



Screening of Pearl Millet [*Pennisetum glaucum* [L.] R. Br.] Germplasm Lines against Drought Tolerance Based on Biochemical Traits

M. L. Choudhary¹, M. K. Tripathi^{1*}, Neha Gupta¹, Sushma Tiwari¹, Niraj Tripathi², Prerana Parihar³ and R. K. Pandya³

¹Department of Plant Molecular Biology and Biotechnology, College of Agriculture, RVS Agricultural University, Gwalior, 474002 M.P., India.

²Directorate of Research Services, JN Agricultural University, Jabalpur 482004, India.

³Department of Plant Pathology, College of Agriculture, RVS Agricultural University, Gwalior, 474002 M.P. India.

Authors' contributions

This work was carried out in collaboration among all authors. All authors read and approved the final manuscript.

Article Information

DOI: 10.9734/CJAST/2021/v40i2331483

Editor(s):

(1) Dr. Alessandro Buccolieri, Università del Salento, Italy.

Reviewers:

(1) Mohammad Hossain, Bangladesh.

(2) Khushboo Dubey, India.

Complete Peer review History: <https://www.sdiarticle4.com/review-history/73335>

Original Research Article

Received 26 June 2021
Accepted 06 September 2021
Published 10 September 2021

ABSTRACT

Aim: The current investigation was commenced to investigate genetic miscellany among pearl millet genotypes based on diverse biochemical parameters interrelated to drought tolerance.

Study Design: In investigation, 96 pearl millet germplasm lines were screened against drought using diverse biochemical traits.

Place and Duration of the Study: The present study was conducted at College of Agriculture, Gwalior, Rajmata Vijayaraje Scindia Krishi Vishwa Vidyalaya, Gwalior, M.P., India during July 2019 to December, 2020.

Methodology: Five biochemical parameters viz., chlorophyll content, carotenoid content, total soluble sugars, proline and protein were investigated for explanation of differences among 96 pearl millet germplasm lines in respect to drought tolerance.

Results: Data of present investigation revealed the mean leaves chlorophyll at 30DAS was 2.90

mgg⁻¹ with the range of 1.31-4.69 mgg⁻¹, whereas chlorophyll at 60DAS was arrayed between 1.46-3.84 mgg⁻¹ with an average of 3.02 mgg⁻¹. Carotenoid at 30DAS was ranging from 4.5-11.44 mgg⁻¹ with an average value of 7.23 mgg⁻¹, while carotenoid at 60DAS was recorded in range of 5.01 to 10.10 mgg⁻¹ with an average of 6.66 mgg⁻¹. TSS was ranged between 1.10-2.20 mgg⁻¹, proline 0.10 to 0.17 mg g⁻¹ and protein content 9.2-16.60 mgg⁻¹.

Conclusions: According to the biochemical data a total of 16 pearl millet genotypes were found to be grouped distantly among all the genotypes. Possibility existed to be drought tolerance of these genotypes.

Keywords: Drought; bio-chemical; crop improvement; pearl millet.

1. INTRODUCTION

Pearl millet [*Pennisetum glaucum* (L.) R. Br.] is cultivated under both in arid and semi-arid circumstances in India, where other cereals are hard to grow [1]. It is the most imperious cultivated cereals in the world, standing sixth following rice, wheat, maize, barley and sorghum in rappers of expanse. It is cultivated on around 30 mha in more than 30 countries [2]. The mainstream of this acreage is in Asia, Africa and Americas [3]. In India, pearl millet is the fourth most widely cultivated edible crop after rice, wheat and maize [4]. It occupies 7.48 million hectares with an average production of 9.21 million tonnes and the productivity of 1231 kgha⁻¹ during 2017-18 [5]. In Madhya Pradesh area under cultivation is 0.31 million hectares with 0.76 million tonnes production and 2435 kgha⁻¹ productivity.

Nutritionally, pearl millet is a good source of energy and high levels of minerals vitamins, lipids, crude fibres and high-quality protein 9-13% [6]. Abdalla et al. [7] analysed pearl millet grain and reported 4.31-5.30 per cent crude fibre, 1.53-2.00 per cent ash, 450-990 mg phosphorus, 10-80 mg calcium, 7-18.0 mg iron, 5.3-7.0 mg zinc, 1.0-1.8 mg copper and 1.8-2.3 mg manganese content. Carbohydrates are the main component of *Pennisetum glaucum* grains varying from 71.82 to 81.02 per cent [8]. Fat content of pearl millet varieties vary from 4.32 to 5.11 per cent [7]. The total sugars in pearl millet ranges from 2.55 to 2.93 per cent, non-reducing sugars between 2.15 to 2.57 per cent and reducing sugars from 0.34 to 0.39 per cent [9,10]. Due to the nutritional superiority and climate-resilient nature of pearl millet over other crops, it has given the tag of “nutricereals” by the Ministry of Agriculture and Farmers Welfare, Government of India. Over 170 million individual's dependents on pearl millet as an essential from India. It is mostly used in deprived countries and by the poorest peoples. So, also known as the “Poor man's cereal crop” [11].

Although, pearl millet is a resilient crop proficient of growing in punitive and peripheral environments where no other cereals may give economic returns but same time its yield reduced drastically due to drought conditions substantially. Drought itself is a complex phenomenon and several parameters influencing it were found to be under genetic control. Drought stress is the most important environmental constraint limiting factor for crop production worldwide [2, 12,13,14,15]. Drought is the most damaging abiotic stress affecting crop productivity, which is caused by insufficient rainfall and/or altered precipitation patterns [16]. The seriousness of drought stress depends on its timing, period and intensity [17]. Drought limits the agricultural production by preventing the crop plants from expressing their full genetic potential [18]. Out of those, terminal drought is shown to contribute to the foremost severe yield losses because it affects spikelet establishment and reduces its fertility [19].

