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Genotypic Response of Common Bean (*Phaseolus vulgaris* L.) to Moisture Stress Conditions in Kenya

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Abstract: The aim of this study was to determine drought response of thirty bean genotypes under glasshouse and field conditions. The experiment was laid in a Randomized Complete Block in split-plots design with three replications. Significant (p<0.05) genotypic differences in drought resistance were observed under moisture stress conditions. Significant interaction (p<0.01) between variety and moisture level were observed both in the glasshouse and in the field. Drought resistant genotypes showed high relative water content in stress environment as compared to susceptible ones. They also took relatively longer time to wilt, consequently showing low soil water content at wilting and lower biomass reductions under moisture stress conditions. A significant negative correlation was obtained between days to permanent wilting and relative water with biomass reduction. Relative water content was positively and significantly correlated to stomatal conductance and days to permanent wilting. Correlation between stomatal conductance and biomass yield reduction was not significant. The responses relative water content and stomatal conductance, in certain conditions, were recognized as beneficial drought resistance indicators and may be used as selection criteria in bean breeding programme.

Key words: Drought, beans, drylands, relative water content, selection

INTRODUCTION

Common bean (Phaseolus vulgaris L.) is one of the most important crops grown in the semi-arid areas of Kenya. It is the most important source of proteins in Kenya and ranks second to maize in importance as source of food, especially amongst rural communities. Drought conditions in Kenya are widespread covering up to 80% of landmass, therefore beans are frequently exposed to drought stress conditions. Studies on the crop in these regions indicate a wide genetic variability in drought resistance among the unimproved bean cultivars^[1]. Information on drought resistance amongst improved cultivars was reported by Nyabundi et al.[2], however, the information is inadequate and therefore there is need to test various genotypes to ensure that losses due to drought are reduced. To achieve this, identification of traits associated with drought is necessary. Runkulatile et al.[3] reported that land races from dry areas maintain a consistently higher stomatal conductance, transpiration rates, net photosynthesis and relative water content. Alternatively, Mcree and Richardson^[4] suggested that the ability of a drought adapted variety to root deeper in the soil horizon and extract more water

from the receding waterfronts may help maintain high relative water content of the plant. This is associated with high photosynthetic activity without a significant reduction in stomatal closure and transpiration rate. Cultivars with these abilities tend to exhibit delay in dehydration as indicated by the number of days to permanent wilting with the onset of drought^[5]. In common beans, information on responses of these traits to drought stress is scanty, especially in tropical regions.

Relative Water Content (RWC) has been suggested to be a better measure for plant's water status than the thermodynamic state variables (water potential, turgor potential and solute potential). Slatyer^[6] showed that when the water potential was -20 bars, the RWC was 50% in tomato leaves, but about 90% in *Acacia anaeura phyllodes*, implying that the tomato plant had a higher water deficit than *Acacia*. Similarly, Fischer^[7] found that RWC was directly related to soil water content and suggested that RWC might also be used to indicate soil water. Leaf RWC in plants decrease as water stress levels increase. It would be expected that under non-stress conditions, RWC of the plant would be near 100% with water potential values approaching zero. Bennet *et al.* ^[8] and Schonfeld *et al.* ^[9] noted that superior performance of

drought tolerant soybeans, maize and wheat varieties under water stress environment is attributed to osmoregulation and stomatal closure when stress sets in. These cultivar differences in RWC could be used to select high yielding genotypes that maintain cell turgor under water stress environment to give high relative yield[10]. Water stress reduces plant growth more than all other stresses combined[11]. Bradford and Hsiao[12] demonstrated that small reductions in growth rates in the early growth stage due to water stress would, because of exponential nature, compound with time into large reductions in biomass productions. Water stress depresses dry matter production in most crops^[13,14] but it is the stress intensity, duration of application and crop's growth stage that determines the degree of biomass depression^[15]. There is little information on the response of Kenyan land races and their crosses to drought stress. Information on selection criteria is also scanty. Therefore, the objective of this study was to evaluate the value of morphological and physiological traits as indicators that can be used in selection for drought resistance in common beans. The hypothesis tested was that common bean response to water stress was similar across all the commercial check land races and their crosses and that no trait was associated with drought in beans.

MATERIALS AND METHODS

Site description: The study was conducted between March 1994 and March 1994 in Central province of Kenya, at Kabete Campus, University of Nairobi (1°45 20 S and 36°45 20 E). The site is in agriculturally high potential zone, LH3, within the Kenya highlands^[16]. Kabete is at an altitude of 1829 m above sea level and receives annual rainfall of about 1000 mm with a mean monthly maximum temperature of 32°C and minimum of 12°C. The soils are well-drained friable clay with humic Nitosols. The mean minimum and maximum glasshouse temperature during the experimental period was 14.8 and 36.2°C, respectively.

