# GENETIC DIVERSITY IN POTATO (SOLANUM TUBEROSUM L.)

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## Abstract

Genetic diversity in 31 potato genotypes (parents and their hybrid progenies) was determined using multivariate analysis. Cluster analysis revealed that the parents and their hybrid progenies could be grouped into five different clusters. The maximum number of genotypes were included in clusters II and V. Cluster V had maximum and cluster I had minimum intra-cluster distance. Cluster mean showed wide range of variation for several characters among single as well as multi-genotypic clusters. Considering diversity pattern, parents should be selected from clusters I, III and V for the improvement of potato.

## Introduction

Potato is the fourth most economically important food crop after wheat, rice and maize in the world. The crop has high nutritional value as well as great yield potential. The existence of variability in a particular trait is an important prerequisite for its heritable improvement. For improving the yield potential of varieties and hybrids the decision should be made about the choice of right type of parents for hybridization. Since potato is a vegetatively propagated crop, variation among the existing commercial cultivated varieties seldom occur. Therefore, induction of variability in potato is urgently needed for ultimate use in any crop improvement programme.

It has been found that the progenies derived from crossing between divergent parents give divergent and useful trait. It has been often postulated by the breeders that geographical distribution reflects genetic diversity in selecting parents for hybridization. A limited study has been made on genetic divergence in potato either at tetraploid (Gaur *et al.* 1978, Sidhu *et al.* 1981 and Singh *et al.* 1988) or at diploid level (Grag 1988). An understanding of the nature and magnitude of variability among the genetic stocks is of prime importance to the breeders. Genetic diversity is one of the important tools to qualify genetic variability in both cross- and self-pollinated crops (Murty and Arunachalam 1966, Gaur *et al.* 1978). Such a study also permits to select the genetic divergent parent to obtain the desirable recombinant in the segregating generations. Therefore, the present study was undertaken to analyze the genetic divergence in 31 potato genotypes.

#### **Materials and Methods**

Thirty one genotypes of potato were grown in the research field of Institute of Biological Sciences, Rajshahi University, Rajshahi. The experimental design was RCBD with four replications. Each replication consists of 31 plots. Each plot having two rows of 40 plants. Row to row and tuber to tuber distance was 60 and 20 cm, respectively. Recommended dose and application methods of fertilizers were used. To get good crop conditions irrigation, intercultural operations, spraying of insecticide and fungicides were performed. The planting was done on 15 November 2001 and the harvesting was made after 90 days of planting. Observations were recorded and calculated on ten randomly selected plants from each plot on days to emergence (DE), plant height (PH), number of stems/plant (NS), number of tubers/plant (NT), tuber weight/ plant (TW), individual tuber weight per plant (ITW), tuber dry-matter content (DM%) and tuber

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weight loss due to respiration 150 days after harvest (TWL%). Data were subjected to principal component and Mahalanobis (1936)  $D^2$  analysis extended by Rao (1952) using GENSTAT 513 Computer programme.

# **Result and Discussion**

Analysis of variance revealed that the differences in 31 potato genotypes/varieties were significant for all the characters indicating the presence of notable genetic variability among them. The  $D^2$  values ranged from 3.823 to 23.694 and principal component scores also indicated a high degree of genetic diversity among the genotypes.

Cluster analysis: By application of non-hierarchical clustering using co-variance matrix, 31 genotypes (parents and their hybrids) of potato were grouped into five different clusters (Table 1). It was revealed that clusters II and IV had the maximum number of genotypes (nine) followed by cluster V and III having seven and five genotypes each, respectively. Cluster I had only one genotype and it was the lowest. Clustering pattern of parental genotypes under this study reveals that parents showed considerable genetic diversity among themselves by occupying four different clusters. Similar results were reported by Gaur et al. (1978) in potato, Masud et al. (1995) in pumpkin, Mannan et al. (1993) in Colocasia esculenta and Singh and Singh (1979) in okra. The 21 hybrids were distributed into four different clusters having more than one hybrid in each cluster. In some cases, the hybrid and one of its parents occupied the same cluster as in clusters II, IV and V. Similar results were reported by Main and Bahl (1989) and Singh and Prasad (1991). Intercluster hybrids were more frequent among hybrids with significant heterosis though the expression of heterosis was better in intercluster hybrids because of high mean performer of the parents and their different origin. Shanmugam and Rangasmy (1982) reported that falling materials of same origin into different clusters was an indication of broad genetic base of the genotypes belonging to that origin. Most of the female parents were grouped in cluster V indicating low genetic diversity among these parents. This could be due to their narrow genetic background.

