

# PHENOLOGY, GROWTH, FIBRE YIELD AND QUALITY STUDIES IN KENAF (HIBISCUS CANNABINUS L.)

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#### Abstract

Six kenaf genotypes were evaluated in randomized block design with four replications during kharif- 2022-23 under irrigated conditions at Cotton Improvement Project, MPKV, Rahuri (Ahmednagar) to study the phenology, growth, fibre yield and quality studies in kenaf (Hibiscus cannabinus L.). The principal growth stages for fibre yield are observed through BBCH scale (Biologische Bundesanstalt, Bundessortenamt and CHemische Industrie). It is observed that genotypes JRK-2019-2 and JRK-2019-1 required higher number of days to 50% flowering flowering. Genotype JRK-2019-3 shows superior performance in growth parameters. Genotype JRHC-15 shows superior performance in absolute growth rate, relative growth rate and leaf area duration. Genotype JRK-2019-2 shows good performance in leaf area index and unit area efficiency. Highest canopy temperature at top, middle and bottom and chlorophyll content is observed in JRK-2019-3. The genotype AMC 108 has better light extinction coefficient while, highest light intensity at top, middle and bottom and light transmission ratio is observed in JRHC-15. Genotype JRK-2019-1 was reported to be the highest green weight (q/ha) at harvesting. JRK-2019-2 showed better performace in yield and yield contributing characters. Genotypes JRK-2019-2 and JRK-2019-1 was reported to be the highest root content (wt.%). Genotypes JRK-2019-2 and JRK-2019-3 was reported to be the lesser defects (wt.%). Good tenacity is observed in JRK-2019-3 Genotypes followed by JRK-2019-2. Highest fineness is observed in JRK-2019-1. Genotypes JRK-2019-2, JRK-2019-1, JRHC-15 shows heavy bodied bulk density and HC 583, JRK-2019-3, AMC 108 shows medium bodied bulk density. The fibre of all genotypes appeared creamy white in colour. The yield contributing characters were significantly associated with fibre yield per plant.

Keywords: Phenology, growth parameters, canopy temperature, light intensity, fibre quality

### **1. INTRODUCTION**

Kenaf (*Hibiscus cannabinus* L.) is an herbaceous annual fibre crop belongs to *Malvaceae* family. It grows to 1.5-4.5 m tall with woody base. The prickly unbranched stems are mostly 1.5-2.5 cm in diameter. The 10-15 cm long leaves are alternate from side to side on stalk and branches. The younger leaves on all kenaf plant are very simple, entire and cordate. The leaves near the base of stems are deeply lobed with 3-7 lobes, while leaves near the top of stem are shallowly lobed. Kenaf plant produces large showy, white, light yellow, creamy colour flowers with maroon or purple colour center. The flowers are bell shaped, widely open, solitary, auxiliary having 8-15 cm diameter with 5 petals, 5 sepals and numerous stamens [1]. Kenaf plant contains three type of fibres i.e. bast, core and pith [2]. Kenaf requires a warm, moist and tropical-subtropical climate and thrives with abundant solar radiation and high rainfall. Kenaf grows at latitude  $30^{0}$ N and  $30^{0}$ S and at altitude upto 1.25 m with mean relative humidity are 68%-85% and the optimum range of temperature ranging from  $20^{0}$ C- $30^{0}$ C. Kenaf is a versatile plant that offers a wide range of uses across various industries, making it a valuable resource for fibre, food, medicine, oil, chemical absorbent, mushroom cultivation, natural

fibre, textile application, construction, housing industries, environmental cleaning, and bioenergy production. In the fibre industry, kenaf's strong and durable fibres are utilized for making products such as ropes, twines, canvas, and burlap sacks. Its fibres can also be blended with other materials to create composite materials for construction and automotive parts. Phenology in plants refers to the study of cyclic and seasonal patterns of plant life events and their relationships with environmental factors, such as temperature, light, and precipitation [3]. In essence, the BBCH scale serves as a valuable tool for monitoring and understanding the phenological development of crops, enhancing agricultural practices, and contributing to the advancement of crop science and management.

# 2. MATERIAL AND METHODS

#### 2.1 Location

The experiment was conducted at Cotton Improvement project, MPKV, Rahuri during *kharif* 2022. The centre is located in the Agro-Ecological region (AER) 6.0 [Deccan Plateau, Hot Semi-Arid Eco-Region] [Agro-Ecological Sub Region 6.1]. It is situated 33 km away from Ahmednagar on Ahilyanagar- Manmad state Highway No.14. It lies between 19°-48° N and 19°-57° N latitude and between 74°-32°E and 76°-19° E longitude and at altitude of 657 meters above mean sea level.