Bean genotypes: Thirty bean genotypes (Table 1) obtained from the Bean Project of the Department of Crop Science; University of Nairobi comprised five seed types namely Rose Coco (RC), Canadian Wonder (CW), Mwitemania (MW), Mwezi Moja (MM) and Red Haricot (RH). Each seed type had a number of nearly homozygous F7, F8 and F9 generation bean lines selected from populations created from crosses among seven parents including five local commercial cultivars GLP-2, GLP-24, GLP-92, GLP-585 and GLP-1004 which were included in this study. Based on the days to maturity, the genotypes

Table 1: Bean cultivars used in this study

Cultivar	Seed type
E1	RC+
E3	RC
E4	RC+
E5	RC
E6	RC+
E8	RC
E9	RC
E10	RC+
M11	CW+
M13	CW+
M14	RC
M15	RC
M16	\mathbf{MW}^+
M18	$\mathbf{M}\mathbf{M}^{+}$
M19	\mathbf{MW}^+
M20	CW
M23	RC+
M26	RH+
L37	RC
L38	CW+
L42	CW
L44	$\mathbf{M}\mathbf{M}^{+}$
L46	RH+
L49	CW+
L50	CW
GLP-2**	RC+
GLP-24**	CW+
GLP-92**	MW+
GLP-585**	RH+
GLP-1004**	MM+

 $\begin{array}{llll} \hline RC = Rose \ Coco, & CW = Canadian \ Wonder, & MW = Mwitemania, \\ MM = Mwezi \ Moja, & RH = Red \ Haricot, & ** Commercial check cultivars, \\ E = Early, & M = Medium \ and \ L = Late \ maturing, + Cultivars \ used \ in \\ \hline determination \ of \ RWC \ in the \ field & \hline \end{array}$

were designated as Early (E), Medium (M) or late (L) maturing with days to maturities of 80 to 85, 86 to 96 and more than 96 days, respectively.

Experimental design and sampling procedures: Thirty genotypes were planted in the glasshouse in March to May 1994 in 8L pots. Each pot comprised of topsoil mixed with gravel and sand in a ratio of 2:1:1 and DAP (18:46:0) at the rate of 100 kg ha⁻¹. A 2 x 30 factor experiment was arranged as a split-plot in Randomized Complete Block Design replicated three times. Two watering levels (water stress and water non-stress) were assigned as whole plots while the genotypes were ascribed to the sub-plots. Six seeds were planted in each pot and watered to field capacity. The seedlings were thinned to four per pot seven days after emergence. Stress treatment was initiated at the third trifoliate leaf stage (at about 28 Days After Emergence (DAE)) by bringing all the pots to field capacity in the morning of day zero. There after, water was withheld until dawn time (6 am) wilting was evident in each genotype. This was assumed to be the permanent wilting point^[17]. Data in the glasshouse were taken on three randomly selected plants from each sub-plot (genotype) for the leaf RWC, Days to Permanent Wilting (DPW), Soil Moisture Content at wilting (SMC) and stomatal conductance at wilting. The RWC values were determined as the water content at sampling relative to that at full turgor as described by Bennet et al.[8], thus percent RWC = (Fresh weight-dry weight/turgid weightdry weight) x 100. Days to permanent wilting represented the number of days from when water was withheld till dawn time wilting was evident in each genotype. Stomatal conductance (cm sec⁻¹) was determined on the abaxial leaf surface of wilted plants using a steady state porometer (Li-1600). Soil moisture content at wilting was determined gravimetrically by oven-drying the soil sample at 105°C for 24 h to get the dry weight for each genotype. The soil moisture content was expressed as percent by weight using the formula: % Water by weight = (Wet weight-dry weight/dry weight) x 100^[18].