Group/cluster No.	No. of genotypes	Genotypes in different cluster
Ι	1	Dheera
Π	9	Shill Bilati × TPS-67, Shill Bilati × Dheera, Lal Pakri × Dheera, Sada Gutti × TPS-67, Ausha × TPS-67, Patnai × TPS-67, Patnai × TPS-13,
III	5	TPS-67, TPS-13 Lal Shill × TPS-67, Lal shill × Dheera, Lal Pakri × TPS-13, Ausha × TPS- 13, Challisha × TPS-67
IV	9	Lal Shill × TPS-13, Shill Bilati × TPS-13, Lal Pakri × Dheera, Sada Gutti × TPS-13, Sada Gutti × Dheera, Ausha × Dheera, Patnai × Dheera,
V	7	Challisha $\times$ TPS-13, Ausha Challisha $\times$ Dheera, Lal Shill, Shill Bilati, Lal Pakri, Sada Gutti, Patnai, Challisha

Table 1. Distribution of 31 potato genotypes among five clusters.

TPS = True potato seed.

The maximum inter-cluster divergence (Table 2) was observed between the clusters I and V, and it was minimum between clusters II and III. The maximum intra-cluster distance was observed in cluster V and minimum in cluster I. Cluster I had only a single genotype. The crosses involving parents from most divergent clusters are expected to manifest maximum heterosis and generate wide variability in genetic architecture. Intracluster distance was being much lower than the

intercluster one, suggested, heterogeneous and homogeneous nature between and within groups, respectively. This was further supported by an appreciable variation observed for cluster means (Table 3). ITW, TW and WL% were the highest in cluster I; NS and TN in cluster III; PH in cluster II and DM% in cluster V; while ITW and TW were lowest mean values in cluster V. A wide range of variation for several characters among single as well as multigenotypic cluster was observed. However, the difference was more clear for PH, NS, TW, ITW, TW, DM% and TWL% which has contributed largely to the total divergence. Similar results have also been reported by Desai and Jaimini (1997), Gaur *el al.* (1978), Sidhu and Pandita (1980) and Sidhu *et al.* (1981) for TW, PH, ITW and TN towards total divergence in cluster. Hence, for the improvement of different characters viz. TN, ITW, TW, DM% and TWL% under the present study, parents should be selected from cluster I, III and V.

Cluster	Ι	II	II	IV	V
Ι	0000	10.457	13.799	18.706	23.694
II		0.818	3.823	8.798	14.281
III			0.557	5.015	10.458
IV				0.762	5.169
V					0.930

Table 2. Average intra-	(bold face) and intercluster	distance (D <sup>2</sup> ) of 31 potato genotypes.
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Characters	Clusters				
	Ι	II	III	IV	V
Days to emergence	8.69	10.48	10.83	9.77	11.78
Plant height	42.47	55.00	51.34	53.29	51.42
Number of stems/plant	5.25	5.40	5.71	5.36	4.88
Tuber numbers/plant	9.13	20.66	20.78	19.69	13.33
Individual tuber weight/plant	41.93	17.48	14.58	16.43	8.67
Tuber weight/plant	365.00	290.30	244.28	186.96	119.13
Tuber dry-matter content (%)	19.89	21.06	21.61	20.45	24.04
Tuber weight loss due to respiration (%)	24.60	16.40	14.58	16.43	8.67

Table 3. Cluster means for eight characters in 31 potato genotypes.

The principal component analysis revealed that in major vector 1 the important characters responsible for genetic divergence in the major axis of differentiation were PH and TN (Table 4). In vector II which was the second axis of differentiation PH, TN and TW were important. The role of PH and TN for both the vectors was positive across two axes which is the indication of the important components of genetic divergence in these materials.

Table 4. Latent vectors f	or eight characters o	f 31	potato genotypes.

Characters	Vector I	Vector II
Days of emergence	- 0.0450	-0.0057
Plant height	-0.0064	0.0210
Numbers of stem/plant	-0.5181	- 0.4344
Tuber numbers/plant	0.1048	0.1968
Individual tuber weight/plant	0.0056	0.0521
Tuber weight/plant	- 0.0823	0.0030
Tuber dry matter content (%)	- 0.6129	- 0.9156
Tuber weight loss due to respiration (%)	- 0.3334	-0.4254

Group consultation was also independently derived by principal component analysis to verify grouping obtained through  $D^2$  statistic in a two dimensional chart ( $Z_1$ - $Z_2$ ). Therefore, scores obtained for the first two components were plotted against two main axis and then supper imposed with clustering (Fig. 1). This clustering pattern confirmed the results obtained by  $D^2$  analysis.

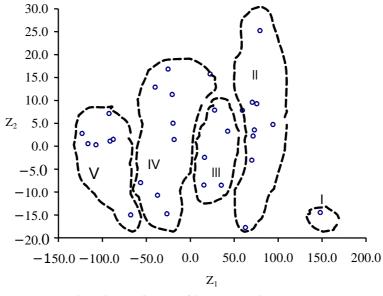


Fig.1. Scatter diagram of 31 genotypes in potato.

The crosses involving parents belonging to the maximum divergent clusters were expected to manifest maximum heterosis and also wide variability in genetic architecture. Thus crosses among the genotypes of clusters I, III and V would exhibit high heterosis and is also likely to produce new recombinants with desired characters in potato.

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