#### **2.2 Experimental Design and Procedures**

Six genotypes of kenaf (JRHC-15, JRK-2019-1, JRK-2019-2, JRK-2019-3, HC 583, AMC- 108 were evaluated in randomized Block Design with four replications at Cotton Improvement to study the phenology, growth, fibre yield and quality. The 30 x 5-7 cm spacing was adopted. The gross and net plot sizes were 6.00 x 4.50 m and 5.40 x 3.90 m, respectively. The fertilizer dose was applied as per recommended dose of 60:30:30 N, P2O5 and K2O kg/ha, the first half dose of nitrogen and full dose of P and K was given at the time of sowing and remaining dose of nitrogen was applied in 2 equal splits as top-dressing at 35-40 and 65-70 days of crop-age. Protective irrigations were given as when required. Five plants in each plot were randomly selected in a net plot area and tagged for recording the various growth traits, yield and other physiological parameters. The fibre quality parameters were estimated in laboratory after the harvesting of genotypes. Principal growth stages (days) were recorded as per the BBCH-scale [4]. The growth parameters viz. plant height, basal diameter, number of leaves and leaf area was recorded at 20 days interval from sowing. The physiological parameters viz. chlorophyll content, canopy temperature at top, middle and bottom level and light intensity at top, middle and bottom bevel was recorded at initiation of flowering by using SPAD meter, infra red thermometer and lux meter, respectively at initiation of flowing. The growth rates and physiological parameters were calculated by using following formulae.

Parameter	Formulae
Absolute Growth Rate (cm/day) [5]	$AGR = \frac{h1 - h2}{t2 - t1}$
Relative Growth Rate (cm/cm/day) [6]	$RGR = \frac{Logeh1 - Logeh2}{t2 - t1}$
Leaf Area Index [6]	Total leaf area per plant
Leaf Area Duration [7]	$LAD = \frac{(L1 + L2)}{2}X (t2 - t1)$
Light Extinction Coefficient	$K = \frac{LogeI/I0}{LAI}$
Light Transmission Ratio (LTR)	$LTR = \frac{I}{I0}$
Unit Area Efficiency (UAE)	$UAE = \frac{Fibre \ yield}{Land \ area} X \frac{1}{Duration \ of \ crop}$

Where, h1 and h2 are the plant height at t1 and t2 times, respectively, L1 and L2 = LAI at the first stage and second stage, Io and I are the light intensity at top and bottom of a population with LAI. Fibre yield and yield attributes and fibre quality characters were recorded after harvesting and retting. The data analyzed for coefficient of variation through randomized block design by Panse and Sukhatme [8].

# **3. RESULTS AND DISCUSSION**

#### **3.1 Crop Phenology**

The data on various principle growth stages such as germination, leaf and shoot development, inflorescence emergence and flowering in days through BBCH scale are presented in table 1 and depicted in figure 1. The extended BBCH scale considers 10 principal growth stages in roselle is evidenced and numbered from 0 to 9 [9]. Starting at germination (stage 0) and ending at the beginning of the rest period (stage 9). Based on BBCH scale, nine phenological stages were recorded for roselle that included: (0) Germination, (1) Leaf development, (2) Formation of side shoots, (3) Main stem elongation, (5) Inflorescence emergence, (6) Flowering, (7) Development of bolls, (8) ripening of sepals and (9) seeds Senescence.



Figure 1: Phenological development influenced by kenaf genotypes



# 3.1.1 Germination, sprouting and bud development

Days required for germination, sprouting and bud development are 1 to 6. The genotype JRK-2019-1 (6 days) required highest number of days while genotype JRHC-15 (4 days) required less number of days for germination, sprouting and bud development among 6 kenaf genotypes.

#### **3.1.2 Leaf development (main shoot)**

Days required for leaf development (main shoot) are 7 to 38. The genotype JRK-2019-1 required highest number of days (9.25 days) for first true leaf emergence, cotyledons completely unfolded and first leaves separated and (38 days) to more true leaves unfolded among 6 kenaf genotypes. The genotype JRHC-15 required less number of days (7.75 days) for first true leaf emergence, cotyledons completely unfolded and first leaves separated and (33 days) to more true leaves unfolded among 6 kenaf genotypes.

#### **3.1.3 Stem elongation, shoot development (main shoot)**

Days required for stem elongation and shoot development (main shoot) are 45 to 71 days. The genotype JRK-2019-1 required highest number of days (56 days) for stem 10% of final length (diameter); 1 node detectable and (71 days) to maximum stem length diameter reached or more nodes detectable among 6 kenaf genotypes. The genotype JRHC-15 required less number of days (45 days) for stem 10% of final length (diameter); 1 node detectable and (60 days) to maximum stem length diameter reached or more nodes detectable among 6 kenaf genotypes.