The field experiment was conducted during the dry spell between January and March 1995. Sixty treatments were arranged as a split-plot in a RCBD replicated three times. Three levels of watering (high, medium and low) were assigned to whole plots measuring 2x30 m. The subplots measured 1x2 m and the treatments consisted of the genotypes, each in five rows spaced 0.5 m apart. Seeds were planted at an intra-row spacing of 10 cm giving a plant population of 200,000 plants ha⁻¹. Seedlings were thinned to one per hill ten days after emergence. Irrigation water was applied using a line source sprinkler irrigation set that provided a moisture gradient, decreasing with increase in distance from the sprinkler line. Sprinkler stands were spaced at 30 m and sprinkler throw of about 15 m was achieved. The high, medium and low level of watering treatments received a total of 413, 312 and 174 mm of water, respectively. These amounts included little rainfall (144 mm) that fell during the experimental period. Field data were collected from each sub-plot region receiving designated amounts of water (excluding double irrigated overlap areas) and included the following response variable: Total plant biomass accumulation, leaf relative water content and biomass yield reductions from the high, medium and low watering levels. Total aboveground biomass was determined on five plants per sub-plot at fortnightly interval starting 14 DAE. The plants were cut at ground level and dried at 70°C for 72 h to constant weight. Leaf RWC was measured as described for the glasshouse experiment at 14 day interval, beginning 15 DAE up to 43 DAE. The biomass yield reductions for the medium and low water treatments were expressed as percentage of the high water treatment biomass. The high water level biomass was assumed to be non-stress while the medium and low water levels were differentially stressed, thus biomass yield reduction was computed as follows: % Biomass reduction = (Biomass stress/biomass non-stress) x 100.

Data analysis: Data were subjected to analysis of variance using the General Linear Model (GLM) and means separated by Duncan's Multiple Range Test (p<0.05) on SAS software^[19]. Correlation analysis was conducted for the data according to Steel *et al.*^[20].

RESULTS

The mean RWC across the genotypes in the glasshouse water stress treatment generally decreased with increasing number of Days After Withholding Water (DAWW) (Table 2) with the highest mean value being 87.59% (0 DAWW) and lowest value of 68.92% at wilting. The mean leaf RWC across the genotypes remained relatively stable throughout the first three days after watering was withheld, however, by the sixth day, the mean leaf RWC across the genotypes had dropped to 80.88%. Further drops to means of 73.6 and 69.31% were similarly observed for nine and twelve DAWW and at wilting, respectively. At dawn time wilting, cultivars M19, M16, M18, GLP-1004 and GLP-92 had the highest RWC means while E4, L46, M15, L37 and GLP-585 had the lowest mean values at the same time. Cultivar GLP-1004 took the longest period to permanent wilting (22 days) followed by GLP-92 and M18 (21 days each) and M19 that wilted after 20 days (Table 3). These genotypes appeared to maintain cell turgidity even at comparatively low soil moisture contents at which wilting occurred. These genotypes maintained their stomatal conductance at high values (> 0.30 cm sec⁻¹) despite the low moisture contents at wilting. In contrast, E6, E10, M11, M14, M15, M26 and L37 took minimum time to loose turgidity, wilting after 12 days. At this time, the soil moisture contents were comparatively high and the stomatal conductance values were low compared with the seemingly drought resistant genotypes such as GLP-1004, GLP-92, M16 and L44, which had lower SMC and higher stomatal conductance under the same treatment conditions. Days to Permanent Wilting (DPW) were significantly negatively correlated to soil water content at wilting (r= 0.88**)(Data not shown). It was however significantly and positively correlated to leaf relative water content at wilting (r=0.81*). Significant negative correlation was found between soil moisture content at wilting and leaf RWC at wilting (r=-0.65*). Relative water content at zero DAWW associated significantly positively with DPW but negatively with SMC in the glasshouse (r=0.54* and -0.51*, respectively).

Table 2: Relative water content (%) for stressed bean genotypes at different DAWW

