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ABSTRACT

Under the present study, assessment of genetic variability of 60 numbers of rice landraces was carried out under morphological and molecular level for selection of promising drought tolerant genotypes under upland ecosystem of North East India. Predominance of additive gene action on characters like leaf rolling index, root: culm ratio, grain yield and other yield contributing traits like plant height came out as effective parameters for selection of drought stress tolerant landraces. Molecular analysis revealed moderate genetic diversity with average Polymorphic Information Content values of 0.44 across the rice landraces. No specific clustering patterns of landraces against some morphological traits numbers of productive tillers per plant, numbers of filled grains per plant etc were found. This signifies the polygenic nature of the quantitative characters and influence of environment on them. Precise correlation between the morphological performance of the landraces and their clustering pattern under molecular analysis was found to be effective in identifying suitable landraces like Chikanswarikabar as a promising parent for future breeding programme and also to formulate efficient strategies for sustainable management of rice landraces under rain fed ecosystem.

Keywords: Rice, Landraces, variability, drought, molecular marker.

INTRODUCTION

Rice (Oryza sativa L) is the most staple food of more than half of the world's population (Rasheed et al., 2020) and 90% of it is being produced and consumed in Asia (Fukagawa and Ziska, and share maximum in grain 2019) production. Being a semi aquatic crop, rice can be grown in diverse range of ecosystems such as rain fed uplands and lowlands, irrigated, and deep-water ecosystems. But, in present day scenario, the entire ecosystems suffer from the threat of the climate change like drought stress and to be more specifically, the rain fed uplands are the most severely affected by drought stress because of its lighter soil texture and quick percolation tendency of rain water. Widespread drought occurrence in rice-growing areas results in severe decline in yield and thus the development of drought-tolerant varieties that maintain good vield under moisture stress is a priority area of rice research for rice production sustainable and improvement. The success of a crop improvement programme depends on the availability of genetic variability in the population. Rice landraces, because of their effectual evolution process and enormous in-built genetic variability can grow in a wide range of ecosystem and climatic conditions and can be treated as promising source towards development of drought tolerant population.

India is considered to be one of the prime centres for rice diversity and Tripura, being a state of north east India is endowed with a great diversity of rice landraces in its varied topographical rain fed upland ecosystem. Many of the rice landraces grown in the hilly region of this state are found to be effective as sources of drought tolerance. But, over the last few decades, rice has faced a severe diversity loss like other locally available landraces of different crop plants that are often ignored in existence of high-yielding varieties. however, have survived environmental turmoil over the ages (Pradhan et al., 2020). Along with, it is also undoubtedly proven fact that. information on the amount of genetic variation in the germplasm and genetic relationships between them is imperative for the study and designing of breeding programs (Farahani et al., 2019) towards development identification and of germplasm and defining the existence of varieties (Eid, 2019). In any breeding programs, determination of the level of genetic diversity in crop species is of great importance for the selection of suitable parents and maximum utilization of heterosis (Luo et al., 2019). Therefore, the present study was undertaken with the aim to assess the trend in genetic diversity in sixty numbers of rice landraces under both morphological and molecular level and to generate comparative information for selection of appropriate drought tolerant promising genotypes in future rice breeding programme.

MATERIALS AND METHODS Field experiment

The field experiment was conducted in the farm complex of Indian Council of Agricultural Research (ICAR) Complex for North Eastern Hilly Region, Tripura Centre. Lembucherra, West Tripura (23°90'E, 92°29'N), India, during the years 2015-16 and 2016-17. 60 numbers of local rice land races (Table1), collected from different hill ranges of Tripura evaluated through were Randomised Complete Block Design with 3 replications. Artificial drought conditions were imposed in the experiment field during the 'Boro' seasons of 2015-16 and 2016-17 as per the standard procedure of IRRI Knowledge bank (http://www. knowledgebank.irri.org/rice breeding course). Seeds were directly sowed at a distance of 25 cm (row-row) x 25 cm (plant -plant) in dry soil. Regular furrow type of irrigation has been provided up to 30 days after sowing to ascertain optimum seedling growth. After that, the frequencies of irrigation were reduced. No irrigation was provided until or unless field surface is completely dry or plants under the study showed severe wilting symptoms. When the target level of soil dryness and plant stress are reached, the field was liberally irrigated to saturate the root zone. This irrigation pattern was repeated until harvest. Fertilizers were applied 60:40:40 Kg N: P: K ha⁻¹. The same pattern of management practices, irrigation schedules, inter culture operation other pest control measures were followed as and when necessary during the whole growing periods for the investigated years.

Phenotypic analysis based on morphological traits

Pulled mean values of 15 numbers of morphological characters as listed in Table (2) were recorded from five numbers of competitive plants preferably from the replications. rows over the middle Standard evaluation system of Rice (IRRI, 2002) was considered as reference while recording the characters. The mean data were also subjected to heritability and genetic advance analysis as per Johnson et al. (1955) using Windostat Version 9.2 from Indostat service. In order to arrange the landraces in various groups and subgroups and also to find out the behavioural similarity of them. Hierarchical cluster analysis for all the morphological traits in 60 rice land races was carried out using un-weighted pair group linkage type through NCSS statistical software 2019. The Euclidean distance method and standard deviation scaling method were used with cluster cut off value of 1.0 to form a horizontal dendrogram.

Molecular analysis

DNA was extracted from the tender leaf tissues of 21 days old seedlings (a

healthy single seedling per genotype) of the rice land races, based on Cetyl trimethyl ammonium bromide (CTAB) method described by Muray and Thompson (1980) with some modification. A set of 40 numbers of simple sequence repeat (SSR) markers, distributed among different rice chromosomes were screened and subsequently utilised for genetic diversity analysis of the rice land races. Polymerase chain reactions were carried out in 12.5 µl of total reaction volume containing 1.0 µl genomic DNA, 4.25 µl nano pure H_2O , 6.25 µl of PCR master mix (Hi Media) containing Thermus aquaticus (Taq) DNA polymerase, dNTPs, MgCl₂ and reaction buffers at optimal concentrations and 0.5µl from microsatellite (SSR) markers (forward and reverse primers). Amplifications were performed in Eppendorf nexus gradient Master cycler with the temperature profile consisting of Initial denaturation stage of 94[°]C for 5 minutes and 35 cycles of [(i) Denaturation stage : 94° C for 1 minute (ii) Annealing stage : 55° C for 45 seconds (iii)Extension stage : $72^{\circ}C$ for 2 minutes] followed by final extension stage of $72^{\circ}C$ for 10 minutes. The ultimate PCR products were then diluted in 50 µl of distilled water kept in 4°C for further Gel and electrophoresis. Subsequently, around 2 µL aliquot of amplified PCR products of each of the landraces were mixed gently with loading buffer and loaded in 2% agarose gel prepared in Tris- Acetate (TAE) buffer along with 100 bp and 50 bp DNA marker ladder. For gel electrophoresis of SSR amplified PCR products, agarose gel were also used by several previous workers (Anupam et al., 2017; Ravikiran et al., 2018; Hamidah et 2020) with al.. resultant significant polymorphism. The amplified PCR products were then size fractioned by electrophoresis and visualized by staining with ethidium bromide (0.5µg/ml) in MultiDoc-It Imaging System of UVP.