#### 3.1.4 Inflorescence emergence (main shoot) / heading

Days required for inflorescence emergence or heading are 63 to 93 days. The genotype JRK-2019-1 required highest number of days (76 days) for beginning of heading and (93 days) to first flower petals visible and inflorescence fully emerged. The genotype JRHC-15 required less number of days (63 days) for beginning of heading and (81 days) to first flower petals visible and inflorescence fully emerged among 6 kenaf genotypes.

#### 3.1.5 Flowering (main shoot)

Days required for inflorescence emergence and flowering are 86 to 108 days. The genotype JRK-2019-1 required highest number of days (98 days) for first flowers open (sporadically) and (108 days) to full flowering i.e. 50% of flowers open, first petals may be fallen. The genotype JRHC-15 required less number of days (86 days) for first flowers open (sporadically) and (100 days) to full flowering i.e. 50% of flowers open, first petals may be fallen among 6 kenaf genotypes. There were significant differences in the number of days to reach 50% flowering, 50% fruiting, and physiological maturity among the treatments [10].

	Principle growth stages (days)		IPK-		HC	IDK-	AMC
	Therpie growin stages (days)	2019-2	2019 <b>.</b> 1	JKIIC- 15	583	2019-3	108
1	Germination Sprouting Bud development	2017 2	2017 1	10	202	2017 0	100
01	Beginning of seed imbibition.	1	1	1	1	1	1
01	Beginning of bud swelling (P.V)	-	-	-	-	-	-
03	Seed imbibition complete: End of	2	2	2	2	2	2
00	bud swelling (P. V)	-	-	-	-	-	-
05	Radicle (root)emerged from seed:	3	3	2	3	3	3
	Perennating organs forming roots	-	-		-	-	-
	(P. V)						
06	Elongation of radicle, formation	3	4	3	3	3	3
	of root hairs and lateral roots						
07	Coleoptile emerged from caryopsis (G);	4	4	3	4	4	4
	Hypocotyl with cotyledons or shoot breaking						
	through seed coat (D, M), Beginning of						
	sprouting or bud breaking (P, V)						
08	Hypocotyl with cotyledons growing towards	4	5	4	4	4	4
	soil surface (D), Shoot growing towards soil						
	surface (P, V)						
09	Emergence: Coleoptile breaks through soil	5	6	4	5	5	5
	surface (G), Emergence: Cotyledons break						
	through soil surface (except hypogeal						
	germination D, M); Emergence: Shoot/leaf						
	breaksthrough soil surface (D, V); Bud						
	shows green tips (P)						
2	Leaf development (main shoot)						
10	First true leaf emerged from coleoptile (G);	8.5	9.25	7.75	8.5	8	8
	Cotyledons completely unfolded (D, M); First						
	leaves separated (P)						
11	First true leaf, leaf pair or whorlunfolded;	11	12	9	11	10.75	11
	First leaves unfolded(P)						
12	true leaves, leaf pairs or whorls	14	15	10	13	13.25	14
	unfolded						
13	true leaves, leaf pairs or whorls	16.75	18	13	16	17	17
	unfolded						
19	more true leaves, leaf pairs or	37	38	33	36	37	35.5
	whorls unfolded						
3	Stem elongation, shoot development (main s	hoot)			-		-
31	Stem 10% of final length	53	56	45	50	53	50
	(diameter); I node detectable (G)		50	10	50 75	~ ~	50
32	Stem 20% of final length	55	58	48	52.75	55	53
20	(diameter); 2 nodes detectable(G)	50	<i>(</i> 1 <i>)</i>	<b>C</b> 1	- 7	50	<b>C</b> 0
33	Stem (rosette) 30% of final length	59	61.5	51	57	59	60
	(diameter); 3 nodes detectable(G), Stages						
20	continuous till.	70	71	<u>(</u> )	<b>65</b>	70	66
39	continuous till. Maximum stem length diameterreached; or	70	71	60	65	70	66
39	continuous till. Maximum stem length diameterreached; or more nodes detectable	70	71	60	65	70	66
39	continuous till. Maximum stem length diameterreached; or more nodes detectable (G 9)	70	71	60	65	70	66
39 4 51	continuous till. Maximum stem length diameterreached; or more nodes detectable (G 9) Inflorescence emergence (main shoot) / hea Inflorescence or flower buds	70 nding 74	71	60	65 70	70	66