	Days after W					
Cultivars	0	3	6	9	12	RWCW
E1	81.62ef	78.42d	71.47hi	69.54f-h	67.90cd	66.66j-m
E3	83.33d-f	84.19a-d	76.38f-i	73.08d-h	69.11cd	68.57f-k
E4	87.30a-f	86.11a-d	74.24g-i	69.74f-h	67.28cd	67.49h-m
E5	86.28a-f	86.75a-d	79.60a-d	71.33e-h	68.95cd	69.27f-j
E6	87.08a-f	88.38a-c	82.32a-f	74.45d-g	68.95cd	67.40h-m
E8	83.26d-f	82.17cd	77.94e-I	71.89eh	70.30b-d	71.43d-f
E9	87.76a-f	86.54a-d	86.25a-d	70.42e-h	66.71cd	68.54f-k
E10	87.75a-f	85.60a-d	76.83e-I	73.97d-g	68.75cd	67.13i-m
M11	87.21a-f	85.46a-d	78.40e-h	74.23d-g	67.10cd	66.59j-m
M13	86.54a-f	86.75a-d	79.60c-g	69.10f-h	67.10cd	66.18j-m
M14	91.62a-c	89.31a-c	84.98a-e	76.83c-e	69.38cd	70.53e-h
M15	88.60a-e	85.46a-d	78.88d-h	72.30d-h	67.80cd	66.07j-m
M16	89.10a-e	85.64a-d	85.46a-e	78.97b-d	73.22a-c	76.46ab
M18	93.02a	92.23ab	89.89a	85.76a	76.83ab	75.35a-c
M19	91.16a-c	89.10a-c	86.75a-c	82.57a-c	78.85a	77.25a
M20	88.60a-e	83.69b-d	77.02f-I	72.69d-h	67.10cd	65.09lm
M23	89.10a-e	87.21a-c	88.15ab	73.22d-h	67.97cd	68.47f-k
M26	85.98a-f	83.26b-d	78.27e-h	72.11e-h	69.74b-d	68.22f-I
L37	80.48f	82.42cd	73.97g-I	67. 8 0gh	67.63cd	65.60k-m
L38	84.55c-f	83.26b-d	78.97c-h	74.45d-g	68.22cd	68.25f-1
L42	92.28ab	89.12a-c	78.33e-h	72.30cd	66.77cd	67.68g-m
L44	91.23a-c	92.23ab	87.54ab	67.67cd	67.67cd	70.78e-g
L46	87.11a-f	86.03a-d	76.83f-I	70.91e-h	66.77cd	65.54k-m
L49	89.53a-d	86.11a-d	84.19a-f	74.45d-g	71.89a-d	70.16e-I
L50	89.90a-d	86.90a-c	83.26a-f	74.45d-g	69.54b-d	67.03i-m
GLP-2	84.70b-f	82.83b-g	80.93b-g	72.57d-h	69.80b-d	66.44j-m
GLP-24	91.62a-c	89.43a-c	86.75a-c	75.53d-f	68.22cd	68.19f-l
GLP-92	87.54a-d	85.82a-d	83.76a-f	70.00e-h	64.16d	72.74c-e
GLP-585	83.26d-f	83.81b-d	70.48I	66.77h	66.77cd	64.54m
GLP-1004	91.23a-c	92.28a	89.10a	83.44ab	78.97a	73.85c-d
Mean	87.59	86.15	80.88	73.60	69.31	68.92
SE	2.22	2.43	2.31	1.98	2.27	1.36
CV (%)	4.39	4.88	4.95	4.66	5.67	1.97

RWCW = Relative Water Content at Wilting

Table 3: Days to permanent wilting, stomatal conductance and percent soil water content at wilting, for bean cultivars in the glasshouse experiment

Cultivars	Days to permanent wilting	Stomatal conductance at wilting	Soil water content at wilting
E1	13i	0.14n	25.00a-f
E3	13i	0.18j-m	26.87a-d
E4	15h	0.16ln	24.98a-f
E5	13i	0.15l-n	27.54ab
E6	12i	0.16l-n	28.03a
E8	17e-h	0.25f	24.53b-g
E9	15h	0.15mn	23.28e-i
E10	12i	0.17 j- m	25.18a-f
M11	12i	0.17 j-m	23.80d-h
M13	13i	0.1 <i>6</i> k-n	24.06c-g
M14	12i	0.17k-n	27.38a-c
M15	12i	0.15l-n	25.49a-f
M16	18c-e	0.32b-d	17.681
M18	21ab	0.30de	19.94i-l
M19	20bc	0.34ab	18.58kl
M20	17e-h	0.20h-i	22.94f-i
M23	19cd	0.29e	22.36f-i
M26	12i	0.20h-j	26.52a-e
L37	12i	0.18i-l	27.78ab
L38	16gh	0.1 <i>7</i> k-m	24.37b-g
L42	17 e-h	0.24f	23.20e-i
L44	18c-f	0.35a	20.67h-l
L46	13i	0.18i-k	24.48b-g
L49	19cd	0.22gh	22.48f-i
L50	18c-e	0.23fg	21.51g-k
GLP-2	16gh	0.18i-k	23.29e-i
GLP-24	17 d- g	0.20hj	21.36g-k
GLP-92	21ab	0.31с-е	17.851
GLP-585	16gh	0.15mn	22.28f-j
GLP-1004	22a	0.33bc	19.04j-l
Mean	16	0.21	23.42
SE	0.54	0.011	1.02
CV (%)	6.00	5.29	7.55

Table 4: Varietal differences in percent leaf relative water content under different watering levels at different DAE of field experiment