Acclimation of plants to water deficit is the results of different events, which result in adaptive changes in plant growth and physio-biochemical processes [20]. Total soluble sugars and proline are also increased [21,22]. Increase in proline which are closely linked with drought tolerance and have the potential to improve crop yield [23]. Chlorophyll a, b, total chlorophyll and carotenoid were significantly reduced under water stress treatments. Water stress increased the proline [24]. Drought stress generally causes decrease in the total chlorophyll content [25,26,27] while the Chlorophyll a/b ratio usually increases [28].

No systematized works on screening of these pearl millet germplasm lines on the basis of biochemical parameters carried out so far. Consequently, in current study an effort has been made to judge the extent of genetic diversity present among diverse pearl millet germplasm lines putative tolerant to drought by means of different biochemical indices.

2. MATERIALS AND METHODS

The present investigation was consisted of 96 genotypes (Table) with diverse responses to drought *viz.* susceptible and tolerant. The seeds of germplasm lines were acquired from AICRP on pearl millet, College of Agriculture, Gwalior, RVSKVV, Gwalior, M. P., India collected from different sources. The experimental material was scrutinized in randomized block design with two replications. The seeds were sown by hand dibbling. Rainfall augmented generally the irrigation necessities. Fortunately, no rainfall was chronicled during period between 50-70 days afterward sowing which was proved helpful for enhancing drought conditions and thereby estimating biochemical parameters. The sampling was done at 30 and 60 days after sowing (DAS). Five plants were arbitrarily selected from each treatment per replication for conducting biochemical analysis and subsequently recording data. Data were recorded for various biochemical parameters *viz.*, chlorophyll, carotenoid, proline, protein and sugar to efficiently screen drought tolerant and susceptible genotypes.

2.1 Biochemical Screening

In the current study, different biochemical parameters like total chlorophyll, carotenoid, proline, soluble sugar contents and protein percentage in immature seeds (under developmental stage) were unhurried. Photosynthetic pigments were estimated by Arnon's method [29]. Free proline content in leaves was determined according to the method proposed by Bates et al. [30] based on the formation of red colour by proline with ninhydrin in acidic medium. The total sugar was estimated as per protocol described by Dubois et al. [31]. The protein content was calculated as per method given by Lowry et al. [32].

2.2 Data Analysis

Data were analysed as suggested by Snedecor & Cochran [33]. Dendrogram analysis of 96 pearl millet germplasm lines based on similarity between these germplasm lines on the basis of different biochemical parameters *i.e.*, proline, sugar and protein were depicted by using NTSYS ver 2.0 software.

3. RESULTS AND DISCUSSION

3.1 Biochemical Variations among Pearl Millet Germplasm Lines

In the present investigation, biochemical traits such as chlorophyll, carotenoid, total soluble sugars, proline and protein contents were investigated on explanation of drought stress. Present study showed that drought increased level of sugar, proline and protein. Analysis of variance was found significant for most of the traits that suggested existence of substantial sum of variability in studied materials for further improvement (Table 2).

3.2 Chlorophyll Content (mgg^{-1} Fresh Weight)

Data depicted reveals that there was presence of significant magnitude of variations in chlorophyll content among different germplasm lines. Chlorophyll content taken at 30 days ranged from 1.31 mgg^{-1} to 4.69 mgg^{-1} with an average value of 2.90 mgg^{-1} . Out of the ninety-six pearl millet germplasm lines studied, genotype IP194 (4.69 mgg^{-1} fresh weight) contained highest chlorophyll content followed by genotypes IP168 (4.35 mgg^{-1} fw), IP161 (4.13 mgg^{-1} fw) whereas minimum chlorophyll content displayed by genotype IP127 (1.31 mgg^{-1} fw). Chlorophyll content at 60 days varied between 1.46 mgg^{-1} to 3.84 mgg^{-1} with a mean value of 3.02 mgg^{-1} . Genotypes IP161 (3.84 mgg^{-1} fw) portrayed highest chlorophyll content tracked by germplasm lines IP165 (3.83 mgg^{-1} fw) and IP143 (3.80 mgg^{-1} fw) whilst minimum displayed by genotype IP127 (1.46 mgg^{-1} fw) (Table 1). Usually, the level of chlorophyll content in leaves regulates the proportion of photosynthesis. Generally, a reduction in chlorophyll component was found in susceptible genotypes in comparison to tolerant genotypes [34]. Abridged level of chlorophyll synthesis in susceptible genotypes may be the motive of less bustle of the photosynthetic elements. Formerly, forfeiture of chloroplast membranes under drought stress has also been reported by Anjum et al. [35]. Analogous reduction in chlorophyll levels in many other plant species *viz.*, soybean, maize, rice, chickpea, pearl millet *etc.* have also been reported [15, 36,37,38]. Higher reduction in chlorophyll content was evidenced in drought susceptible genotypes in the current investigation.

