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Research Article

Genetic Dissection of Variability Using Morphological Traits in Drumstick (Moringa oleifera Lam.) Genotypes

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ABSTRACT

Twenty one moringa genotypes that were collected from different geographical regions of India were evaluated for variability, heritability and genetic advance for growth, yield and earliness characters and significant variations were recorded for the investigated traits. Widest range of variation was observed in pod length, number of pods per plant, single pod weight and pod yield per plant. Maximum genotypic and phenotypic coefficient of variation was observed for number of primary branches, pod length, number of pods per cluster, single pod weight and pod yield per plant. Higher value of heritability associated with high genetic advance as percent mean was observed for pod yield per plant, single pod weight, pod length, plant height, leaf length, pod girth, number of pod per cluster and number of branches per inflorescence. High heritability and genetic advance indicating additive gene action suggested that simple selection could be effective for the improvement of these traits.

Keywords: Moringa, Variability, Heritability, Genetic advance.

INTRODUCTION

Drumstick [*Moringa oleifera* Lam. (Syn. *M. pterygosperma Gaertn.*)] is one of the best known and most widely distributed and naturalized species of a monogeneric family Moringaceae with true diploid chromosome 2n=28. The genus *Moringa* has about 13 species of which two species viz., *M. oleifera* Lam. (syn. *M. pterygosperma* Gaertn.) and *M. concanensis* Nimmo occur in India and the

former being the vegetable type (Panday et al., 2011). Drumstick is an important food commodity which has had enormous attention as the 'natural nutrition of the tropics'. It is an exceptionally nutritious vegetable tree with multiple uses and beneficial properties and has therefore been called a "miracle tree" (Fuglie, 1999) or "one of the world's most useful trees" (Lea, 2010).

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The miracle tree is a perennial softwood crop native to the sub-Himalayan ranges of India, Pakistan, Bangladesh and Afghanistan (Fahey, 2005). It is widely grown in the tropics of Asia, Latin America, the Caribbean, Florida and the Pacific Islands to West, East, and South Africa. India is the prime producer of Moringa (Drumstick) with an annual production of 2.20 to 2.40 million tonnes of tender fruits from an area of 38,000ha leading to the productivity of around 63 tonnes per ha.5 Among the different states, Andhra Pradesh leads in both area and production (15,665ha) followed by Karnataka (10,280ha) and Tamil Nadu (13250ha). In other states, it occupies an area of 4,613ha.

Moringa oleifera is a fastgrowing, drought-resistant tree with many medicinal properties. Common names include moringa, drumstick tree (from the long, slender. triangular seedpods), horseradish tree (from the taste of the roots, which resembles horseradish), and ben oil tree or benzolive tree. Flower favors cross pollination due to delayed stigma receptivity. The successful pollination requires large number of insect population, among which *Xylocopa* is most important. The zygomorphic gullet type flowers show a forenoon (6.00 h -12.00 h) pattern of anthesis after which pollen anthesis takes place (7.00 h - 13.00 h)(Bhattacharya and Mandal, 2004). It is widely cultivated for its young seed pods and leaves, as vegetables, for traditional herbal used medicine and also used for water purification.

Moringa has many medicinal properties. Almost all parts viz., root, bark, gum, leaf, fruit (pod), flower, seed and seed oil have been used for treatment of various inflammation and infectious diseases along with cardiovascular, gastrointestinal, haematological and hepatorenal, disorders (Singh et al., 2011). Flowers are used as stimulant, tonic and diuretic. They are useful in increasing the flow of bile. The seeds of Moringa are considered to be antipyretic, acrid, bitter (Oliveira et al., 1999) and reported to show antimicrobial activity. The seeds of Moringa are used as water purifier. The oil extracted from the seed

known as "Ben" is used as lubricant in watches, for edible purpose and in cosmetics. The kernel of the seed is rich in crude protein, fatty oil and fiber. Leaves contain 4.0per cent moisture, 38.4per cent crude protein, 34.17per cent fatty oil, 3.5 per cent fibre and 3.2 per cent mineral matter.