For diversity analysis at molecular level, only consistent and reproducible bands were scored in binary format as '1' for presence and '0' for absence. Smeared and weak bands were excluded. Fragments of the same molecular weight were considered to represent the same allele. Polymorphic information content (PIC) is a measure of a marker's ability to detect polymorphism in a population, based on the number of alleles detected and their frequency distribution; hence, it provides an estimate of the discriminating power of a marker.PIC is calculated for each marker using the formula: $PIC_i = 1 - \Sigma P^2 i j$, where Pij is the frequency of the jth allele in genotype (i). The polymorphism information content for each locus and heterozygosity was calculated using PIC calculator (Jan. 2002). Di-nucleotide repeat motifs, expected amplified band sizes and annealing temperatures of microsatellite primers under study were availed from SSR marker resources of Gramene microsat database (https://archive.gramene.org/markers/micr osat/all-ssr.html), whereas, major allele frequency of a specific primer was calculated out considering the numbers of occurrence of that particular allele and total numbers of alleles. SSR allelic binary (0 and 1) data generated for 60 rice land races was converted to text tab limited format and imported to DARwin software VERSION 6 (Perrier and Jacquemoud-Collet, 2006). Dissimilarities of allelic data were calculated with 1000 bootstraps option using simple matching dissimilarity index. Unweighted Neighbour joining method was adopted while constructing the horizontal dendrogram based on the allelic dissimilarity.

RESULTS AND DISCUSSION Phenotypic analysis based on morphophysiological traits:

Under the study, the differences between the phenotypic coefficient of variations (PCV) and genotypic coefficient of variations (GCV) for some of the studied characters were found to be comparatively thin indicating sensitivity to environment and consequently greater role of drought stress influencing the expression of those characters. The magnitude of PCV and GCV were found to be highest in case of leaf rolling Index (LRI) followed by leaf tip drying index (LDI). High percentile of coefficient of variations values of LRI and LDI are in agreement with the research works of Haider et al. (2012) and Kumar et al. (2015).

High heritability coupled with moderate to high genetic advance have been observed in LRI, Root: culm ratio, numbers of filled grain per panicle, numbers of spikelet per panicle, grain yield per plant and 100 seed weight which implies predominance of additive gene action for those character. Thus, selection of land races based on those characters would be effective. Similar types of findings under drought stress condition have also been reported by Haider et al. (2012) for LRI, Panja et al. (2017) for root:culm ratio, and Perween et al.(2020) for numbers of filled grains per panicle. High value of heritability along with moderate to high magnitude of genetic advance of traits viz. numbers of spikelet per panicle and grain yield and test weight were also observed by Perween et al. (2020) and Nithya et al. (2020). The estimates of GCV, PCV, Heritability and Genetic advance etc are presented in Table (3).

For effective selection of promising rice landraces, grouping of the land races had also been carried out based on the mean value analysis of some significant morphological traits on the basis of Standard evaluation system of Rice, IRRI 2002 (Table 4), wherein, landraces namely ChikanswariKabar, Garo Malati, Galong, Turkey, Tarkol, Dhalabalam, Maimi Hungar were found to be better performer as far as the grain yield under moisture stress condition is considered. Thus, those landraces may be opted as parent materials for subsequent breeding programme.

Molecular analysis based on microsatellite-based rice marker:

Out of 40 micro satellite or SSR primers, 39 numbers of primers generated polymorphic alleles, while only one marker (RM 172) produced monomorphic allele. A total number of 119 alleles were detected at the loci of the 40 numbers of SSR markers across the sixty rice landraces with an average count of 2.97 alleles per locus. This is an indication of moderate level of molecular diversity among the landraces under the study.

The highest numbers of total alleles per locus were observed for RM252 (6), RM254 (5), followed by RM505, RM285, RM320, RM175, RM339, RM276 and RM592 with 4 numbers of alleles per locus for each primer. The result is confirmatory with the studies of Yadav *et al.* (2013) against RM252 and Anupam *et al.* (2017) against RM252 and RM3. As far as the average number of alleles per locus is concern, to some extent the same range of results were found by Ramadan *et al.* (2015), Nachimuthu *et al.* (2015) and Ngangkham *et al.* (2019).

Polymorphic information content (PIC) is the comparative measure of how much a marker is informative to distinguish between different populations and it depends upon the number of alleles detected by this marker and their relative frequency; thus, it provides an estimate of the discriminating power of the marker (Nagy et al., 2012). PIC values for the microsatellite markers used in this study varied from 0.00 to 0.743 with an average of 0.44. Markers with PIC values of 0.5 or higher are more useful in distinguishing the polymorphism rate of a marker at a specific locus (DeWoody et al. 1995), however, the mean polymorphism rate (0.44) is found to be moderate in this study which are in agreement with the studies of Becerra (2015) and Adak (2020). The most appropriate explanation to this might be the similar origin, ecotype and speciation of the land races. The highest PIC values observed for RM252 were (0.743)followed by RM505 (0.7), RM285 (0.648), RM254 (0.630), RM175 (0.613). In the study according to present the classification of Botstein et al. (1980), there was 14 numbers (35% of total numbers of markers) of highly informative markers with PIC value range > 0.50. On the other hand, 21 numbers of SSR markers with a PIC value (0.50 < PIC <0.25) and 5 numbers of SSR markers with PIC value (< 0.25) proved themselves as informative moderately and slightly informative markers respectively.

