.93 46.32
Dry	plant	biomass (g)	155.23	224.57	260.67	293.11	80.74 187.81
Plant	spread	(cm)	69.20	64.52	69.84	73.91	80.74
No. of Length of Plant Dry	inflores- inflores- spread plant	cences (cm)	16.46	11.59	14.91	8.39	13.20
No. of	inflores-	cences branch ⁻¹	3.87	4.13	3.58	2.56	3.33
No. of	height primary secondary	(cm) branches branches plant ⁻¹ plant ⁻¹	8.00	3.20	9.58	5.00	5.75
Plant No. of	primary	branches plant ⁻¹	13.07	9.67	11.33	6.11	13.42
Plant	height	(cm)	66.32	77.49	77.17	97.18	70.72
Days to	maturity		129.27	178.67	155.17	129.89	124.50
No. of Days to Days to	50%	flowering	62.00	84.80	75.92	51.44	52.08
No. of	Cluster genotypes 50%		Ŋ	Ŋ	4	3	4
	Cluster		I	Π	Ш	IV	Λ