Germplasm evaluation and characterization for economically important traits are basic prerequisite for crop improvement. For any crop improvement programme, evaluation of germplasm to assess the existing variability is the first step. Greater the variability present in the initial material better would be the chances for evolving desired types. A clear understanding of variability of various characters of the breeding materials is an asset to the plant breeder for selecting superior genotypes on the basis of their phenotypic expression. The diversity in moringa genotypes presents an opportunity for selection of superior types and improvement of different quantitative and qualitative characters. The determination of genetic variability and its partitioning into various components is necessary to have an insight into the genetic nature of yield and its components. In this regards, estimates of genotypic and phenotypic variance for various quantitative characters along with heritability and genetic advance expected by selection for yield and its components are useful in designing an effective breeding programme.

MATERIALS AND METHODS

The study has taken up in the education and research block of Department of biotechnology and crop improvement, College of horticulture, University of horticultural sciences, Bagalkot. Twenty-one genotypes of Moringa oleifera used in the present study were collected from different eco-geographical locations covering different states of India were studied in Randomized block design in three replications (Table 1). The observation were recorded for twenty one different quantitative morphological parameters viz., plant height, tree girth, number of primary branches, leaf length, petiole length, days to

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flower initiation, inflorescence length, number of branches per inflorescence, flower size, number of pods per plant, pod length, pod girth, number of pods per cluster, number of pods per branch, days to pod initiation, days to pod maturity, single pod weight, number of seeds per pod, seed size, hundred seed weight and pod yield per plant. The data collected on all the characters were subjected to standard methods of analysis of variance (ANOVA) (Panse & Sukhatme, 1985). Genotypic and phenotypic co-efficient of variation were estimated according to Burton (1952) GCV and PCV were classified by Sivasubramanian and Madhamenon (1973). Heritability and genetic advance as per cent of mean was calculated based on Johnson et al. (1955).

Sr. No.	Name of the accession	Code	Area of collection	State of collection	
1	РКМ-01	MO_1	Periyakulam	Tamil Nadu	
2	РКМ-02	MO_2	Periyakulam	Tamil Nadu	
3	Dhanraj	MO_3	Dharwad (UASD campus)	Karnataka	
4	Bhagya (KDM-01)	MO_4	Bagalkot(UHSB campus)	Karnataka	
5	Karwar-01	MO_5	Kumburda village, Karwar	Karnataka	
6	Badami-02	MO_6	Badami	Karnataka	
7	Mysore-01	MO_7	Mysore	Karnataka	
8	Mysore-02	MO_8	Mysore	Karnataka	
9	Mysore-03	MO_9	Mysore	Karnataka	
10	Yelwala-01	MO_10	Shettinayakanahalli, Mysore	Karnataka	
11	Shirdi-01	MO_11	Shirdi	Maharashtra	
12	Tangi-01	MO_12	Tangi village	Orissa	
13	Bhubaneshwar-01	MO_13	Bhubaneshwar	Orissa	
14	Cuttack-01	MO_14	Cuttack	Orissa	
15	Bagalkot-01	MO_15	Bagalkot	Karnataka	
16	Thar-harsha	MO_16	Vejalpur, Godhra	Gujarat	
17	Mandya-01	MO_17	Mandya	Karnataka	
18	BG-1	MO_18	Bagalkot	Karnataka	
19	BG-2	MO_19	Bagalkot	Karnataka	
20	BG-3	MO_20	Bagalkot	Karnataka	
21	BG-4	MO_21	Bagalkot	Karnataka	

Table 1: Details of the	drumstick gen	otypes used in	the experiment
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RESULT AND DISCUSSION

The analysis of variance revealed the existence of significant differences among the genotypes for all the traits, indicating the presence of considerable genetic variability among the experimental material under study (Table 2). Thus, there is a plenty of area and scope for the improvement of different quantitative and qualitative traits through selection. The tendency of individual genetic characteristics in a population to vary from one another. Similarly highly significant variation for all characters studied was reported by Raja and Bagle (2003), Selvakumari et al. (2013), Varma et al. (2019), Priya et al. (2019) and Balaguru et al. (2020). The mean performance, range and general mean of twenty one genotypes of drumstick for twenty one quantitative characters studied are presented in Table 3. Range of variation observed for all Nair et al.Ind. J. Pure App. Biosci. (2021) 9(1), 442-449ISSN: 2582 - 2845the traits in the present study indicated the
presence of sufficient amount of variationamong the genotypes for all the characters