Details of the microsatellite primers used in present study along with their allele size (bp), allele diversity, gene diversity and polymorphism information content (PIC) is depicted in Table (5). A significant correlation between PIC value and the total number of repeat motifs counts of di-nucleotide sequence per microsatellite marker was also revealed in the study. Loci amplifying di-nucleotide repeat motifs were found to be more polymorphic than those with tri-nucleotide and tetra-nucleotide repeat motifs. Most of the high values of PIC (> 0.5) were found to be associated with high numbers of repeated values of (GA) or (CT) disequences. nucleotide These results. suggest that the total repeat count of SSR or microsatellite loci is associated with the high PIC value of particular marker. The repeat numbers larger the in the microsatellite DNA, the larger the number of identified alleles and PIC values. These results were consistent with those reported by Ramadan et al. (2015) and Melaku et al. (2018). Among the set of 40 numbers microsatellite primers tested in the study, RM 252 (PIC value of 0.743) with [(CT)19] repeat motifs, RM 505 (PIC value of 0.7) with [(CT)12] repeat motif and RM285 (PIC value of 0.648) with [(GA)12] repeat motifs came out to be more informative as they could reveal more polymorphism in comparison to other primers. PCR amplified fragments produced by the some of the polymorphic markers in the current study are given in Figure 1(a, b) to Figure 3(a, b).

Comparative analysis of molecular and morphological study:

For carrying out a reproducible comparative analysis of the rice landraces under both morphological and molecular level, efforts have been rendered to analyse maximum possible numbers of small subgroups. In general, the dendrogram generated against the microsatellite based binary data produced more numbers of sub clusters (11 numbers) in comparison to morphological traits based dendrogram with 6 numbers of clusters (Fig. 4 and Fig. 5). Often, it has been observed, landraces, clustered in a single subgroup under morphological analysis are categorised into different subgroups under molecular analysis which signifies the better capacity discrimination of the microsatellite based molecular analysis the morphology-based diversity over analysis, however some similarities in clustering pattern was also been observed between those analyses. Four numbers of aromatic rice landraces viz. Kalikhasa, KhasaKasam, Kala Jira and American Ration clustered in a same group of B_1B under morphological analysis, while under molecular analysis, only Kalikhasa and Khasakasam grouped together in a single cluster. Morphologically, there were ample similarities between the landraces of Kalikhasa and Kala Jira to raise the doubt on uniqueness of the landraces but better discrimination capacity of SSR based molecular analysis proved themselves as separate landraces under the study. The 'Fazu' group of landraces [FazuVom and Fazu Sen] grouped themselves in a single cluster under both morphological and which probably molecular analysis, signifies the same origin of the said land races.

Based on the mean value analysis of different significant yield governing

morphological traits and molecular analysis, a confirmatory analysis was also carried out to find out the correlation between the mean performance of the landraces and their clustering pattern under molecular analysis (Table 6). It has been revealed that the landraces viz. Kaporok, Releng, Tarkol, TarikolKolte, with more than 20 cm of total root length and Root: Culm ratio value of > 0.3 clustered themselves into a single sub group of B_1 . Yang Dhan and Badaya with moderate to high leaf drying index clustered themselves into a specific sub group of C₂A. Kaporok, Releng, Tarkol, Beti are found to be clustered in a single sub group of B_1 with low to moderate leaf rolling capacity.

But in this study, the main concern was to trace out the high yielder landraces and other yield contributing traits under moisture stress and as far as the grain yield under moisture stress condition is considered the landraces namely ChikanswariKabar, Garo Malati, Galong, Turkey, Tarkol, Dhalabalam, Maimi Hungar were found to be better performer under study. out of which, the ChikanswariKabar, the tallest and the highest vielder landrace settled itself in a single sub group of C₂C under molecular analysis. Significantly, under morphological evaluation also, the said landrace grouped itself with only one high yielder landrace viz. Garo Malati. This certainly signifies the uniqueness and diverse nature of Chikanswarikabar from rest of the landraces and thus, can justifiably be utilised as a promising parent with expected heterosis effect in drought stress breeding programme. Interestingly, some landraces viz. Chikanswarikabar, Maimi Hungar, Garomalati, Kaporok and Tarkol which found to be better yielder under drought stress also showed comparatively deeper root length and higher root : culm ratio. This phenomenon probably justifies the better moisture utilising capacity of the landraces with improved root traits which ultimately

resulted in better yield. Thus, those landraces also create a scope for further utilisation of those in future drought stress breeding programme.

It has also been found that, the landraces which shows slight to no leaf tip drying characters and slight to moderate leaf rolling capacity under moisture stress are basically good yielder. Along with, under this study, land races with intermediate to tall plant height also performed better as far as the grain yield is concern. Thus, along with grain yield, the better root attributes, leaf tip drying index (LDI), leaf rolling index (LRI) and plant height (PH) may also be considered for selection of promising landraces under drought stress. No specific clustering patterns of landraces against the rest of the morphological traits numbers of productive tillers per plant, numbers of filled grains per plant etc were found. This signifies the polygenic nature of the quantitative characters and influence of environment on them.

Overall, the results obtained would help in the identification and differentiation of various cultivars being cultivated in this region to broaden the landrace base in the future of rice breeding programs. In addition, it will help in identifying efficient strategies for the sustainable management of the genetic resources of rice crops in the particular rain fed ecosystem of North East India

ABBREVIATIONS

(CT) - Cytosine and Thymine base, (GA) -Guanine and Adenine base, dNTPs-Deoxyribonucleotide triphosphate, RM: Rice Marker, TAE: Tris- Acetate, (Ethylenediaminetetraacetic Acid), Taq: Thermus aquaticus

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Table 1: List of the 60 numbers of rice land races collected under the study and their collection sites