Table 1. Complete Biochemical data of 96 pearl millet germplasm lines

Sr. No.	Name of genotype	Chlorophyll content [30 days] [mg/g]	Chlorophyll content [60 days] [mg/g]	Carotenoid content [30 days] [mg/g]	Carotenoid content [60 days] [mg/g]	Proline [mg/g]	Sugar [mg/g]	Protein [mg/g]
1	IP 132	3.57	3.46	8.29	7.50	0.10	1.60	12.40
2	IP 118	3.63	3.47	8.13	7.56	0.12	2.10	11.20
3	IP 152	4.06	3.64	9.35	8.12	0.16	1.20	11.80
4	IP 175	2.75	3.09	5.91	6.35	0.10	1.30	12.80
5	IP 133	2.43	2.69	4.54	5.94	0.11	2.20	14.20
6	IP 173	3.43	3.44	6.00	5.02	0.14	1.10	12.70
7	IP 199	1.76	2.06	6.22	7.46	0.16	1.40	11.90
8	IP 127	1.31	1.46	6.03	7.00	0.11	1.70	13.60
9	IP 198	1.98	2.04	6.74	7.92	0.12	1.80	11.40
10	IP 177	3.47	3.26	7.96	5.51	0.14	1.90	14.20
11	IP 182	2.46	2.70	5.92	7.34	0.10	1.70	13.90
12	IP 147	2.84	3.59	7.12	5.46	0.17	1.40	11.20
13	IP 107	3.19	3.44	7.81	7.28	0.12	1.80	15.90
14	IP 140	3.00	3.50	5.83	5.38	0.10	2.20	16.20
15	IP 164	3.33	3.21	8.50	10.10	0.12	1.40	15.40
16	IP 142	3.40	3.10	6.68	6.58	0.16	1.50	14.20
17	IP 180	3.11	3.47	6.31	6.60	0.10	1.40	11.00
18	IP 188	2.90	3.10	6.35	6.42	0.11	1.80	9.20
19	IP 181	1.71	2.58	4.85	5.41	0.14	2.20	12.70
20	IP 129	1.51	1.81	5.77	5.68	0.16	1.40	11.90
21	IP 119	2.36	2.23	6.13	6.78	0.11	1.50	13.60
22	IP 150	2.31	2.63	5.62	5.42	0.12	1.60	11.40
23	IP 120	2.50	3.04	6.36	6.71	0.14	2.10	14.20
24	IP 111	1.61	2.18	6.14	6.38	0.10	1.20	13.90
25	IP 160	2.73	3.28	5.56	5.78	0.17	1.30	16.10
26	IP 136	1.87	2.36	5.86	7.75	0.12	2.20	15.20
27	IP 171	2.63	3.02	7.26	5.95	0.10	1.10	15.50
28	IP 130	3.16	3.38	7.71	5.38	0.12	1.40	16.60
29	IP 166	3.50	3.34	8.67	7.86	0.16	1.70	12.70
30	IP 128	3.58	3.66	8.12	5.29	0.10	1.80	13.50
31	IP 183	3.44	3.46	7.85	6.85	0.11	1.90	14.70
32	IP 165	3.63	3.83	9.96	6.68	0.14	1.70	12.80
33	IP 192	3.24	3.54	10.57	6.59	0.16	1.20	14.20

Sr. No.	Name of genotype	Chlorophyll content [30 days] [mg/g]	Chlorophyll content [60 days] [mg/g]	Carotenoid content [30 days] [mg/g]	Carotenoid content [60 days] [mg/g]	Proline [mg/g]	Sugar [mg/g]	Protein [mg/g]
34	IP 122	3.99	3.54	7.23	5.68	0.11	1.30	12.70
35	IP 143	4.00	3.80	9.45	5.30	0.12	1.50	11.90
36	IP 167	3.65	3.38	8.78	6.21	0.14	1.40	13.60
37	IP 172	2.91	3.34	6.14	7.89	0.10	1.80	11.40
38	IP 106	3.27	3.22	8.20	6.03	0.17	1.90	14.20
39	IP 137	3.39	3.63	7.58	5.82	0.12	2.00	13.90
40	IP 116	3.51	3.59	7.20	6.34	0.10	2.10	11.20
41	IP 194	4.69	3.77	11.44	5.01	0.12	2.20	12.60
42	IP 195	3.76	3.70	9.02	7.99	0.16	1.90	15.50
43	IP 126	3.08	3.09	4.73	5.06	0.10	2.10	16.60
44	IP 155	2.64	2.77	6.35	6.02	0.12	1.60	12.70
45	IP 149	3.32	3.22	5.90	5.85	0.14	1.70	13.50
46	IP 185	3.64	3.57	7.84	5.16	0.16	1.80	14.70
47	IP 161	4.13	3.84	7.55	5.08	0.14	1.90	12.80
48	IP 168	4.35	3.80	5.66	5.13	0.12	1.90	14.20
49	IP 190	3.64	3.40	7.85	5.31	0.14	1.70	12.70
50	IP 156	1.66	2.01	7.86	7.36	0.10	1.80	11.90
51	IP 187	4.10	3.46	8.63	8.09	0.17	1.90	12.40
52	IP 159	3.12	3.00	10.61	8.00	0.12	1.70	11.50
53	IP 139	2.81	2.96	7.81	5.51	0.10	1.80	11.00
54	IP 146	3.12	3.47	6.38	5.78	0.12	1.60	13.10
55	IP 196	2.86	2.83	9.49	8.28	0.16	1.40	13.20
56	IP 186	2.01	2.52	7.43	8.59	0.10	1.20	11.40
57	IP 158	2.58	2.75	7.55	6.64	0.11	2.10	12.30
58	IP 151	2.92	3.29	4.61	5.78	0.14	1.90	13.40
59	IP 193	2.79	3.02	9.36	8.17	0.16	1.50	11.60
60	IP 105	2.71	3.15	6.23	6.39	0.13	2.00	10.60
61	IP 123	2.84	3.13	8.63	7.99	0.12	1.20	9.90
62	IP 131	3.11	3.01	7.86	7.26	0.14	1.40	11.20
63	IP 178	3.15	3.13	6.68	5.86	0.12	2.10	14.80
64	IP 121	2.75	2.78	5.90	5.68	0.17	1.60	15.20
65	IP 104	2.65	2.98	6.87	5.97	0.12	1.70	11.60
66	IP 134	2.66	3.05	7.46	5.49	0.10	1.80	12.90
67	IP 112	2.38	2.62	5.44	5.45	0.12	1.90	11.80
68	IP 141	2.52	2.62	7.07	8.26	0.13	1.90	12.80