Table 1	: Phenological	developmen	nt influenced	by kenaf	genotype

55 First individual flowers visible (still closed); Half of inflorescence emerged (G)	82	85	73	78	83	80
59 First flower petals visible (inpetalled	91	93	81	86	90	89
forms); Inflorescence						
fully emerged (G)						
5Flowering (main shoot)						
60 First flowers open (sporadically)	97	98	86	92	96	93.67
61 Beginning of flowering: 10% of	99	101	89	94	98	95
flowers open						
62 20% of flowers open	102	102	93	96	101	97
63 30% of flowers open	104	104	95	98	103	99
64 40% of flowers open	106	106	98	100	105	101
65 Full flowering: 50% of flowers	108	108	100	102	107	102
open, first petals may be fallen						

#### **3.2 Growth parameter**

The data on various growth characters viz. plant height (cm), basal diameter (cm), number of leaves and leaf area/plant (dm<sup>2</sup>) influenced by kenaf genotypes at various stages of growth are presented in Table 2. The differences among the genotypes were statistically significant for all the growth characters at various stages of growth. Kenaf has an indeterminate type of growth, with rapid growth rate increases until the appearance of the first flower and gradual decreases afterwards [11].

In the present study, the plant height and basal diameter increased progressively with the advancing age of the crop, whereas, number of leaves and leaf area increased progressively upto 80 DAS and thereafter it was decreased towards advancing age of the crop due to defoliation. The genotype, JRK-2019-3 maintained higher plant height (387.50 cm) and basal diameter (2.35 cm) at harvesting as well as at various stages of growth. The genotypes, JRK-2019-2 (386.25 cm, 2.28 cm) and JRHC-15 (385.75 cm, 2.21 cm) were also promising for maintaining plant height and basal diameter at various stages of growth. On an average, genotype JRK-2019-2 recorded the highest number of leaves (366.70) and leaf area (425.37 dm<sup>2</sup>) at 80 DAS as well at various stages of growth. In addition to this, JRK-2019-3 maintained higheer number of leaves (364.79) and leaf area (386.75 dm<sup>2</sup>) at 80 DAS as well at various stages of growth. The kenaf varieties has significant difference between the, in terms of height, basal diameter and biomass [12].

	Table 2: Growth cl	haracters influe	enced at vario	us stages of gr	owth	
Genotype	20 DAS	<b>40 DAS</b>	60 DAS	80 DAS	100 DAS	120 DAS
Plant height (cm)						
JRK-2019-2	50.50	146.91	261.40	315.75	358.95	386.25
JRK-2019-1	40.20	133.68	239.47	299.62	346.25	374.00
JRHC-15	43.28	137.53	251.90	307.50	354.00	385.75
HC 583	34.33	122.58	234.18	288.95	343.50	366.00
JRK-2019-3	52.70	150.03	266.75	317.00	362.45	387.50
AMC 108	37.75	128.20	244.50	301.00	348.05	371.50
SE <u>+</u>	0.98	1.45	2.03	2.26	1.68	1.75
CD at 5 %	2.96	4.38	6.13	6.82	5.07	5.23
Basal diameter (cm)						
JRK-2019-2	0.80	1.38	1.71	1.97	2.11	2.28
JRK-2019-1	0.65	1.17	1.55	1.83	1.98	2.19
JRHC-15	0.73	1.25	1.62	1.89	2.02	2.21
HC 583	0.63	1.13	1.50	1.76	1.90	2.17
JRK-2019-3	0.90	1.51	1.74	2.02	2.16	2.35
AMC 108	0.77	1.31	1.61	1.89	2.05	2.24
SE <u>+</u>	0.069	0.099	0.019	0.023	0.028	0.017
CD at 5 %	NS	NS	0.058	0.071	0.08	0.053

No. of leaves						
JRK-2019-2	52.30	201.50	357.60	366.70	271.00	133.25
JRK-2019-1	42.75	193.75	323.75	341.00	235.75	118.75
JRHC-15	48.25	187.75	260.50	298.75	213.25	90.25
HC 583	36.25	176.25	258.50	315.50	220.00	98.25
JRK-2019-3	57.00	207.00	364.00	364.79	277.65	138.60
AMC 108	46.25	198.00	286.50	316.00	222.50	109.75
<u>SE+</u>	1.926	2.519	2.351	1.993	2.396	1.923
CD at 5 %	5.80	7.59	7.08	6.007	7.22	5.81
Leaf area ( dm <sup>2</sup> )						
JRK-2019-2	19.35	130.95	343.30	425.37	308.94	147.90
JRK-2019-1	9.83	93.00	229.83	310.31	209.81	102.12
JRHC-15	16.88	114.52	231.85	331.61	228.25	93.75
HC 583	10.87	96.93	206.83	318.65	215.60	92.35
JRK-2019-3	18.81	122.13	309.44	386.75	285.97	139.98
AMC 108	12.16	102.95	217.75	304.62	207.59	99.12
SE <u>+</u>	2.1041	2.8159	2.7658	2.2779	3.0369	2.7162
CD at 5 %	6.6060	8.5078	8.3576	6.8939	9.1708	8.2090