High (413 mm) DAE			`	Medium (312 mm) DAE			Low (174 mm) DAE		
Cultivars	15	29	43	15	29	43	15	29	43
E1	84.53a	77.82b	81.20ad	71.08be	75.80be	62.88ef	62.88ef	66.58h	65.02ac
E4	87.30a	89.15ab	76.11bd	74.24bd	79.45ae	77.95ae	65.20df	74.24cg	69.11ac
E6	88.73a	82.90ab	78.63ad	75.80bd	71.80ce	68.44e	61.93f	67.24cg	61.59c
E10	85.45a	81.54ab	81.25ad	75.06bd	69.88e	70.63ce	62.41ef	68.98gh	63.88bc
M11	86.63a	80.62ab	86.68ac	72.66cd	78.67ae	71.95ce	64.81df	76.53ae	68.09ac
M13	87.09a	82.99ab	79.94ad	76.93bd	70.31e	81.48ac	65.79df	68.97gh	69.48ac
M16	85.70a	88.13ab	85.65ac	76.20bd	84.45ab	75.44be	72.70ae	79.11ac	72.88ac
M18	92.60a	88.45ab	87.54ab	85.85ab	87.86a	80.92ad	77.95ab	72.34dh	70.00ac
M19	88.05a	91.90a	85.90ac	79.54bd	86.15ab	85.55ab	73.04af	81.70a	77.35a
M23	89.56a	86.70ab	78.70ad	77.48bd	71.48de	80.33ae	73.48ae	78.19ad	76.63ab
M26	87.49a	83.81ab	77.05bd	69.54d	71.72ce	68.52e	72.11bf	71.37eh	74.38ac
L38	83.62a	81.35ab	87.95ab	76.84bd	76.45be	70.06ce	69.74bf	68.63gh	71.19ac
L44	93.42a	89.45ab	83.33ad	81.46ac	85.56ab	78.29ae	77.29ae	76.36ae	69.86ac
L46	83.70a	76.96b	86.87ac	76.83bd	81.48ad	78.24ae	66.77cf	69.58fg	71.85ac
L49	88.26a	78.05b	83.68ad	81.14ad	71.41 de	70.60ce	75.15ad	75.38bf	70.05ac
GLP-2	84.96a	79.60ab	74.65cd	80.19bd	82.26ac	71.33ce	69.80bf	74.70cg	67.70ac
GLP-24	74	78.90b	82.53ad	72.30cd	78.96ae	69.38ae	73.38ae	79.08ac	71.11ac
GLP-92	94.23a	88.92ab	90.32a	77.06bd	83.26ab	85.46ab	78.90ab	81.20ab	73.21ac
GLP-585	85.40a	79.98ab	72.56d	75.73bd	69.88e	70.22ce	69.93bf	68.64gh	73.33ac
GLP1004	89.10a	87.58ab	90.35a	91.34a	88.64a	88.46a	83.33a	77.88ad	73.20ac
Mean	86.99	83.74	82.54	77.36	78.27	75.89	70.82	73.82	70.54
SE	3.09	3.71	3.56	3.41	3.16	3.49	3.25	3.66	3.87
CV (%)	5.02	6.27	6.11	6.23	5.71	6.50	6.49	7.02	7.76

Table 5: Varietal differences in biomass accumulation (g m^{-2}) under different watering levels at different DAE