SI.	Name of the Land races	Collection site	District	Lattitude/ Longtitude	SI.	Name of the Land races	Collection site	District	Lattitude/ Longtitude
1	Kaporok	Thamsarai para	N	24°14'N /92°17'E	31	Darka Sona	Thamsarai para	N	24°14'N/ 92°17'E
2	Releng	Boithangbar i	Ν	24°19'N /92° 8'E	32	Tarikol Kolte	Boithangbar i	Ν	24°20'N /92°12'E
3	Beti	Boithangbar i	Ν	24°19'N /92° 8'E	33	SadaBiroin	Ganganagar	Ν	24°20'N /92°12'E
4	MaimiUzr a	PurbaJalaiba ri	G	23°30'N /91°30'E	34	MaimiTauk ha	PurbaJalaiba ri	G	23°12'N /91°36'E
5	Kalikhasa	Ganganagar	Ν	24°20'N /92°12'E	35	Saluma	PurbaJalaiba ri	G	23°12'N /91°36'E
6	Chinal	Uttar Gakul Nagar	W	23°42'N /91°15'E	36	Tarkol	Noagang	Ν	24°19'N/ 92°13'E
7	American	Rajnagar	Ν	24°19'N /92° 8'E	37	Madoop	Dalapati	D	23°37'N/ 91°51'E
8	KhasaKas am	Kairai	W	23°51'N /91°28'E	38	Waibang	Bilaiham para	W	23°52'N/ 91°41'E
9	Biroin	Rajnagar	Ν	24°20'N /92°12'E	39	Jhum Bini	Garjeecherra	G	23°26'N /91°30'E
10	Galong	Boithangbar i	N	24°20'N /92°12'E	40	Fazu Sen (White)	Athramura range	K	23°51'N/ 91°44'E
11	FazuVom	Boithangbar i	Ν	24°20'N/ 92°12'E	41	Bongbu	Noagang	Ν	24°19'N /92°13'E
12	Garo Malati	Uttar Gakul Nagar	W	23°42'N/ 91°15'E	42	Sadok	Noagang	Ν	24°19'N /92°13'E
13	Maimi Usha	PurbaJalaiba ri	G	23°12'N /91°36'E	43	Kala Jira	Kameswar	Ν	24°22'N/ 92°11'E
14	Maimi Red	Thamsarai para	Ν	24°14'N /92°17'E	44	Gaigash	Murasingh Para	G	23°42'N/ 91°43E
15	Suri	Aswinirojap ara	D	23°34'N/ 91°54'E	45	Vanbang	Killa	S	23°36'N /91°31'E
16	Lebuka	Thamsarai para	Ν	24°14'N /92°17'E	46	Makajaria	Rajnagar	Ν	24°19'N /92° 7'E
17	Aaduma	Aswinirojap ara	D	23°34'N /91°54'E	47	Jilong	Bilaiham para	W	23°42'N/ 91°15'E
18	Fazu Sen	Boithangbar i	Ν	24°20'N /92°12'E	48	American Ration	Thamsarai para	Ν	24°14'N/ 92°17'E
19	FazuNgoi	Boithangbar i	Ν	24°20'N/ 92°12'E	49	Kala Dhan	Bilaiham para	W	23°42'N/ 91°15'E
20	BetiKalai	Brahmacharr a	K	23°48'N /91°39'E	50	Turkey	Dalapati	D	23°37'N/ 91°51'E
21	Saanki ka Phool	Dalapati	D	23°37'N/ 91°51'E	51	SaankiKach ak	Dalapati	D	23°37'N/ 91°51'E
22	Bihar	Brahmacharr a	K	23°48'N /91°39'E	52	MaimiUkhl ao	PurbaJalaiba ri	G	23°12'N /91°36'E
23	Chikansw ariKabar	Harbang	W	23°54'N /91°31'E	53	Maiwasha	Old Dalapati Para	D	23°37'N/ 91°51'E

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24	Bangbu Jhum	Nunnacharra Range	K	23°46'N/ 91°44'E	54	MaimiWato lok	Dalapati	D	23°37'N/ 91°51'E
25	Lal Biroin	Kameswar	Ν	24°22'N /92°11'E	55	Santinmaw Wakhum	Killa	S	23°36'N /91°31'E
26	DhalaBala m	Rajnagar	N	24°19'N/ 92° 7'E	56	Yang Dhan	Killa	S	23°36'N /91°31'E
27	Goria	Char Garia Para	W	23°54'N/ 91°31'E	57	Badaya	Killa	S	23°36'N/ 91°31'E
28	Bahadur	South Ganga Nagar	Ν	24°20'N /92°12'E	58	Kanchali	Killa	S	23°36'N /91°31'E
29	Maimiwat oklokMan doori	Brahmacharr a	K	23°48'N/ 91°39'E	59	Australian Biroin	Kameswar	N	24°22'N /92°11'E
30	MaimiHu ngar	Twisarangch ak para	W	23°54'N/ 91°31'E	60	Assam Paisom	Rajnagar	N	24°19'N /92° 7'E

D-Dhalai, G- Gomati,K-Khowai,N- North Tripura, S-South Tripura,W-West Tripura Source: Acharjee *et al.*(2019)

Table 2:	Pulled mean	values of 1	5 (Fifteen)) numbers	of morp	hological	characters	of 60
(sixty) lan	nd races unde	r drought st	ress condi	tion				