# **3.3 Growth rates**

The data on various growth rates viz. AGR (cm/day), RGR (cm/cm/day), LAI and LAD (days) influenced by kenaf genotypes at various stages of growth are presented in Table 3. It revealed that, the AGR increased progressively upto 40-60 DAS, rate was declined towards reproductive growth. However, RGR declined with the advancing age of the crop. LAI and LAD increased progressively upto 60-80 DAS, rate was declined towards reproductive growth due to defoliation. The genotype JRK-2019-3 maintained higher AGR at 0-20 DAS (32.70 cm/day), 20-40 DAS (77.33 cm/day) and 40-60 DAS (96.72 cm/day), whereas, JRK-2019-1 (40.15 cm/day) at 60-80 DAS, HC 583 (34.55 cm/day) at 80-100 DAS and JRHC-15 at 100-120 DAS (11.75 cm/ day) recorded higher AGR. The genotype HC 583 (0.027 cm/cm/day) at 20-40 DAS, HC-583 and AMC 108 (0.014 cm/cm/day) at 40-60 DAS, JRK-2019-1 (0.0048 cm/cm/day) at 60-80 DAS, HC 583 (0.0037 cm/cm/day) and JRHC-15 (0.0018 cm/cm/day) recorded higher RGR at respective growth stages. The genotypes JRK-2019-2 (236.31) and JRK-2019-3 (214.86) recorded higher LAI at 80 DAS as well as various stages of growth. Similarly, JRK-2019-2 (4270.38) and JRK-2019-3 (3867.72) recorded higher LAI at 60-80 DAS as well as various stages of growth.

	Table 3: Growth rates in	fluenced by l	kenaf genotypes	at various sta	ges of growth	
Genotypes	0-20 DAS	20-40	40-60	60-80	80-100	100-120
		DAS	DAS	DAS	DAS	DAS
		Absolute g	rowth rate (cm/o	day)		
JRK-2019-2	30.50	76.41	94.49	34.35	23.20	7.30
JRK-2019-1	20.20	73.48	85.79	40.15	26.63	7.75
JRHC-15	23.28	74.25	94.37	35.60	26.50	11.75
HC 583	14.33	68.25	91.60	34.77	34.55	2.50
JRK-2019-3	32.70	77.33	96.72	30.25	25.45	5.05
AMC 108	17.75	70.45	96.30	36.50	27.05	3.45
Mean	23.12	23.12	73.36	93.21	35.27	27.23
		Relative gro	owth rate (cm/cn	n/day)		
JRK-2019-2		0.023	0.012	0.0041	0.0027	0.0015
JRK-2019-1		0.026	0.012	0.0048	0.0031	0.0016
JRHC-15		0.025	0.013	0.0043	0.0030	0.0018
HC 583		0.027	0.014	0.0045	0.0037	0.0013
JRK-2019-3		0.022	0.012	0.0037	0.0029	0.0014
AMC 108		0.026	0.014	0.0045	0.0031	0.0014

Mean		0.025	0.013	0.0043	0.0031	0.0015
		Leaf ai	ea index			
JRK-2019-2	10.75	72.75	190.72	236.31	171.63	82.16
JRK-2019-1	5.46	51.66	127.68	172.39	116.56	56.73
JRHC-15	9.37	63.62	128.80	184.22	126.80	52.08
HC 583	6.038	53.85	114.90	177.02	119.77	51.30
JRK-2019-3	10.45	67.85	171.91	214.86	158.87	77.76
AMC 108	6.75	57.19	120.97	169.23	115.33	55.07
Mean	8.13	61.15	142.50	192.34	134.83	62.52
		Leaf are	a duration			
JRK-2019-2		835.00	2634.72	4270.38	4079.50	2538.00
JRK-2019-1		571.27	1793.50	3000.77	2889.55	1732.94
JRHC-15		730.00	1924.27	3130.33	3110.33	1788.88
HC 583		598.88	1687.55	2919.33	2968.05	1710.83
JRK-2019-3		783.00	2397.61	3867.72	3737.33	2366.38
AMC 108		639.51	1781.66	2902.08	2845.66	1704.00
Mean		692.94	2036.55	3348.43	3271.74	1973.50