	High (413 mm) DAE			Medium (312 mm) DAE			Low (174)	·		
Cultivars	14	28	42	14	28	42	 14	28	42	
E1	26.1df	65.3kl	103.1kl	20.0bc	48.5jk	70.3fh	16.6be	39.7fh	46.8jl	
E3	29.8af	76.6fi	107.7hk	22.6ac	55.4ej	76.4dh	19.1ae	29.6i	50.5ik	
E4	33.5ac	83.8ad	122.4e	25.3b	62.2ag	82.5cg	21.5ac	47.1bf	54.1gj	
E5	31.7ae	79.2bh	176.8b	24.5ab	61.8ae	117.6b	19.9ae	43.9ch	84.9c	
E6	30.5af	76.3fi	136.5d	23.7ac	57.8ch	93.3c	19.3ae	42.0eh	62.8dg	
E8	29.3af	73.3hj	96.1 l m	23.0ac	53.8gk	68.9gh	18.6ae	40.1fh	40.8lm	
E9	32.2ad	80.7bh	105.5im	24.8ac	60.8bg	76.3dg	21.1ad	48.9be	56.2fi	
E10	30.5af	76.2fi	104.5jl	23.1ac	57.9ch	72.7dh	19.6ae	45.7bf	57.0ei	
M11	32.4ad	81.0bg	117.5eh	25.0ac	61.9af	86.0ce	21.1ad	50.0be	62.0dg	
M13	31.1af	77.6di	113.3ej	23.9ac	58.8bh	84.5cf	19.9ae	48.5be	62.0dg	
M14	30.5af	76.3fi	110.7gk	23.8ac	56.6di	72.5eg	18.8ae	44.4ch	62.6dg	
M15	31.3af	78.2ci	114.6ei	24.0ac	57.6ch	77.3dg	19.5ae	45.0cg	56.9ei	
M16	31.0af	77.5di	105.5im	24.2ac	62.8ae	76.0dg	20.8ad	47.3bf	57.0ei	
M18	29.7af	74.2gj	113.7ej	23.4ac	60.1bg	84.2cf	20.2ae	46.6bf	63.3df	
M19	32.5ad	79.6bh	166.6c	25.7ab	65.8ab	118.8b	21.4ac	49.6be	91.1bc	
M20	32.6ad	81.9bf	117.0eh	25.0ac	61.5af	84.1cf	20.3ae	53.4ab	65.2de	
M23	32.0ae	80.0bh	118.5eg	24.1ac	58.6bh	82.1cg	20.3ae	45.7be	51.2ik	
M26	27.0cf	67.5jl	117.5eh	20.0bc	51.8hk	79.0cg	16.5ce	37.5gh	53.3hk	
L37	35.7a	89.4a	113.4ej	26.7ab	64.9ac	78.0dg	22.3a	50.2bd	47.2j1	
L38	34.5ab	86.2ab	120.6ef	26.6ab	64.3ad	83.7cg	22.8a	58.2a	68.4d	
L42	33.7ac	84.4ad	121.6ef	26.0ab	65.0ac	87.5cd	22.3a	51.5ac	62.0dg	
L44	27.0cf	67.6j-l	171.2bc	21.6ac	53.8gk	128.0b	18.4ae	43.0dh	96.9b	
L46	25.0ef	62.51	93.0m	18.5c	47.2k	60.8h	15.2e	36.5h	37.8m	
L49	30.7af	76.9ei	116.0eh	23.4ac	59.3bg	79.8cg	20.0ae	45.5bf	58.4ei	
L50	31.5ae	79.1bh	112.4bh	24.3ac	60.2bg	78.0dg	20.5ae	46.3bf	60.3dh	
GLP-2	28.7bf	71.6ik	103.5kl	21.4ac	54.9fj	69.1gh	18.0ae	42.5dh	57.5ei	
GLP-24	32.3ad	80.6bh	108.9gk	25.3ac	61.1bg	76.3dg	20.9ad	47.1bf	52.6hk	
GLP-92	34.0ac	85.0ac	227.8a	27.2a	69.9a	161.7a	22.1ab	51.9ac	125.3a	
GLP-585	26.0df	65.0kl	105.3il	20.4ac	49.5ik	69.9fh	15.8de	37.0h	45.0km	
GLP-1004	24.4f	61.0l	228.6a	19.8bc	47.6k	171.9a	16.6be	39.4fh	130.5a	
Mean	30.6	76.4	125.7	23.6	58.3	88.2	19.7	45.1	64.0	
SE	2.04	2.2	2.9	2.1	2.2	4.3	1.6	2.4	2.6	
CV (%)	11.54	4.88	4.0	15.10	6.66	8.46	14.04	9.08	7.00	

Table 6: Biomass yields reduction of twenty bean cultivars at different DAE under medium and low watering levels

Cultivars	Medium (312	mm) DAE		Low (174 mm) DAE			
	14	28	42	14	28	42	
E1	23.5	25.7	31.8	36.32	39.2	54.6	
E4	24.6	25.8	29	35.91	43.8	51.87	
E6	22.26	24.28	31.67	36.85	44.99	53.97	
E10	24.23	24.04	30.47	35.82	40.01	45.61	
M11	22.99	23.62	26.74	34.97	38.28	47.21	
M13	23	24.29	28.68	36.01	37.5	45.35	
M16	22	18.99	28	33	38.99	46	
M18	21	18.99	26	32.09	37.19	44.3	
M19	20.95	17.3	25.4	35.98	37.72	45.3	
M23	24.69	26.7	30.7	36.69	42.9	56.8	
M26	25.89	23.3	32.8	38.81	44.4	54.7	
L38	22.89	25.4	30.6	33.91	32.5	43.3	
L44	20.12	20.34	25.24	31.99	36.43	43.4	
L46	26	24.5	34.6	39.4	41.6	60.5	
L49	24.01	22.8	31.2	34.81	40.79	49.6	
GLP-2	25.31	23.4	33.3	37.31	40.7	44.5	
GLP-24	21.61	24.2	29.9	35.19	41.36	51.7	
GLP-92	20	19	26.6	35	39	45	
GLP-585	21.42	23.88	33.59	39.08	43.05	57.25	
GLP-1004	19.02	22	24.8	31.89	35.49	42.9	
Mean	22.77	22.93	29.55	35.55	39.80	49.19	