SL. No.	GENOTYPES	VGR	LRI	ICLI	PL HT	PR TLR	PN LN	PRI BR	RTLN	RT: CLM	Nd /LTNdS	FLD GRN / PN	SPKLT FRT	100 SD WT	IH	GR Y IPLT
1	Kaporok	5.50	2.00	0.25	94.67	4.33	23.33	7.50	22.15	0.31	103.50	90.42	87.63	2.22	26.10	9.31
2	Releng	7.00	1.00	0.00	87.08	3.75	23.75	7.91	21.23	0.34	103.17	72.59	70.35	2.60	26.51	7.53
3	Beti	5.00	3.00	0.00	85.33	3.92	20.42	6.84	19.26	0.30	91.00	80.75	88.75	2.43	24.90	7.87
4	MaimiUzra	7.00	0.00	1.25	102.08	3.25	23.75	8.08	15.01	0.19	84.08	68.50	81.54	2.22	24.06	5.51
5	Kalikhasa	7.50	2.00	2.00	84.92	3.92	17.08	5.67	10.58	0.16	80.17	69.33	86.49	1.10	13.63	3.23
6	Chinal	6.50	0.00	3.50	80.50	3.58	24.42	6.33	11.51	0.21	65.08	54.75	84.07	2.19	16.42	4.65
7	American	8.00	0.00	0.00	85.25	2.83	24.92	6.58	13.42	0.22	114.34	88.08	77.11	2.33	25.84	6.31
8	KhasaKasam	8.00	0.50	0.50	75.75	3.33	16.83	5.58	10.32	0.19	80.17	66.08	82.18	1.26	13.88	2.97
9	Biroin	6.50	0.00	0.00	85.25	3.92	25.25	7.75	21.08	0.35	90.67	78.67	86.74	2.35	27.49	7.48
10	Galong	6.50	3.00	0.25	100.91	3.42	26.17	7.92	21.62	0.29	164.33	147.42	89.51	2.69	39.07	14.08
П	FazuVom	7.50	2.00	1.00	69.75	3.42	23.75	7.17	10.76	0.24	64.42	51.67	80.19	2.39	21.60	4.58
12	Garo Malati	5.00	1.00	0.00	124.16	4.42	29.17	9.33	26.53	0.28	162.83	147.75	90.84	2.75	34.85	18.63
13	Maimi Usha	8.00	1.00	1.50	61.25	3.50	23.25	6.92	8.69	0.23	94.75	80.75	85.24	2.19	28.88	6.66
14	Maimi Red	7.50	1.00	0.50	94.50	3.50	23.16	6.42	18.58	0.26	77.83	65.02	83.54	2.89	23.72	6.65
15	Suri	8.50	0.00	1.00	73.00	3.58	20.50	6.00	10.62	0.20	76.67	63.09	82.23	2.10	22.12	5.39
16	Lebuka	7.00	0.25	0.00	83.33	3.50	26.08	7.25	8.24	0.15	85.08	67.83	79.72	2.05	20.60	5.58
17	Aaduma	7.00	0.00	2.00	79.33	4.25	22.58	6.67	15.78	0.28	78.91	65.91	82.28	1.67	19.07	5.00
18	Fazu Sen	8.00	1.00	1.00	83.92	3.08	22.34	5.92	14.17	0.23	65.25	49.67	76.35	2.73	19.45	4.91
19	Fazu N'G	9.00	2.50	2.00	77.25	2.83	20.08	7.92	14.96	0.27	63.33	54.09	85.06	2.35	19.62	4.29
20	BetiKalai	6.00	0.50	0.50	91.08	3.67	23.75	6.33	16.03	0.24	95.92	83.34	87.02	2.75	28.95	8.88
21	Saanki ka Phool	8.00	0.00	0.00	79.83	3.58	22.42	6.33	8.40	0.15	62.00	49.75	79.95	2.09	16.07	4.19
22	Bihar	7.00	1.00	0.50	86.58	3.58	20.17	5.25	12.12	0.18	69.92	55.92	80.40	1.78	16.17	4.56
23	ChikanswariK abar	5.50	0.50	0.50	125.42	4.50	29.50	10.25	24.76	0.26	195.33	178.58	91.43	2.52	40.86	21.66
24	Bangbu Jhum	6.50	0.00	1.00	77.33	2.92	22.25	6.50	12.82	0.23	57.50	45.42	79.53	2.78	21.39	3.94
25	Lal Biroin	7.00	0.00	2.50	78.75	3.17	17.67	7.42	11.71	0.19	80.25	66.50	82.89	2.55	25.46	5.75
26	DhalaBalam	6.50	0.50	0.25	83.42	4.08	22.58	7.75	12.30	0.20	96.83	83.92	86.64	2.77	30.49	10.00
27	Goria	7.00	0.00	0.50	94.92	3.41	19.67	5.33	9.73	0.13	68.67	56.92	83.03	2.44	22.30	5.18
28	Bahadur	9.00	0.00	1.50	95.00	4.00	25.16	6.42	14.08	0.20	65.67	51.00	77.72	1.82	12.80	3.76
29	M.W.Mand oori	7.00	0.00	2.00	91.83	3.83	27.50	8.58	9.65	0.15	98.42	79.42	80.05	2.16	27.07	6.96
30	Mami Hungar	5.50	0.50	0.50	98.42	3.92	28.92	8.67	23.13	0.33	103.33	91.50	88.49	2.99	33.76	11.04

VGR- Seedling vegetative vigour, LRI- Leaf rolling index, LDI-Leaf tip drying index, PL HT - Plant Height (cm.), PR TLR-Numbers of Productive Tiller per plant, PN LN -Panicle length (cm.), PRI BR-Numbers of Primary branch per panicle, RT LN-Root length (cm.), RT:CLM-Root- Culm ratio, SPKLT/PN-Numbers of Spike lets per panicle, FLD GRN/ PN-Numbers of filled grain per panicle, SPKLT FRT-Spikelet fertility (%),100 SD WT-100 Seed's weight (gm.), HI-Harvest Index (%),GR Y /PLT-Grain yield per plant (gm.)

Table 2: cont. Pulled mean value of 15 (Fifteen) numbers of morphological characters of 60 (sixty) land races under drought stress condition