# 3.4 Chlorophyll content and canopy temperature

The SPAD meter (Soil Plant Analytical Development) is a simple hand held and portable instrument that is widely used for the rapid, accurate and non-destructive measurement of leaf chlorophyll concentrations. Canopy temperature (IRT) is an integrative trait that reflects the plant water status or the resultant equilibrium between the root water uptakes and shoots transpiration [13]. The SPAD chlorophyll meter reading at initiation of flowering in kenaf genotypes ranges from 38.8 (HC 583) to 47.63 (JRK-2019-3). At initiation of flowering, canopy temperature at top, middle and bottom level ranges from 25.2 to 35.1 (<sup>0</sup>C). Genotype JRK-2019-3 at top (35.10°C), middle (30.15°C) and bottom (27.7°C) level recorded significantly highest canopy temperature. Genotype JRHC-15 showed lowest canopy temperature at top (32.50°C) and middle (27.50°C) level, while at bottom level genotype JRK- 2019-1 (25.2°C) showed lowest canopy temperature at initiation of flowering (Table 4).

Genotype	Chlorophyll	Canopy Temperat	ure (IRT)	
	Content (SPAD)	Тор	Middle	Bottom
JRK-2019-2	42.63	34.40	30.15	25.6
JRK-2019-1	42.75	33.70	29.43	25.2
JRHC-15	44.83	32.50	27.50	26.4
HC 583	38.80	33.20	28.50	25.5
JRK-2019-3	47.63	35.10	30.15	27.7
AMC 108	41.89	34.12	30.10	26.2
SE <u>+</u>	1.5178	0.4298	0.5637	0.5117
CD at 5%	4.5752	1.2955	1.6992	1.5424

Table 4: Chlorophyll content, canopy temperature influenced by kenaf genotypes

# 3.5 Light intensity and unit area efficiency

A genetic, molecular, physiological, biochemical, and functional genomics approach, significant developments have been made in identifying genes and molecular mechanisms underlying the relationship of light intensity and photosynthesis [14]. In simple crop models such as LINTUL, extinction coefficient of light (KL) is widely used to calculate light interception by the canopy and to predict biomass yields based on the light use efficiency concept [15]. At initiation of flowering, light intensity by using LUX meter at top, middle and bottom level ranges from 8000 to 72000 LUX. At top, middle and bottom level genotype JRHC-15 recorded significantly highest light intensity. Genotype JRK-2019-2 shows lowest lux meter reading at top and bottom level while genotype JRK-2019-3 shows lowest lux meter reading at middle level at initiation of flowering. The mean light extinction coefficient (K) was of 0.004353. The genotype AMC 108 (0.004901) JRK-2019-2 (0.003429) recorded highest and lowest light extinction coefficient (K), respectively. The mean

light transmission ratio was of 0.14. The genotypes, JRHC-15 (0.167) JRK-2019-2 (0.123) recorded highest and lowest light transmission ratio, respectively. The mean unit area efficiency was 0.001391. The genotype JRK-2019-2 (0.001610) and HC 583 (0.001293) recorded the highest and lowest unit area efficiency, respectively (Table 5).

Table 5: Light intensity and unit area efficiency influenced by kenar genotypes								
Genotype	Light Intensi	ty (LUX Meter	r)	Light extinction	Light	Unit area		
	Тор	Middle	Bottom	coefficient (K)	transmission	efficiency		
	-				ratio			
JRK-2019-2	65000	26000	8000	0.003429	0.123	0.001610		
JRK-2019-1	70000	30000	10000	0.004791	0.143	0.001308		
JRHC-15	72000	38000	12000	0.004558	0.167	0.001522		
HC 583	71000	30000	10000	0.004657	0.141	0.001293		
JRK-2019-3	68000	25000	8500	0.003783	0.125	0.001350		
AMC 108	70000	35000	10500	0.004901	0.150	0.001264		
SE <u>+</u>	859.08	896.13	617.59					
CD at 5%	2589.55	2701.24	1861.64					

**Table 5:** Light intensity and unit area efficiency influenced by kenaf genotypes

# 3.6 Yield and yield contributing characters

The data on yield and yield contributing characters are presented in table 6. The differences among the genotypes were statistically significant for all the yield and yield contributing characters (Table 6). The genotype JRK-2019-1 (539.13 q/ha) and HC 583 (485.89 q/ha) produced highest and lowest green weight, respectively. The highest and lowest fibre yield was recorded by the genotype JRK-2019-2 (34.93 q/ha) and AMC 108 (27.44 q/ha), respectively. The genotype JRK-2019-2 (365.72 q/ha) showed significantly highest stick yield followed by JRHC-15 (353.2 q/ha), JRK-2019-3 (325.88 q/ha). The highest fibre recovery was recorded by the genotype JRK-2019-2 (8.71 %) and the lowest fibre recovery was recorded by the genotype JRK-2019-3 (8.29 %).