Table 2: One- factor analysis of variance across 21 quantitative characters in drumstick genotypes

Sl. No.	Traits	MSS (genotypes)	Error	F-Value (Genotypes)	
	d. f	20	40		
1	Plant height (cm)	9849.001	300.0813	32.82111*	
2	Tree girth (cm)	52.52569	4.18565	12.54899*	
3	Number of primary branches	1.337714	0.529429	2.526713*	
4	Leaf length (cm)	92.09041	3.177532	28.981743*	
5	Petiole length (cm)	4.734206	0.316497	14.95814*	
6	Days to flower initiation	208.9549	6.403143	32.63317*	
7	Inflorescence length (cm)	10.18029	2.799306	3.63672*	
8	Number of branches per inflorescence	6.603302	0.276063	23.9195*	
9	Flower size (mm)	6.11591	0.996367	6.138211*	
10	Number of pods per plant	717.4665	20.51368	34.97502*	
11	Pod length (cm)	612.8009	10.92919	56.07009*	
12	Pod girth (mm)	14.73018	0.440962	33.40467*	
13	Number of pods per cluster	0.577206	0.020063	28.76899*	
14	Number of pods per branch	13.6313	0.971111	14.03681*	
15	Days to pod initiation	17.89349	1.449302	12.34629*	
16	Days to pod maturity	59.09283	8.577206	6.889519*	
17	Single pod weight (g)	4283.26	32.17838	133.1099*	
18	Number of seeds per pod	27.20971	2.041619	13.32752*	
19	Seed size (mm)	1.745071	0.375123	4.651999*	
20	Hundred seed weight (g)	4.41905E-05	1.23582E-05	3.575819*	
21	Pod yield per plant (Kg)	31.59736	0.770268	41.02127*	

*5 per cent significance

Comparative variability of traits is evaluated by estimating the genotypic coefficient of variation (GVC) and the phenotypic coefficient of variation (PCV). Greater the variability in a population, there are the greater chances for effective selection for desirable types. The estimates of variances due to these three components for twenty one quantitative characters are represented in table 4. In the present study, the phenotypic variance ranged between 0.206 (number of pods per cluster) to 3483.055 (plant height). The genotypic variance ranged between 0.186 (number of pods per cluster) to 3182.973 (plant height). The environmental variance ranged between 0.020 (Number of pods per cluster) to 300.081 (plant height). In general, phenotypic variance was higher in magnitude than genotypic variance for all characters and the genotypic variances were higher in magnitude over respective environmental variances for all the characters. The magnitude of phenotypic coefficient of variation was higher than corresponding genotypic coefficient of variation for all the 21 quantitative parameters, which indicated prime role of environment on the character expression. With respect to the genetic variability components such as PCV and GCV, the value of 0-10 per cent indicate low genetic variation which is represented by the traits flower size (9.176) and days to pod maturity (8.791) (Table 4). 10-20 per cent indicates moderate variation which is present in most of the traits under study. The value of >20 per cent indicate higher genetic variation which is present in the traits like number of primary branches (20.269), pod length (27.643), number of pods per cluster (28.636), single pod weight (34.858) and pod yield per plant (37.144). This showed similarity with the studies of Raja and Bagle (2003), Selvakumari et al. (2013), Varma et al. (2019), Priya et al. (2019) and Balaguru et al. (2020).

Heritability is the portion of phenotypic variation which is transmitted from parent to the progeny. High heritability was recorded for all the characters except number of primary branches, inflorescence length, seed size and hundred seed weight (Table 3). The higher broad sense heritability was recorded for pod length (94.8 per cent) followed by pod yield per plant (93.0 per

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cent), number of pods per plant (91.9 per cent), pod girth (91.5 per cent), plant height (91.4 per cent), days to flower initiation (91.3 per cent) etc., Higher heritability in broad sense showed that large proportion of phenotypic variance was attributable to the genotypic variance and these traits were less affected by the environment. The least heritability was reported for number of primary branches (33.7 per cent) followed by hundred seed weight (46.2 per cent) and inflorescence length (46.8 per cent).