Sr. No.	Name of genotype	Chlorophyll content [30 days] [mg/g]	Chlorophyll content [60 days] [mg/g]	Carotenoid content [30 days] [mg/g]	Carotenoid content [60 days] [mg/g]	Proline [mg/g]	Sugar [mg/g]	Protein [mg/g]
69	IP 145	1.98	2.34	4.87	7.09	0.11	1.70	14.20
70	IP 144	2.07	2.12	5.69	8.30	0.11	1.80	12.70
71	IP 138	2.23	2.60	7.53	5.77	0.14	1.90	11.90
72	IP 179	2.38	2.65	7.70	6.05	0.16	1.70	12.40
73	IP 153	2.69	2.83	6.84	6.03	0.13	1.80	11.50
74	IP 101	2.50	2.24	5.94	6.14	0.12	1.60	11.00
75	IP 135	2.52	2.87	6.30	6.12	0.14	1.40	13.10
76	IP 162	2.72	3.15	7.90	8.14	0.10	1.20	13.20
77	IP 115	2.46	2.77	6.21	5.96	0.13	2.10	11.40
78	IP 170	3.10	3.23	7.73	5.99	0.12	1.90	12.30
79	IP 109	2.72	2.71	7.45	8.16	0.10	1.50	13.40
80	IP 154	2.19	2.78	4.50	5.57	0.12	2.00	11.60
81	IP 174	2.90	2.84	7.54	8.20	0.16	1.20	13.90
82	IP 108	2.68	2.88	7.84	6.60	0.12	1.40	11.20
83	IP 189	3.59	3.35	9.55	8.23	0.11	1.70	15.90
84	IP 110	3.09	3.01	7.63	8.27	0.14	1.80	16.20
85	IP 117	3.26	2.99	7.82	7.26	0.16	1.90	15.40
86	IP 169	2.65	3.04	7.52	6.83	0.11	1.70	14.20
87	IP 114	2.25	2.35	8.61	8.44	0.12	1.40	11.00
88	IP 163	2.04	2.41	5.55	5.80	0.14	1.80	9.40
89	IP 274	2.99	3.32	9.11	8.82	0.10	2.20	11.60
90	IP 283	2.10	2.76	6.70	6.47	0.17	1.40	11.90
91	IP 236	2.76	2.95	6.36	6.89	0.10	1.50	12.40
92	IP 291	2.52	2.66	7.62	7.15	0.10	1.40	11.40
93	IP 230	2.69	2.78	7.39	7.69	0.12	1.80	13.10
94	IP 262	3.18	2.68	7.73	5.72	0.16	2.20	13.40
95	IP 231	2.98	2.92	7.65	6.06	0.14	2.00	16.10
96	THAK 1827	3.62	3.79	8.17	8.93	0.11	1.80	15.20
Mean		2.90	3.02	7.23	6.66	0.13	1.70	13.02
Range	Min	1.31	1.46	4.50	5.01	0.10	1.10	9.20
	Max	4.69	3.84	11.44	10.1	0.17	2.20	16.60
CD_{0.05}		0.0928	0.1396	0.3953	0.3183	0.0032	0.0835	0.4685

Table 2. Correlation coefficient among different biochemical parameters of pearl millet germplasm lines

Parameters	Correlations						
	CH ₃₀	CA ₃₀	CH ₆₀	CA ₆₀	Proleine	Sugar	Protein
CH ₃₀	1	.537**	.885**	-.097	.131	.097	.213*
CA ₃₀		1	.425**	.354**	.157	-.137	-.016
CH ₆₀			1	-.173	.075	.073	.197
CA ₆₀				1	-.072	-.191	-.015
Proleine					1	-.099	.085
Sugar						1	.111
Protein							1

** Correlation is significant at the 0.01 level [2-tailed].

* Correlation is significant at the 0.05 level [2-tailed].

CH30= Chlorophyll 30 days, CA30= Carotenoid 30days, CH60= Chlorophyll 60 days, CA60= Carotenoid 60days

3.3 Carotenoid Content (mgg⁻¹ Fresh Weight)

Carotenoid content at 30 days ranging between 4.5 mgg⁻¹ to 11.44 mgg⁻¹ with an average of 7.23 mgg⁻¹. The highest carotenoid content at 30 days was evidenced in the genotype IP194 (11.44 mgg⁻¹) trailed by genotypes IP159 (10.61 mgg⁻¹ fresh weight) and IP192 (10.57 mgg⁻¹ fw), while minimum demonstrated by genotype IP154 (4.5 mgg⁻¹ fw). Carotenoid content at 60 days differed from 5.01 mgg⁻¹ to 10.10 mgg⁻¹ with mean worth of 6.66 mgg⁻¹. The highest carotenoid content at 60 days was recorded in the genotype IP164 (10.10 mgg⁻¹ fw) intimately tracked by Thak1827 (8.93 mgg⁻¹ fw), IP 274 (8.82 mg g⁻¹ fw). However, least carotenoids content was documented in the genotype IP154 (5.01 mgg⁻¹ fw) (Table 1).

3.4 Total Soluble Sugars (TSS) Content (mgg⁻¹ Fresh Weight)

Sugar content was documented in range of 1.10 to 2.20 mgg⁻¹ with the highest in genotypes IP133, IP144, IP181, IP136, IP196, IP262 (2.20 mgg⁻¹ fresh weight) and minimum TSS content was recorded in genotype IP173 (1.10 mgg⁻¹ wt.) (Table 1). The accumulation of soluble sugars in plants response to drought stress is well documented. The role of soluble sugars in plant metabolism as typical osmo-protectants, stabilizing cellular membranes and maintaining turgor pressure. It was claimed that under drought conditions, sugar fluidity may even be a signal for metabolic directive. Soluble sugars are the key osmotic adjustment substances and important indicators of drought tolerance [15,27, 38, 39, 40]. During present study germplasm lines possessing higher levels of sugar might be drought tolerant.

3.5 Proline Content (mgg⁻¹ Fresh Weight)

The mean value of proline content ranges from 0.10 to 0.17 mgg⁻¹ fresh weight with an average of 0.13 mgg⁻¹ fr.Wt (Table 1). Highest proline content was recorded in genotypes IP147 tracked by genotypes IP160, IP106, IP187 and IP121 and minimum in genotypes IP134 trailed by germplasm lines IP162, IP274, IP2426 and IP291 (0.10 mg g⁻¹ fr. Wt). Proline is whispered as an imperative drought tolerance pointer and estimated in ninety-six genotypes during the present investigation. Role of proline in osmotic regulation under water stress has been monitored in various plant species [15, 38, 41-43]. Pearl millet genotypes with significant rise in proline contents have been considered as drought tolerant. Increased proline content maintains cell water level under drought [15, 38,44]. Further, George et al. [45] suggested that increased proline has osmo-protective functions by preventing separation of enzymes during metabolic activities. It seems to proline may play a role in minimizing the injury caused by dehydration. Similar results were also documented by Mohammad and Heidari [46].