SL. No	GENOTYP ES	VGR.	LRI	IDI	PL. HT.	PR TLR	PN LN	PRI BR	RTLN	RT:CLM	NALLINAS	FLD GRN / PN	SPKLT FRT	100 SD WT	Ħ	GR Y / PLT
31	Darka Sona	6.00	1.00	1.00	86.83	3.33	23.00	5.50	14.51	0.23	95.58	83.34	87.04	2.38	27.85	7.01
32	Tarikol Kolte	7.00	0.00	0.75	102.42	3.67	23.58	6.83	22.12	0.28	80.66	70.75	87.62	2.58	27.13	7.31
33	SadaBiroin	8.00	1.00	3.00	73.50	3.41	20.00	7.08	10.23	0.19	75.91	56.16	74.15	2.32	24.99	4.73
34	Maimi Taukha	7.50	4.50	0.50	78.83	2.92	26.17	7.91	18.27	0.35	100.67	84.59	84.00	2.48	29.40	6.52
35	Saluma	9.00	2.00	2.00	61.25	3.00	24.33	6.25	11.57	0.31	64.33	50.91	78.98	2.22	16.74	3.46
36	Tarkol	5.50	5.00	1.00	92.58	3.58	24.00	7.16	24.54	0.36	133.00	108.92	81.83	2.69	32,45	11.14
37	Madoop	6.50	0.00	0.50	98.25	3.75	21.59	7.92	21.71	0.28	148.84	122.25	82.19	1.91	24.24	9.14
38	Waibang	9.00	5.00	1.50	73.92	3.42	19.08	6.92	13.09	0.24	65.58	56.92	86.22	2.61	19.78	5.26
39	Jhum Bini	7.00	2.00	1.00	93.83	3.50	24.17	9.75	21.59	0.31	97.08	82.76	85.27	2.69	29.65	8.66
40	Fazu Sen (White)	7.00	0.00	1.00	67.17	3.16	19.92	6.75	14.50	0.31	59.25	51.50	87.19	2.28	17.35	3.91
41	Bongbu	7.50	0.00	0.00	98.58	3.33	22.92	6.50	12.55	0.17	77.67	60.75	78.31	2.74	21.57	5.75
42	Sadok	8.50	1.00	1.00	62.33	2.75	21.83	5.92	8,78	0.22	63.92	50.83	79.52	2.39	21.38	3.61
43	Kala Jira	9.00	2.00	3.00	85.83	3.33	19.92	6.25	11.46	0.18	73.42	61.75	84.57	1.17	13.20	2.54
44	Gaigash	6.00	1.00	2.00	82.41	3.75	24.58	8.00	10.26	0.18	83.17	70.58	85.03	2.97	26.11	8.34
45	Vanbang	7.00	0.00	2.00	65.17	3.08	17.33	6.42	8.65	0.18	69.33	57.58	82.96	2.19	19.22	4.13
46	Makajaria	8.00	0.00	1.00	101.50	3.25	26.41	7.58	10.87	0.15	79.08	67.83	85.64	1.74	17.24	4.13
47	Jilong	7.50	1.00	1.00	80.17	3.42	21.50	5.75	11.24	0.19	71.66	63.33	88.23	2.03	28.06	5.12
48	American Ration	8.00	0.00	1.00	83.58	3.50	25.00	6.33	9.38	0.16	85.17	70.50	82.83	1.85	19.63	4.70
49	Kala Dhan	6.00	0.50	0.50	89.42	3.83	22.33	7.00	13.84	0.21	76.75	66.00	86.02	2.74	25.20	7.09
50	Turkey	7.50	0.00	0.00	107.58	3.83	22.83	7.67	14.87	0.18	133.41	119.75	89.38	2.64	34.88	12.39
51	SaankiKachak	5.50	0.00	0.50	76.83	3.16	21.42	9.16	9.76	0.18	99.33	84.59	85.11	2.24	29.95	6.50
52	MaimiUkhlao	7.00	1.00	1.00	81.83	3.00	27.42	8.17	9.60	0.18	88.75	71.09	79.45	1.91	20.02	4.34
53	Maiwasha	7.50	0.00	0.50	81.33	3.16	19.75	6.08	14.10	0.23	72.50	53.17	73.47	2.29	20.55	4.20
54	Maimi Watolok	5.50	3.00	0.00	100.00	3.16	22.17	8.92	20.53	0.27	113.25	97.34	85.95	2.91	35.03	9.25
55	Santin Wakhum	7.00	0.00	1.00	91.08	3.00	21.25	6.92	9.76	0.14	62.25	49.41	79.32	2.49	20.35	3.79
56	Yang Dhan	9.00	0.00	3.00	74.17	2.92	18.75	5.83	7.14	0.13	58.83	47.25	80.61	2.28	21.34	3.38
57	Badaya	9.00	0.00	4.00	62.16	2.75	20.42	5.72	7.11	0.17	67.17	56.09	83.24	2.60	21.53	4.28
58	Kanchali	9.00	0.00	4.50	77.83	3.00	20.08	7.25	12.77	0.22	76.83	61.25	79.61	1.81	17.00	3.51
59	Australian Biroin	6.50	0.00	1.00	112.50	3.33	29.42	9.00	18.49	0.22	90.92	67.92	74.75	2.20	20.01	5.79
60	Assam	7.00	0.50	1.00	96.67	4.17	24.75	7.67	14.86	0.21	107.17	95.25	88.64	1.42	23,91	5.88

VGR- Seedling vegetative vigour, LRI- Leaf rolling index, LDI-Leaf tip drying index, PL HT - Plant Height (cm.), PR TLR-Numbers of Productive Tiller per plant, PN LN -Panicle length (cm.), PRI BR-Numbers of Primary branch per panicle, RT LN-Root length (cm.), RT:CLM-Root- Culm ratio, SPKLT/PN-Numbers of Spike lets per panicle, FLD GRN/ PN-Numbers of filled grain per panicle, SPKLT FRT-Spikelet fertility (%), 100 SD WT-100 Seed's weight (gm.), HI-Harvest Index (%),GR Y /PLT-Grain yield per plant (gm.)

Table 3: Estimates of Genetic parameters for 15 (Fifteen) numbers of Morpho-Physiological characters of 60 (sixty) land races under drought stress condition

Character	Range Min. Max.		Grand Mean	Expected mean next	CV	GCV	PCV	H^2	GA as % of mean
	•			generation		·		·	5%
VGR.	5.0	9.0	7.17	8.76	12.06	14.15	18.60	0.57	22.19
LRI	0.00	5.00	0.89	3.28	50.81	137.95	147.01	0.88	266.66
LDI	0.00	4.50	1.11	2.62	71.99	86.06	112.20	0.58	135.99
PL. HT.	61.24	125.41	86.17	108.39	10.29	15.14	18.31	0.68	25.79
PR TLR	2.74	4.5	3.47	3.95	12.44	10.51	16.29	0.41	13.98
PN LN	16.83	29.49	22.87	27.51	9.45	12.40	15.59	0.63	20.31
PRI BR	5.25	10.24	7.11	8.84	11.51	14.95	18.87	0.62	24.41
RT LN	7.11	26.52	14.28	23.56	15.17	34.44	37.64	0.83	64.94
RT: CLM	0.13	0.35	0.22	0.34	8.22	26.76	27.99	0.91	52.69
SPKLT/PN	57.49	195.33	88.51	139.88	13.25	30.68	33.42	0.84	58.03
FLD GRN/ PN	45.41	178.58	74.08	122.64	14.30	34.45	37.30	0.85	65.55
SPKLT FRT	70.34	91.43	83.06	90.27	3.52	5.117	6.21	0.67	8.67
100 SD WT	1.10	2.99	2.29	3.13	6.36	18.60	19.65	0.89	36.25
HI	12.8	40.86	23.81	34.74	14.50	25.61	29.43	0.75	45.90
GR Y /PLT	2.53	21.66	6.53	13.03	23.40	52.77	57.72	0.83	99.37

Min-Minimum, Max- Maximum, CV- Coefficient of variance, GCV - genotypic coefficient of variations PCVphenotypic coefficient of variations, H^2 – Broad sense heritability in percentage, GA – Genetic advance