There were varietal differences in percent leaf RWC of the three watering levels at different DAE for the field experiment (Table 4). At 43 DAE of the high water level, varieties GLP-1004, GLP-92, L38 and M18 had relatively higher mean absolute RWC values, while comparatively lower values were scored by the cultivars GLP-585, GLP-2, M26 and E4. Higher mean values were noted with cultivars GLP-1004, GLP-92, M19 and M18 at 43 DAE of the medium watering level. Among the cultivars that exhibited relatively higher RWC values at 43 DAE of the low water treatment were M19, M23, M26, GLP-92 and GLP-1004. Significant watering x variety interaction was observed at 15 DAE (p<0.05) and 29 DAE (p<0.01). It can be deduced from the above observations that the cultivars GLP-1004, GLP-92, M18 and M19 maintained fairly high relative water content under the three watering levels while genotypes E6, M26, GLP-2 and GLP-585 were among those with lowest values under similar conditions.

Plant biomass was significantly (p<0.01) reduced by water stress at 28 and 42 DAE in all cultivars (Table 5). Significant watering x variety interaction was observed at 14 DAE (p<0.05) and at 28 and 42 DAE (p<0.01). The mean across the genotypes reveals that biomass accumulation was lower at low irrigation level followed by medium level at similar number of days after emergence. Though there was a general increase in biomass accumulation with the number of days after emergence, the increase was affected by the irrigation level, being highest in the high irrigation level and lowest in the low level. At 42 DAE, genotypes GLP-1004, GLP-92, L44, M19 and E6 had significantly more biomass under all the watering levels. GLP-1004 had the

highest biomass followed by GLP-92 for all watering levels at 42 DAE. Varieties E1, L46 and GLP-2 were among those with the least values at this time under the three watering levels. The mean percent biomass reduction in the medium water treatment was lower than that of the low treatment at all DAE (Table 6). The reduction tended to increase with the number of days after emergence under both treatments, but the reduction was marginally small between 14 and 28 DAE under medium water treatment, being 0.16%. The margin under low water treatment between these two samplings was comparatively large. Greatest mean yield reduction of 49.19% was observed at 42 DAE under low watering level. At 42 DAE in medium water treatment, cultivars L46, GLP-585, GLP-2, M26, E1 and E6 suffered the greatest biomass yield reductions. However, least biomass reduction was observed in GLP-1004, L44, M19, M11, GLP-92 and M18. Depression of biomass yield at 42 DAE in low water treatment was more pronounced. Genotypes L46, GLP-585, M23, E1, E6 and M26 experienced the greatest reductions. Of the cultivars that suffered least biomass reductions at 42 DAE of medium water treatment, GLP-1004, L44, M19 and M18 still had relatively least percent reductions at 42 DAE in the low water treatment. Days to permanent wilting correlated negatively and significantly with percent biomass reduction at 42 DAE, low watering level (r = -0.63*) of the field experiment. However, SMC and RWC at wilting respectively associated positively (r = 0.58*) and negatively (r =-0.53*) with percent biomass reduction at similar DAE of low watering level.

DISCUSSION

The bean genotypes tested in this study showed variability in phenology and physiological responses to water deficits. Water stress decreased leaf relative water content and biomass in all the tested genotypes. Similar observations have been reported in blackgram, greengram, cowpeas and soybeans^[21], in common bean^[3] and in sugar beet^[4]. Variety differences were observed in leaf RWC in the glasshouse and in the field. Varieties M18, L44, GLP-24, M19, GLP-1004 and GLP-92 had consistently higher RWC values in the glasshouse and in the medium and low watering levels in the field. This observation indicated that these genotypes had a better plant water status and hence better yields as compared with those that had generally lower scores for the traits under similar conditions.