3.6 Protein Content (mgg⁻¹ Fresh Weight)

Protein content varied significantly in array of 9.2-16.60 mgg⁻¹ with an average of 13.02 mgg⁻¹, highest in the genotype IP130 (16.60 mgg⁻¹ fresh weight) and IP126 (16.6 mgg⁻¹) intimately tracked by genotype IP110 (16.2 mgg⁻¹), whilst the least count was evidenced with genotype IP188 (9.2 mgg⁻¹). Protein synthesis responds to drought stress. Late embryogenesis abundant (LEA) proteins play an important role in the protection of plants under drought. Comparable study has also conducted by Sahu et al. [37], Hadimani et

al. [47] and Gupta et al. [48] in different crop species.

3.7 Correlation Coefficient Analysis

Correlation coefficient are presented in Table 2. Chlorophyll at 30 days is highly, positively and significantly correlated with carotenoid 30 days ($r= 0.537$) and chlorophyll at 60 days ($r= 0.885$) at 1% level of significance and protein ($r= 0.213$) at 5 % probability level. Carotenoid at 30 days had positive and significant correlation with chlorophyll at 60 days ($r= 0.425$) and carotenoid at 60 days ($r= 0.354$).

3.8 Biochemical Activities Based Principal Component Analysis (PCA)

Principal component analysis (PCA) was drawn by considering biochemical variables instantaneously. The pattern of variations displayed by the PCA designated by correlation coefficients explained for pair-wise suggestion of the traits. The PCA correlation illustrated that genotype possessed higher and lower content occupying unique position towards the graph (Fig. 1). On the basis of highest and lowest content of the biochemical attributes genotypes *i.e.*, IP189, IP192, IP194, IP159, IP123, IP188, IP163, IP126, IP133, IP110, IP140, IP-160 and IP126 showed distinctive position on the plot.

3.9 Dendrogram Based on Different Biochemical Parameters

On the basis of dendrogram pearl millet germplasm lines grouped into two clusters one major and one minor. Minor cluster consisted of one genotype *i.e.*, IP173 and major cluster consist 95 germplasm lines that further divided into two groups one major and one minor. Minor cluster consisted of sixteen genotypes *viz.*, THAK1827, IP107, IP231, IP140, IP291, IP139, IP283, IP236, IP274, IP196, IP146, IP230, IP163, IP198, IP147 and IP114 and major cluster had 79 germplasm lines and again divided into two groups one major and one minor. Minor cluster had 21 germplasm lines *namely*, IP193, IP154, IP153, IP108, IP188, IP105, IP101, IP119, IP167, IP129, IP156, IP152, IP122, IP165, IP143, IP141, IP149, IP109, IP120, IP169 and IP183. While major cluster had 58 germplasm lines which further grouped into two clusters one major and one minor. Minor cluster contain 22 germplasm lines including IP115, IP199, IP131, IP123, IP138, IP116, IP159, IP186, IP130, IP189, IP110, IP126, IP162, IP132, IP134, IP142, IP145, IP262, IP177, IP168, IP133 and IP195 and major cluster consist 36 germplasm lines *viz.*, IP160, IP171, IP121, IP111, IP192, IP185, IP106, IP178, IP164, IP136, IP117, IP158, IP112, IP180, IP118, IP104, IP150, IP172, IP166, IP155, IP161, IP137, IP151,

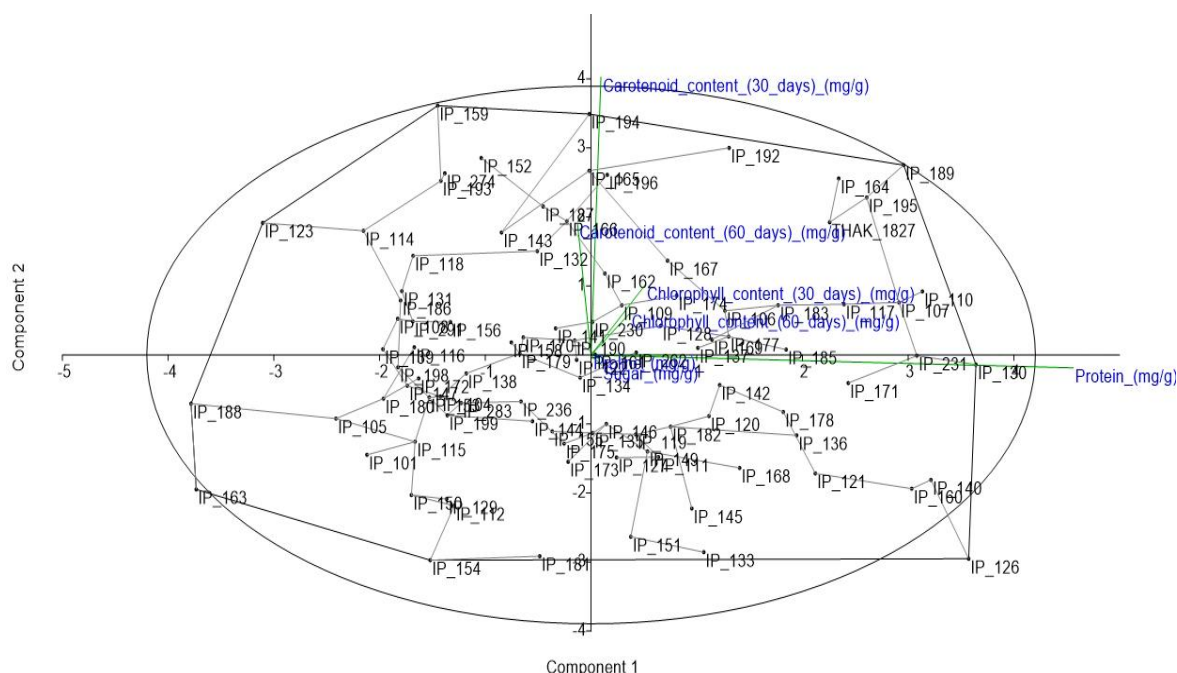


Fig 1. Diagram of pearl millet germplasm lines based on Chlorophyll and carotenoid attributes