	Table 6: Yield and yield contributing characters influenced by kenaf genotypes								
Genotype	Green	Fibre	Stick	Fibre					
	weight (q/ha)	yield (q/ha)	yield (q/ha)	<b>Recovery</b> (%)					
JRK-2019-2	511.67	34.93	365.72	8.71					
JRK-2019-1	539.13	28.38	303.94	8.51					
JRHC-15	532.51	33.02	352.20	8.55					
HC 583	485.89	28.06	300.90	8.55					
JRK-2019-3	526.93	29.29	325.88	8.29					
AMC 108	507.04	27.44	296.55	8.47					
SE <u>+</u>	10.615	0.59	10.17						
CD at 5%	32.289	1.823	30.65631						

#### Fibre quality parameter

The data on fibre quality parameter such as root content, defects, tenacity, fineness, colour, bulk density and BIS Grade are presented in table 7. The genotypes JRK-2019-2 and JRK-2019-1 recorded highest (40 wt. %) root content, while genotypes HC 583, JRK-2019-3 and AMC 108 recorded lowest (12 wt. %) root content. Genotypes JRK-2019-2 and JRK-2019-3 was reported to be the lesser defects (0.5 wt. %), whereas, genotypes JRHC-15 and AMC 108 recorded highest defects (1.5 wt. %). The genotype JRK-2019-2 (26 g/tex) and AMC 108 (23.6 g/tex) showed significantly highest and lowest tenacity, respectively. The mean fineness (tex) was 4.98 and it ranged between 4.5 tex (HC 583) to 5.5 tex (JRK-2019-1) fineness. All genotypes showed creamy white fibre colour. The genotypes JRK-2019-2, JRK-2019-1 and JRHC-15 shown heavy bodied bulk density while other three genotypes HC 583+, JRK-2019-3 and AMC 108+ shown medium bodied bulk density. The genotypes JRK-2019-2, JRHC-15 and AMC 108 recorded M3+70% BIS grade while genotypes JRK-2019-3 and JRK-2019-3 recorded M3+30% BIS grade. The genotype HC 583 recorded M2+10% BIS Grade.

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Genotype	Root	Defects	Tenacity	Fine-	Colour	<b>Bulk Density</b>	BIS
	content	(wt.%)	(g/tex)	ness(tex)			Grade*
	(wt.%)						
JRK-2019-2	40	0.5	25.6	5.1	Creamy white	Heavy Bodied	M3+70%
JRK-2019-1	40	1.0	24.1	5.5	Creamy white	Heavy Bodied	M3+30%
JRHC-15	20	1.5	25.4	4.8	Creamy white	Heavy Bodied	M3+70%
HC 583	12	1.0	25.2	4.5	Creamy white	Medium Bodied	M2+10%
JRK-2019-3	12	0.5	26	4.6	Creamy white	Medium Bodied	M2+30%
AMC 108	12	1.5	23.6	5.4	Creamy white	Medium Bodied	M3+70%

#### 4. CONCLUSIONS

The genotype JRHC-15 required less number of days (100) for 50% of flowering and JRK-2019-1 (108 days) and JRK-2019-3 (107 days) required more number of days for 50% of flowering. Genotype JRK-2019-3 shows superior performance in growth and development. Genotype JRHC-15 shows superior performance in AGR, RGR and LAD, while, JRK-2019-2 shows good performance in LAI and unit area efficiency. Genotype AMC 108 performed better results in light extinction coefficient while, highest light intensity at top, middle and bottom and light transmission ratio is observed in JRHC-15. JRK-2019-2 shows better performance in yield and yield contributing characters. Genotypes JRK-2019-2 and JRK-2019-1 was reported to be the highest root content (wt. %), JRK-2019-2 and JRK-2019-3 was reported to be the lesser defects (wt.%), JRK-2019-3 and JRK-2019-2 has good tenacity and JRK-2019-1 had Higher fibre fineness. Genotypes JRK-2019-2, JRK-2019-1, JRHC-15 shows heavy bodied bulk density and HC 583, JRK-2019-3, AMC 108 shows medium bodied bulk density. All genotypes appeared creamy white fibre in colour. The yield contributing characters were significantly associated with fibre yield per plant.