This genotypic variation in RWC may be attributed to differences in the ability of the varieties to absorb more water from the soil and/or the ability to control water loss through the stomates. These findings are in agreement with those reported by Sinclair and Ludlow^[22]. It may also be due to differences in the ability of the tested varieties to accumulate and adjust osmotically to maintain tissue turgor and hence physiological activity^[9]. Varietal differences in RWC may also be a result of varieties maximizing on soil water reserves by fully extracting water in the existing rooting zone and/or extending rooting depth to increase water reserve for the crop^[23]. Other researchers have investigated the role of increased root length and density in maintenance of guard cell turgidity under water stress. Lorens et al.[24] attributed the cultivar differences in net photosynthetic rates to the differential abilities for the better-adapted cultivars to root deeper maintain higher plant water potential. and Runkulatile et al.[3] found that drought resistant varieties (Ulonzo, GLP-1004 and White Haricot) exhibited faster downward root penetration than the less drought resistant varieties. In the present study, the findings concur as drought resistant genotypes such as GLP-1004, GLP-92, M18 and M19 showed higher RWC and stomatal conductance apart from taking longer periods to wilt and at low pot soil moisture content at wilting under glasshouse compared to the drought susceptible genotypes such as E6, E10, M11, M26 and L37. These genotypes showed similar trends in the field under the medium and low watering levels. The drought resistant genotypes consistently maintain higher RWC and suffered least biomass reductions as compared to their drought susceptible counterparts.

Osmotic adjustment would also explain the maintenance of high RWC values of these cultivars in the

glasshouse, despite the low soil moisture contents at wilting. Martin *et al.*^[25] proposed the mechanism for drought resistant barley cultivars that maintained high RWC under water stress. Similarly, stomatal conductance may be a useful indicator in discriminating drought resistant and non-resistant cultivars in the glasshouse. However, water stress develops quickly in the glasshouse due to limited soil volume in the pots. Therefore, plants may not have time to adjust their stomata as in the field where response to water stress may allow for osmoregulation, which causes a relatively higher RWC and stomatal conductance. Water stress depends on variety and to a greater extent on the length of its exposure, plus that of temperature in the growing conditions^[26].

The delay in dehydration exhibited by the cultivars GLP-1004, GLP-92, GLP-24, M18, M19, L44 and L50 may have important implications in terms of crop productivity. These varieties had comparatively higher RWC values under stress conditions, tended to accumulate higher biomass yields and suffered least depressions in biomass under all levels of water stress. Days to permanent wilting for these genotypes were longer than their counterparts such as E6, E10, M11, M14 and M15 but the soil water content at their permanent wilting was comparatively lower. Since cultivars such as GLP-1004, GLP-92, GLP-24, M16, M18, M19 and L44 had comparatively higher RWC in the glasshouse treatment, it may possibly be concluded that they maintain a relatively higher water potential gradient between the roots and the transpiring leaves. This discrepancy could be due to the differences in leaf solute potentials since the evaporative demand and the initial soil water content was similar for all the varieties. Differences in root length density may also have played part, but this factor is not likely to be important in influencing water uptake by potted plants whose roots ramify extensively into the potting medium. However, it could come into play in the field grown plants as has been reported by Runkulatile et al.[3]. Morgan and Condon[27] and Santamaria et al.[17] also showed that genotypes of wheat and sorghum with high osmotic adjustment produced more root biomass and greater root length than genotypes with low osmotic adjustment. The results of this study therefore may suggest that osmotic adjustment principally enabled the cultivars GLP-1004, M18, M16, M19 and GLP-24 to exploit moisture from relatively dry soils and thus taking comparatively longer periods to wilt.

Differential effect of watering level on biomass accumulation was observed at all days after emergence. But due to inherent genetic differences in biomass accumulation among genotypes, percent decrease in biomass accumulation would provide a better indicator of

drought resistance than differences in total biomass per se. Cultivars such as GLP-1004, L44, M18 and M19 were least affected under both low and medium watering levels compared to GLP-585, E1, E6, or M23. This disparity in their response could also serve as evidence that the former genotypes have higher ability to resist drought. Their higher RWC in the glasshouse and in the field could support their ability to osmoregulate and thus maintained growth under water-limited conditions. Similarly, studies on wheat^[27] and sorghum^[17] have indicated that osmotically non-adjusting varieties are more significantly affected by water stress than the osmotically adjusting.

The combination of both morphological and physiological attributes of drought resistance with high yield potential forms the selection basis for better yield performance under drought conditions. Although common beans are prone to short water stress periods, differences in drought resistance have been observed in this study. There was evidence that the genotypes that showed superior performance in the glasshouse also suffered fairly low biomass reduction under the medium and low watering levels in the field. Strong correlations occurred between traits such as days to permanent wilting, leaf relative water content at wilting and soil moisture content at wilting as well as stomatal conductance at wilting of the glasshouse and biomass reduction in the field. These suggest the possibility of screening the genotypes in the glasshouse and also the likelihood of selecting the cultivars on the basis of their leaf RWC. Therefore, cultivars M16, M18, M19, L44, GLP-24, GLP-92 and GLP-1004 seem to be most adapted to the dry-land conditions among the tested genotypes.

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