LEAF ROLLING AT VEGI	ETATIVE STAGE							
	MaimiUzra, Chinal, American, Biroin, Aaduma, Suri, Saanki ka phool,							
No sign of loof rolling	Bangbu Jhum, Lal Biroin, Goria, Bahadur, Mami WatoklokMandoori,							
No sign of leaf forming	Bangbu, Vanbang, Makajaraia, Turkey, SaankiKachak, SaantinWakhom,							
	Yang dhan, Badaya, Kanchali, Australian Biroin							
	KhasaKasam, Lebuka, BetiKalai, ChikanswariKabar, DhalaBalam,							
Slight leaf rolling	MaimiHungar, Kaladhan, Assam Paisom							
	Tarkol, Waibang, MaimiWatoklok, Galong, Beti, Kaporok, Kalikhasa, Sadok,							
Moderate leaf rolling	FazuVom, Kalajira							
LEAF DRYING AT VEGE	TATIVE STAGE							
	Releng, Beti, American, Biroin, Garomalati, Lebuka, Saanki ka phool,							
No sign of leaf drying	Bangbu, MaimiWatoklok, Turkey							
~~ ~ ~ ~ ~ ~	Maimi Red, BetiKalai, Bihar, ChikanswariKabar, Goria, MaimiHungar,							
Slight leaf drying	MaimiTaukha, Madoop, Waibang, Kaladhan, Maiwasha							
Moderate to high leaf	Chinal, SadaBiroin, Kala Jira, Yang Dhan, Badaya, Kanchali							
drving	·······, ·····························							
ROOT LENGTH								
More than 20 cm under	Kanorok Releng Garomalati ChikanswariKahar MaimiHungar Tarikol							
drought stress	Kolte Tarkol Madoon MaimiWatoklok							
PLANT HEIGHT	Tono, Furkor, Mudoop, Munni Wuoklok							
Tall (More than 125 cm)	ChikanswariKahar							
Tan (Wore than 125 cm.)	Garo malati Kaporok MaimiUzra Galong BetiKalai Goria Bahdur							
Intermediate	M W Mandoori MaimiHungar Tarikolkolte Tarkol Madoon Ihum Bini							
(90, 125) cm	Ranghu Makajaria Turkey MaimiWatoklok SantinWakhum Australian							
(90-125) em.	Biroin Assam Paisom							
Semidwarf	Past of the landraces							
(Less than 90 cm)	Rest of the fandraces							
CDAIN VIELD DED DI AN	<u>т</u>							
ORAIN HELD FER FLAN	ChilcongwariKahan Canomalati Calang Turkay Tarkal							
More than 10 gm	Chikanswan Kabai, Galoinalati, Galoing, Turkey, Tarkor Dhala Dalam Maimi Hungar							
100 SEED WEICHT	,Dharabarani,Maininhungar.							
100 SEED WEIGHT	M.'. 'II							
	MaimiHungar, Gaigash, Maimi watokiok, Maimi Red, Bangbu jnum, Reieng,							
More than 2.5 gm.	Galong, Garo Malan, Fazu Sen, BenKalai, UnikanswariKabar, Lai Biroin,							
C	DhalaBalam, Tarikolkolte, Tarkol, Walbang , Jhum Bini, Bongbu, Kala dhan,							
	Turkey, Badya							
ROOT: CULM RATIO								
More than 0.3	Tarkol, Releng, MaimiHungar, MaimiTaukha, Kaporok, Jhum Bini, Fazu Sen							
	(white)							
NUMBERS OF FILLED G	RAIN PER PANICLE							
More than 150 numbers of	Galong, Garomalati, ChikanswariKabar, Madoop, Tarkol							
filled grain per panicle								
NUMBERS OF PRODUCT	IVE TILLERS							
More than average 4	Kaporok, Garomalati, Chikanswarikabar, DhalaBalam, Assam Pisom, Aaduma							
Numbers of tillers								

Table 4: Grouping of landraces based on Mean value analysis of some significant morphological traits

Table 5: Details of the microsatellite primers used in present study with their Allele size (bp), Allele diversity, Gene diversity and Polymorphic information content (PIC)

Sr. No	Primer	Motif	C hr.	An. Tmp (c ⁰)	Exp. PCR product [bp]	Actual PCR product range[bp]	Na	MA _{fr}	Не	PIC	Database reference (Gramene)
1	RM333	(TAT)19(CTT)19	10	55	191	100-200	2	0.905	0.172	0.158	Tenmykh et al.(2000)
2	RM482	(AT)9	2	55	188	190-210	3	0.667	0.470	0.394	, , , , , , , , , , , , , , , , , , ,
3	RM487	(AC)10	3	55	176	180-190	2	0.828	0.285	0.245	T 11 1(2001)
4	RM491	(AT)14	12	55	263	280-300	2	0.867	0.231	0.204	Tenmykh et al.(2001)
5	RM592	(ATT)20	5	55	270	200-300	4	0.744	0.419	0.388	
6	RM175	(CCG)8	3	67	95	80-180	4	0.464	0.669	0.613	Tenmykh et al. (2000)
7	RM205	(CT)25	9	55	122	120-180	3	0.696	0.469	0.423	Cl (1007)
8	RM252	(CT)19	4	55	216	180-900	6	0.300	0.777	0.743	Chen <i>et al.</i> (1997)
9	RM276	(AG)8A3(GA)33	6	55	149	150-300	4	0.511	0.645	0.591	Tenmykh et al.(2000)
10	RM339	(CTT)8CCT(CTT)5	8	55	148	120-300	4	0.473	0.664	0.608	Tenmykh et al.(2000)
11	RM437	(AG)13	5	55	275	250-300	3	0.643	0.513	0.450	•
12	RM458	(TAG)8	8	55	180	180-200	2	0.510	0.500	0.375	
13	RM484	(AT)9	10	55	299	180-300	3	0.654	0.494	0.426	
14	RM486	(CT)14	1	55	104	150-180	2	0.754	0.371	0.302	
15	RM488	(GA)17	1	55	177	170-180	2	0.655	0.452	0.350	
16	RM505	(CT)12	7	55	199	200-500	4	0.290	0.747	0.700	Tenmykh et al. (2001)
17	RM506	(CT)13	8	55	123	100-130	3	0.413	0.657	0.583	•
18	RM538	(GA)14	5	55	274	280-300	2	0.863	0.237	0.209	
19	RM513	(TC)11	1	55	262	250-260	2	0.553	0.494	0.372	
20	RM570	(AG)15	3	55	208	150-200	3	0.455	0.619	0.539	
21	RM566	(AG)15	9	55	239	230-250	3	0.558	0.582	0.512	
22	RM167	(GA)16	11	55	128	120-200	3	0.456	0.618	0.538	Wu et al. (1993)
23	RM172	(AGG)6	7	55	169	175	1	1.000	0.000	0.000	Tenmykh et al.(2000)
24	RM213	(CT)17	2	55	139	120-200	3	0.646	0.503	0.435	
25	RM215	(CT)16	9	55	148	120-150	3	0.438	0.633	0.556	
26	RM 231	(CT)16	3	55	182	150-200	3	0.527	0.594	0.518	Chen <i>et al.</i> (1997)
27	RM257	(CT)24	9	55	147	140-180	3	0.597	0.525	0.439	
28	RM282	(GA)15	3	55	136	100-150	3	0.588	0.552	0.479	
29	RM287	(GA)21	11	55	118	80-120	3	0.627	0.514	0.440	Tenmykh et al. (2000)
30	RM320	(AT)11GTAT(GT)13	7	55	167	100-400	4	0.429	0.669	0.608	
31	RM3	(GA)2GG(GA)25	6	55	145	100-150	3	0.586	0.556	0.484	Panaud et al. (1996)
32	RM30	(AG)9A(GA)12	6	55	105	70-100	3	0.542	0.577	0.498	Chen et al. (1997)
33	RM52	(AG)19	8	55	240	250-500	3	0.867	0.239	0.225	Tenmykh et al. (2000)
34	RM146	(CT)11-(CT)7	5	55	345	300-350	3	0.607	0.503	0.407	Tenmykh et al. (2000)
35	RM233	(CT)20	2	55	162	150-170	2	0.558	0.493	0.372	Chen et al. (1997)
36	RM254	(TC)6ATT(CT)11	11	55	165	100-800	5	0.453	0.687	0.630	Chen et al. (1997)
37	RM285	(GA)12	9	55	205	200-800	4	0.444	0.697	0.648	Tenmykh et al. (2000)
38	RM315	(AT)4(GT)10	1	55	133	100-130	3	0.623	0.523	0.453	Tenmykh et al. (2000)
39	RM452	(GTC)9	2	55	209	200-230	2	0.964	0.069	0.067	Tenmykh et al. (2001)
40	RM 276	(AG)8A3(GA)33	6	55	149	150-180	2	0.600	0.480	0.365	Tenmykh et al. (2000)
]	TOTAL						119	24.35	19.89	17.34	
A١	/ERAGE						2.97	0.608	0.497	0.433	