Heritability estimates along with genetic advance is more useful than the heritability value alone for selecting the best individual. Higher value of heritability associated with high genetic advance as percent mean was observed for pod yield per plant, single pod weight, pod length, plant height, leaf length, pod girth, number of pod per cluster and number of branches per inflorescence. This is attributed to the additive gene action. Heritability values were high than those of genetic advance for most of the traits which indicated that they were least influenced by the environment and shows that the phenotypes were true representative of their genotypes and selection based on phenotypic performance would be reliable genotypes which is in line with the findings of Karunakar et al. (2018), Lakshmi Narayana Priya et al. (2019) and Varma et al. (2019).

Genotypes	PH (cm)	TG (cm)	NPB	LL	PeL	DFI	IL	NBI	FS	NPP	PL
			(No. s)	(cm)	(cm)		(cm)	(No. s)	(mm)	(No. s)	(cm)
MO_1	300.20	33.95	4.47	45.13	11.97	74.13	20.85	7.73	14.83	101.73	66.62
MO_2	281.00	33.02	4.00	35.20	12.45	64.87	20.31	7.13	16.04	99.47	94.28
MO_3	399.27	38.20	5.13	47.33	11.81	85.87	21.19	11.20	16.70	105.33	29.89
MO_4	421.00	33.63	5.07	40.81	11.16	67.33	19.91	7.33	17.52	96.60	53.89
MO_5	303.67	38.64	3.87	41.50	10.09	87.73	21.98	6.73	15.62	90.60	44.28
MO_6	307.00	36.03	3.40	28.62	9.87	88.13	20.15	4.93	16.95	91.73	45.31
MO_7	352.40	44.93	5.67	45.21	10.03	74.47	18.55	7.07	17.47	90.87	41.19
MO_8	323.33	39.63	5.00	39.41	10.33	82.13	20.27	6.53	18.49	83.73	47.29
MO_9	394.07	38.03	3.87	48.27	12.59	88.93	17.88	8.27	18.11	88.40	68.17
MO_10	400.40	34.64	4.40	39.43	10.79	71.13	22.05	9.80	19.52	77.73	45.26
MO_11	378.93	32.44	3.40	38.70	8.99	78.73	20.80	10.40	17.22	74.53	46.52
MO_12	294.93	35.21	3.93	49.31	12.73	83.87	23.18	8.20	19.55	62.67	44.15
MO_13	382.67	45.80	4.93	35.33	9.63	84.00	20.36	7.07	16.24	41.07	40.38
MO_14	391.33	39.00	5.60	40.33	10.51	83.60	23.01	8.20	18.82	72.93	57.56
MO_15	205.93	33.63	3.40	47.51	12.11	74.67	20.70	9.33	19.06	59.00	71.72
MO_16	409.33	45.53	4.20	42.00	11.74	96.73	17.19	6.13	18.94	79.60	47.57
MO_17	320.00	36.20	4.47	32.04	9.52	70.13	19.32	7.13	19.12	90.73	38.70
MO_18	316.00	37.54	4.53	41.47	9.46	89.33	17.57	7.00	19.96	89.27	54.31
MO_19	413.80	42.03	4.33	46.28	12.85	76.73	19.01	9.20	18.37	89.73	47.86
MO_20	289.00	35.20	4.13	43.03	12.71	81.67	16.92	8.00	18.78	93.93	65.27
MO_21	385.93	42.47	4.80	35.41	10.95	72.00	22.51	7.27	18.95	91.00	54.75
Mean	346.20	37.89	4.41	41.06	11.06	79.82	20.18	7.84	17.92	84.32	52.62
SE(m)	10.00	1.18	0.42	1.03	0.32	1.46	0.97	0.30	0.58	2.61	1.91
C.V %	15.003	15.399	16.5011	14.3409	15.0862	13.1702	8.2924	16.7007	15.5709	15.3716	16.2829
C.D @5%	28.58	3.38	1.20	2.94	0.93	4.18	2.76	0.87	1.65	7.47	5.46
Min.	205.93	32.44	3.40	28.62	8.99	64.87	16.92	4.93	14.83	41.07	29.89
Max.	421.00	45.80	5.67	49.31	12.85	96.73	23.18	11.20	19.96	105.33	94.28