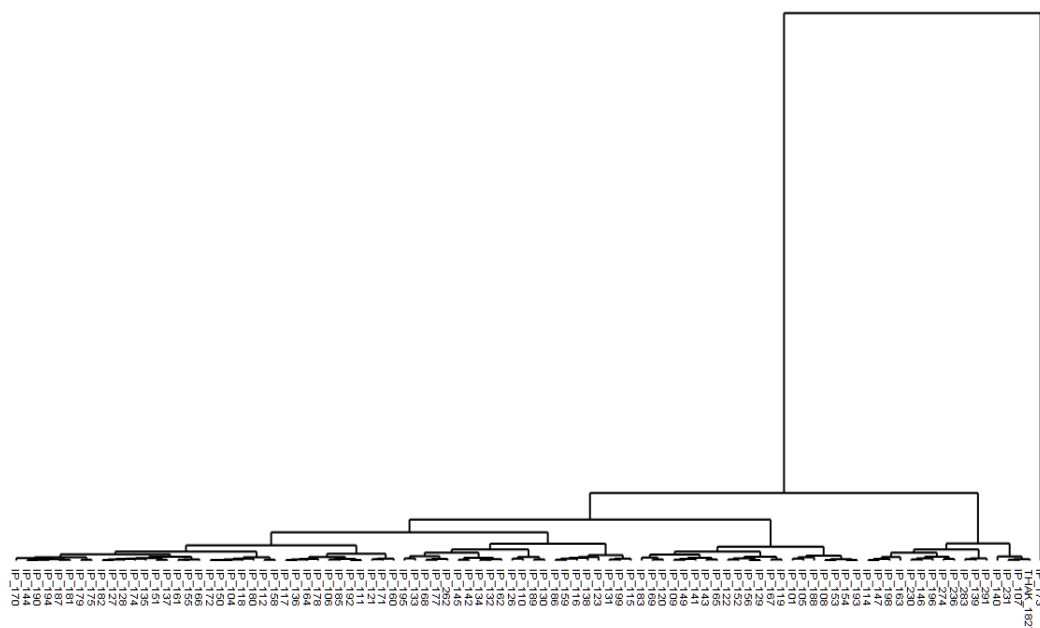


Fig 2. Dendrogram of pearl millet germplasms based on different biochemical parameters

IP135, IP174, IP128, IP127, IP182, IP175, IP179, IP181, IP187, IP194, IP190, IP144 and IP170 (Fig. 2). Cluster analysis based on dendrogram determine relative position of genotypes and decide selection of parents for crossing programme to achieve desired response [2, 49-55].

4. CONCLUSION

In conclusion, genotypes viz: IP133, IP177, IP164, IP142, IP120, IP160, IP136, IP166, IP192, IP195, IP106, IP126, IP121, IP110, IP117 and THAK 1827 made their position in distinct group due to drought by using different biochemical parameters. So, these germplasm lines might be used as a donor parent for future breeding programme for development of drought tolerant genotype(s) identifying QTLs by developing RILs through forward genetics approaches.

COMPETING INTERESTS

Authors have declared that no competing interests exist.

REFERENCES

1. Reddy SP, Satyavathi CT, Khandelwal V, Patil HT, Gupta PC, Sharma LD, Mungra KD, Singh SP, Narasimhulu, R, Bhadarge

HH, Iyanar K, Tripathi MK, Yadav D, Bhardwaj R, Talwar A M, Tiwari VK, Kachole UG, Sravanti K, Shanthi Priya M, Athoni BK, Anuradha N, Govindaraj M, Nepolean T, Tonapi VA. Performance and stability of pearl millet varieties for grain yield and micronutrients in arid and semi-arid regions of India. *Frontiers in Plant Sciences*, 2021;12:670201.

DOI:10.3389/fpls.2021.670201

2. Choudhary ML, Tripathi MK, Tiwari S, Pandya RK, Gupta N, Tripathi N, Parihar P. Screening of pearl millet [*Pennisetum glaucum* (L.) R. Br.] germplasm lines for drought tolerance based on morpho-physiological traits and SSR markers. *Current Journal of Applied Science and Technology*. 2021;40:46-63.

DOI:https://doi.org/10.9734/cjast/2021/v40i531303

3. Gupta SK, Nepolean T, Sankar SM, Rathore A, Das RR, Rai KN, Hash CT. Patterns of molecular diversity in current and previously developed hybrid parents of pearl millet [*Pennisetum glaucum* (L.) R. Br.]. *American Journal of Plant Sciences*. 2015;6:1697-1712.