#### **DISCLOSURE STATEMENT**

The paper is original. No part of the manuscript has been published before, nor is any part of it under consideration for publication in another journal. In addition, we affirm that all the authors have approved the manuscript for submission.

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#### **COMPETING INTERESTS**

Authors have declared that no competing interests exist.

#### REFERENCES

- [1] Webber III CL, Bledsoe VK, Bledsoe RE, Janick J, Whipkey A (). Kenaf harvesting and processing. Trends in new crops and new uses. 2002; 9: 340-347. ASHS Press, Alexandria, VA
- [2] Tahir Paridah Md, Ahmed AB, SaifulAzry SOA, Ahmed Z. (). "Retting Process of Some Bast Plant Fibres and Its Effect On Fibre Quality: A Review" (PDF). BioResources. 2011; 6(4): 5260–5281. DOI:10.15376/BIORES.64.5260-5281
- [3] Meier U, Bleiholder H, Buhr L, Feller C, Hack H, Heß M, Zwerger P. The BBCH system to coding the phenological growth stages of plants-history and publications. *Journal für Kulturpflanzen*, 2009; 61(2):41-52. DOI:10.5073/JFK.2009.02.01
- [4] Hunt R, Hunt R. Absolute growth rates. Basic Growth Analysis: Plant growth analysis for beginners, *pp*:17-24. 1990. DOI:<u>10.5860/choice.28-2138</u>
- [5] Hack H, Bleiholder H, Buhr L, Meier U, Schnock Fricke U, Weber E, Witzenberger A. (). Einheitliche Codierung der phanologischen Entwicklungsstadien mono-Und dikotyler Pfl anzen – Erweiterte BBCH-Skala, Allgemein. Nachrichtenbl. Deut. Pfl Anzenschutzd. 1992; 44: 265-270. © Eugen Ulmer GmbH & Co., Stuttgart
- [6] Williams RF. The physiology of plant growth with special reference to the concept of net assimilation rate. Annals

of Botany. 1946; 10(37): 41-72. https://doi.org/10.1093/oxfordjournals.aob.a083119.

- [7] Power JF, Willis WO, Grunes DL, Reichman GA (). Effect of soil temperature, phosphorus, and plant age on growth analysis of barley. Agronomy Journal. 1967; 59(3): 231-234.
   DOI:10.2134/AGRONJ1967.00021962005900030007X
- [8] Panse VG, Sukhatme PV. Statistical methods for agriculturalresearch. ICAR, New Delhi, pp: 308-318. 1985.
- [9] Javadzadeh SM, Rezvani Moghaddam P, Bannayan Aval M, Asili J. (). Assessment of Required Growing Degree Days for Phenological Stages of Roselle (*Hibiscus sabdariffa* L.) based on BBCH-Scale in Different Cropping Systems. Journal of Agroecology. 2018; 10(2): 368-385. DOI:10.22067/JAG.V10I2.38318
- [10] Kassim HG, Bello NJ, Ufoegbune GC, Makinde AA, Olasantan FO (). Effects of rainfall variability on moisture availability for cultivation of sorghum, kenaf and okra in tropical wet-and dry-climatic Western Zones of Nigeria. Nigerian Journal of Horticultural Sciences. 2022; 26 (4): 169-181. ISSN 1118-2733
- [11] Petrini C, Bazzocchi R, Montalti P. Yield potential and adaptation of kenaf (*Hibiscus cannabinus*) in north-central Italy. Industrial Crops and Products. 1994; 3(1-2): 11-15. https://doi.org/10.1016/0926-6690(94)90073-6
- [12] Shukor NAA, Hamzah MB, Hamid HA, Salleh G, Nasir MFM (). Growth and phenology of kenaf (*Hibiscus cannabinus* L.) varieties. PertanikaJournal of Tropical Agricultural Sciences. 2009; 32(1): 29-33.
- [13] Berger B, Parent B, Tester M. High-throughput shoot imaging to study drought responses. Journal of Experimental Botany. 2010; 61(13): 3519-3528. <u>https://doi.org/10.1093/jxb/erq201</u>
- [14] Wimalasekera R (). Effect of light intensity on photosynthesis. Photosynthesis, productivity and environmental stress, *pp*: 65-73. 2019. <u>https://doi.org/10.1002/9781119501800.ch4</u>
- [15] Van Ittersum MK, Leffelaar PA, Van Keulen H, Kropff M J, Bastiaans L, Goudriaan J (). On approaches and applications of the Wageningen crop models. European Journal of Agronomy. 2003; 18(3-4): 201-234. https://doi.org/10.1016/S1161-0301(02)00106-5