Table 3: Per se performance of 21 genotypes for quantitative traits

PH- Plant height, TG- Tree girth, NPB- Number of primary branches, LL- leaf length, PeL- Petiole length, DFI- Days to flower initiation, IL- Inflorescence length, NBI-Number of branches per inflorescence,

FS- flower size, NPP-Number of pods per plant, PL- Pod length

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Genotypes	PG (mm)	PPC (No. s)	PPB (No. s)	DPI	DPM	SPY (g)	NSP (No.s)	SS (mm)	HSW (g)	PYP (Kg)		
MO_1	14.12	1.53	13.40	19.87	65.20	166.27	20.07	10.21	37.80	16.20		
MO_2	15.37	1.60	8.87	17.47	51.33	179.73	24.27	9.11	27.53	14.56		
MO_3	8.52	1.80	16.07	21.60	57.40	64.87	13.67	8.89	31.27	7.19		
MO_4	13.16	1.27	12.27	15.27	47.80	107.00	22.67	8.49	23.73	10.33		
MO_5	14.11	1.27	11.27	18.33	56.40	91.27	16.20	7.97	25.67	8.07		
MO_6	8.94	2.13	14.13	18.07	58.53	94.60	20.67	10.00	36.67	9.25		
MO_7	13.55	1.27	12.60	23.43	60.27	88.07	19.80	9.18	27.80	7.97		
MO_8	8.21	2.53	11.27	20.87	58.00	102.47	18.87	8.18	25.20	7.78		
MO_9	14.19	1.27	10.93	14.47	62.87	133.13	25.53	8.25	29.00	11.01		
MO_10	11.21	1.33	8.87	18.27	57.87	74.73	19.00	8.81	26.80	5.68		
MO_11	12.69	1.87	15.80	19.87	54.13	79.40	18.33	8.49	31.53	5.26		
MO_12	13.92	2.00	14.07	24.00	52.33	73.27	18.93	9.09	33.53	4.51		
MO_13	15.12	1.13	8.20	22.67	57.13	94.47	18.27	8.49	29.60	4.74		
MO_14	13.24	2.20	14.20	18.33	60.80	86.60	19.80	8.23	27.33	6.58		
MO_15	12.95	2.07	10.13	19.20	49.60	183.67	25.27	9.71	31.93	10.44		
MO_16	12.34	2.07	10.73	20.80	61.00	95.13	24.00	10.20	35.20	8.39		
MO_17	15.63	1.13	12.80	18.87	56.73	67.73	18.47	7.82	27.87	6.62		
MO_18	9.21	1.20	12.87	21.27	54.80	123.87	21.87	9.03	29.53	10.13		
MO_19	11.95	1.20	11.87	18.47	60.47	81.07	22.47	10.07	34.40	7.39		
MO_20	13.65	1.07	13.13	16.87	59.87	166.20	23.00	8.00	28.07	14.30		
MO_21	12.79	1.33	12.80	19.87	61.73	139.87	18.87	8.60	32.00	11.46		
Mean	12.61	1.58	12.20	19.42	57.35	109.21	20.48	8.90	30.12	8.95		
SE(m)	0.38	0.08	0.57	0.69	1.69	3.27	0.82	0.35	2.02	0.51		
C.V %	15.2645	18.9416	18.0753	16.1989	15.1070	15.1942	16.9781	16.8841	11.6724	9.8101		
C.D @5%	1.09	0.23	1.63	1.99	4.83	9.36	2.36	1.01	0.01	1.45		
Min.	8.21	1.07	8.20	14.47	47.80	64.87	13.67	7.82	23.73	4.51		
Max.	15.63	2.53	16.07	24.00	65.20	183.67	25.53	10.21	37.80	16.20		