4. Verma, R, Tripathi: MK, Tiwari S, Pandya, R K, Tripathi: N, and Parihar P. Screening of pearl millet [*Pennisetum glaucum* (L.) R. Br.] genotypes against blast disease on the basis of disease indexing and gene-specific SSR markers. *International*

- Journal of Current Microbiology and Applied Sciences. 2021;10 [02]:1108-1117.
DOI:https://doi.org/10.20546/ijcmas.2021.1001.134.
5. AICRPPM. Annual Report, All India Coordinated Pearl Millet Improvement Project. Mandor, Rajasthan, India:AICPMIP, Indian Council of Agricultural Research; 2019.
 6. Uppal RK, Wani SP, Garg KK, Alagarswamy G. Balanced nutrition increases yield of pearl millet under drought. Field Crop Research. 2015;177:86–97.
 7. Abdalla AA, Ahmed UM, Ahmed AR, Tinay AH, Ibrahim KA. Physicochemical characterization of traditionally extracted pearl millet starch [Jir]. Journal of Applied Sciences Research, 2009;5:2016-2027.
 8. Cheik A T, Aly S, Yaya B, Alfred T S. A comparative study on nutritional and technological quality of fourteen [14] cultivars of pearl millets [*Pennisetum glaucum* (L) Leeke] in Burkina. Faso. Pakistan Journal of Nutrition, 2006;5:512-521.
 9. Rekha. Efficacy of processing techniques in the utilization of pearl millet for value added products. A M.Sc. thesis, CCS Haryana Agricultural University, Hisar, Haryana, India. 1997;125.
 10. Poonam. Effect of acid and heat treatment on nutrient composition and shelf life of pearl millet [*Pennisetum glaucum*] flour. A M.Sc. Thesis, CCSHAU, Hisar, India; 2002.
 11. Alam M K, Ogata Y, Sako Y, Al-Mamun M, Sano H. Intermediary metabolism of plasma acetic acid, glucose and protein in sheep fed a rice straw-based diet. Asian-Australian Journal of Animal Sciences. 2010;23:1333-1339.
 12. Iwuala E, Odjegbab V, Sharmac V, Alame A. Drought stress modulates expression of aquaporin gene and photosynthetic efficiency in *Pennisetum glaucum* (L.) R. Br. genotypes. Current Plant Biology. 2020;21:131.
 13. Mishra N, Tripathi MK, Tiwari S, Tripathi N, Ahuja A, Sapre S, Tiwari S. Cell suspension culture and *in vitro* screening for drought tolerance in soybean using poly-ethylene glycol. Plants. 2021a;10:517-536.
 14. Mishra N, Tripathi MK, Tiwari S, Tripathi N, Gupta N and Sharma A. Morphological and physiological performance of Indian soybean [*Glycine max* (L.) Merrill] genotypes in respect to drought. Legume Research; 2021b.
DOI:10.18805/LR-4550.
 15. Sharma A, Tripathi MK, Tiwari S, Gupta N, Tripathi N, Mishra N. Evaluation of soybean [*Glycine max* (L.)] genotypes on the basis of biochemical contents and antioxidant enzyme activities. Legume Research; 2021.
DOI:10.18805/LR-4678.
 16. Toker C, Canci H, Yildirim T. Evaluation of perennial wild *Cicer species* for drought resistance. Genetic Resources and Crop Evolution. 2006;54:1781–1786.
 17. Serraj R, Hash CT, Rizvi SMH, Sharma A, Yadav RS, Biding FR. Recent advances in marker-assisted selection for drought tolerance in pearl millet. Plant Production Science. 2005;8:334–337.
 18. Mitra J. Genetics and genetic improvement of drought resistance in crop plants. Current Science. 2001;80:758-763.
 19. Bernier J, Kumar A, Venu P R. Characterization of the effect of a QTL for drought resistance in rice, qtl12.1, over a range of environments in the Philippines and eastern India. Euphytica. 2009;166:207–217.
 20. Duan XJ, Deng M, Zhao WZ, Wu K. The prolongation structure of the in homogeneous equation of the reaction–diffusion type. Journal of Physics A:Mathematical and Theoretical. 2007;40:3831–3837.
 21. Jafar SA, Goldsmith A. Transmitter optimization and optimality of beam forming for multiple antenna systems," in IEEE Transactions on Wireless Communications. 2004;3:1165-1175.
 22. Adejare FB, Umebese CE. Somatal resistance to low leaf water potential at different growth stages affects plant biomass in *Glycine max* L. American Journal of Agricultural and Biological Sciences. 2007;2(3):136-141.
 23. Goyal V, Jain S, Bishnoi NR, Munjal R. Leaf water relations, diffusive resistance and proline accumulation in hybrid pearl millet under depleting soil moisture content. Dianj Plant Physiol. 2001;6:41-45.
 24. Manivannan P, Jaleel CA, Kishore KA, Sankar B, Somasundaram R, Sridharan R and Panneerselvam R. Changes in antioxidant metabolism of *Vigna unguiculata* L. Walp. by propiconazole