Chr.: Chromosome location, An. Tmp: Annealing temperature, Exp.PCR product [bp]: Expected PCR product size in base pair, Na: Total number of alleles per locus, MA_{fr}: Major allele frequency, *He*: Gene Heterozygosity, PIC: Polymorphism information content

Traits	Molecular Clustering pattern								
	Name of the landraces	Sub group							
	Kaporok, Releng, Tarkol, Tarikol Kolte	B_1							
Root length (more than 20 cm.)	Maimi Hungar,Chikanswari Kabar Madoop,Garomalati	Scattered in different cluster							
Poot: Culm ratio	Kaporok, Releng, Tarkol, Tarikol Kolte	B ₁							
(more then $(0, 3)$)	Madoop , Maimi Watoklok	A_1							
(more men 0.3)	MaimiHungar	C_1C							
Moderate to high leaf drying index	Chinal, Sada Biroin, Kala Jira, Kanchali	Scattered in different cluster							
	Yang Dhan, Badaya	C_2A							
Low to moderate leaf rolling index	Kaporok, Releng, Tarkol, Beti	B_1							
Moderate to high leaf drying index	Badaya, Yang Dhan	C_2A							
Plant height (more than 125 cm.)	Chikanswari Kabar	C ₂ C							
More than average 4 numbers of productive tillers	Kaporok, Garomalati, Chikanswari Kabar, Dhala Balam, Assam Paisom, Aaduma	Scattered in different cluster							
100 seed weight (more than 2.5 gm.)	Maimi Hungar, Gaigash, Maimi Watoklok, Maimi Red, Bangbu Jhum, Releng ,Galong, Garomalati, Fazu Sen, BetiKalai, Chikanswari Kabar, Lal Biroin, DhalaBalam, Tarikol Kolte, Tarkol, Waibang , Jhum Bini, Bongbu, Kaladhan, Turkey, Badaya	Scattered in different cluster							
Numbers of filled grain	Galong, Garomalati, ChikanswariKabar,	Scattered in							
per panicle (more than 150	Madoop, Tarkol	different							
Numbers)		cluster							
Grain yield per plant	ChikanswariKabar, Garomalati, Galong,	No specific							
(more than 10 gm)	Turkey, Tarkol, Dhalabalam, MaimiHungar	cluster							

Table 6: Similarities in the grouping pattern of landraces under mean value analysis and their clustering pattern under molecular analysis



Fig. 1 a & b: Amplification profile of 60 rice landraces generated through microsatellite primer - RM 257



Fig. 2 a & b: Amplification profile of 60 rice landraces generated through microsatellite primer - RM 538



Fig. 3 a & b: Amplification profile of 60 rice landraces generated through microsatellite primer - RM 254

L- DNA ladder, 1. Kaporok 2. Releng 3. Beti 4. MaimiUzra 5. Kalikhasa 6. Chinal 7. American 8. KhasaKasam 9. Biroin10.Galong 11. FazuVom 12. Garo Malati 13. Maimi Usha 14. Maimi Red 15. Suri 16.Lebuka 17.Aaduma 18.Fazu Sen 19.Fazu N'G 20.Beti Kalai 21.Saanki ka Phool 22.Bihar 23.Chikanswari Kabar 24.Bangbu Jhum 25.Lal Biroin 26.Dhala Balam 27.Goria 28.Bahadur 29.M.W.Mandoori 30.Mami Hungar 31.Darka Sona 32.Tarikol Kolte 33.Sada Biroin 34.Maimi Taukha 35.Saluma 36.Tarkol 37.Madoop 38.Waibang 39.Jhum Bini 40.Fazu Sen (White) 41.Bongbu 42.Sadok 43.Kala Jira 44.Gaigash 45.Vanbang 46.Makajaria 47.Jilong 48.AmericanRation49.KalaDhan 50.Turkey 51.Saanki Kachak 52. Maimi Ukhlao 53. Maiwasha 54. MaimiWatolok 55. SantinWakhum 56. YangDhan 57. Badaya 58. Kanchali 59. Australian Biroin 60. Assam Paisom



Fig 4 Clustering pattern of 60 nos. of rice landraces under morphological analysis



Fig 5 Clustering pattern of 60 nos. of rice landraces under molecular analysis

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