PG- Pod girth, PPC- pods per cluster, PPB- Pods per branch, DPI- Days to pod initiation, DPM- days to pod maturity, SPY-Single Pod weight, NSP- Number of seeds per pod, SS- Seed size, HSW- Hundred seed weight, PYP- Pod yield per plant

Table 4: Genetic variability	and heritability j	parameters for 21	quantitative par	rameters for drumstick
		L		

genotypes PV Sl. No. Traits GV EV PCV GCV ECV h² (Broad sense) GA (%) GAM 3483.055 3182.973 300.081 17.047 16.296 5.004 0.914 111.102 32.092 PH (cm) 1 10.593 5.399 19.443 2 TG (cm) 20.299 16.113 4.186 11.890 0.794 7.367 3 NPB 0.799 0.529 0.269 20.269 16.501 11.771 0.337 0.621 14.083 13.257 4 LL (cm) 32.816 29.637 3.178 13.950 4.341 0.903 10.658 25.954 5 1.789 1.473 0.316 12.093 10.971 5.086 0.823 2.268 20.504 PeL (cm) 73.920 67.517 20.267 6 DFI 6.403 10.771 10.294 3.170 0.913 16.177 5.260 2.385 2.210 2.813 7 IL (cm) 2.799 2.461 11.367 8.292 10.954 7.774 0.468 2.109 18.521 0.276 8 NBI 19.696 6.701 0.88435.877 9 FS (mm) 2.703 1.706 0.997 9.176 7.290 5.573 0.631 2.138 11.930 10 NPP 252.831 232.318 20.514 18.858 18.077 5.372 0.919 30.098 35.696 11 PL (cm) 211.554 200.625 10.929 27.643 26.919 6.283 0.948 28.415 54.002 12 PG (mm) 5.205 4.764 0.441 18.086 17.303 5.264 0.915 4.302 34.101 13 NPC 0.2060.186 0.020 28.636 27.204 8.942 0.902 0.843 53.238 14 NPB 5.191 4.220 0.971 18.671 16.834 8.075 0.813 3.816 31.267 22.085 5.481 15 DPI 6.931 1.449 13.556 12.055 6.199 0.791 4.289 16 DPM 25.416 16.839 8.577 8.791 7.156 5.107 0.663 6.881 11.998 17 SPY (g) 1449.206 1417.027 32.178 34.858 34.469 5.194 70.213 0.978 76.680 NSP 10.431 8.389 2.042 15.773 14.145 6.978 0.804 5.351 26.133 18 19 SS (mm) 0.832 0.457 0.375 10.251 7.595 0.549 1.031 11.590 6.885 20 HSW (g) 22.969 12.358 10.611 15.913 11.672 10.816 0.462 4.561 15.143 11.044 21 PYP(Kg) 10.274 0.770 37.144 35.825 9.808 0.930 71.181 6.369

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Heritability estimates along with genetic advance is more useful than the heritability value alone for selecting the best individual. Higher value of heritability associated with high genetic advance as percent mean was observed for pod vield per plant, single pod weight, pod length, plant height, leaf length, pod girth, number of pod per cluster and number of branches per inflorescence. This is attributed to the additive gene action. Heritability values were high than those of genetic advance for most of the traits which indicated that they were least influenced by the environment and shows that the phenotypes were true representative of their genotypes and selection based on phenotypic performance would be reliable genotypes which is in line with the findings of Karunakar et al. (2018), Lakshmi Narayana Priya et al. (2019) and Varma et al. (2019).

Selection would be depending on pod yield per plant, the number of pods per plant, single pod weight and pod length may bring out the genetic improvement in the moringa because they showed a high value of GCV, PCV, heritability and genetic gain. Hence the selection will be effective for these traits. These traits can be improved through mass selection, progeny selection, or any other modified selection procedures. It can be concluded that PKM-01, PKM-02, Dhanraj, Shirdi-01 and Bagalkot-01genotypes, performances were outstanding for the traits like pod length, number of pods per plant, single pod weight and pod yield per plant, indicating their utilization in the breeding for development of elite varieties or hybrids in the future and genotypes like Bhagya, Tangi-01 and PKM-02 for better performance in relation to earliness parameters.