- under water deficit stress. *Colloids Surf B: Biointerf.* 2007;57:69-74.
25. Terzi R, Kadioglu A. Drought stress tolerance and the antioxidant enzyme system in *Cenante setosa*. *Acta Biologica Cracoviensia Series Botanica.* 2006;48:89–96.
 26. Farooq M, Wahid A, Kobayashi N, Fujita D, Basra SMA. Plant drought stress: Effects, mechanisms and management. *Agronomy and Sustainability.* 2009;29:185-212.
 27. Izanloo A, Condon AG, Langridge P, Tester M, Schnurbusch T. Different mechanisms of adaptation to cyclic water stress in two South Australian bread wheat cultivars. *Journal of Experimental Botany.* 2008;59:3327–3346.
 28. Ashraf M, Rauf H. Inducing salt tolerance in maize (*Zea mays* L.) through seed priming with chloride salts: growth and ion transport at early growth stages. *Acta Physiologia Plantarum.* 2001;23:407–414.
 29. Arnon I. Crop production in dry regions. Leonard Hill, London; 1972.
 30. Bates LS, Waldren RP, Teare ID. Rapid determination of free proline for water-stress studies. *Plant and Soil.* 1973;39:205–207.
 31. DuBois M, Gilles KA, Hamilton JK, Rebers PK, Smith F. Colorimetric method for determination of sugars and related substances, *Analytical Chemistry.* 1956;28:350-356.
 32. Lowry OH, Rosebrough NJ, Farr AL, Randall RJ. Protein measurement with the folin phenol reagent. *The Journal of Biological Chemistry;* 1951.
 33. Snedecor GW, Cochran WG. *Statistical methods.* 1889;VIII Ed. Wiley-Blackwell.
 34. Hossain MM, Lam HM, Zhang J. Responses in gas exchange and water status between drought-tolerant and susceptible soybean genotypes with ABA application. *The Crop Journal.* 2015;3:500-506.
 35. Anjum SA, Xie X, Wang L, Saleem M F, Chen M and Wang L. Morphological, physiological and biochemical responses of plants to drought stress. *African Journal of Agricultural Research.* 2011;6:2026-2032.
 36. Zhang M, Duan L, Tian X, He Z, Li J, Wang B and Li Z. Uniconazole-induced tolerance of soybean to water deficit stress in relation to changes in photosynthesis, hormones and antioxidant system. *Journal of Plant Physiology,* 2007;164:709-717.
 37. Sahu VK, Tiwari S, Gupta N, Tripathi MK, Yasin M. Evaluation of physiological and biochemical contents in desi and kabuli chickpea. *Legume Research,* 2020;10. DOI. 18805/LR-426.
 38. Mishra N, Tripathi M K, Tripathi N, Tiwari S, Gupta N, Sharma A, Shrivastava M K. Changes in biochemical and antioxidant enzymes activities play significant role in drought tolerance in soybean. *International Journal of Agricultural Technology,* 2021c;17:1425-1446.
 39. Gurrieri L, Merico M, Trost P, Forlani G, Sparla F. Impact of drought on soluble sugars and free proline content in selected *Arabidopsis* mutants. *Biology.* 2020;9:367. DOI:10.3390/biology9110367.
 40. Watanabe S, Kojima K, Ide Y, Sasaki S. Effects of saline and osmotic stress on proline and sugar accumulation in *Populus euphratica in vitro.* *Plant Cell Tissue and Organ Culture.* 2000;63:199–206.
 41. Rengasamy P. Transient salinity and subsoil constraints to dryland farming in Australian sodic soils: an overview. *Australian Journal of Experimental Agriculture.* 2002;42:351-361.
 42. Guo P, Baum M, Grando S, Ceccarelli S, Bai G, Li R, Korff M, Varshney RK, Graner A and Valkoun J. Differentially expressed genes between drought-tolerant and drought-sensitive barley genotypes in response to drought stress during the reproductive stage. *Journal of Experimental Botany.* 2009;60:3531-3544.
 43. Singh A K, Rana M K, Singh S, Kumar S, Durgesh K, Arya L. Assessment of genetic diversity among pearl millet [*Pennisetum glaucum* (L.) R Br.] cultivars using SSR markers. *Range Management & Agroforestry.* 2013;34:77-81.
 44. Ghorbanli M, Gafarabad M, Amirkian T, Mamaghani BA. Investigation of proline, total protein, chlorophyll, ascorbate and dehydro-ascorbate changes under drought stress in Akria and Mobil tomato cultivars. *Iranian Journal of Plant Physiology.* 2012;3:651-658.
 45. George S, Minhas NM, Jatoi SA, Siddiqui SU, Ghafoor A. Impact of polyethylene glycol on proline and membrane stability index for water stress regime in tomato (*Solanum lycopersicum*). *Pakistan Journal of Botany.* 2015;47:835-844.

46. Mohammad K N Heidari R. Drought-induced accumulation of soluble sugars and proline in two maize varieties. *World Applied Sciences Journal*. 2008;3:448-453.
47. Hadimani N A, Muralikrisna G, Tharanathan R N, Malleshi N G, Nature of carbohydrates and proteins in three pearl millet varieties varying in processing characteristics and kernel texture. *Journal of Cereal Science*. 2001;33:17–25.
48. Gupta, N, Tiwari, S, Tripathi M K, Bhagyawant, S S. Antinutritional and protein-based profiling of diverse desi and wild chickpea accessions. *Current Journal of Applied Science and Technology*. 2021;40:7-18.
49. Adlak T, Tiwari S, Tripathi M K, Gupta N, Sahu VK. Biotechnology: An advanced tool for crop improvement. *Current Journal of Applied Science and Technology*. 2019; 33(1):1-11.
50. Pramanik A, Tiwari S, Tomar RS, Tripathi MK, Singh AK. Molecular characterization of groundnut [*Arachis hypogaea* L] germplasm lines and varietal set for yield and yield attributing traits. *Indian Journal of Genetics*. 2019;79:56-65. DOI:<https://doi.org/10.31742/IJGPB.79.1.8>
51. Mishra N, Tripathi MK, Tiwari S, Tripathi N and Trivedi H K. Morphological and molecular screening of soybean genotypes against yellow mosaic virus disease. *Legume Research*; 2020. DOI:10.18805/LR4240.
52. Shyam C, Tripathi MK, Tiwari S, Tripathi N, Ahuja A. Molecular characterization and identification of *Brassica* genotype(s) for low and high erucic acid content using SSR markers. *Global Journal of Bioscience and Biotechnology*. 2020;9:56 – 66.
53. Upadhyay S, Singh AK, Tripathi MK, Tiwari S, Tripathi N. Validation of simple sequence repeats markers for charcoal rot and *Rhizoctonia* root rot resistance in soybean genotypes. *IJABR*. 2020b;10:137-144.
54. Pramanik A, Tiwari S, Tripathi MK, Mandloi S, Tomar RS. Identification of groundnut germplasm lines for foliar disease resistance and high oleic traits using SNP and gene-based markers and their morphological characterization. *Legume Research*; 2021. DOI:10.18805/LR-4666.
55. Upadhyay S, Singh AK, Tripathi MK, Tiwari S, Tripathi N, Patel RP. *In vitro* selection for resistance against charcoal rot disease of soybean [*Glycine max* (L.) Merrill] caused by *Macrophomina phaseolina* (Tassi) Goid. *Legume Research*; 2020a. DOI:10.18805/LR-4440

© 2021 Choudhary et al.; This is an Open Access article distributed under the terms of the Creative Commons Attribution License (<http://creativecommons.org/licenses/by/4.0>), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Peer-review history:
The peer review history for this paper can be accessed here:
<https://www.sdiarticle4.com/review-history/73335>