REFERENCES

- Balaguru, P., Sathiyamurthy, V. A., Janavi, G. J., & Santha, S. (2020). Variability in perennial moringa (Moringa oleifera Lam.) genotypes for quantitative and qualitative traits Electron. J. Plant Breed, 11(2), 515-520.
- Bhattacharya, A., & Mandal, S. (2004). Pollination, pollen germination and

stigma receptivity in Moringa oleifera Lam. Grana, 43, 48–56.

- Burton, G. W., & Devane, E. H. (1952). Estimating heritability in tall fescue (Festuca *arundinaceae*) from replicated clonal material. Agron. J., 45, 478-481.
- Fahey, J. W. (2005). Moringa oleifera A review of the medical evidence for its nutritional. therapeutic, and prophylactic properties. Part1. Trees *Life J.*, *1*, 1–15.
- Fuglie, L. J. (1999). The Miracle Tree Moringa oleifera natural nutrition for the tropics. Church World Service, Dakar. 68 p.; revised in 2001 and published as The Miracle Tree: The Multiple Attributes of Moringa, 172 p.
- Johnson, H. W., Robinson, H. F., & Comstock, R. E. (1955). Estimates of genetic and environmental variability in soybean. Agron. J., 47, 314-318.
- Karunakar, J., Preethi, T. L., Manikanta, B. N., Pugalendhi, L., & Juliet, H. S. (2018). Genetic variability, correlation and path analysis in Moringa (Moringa oleifera L.). J. Pharmacogn. Phytochem, 7(5), 3379-3382.
- N. P. C., Paramaguru, Lakshmi, P., Satyamoorthy, V. A., & Bhoopathi, M. N. (2019). Genetic variability of quantitative characters that exist in moringa (Moringa oleifera Lam.) in Tamil Nadu, India. Int. J. Chem, 7(3), 3211-3213.
- Lea, M. (2010). Bioremediation of turbid surface water using seed extract from Moringa oleifera.
- Oliveira, J. T. A., Silveira, S. B., Vasconcelos, I. M., Cavada, B. S., & Moreira, R. A. (1999). Compositional and nutritional attributes of seeds from the multipurpose tree Moringa oleifera Lam. J. Sci. Food Agric., 79, 815-820.
- Pandey, A., Prudheep, K., Gupta, R., Navar, E. R., & Bhandari D. C. (2011). Drumstick tree (Moringa oleifera Lam.): A multipurpose potential

Ind. J. Pure App. Biosci. (2021) 9(1), 442-449

Nair et al. Ind. J. Pure App. Bio species in India. Genet. Resour. Crop Evol, 58, 453-460.

- Panse, V. G., & Sukhatme, P. V. (1961). Statistical methods for agricultural workers 2nd Edition, Indian Council of Agricultural Research, New Delhi, pp. 359.
- Raja, S., & Bagle, B. G. (2005). Variability studies in annual moringa (Moringa pterigosperma Gaertn). Veg. Sci., 32(1), 78-79.
- Selvakumari, P., & Ponnuswami, V. (2017). Correlation and genetic variation of thirty four different genotypes of Moringa (*Moringa oleifera*, Lam.) in Tamil Nadu Condition, India. Int J Curr Microbiol Appl Sci 6, 332-335.

- Sivasubramanian, S., & Madhavamenon, P. (1973). Genotypic and phenotypic variability in rice. *Madras Agric. J* 60(9-13), 1093-1096.
- Singh, B. K., Sharma S. R., Kalia, P., & Singh, B. (2011). Genetic variability for antioxidants and horticultural traits in cabbage. *Indian J. Hortic.*, 68(1), 51-55.
- Varma, L. K., Asati, B. S., Shankar, D., & Chandrashekhar, M. K. (2019).
 Variability and association studies for yield components in drumstick (*Moringa oleifera* L.). J. Pharma cogn. Phytochem, 8(4), 2356-2359.