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EXPLOITATION OF GENETIC POTENTIAL OF SWEETPOTATO FOR END-USER TRAITS IMPROVEMENT

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ABSTRACT

Sweetpotato (*Ipomoea batatas* Lam) is a staple food globally, but it has remained underutilised resource in Ghana due to lack of consumer preferred cultivars. There is the need to develop staple-type sweetpotato cultivars which are preferred by consumers to increase sweetpotato use as a food security, health and industrial crop commodity in Ghana. This study was conducted to evaluate the breeding potential of sweetpotato germplasm for the development of farmer and consumer preferred varieties in Ghana. A total of 115 sweetpotato accessions were evaluated for genetic variability. Significant ($P < 0.01$) differences were observed indicating genetic diversity. $G \times E$ was significant for all traits, except dry matter, sucrose, total sugar, and starch content. Phenotypic Coefficient of Variation (PCV) ranged from 4.78% for starch content to 63.40% for marketable root weight. Genotypic Coefficient of Variation (GCV) ranged from 4.07% for starch content to 55.35% for marketable root weight. Broad-sense heritability estimates varied from medium (0.61) to high (0.90) for all the traits, except for sucrose content. Predicted improvement over the means is 10 up to 105% for all traits, except starch (7.13%). This indicates sufficient useful genetic variation prospect which could be used to provide substantial improvement through selection of superior genotypes. The strong positive genetic association between dry matter and starch ($r = 0.71$), and strong negative relationship for sugar and dry matter ($r = -0.77$) and starch content ($r = -0.99$) indicates the possibility of developing non-sweet high dry matter sweetpotatoes, which are the preferred varieties in Ghana.

Key Words: Beta-carotene, genotypic coefficient of variation, heritability, *Ipomoea batatas*, non-sweet sweetpotato, phenotypic coefficient of variation

RÉSUMÉ

La patate douce (*Ipomoea batatas* Lam) est un aliment de base à l'échelle mondiale, mais c'est une ressource sous-utilisée au Ghana en raison du manque de cultivars préféré par les consommateurs. Il est nécessaire de développer des cultivars de patate douce de type de base qui sont préférés par les consommateurs pour augmenter l'utilisation de la patate douce comme la culture de la sécurité alimentaire, la culture de santé et le produit industriel au Ghana. Cette étude a été menée pour évaluer le potentiel du matériel génétique du germeplasma de la patate douce pour le développement des variétés préférées par les agriculteurs et les consommateurs au Ghana. Un total de 115 germeplasmes de patate douce ont été évalués pour la variabilité génétique. Des différences significatives de ($P < 0,01$) ont été observées indiquant la diversité génétique. $G \times E$ était important pour tous les traits, à l'exception de matière sèche, le saccharose, le sucre total et la teneur en amidon. Coefficient Phénotypique de variation (CPV) variait de 4,78% pour la teneur en amidon à 63,40% pour le poids de racine commercialisables. Coefficient génotypique de variation (CGV) variait de 4,07% pour la teneur en amidon à 55,35% pour le poids de racine

commercialisables. Coefficient génotypique de variation (CGV) variait de 7,60% pour la teneur en matière sèche à 55,35% pour le poids de racine commercialisables. Estimations de l'héritabilité au sens large variait de (0,61) moyen à (0,90) haut pour tous les traits, sauf pour la teneur en saccharose. L'amélioration prédite sur le moyen est de 10 à 105% pour tous les traits, à l'exception de l'amidon (7,13%). Cela indique suffisamment la perspective de la variation génétique utile qui pourrait être utilisée pour fournir une amélioration substantielle par la sélection de génotypes supérieurs. Une forte relation génétique positive entre la matière sèche et l'amidon ($r = 0,71$), et une forte relation négative pour le sucre et la matière sèche ($r = -0,77$) et la teneur en amidon ($r = -0,99$) indiquent la possibilité de développer la patate douce qui est non sucrée, haute de matière sèche et qui sont les variétés préférées au Ghana.

Mots Clés: Bêta-carotène, coefficient génotypique de variation, héritabilité, *Ipomoea batatas*, patate douce non-sucré, coefficient phénotypique de variation

INTRODUCTION

Sweetpotato (*Ipomoea batatas* Lam) is a dicotyledonous plant of the botanical family Convolvulaceae (Watson and Dallwitz, 2000). It is one of the most important root crops in the world (Waramboi *et al.*, 2011). Sweetpotato is a staple food for millions of people and the seventh most abundant crop globally (Bouville-Benjamin, 2007; Devi *et al.*, 2014). The potential of sweetpotato in food security and global well-being has been well recognised (Van Hal, 2000; Lebot, 2010).

In spite of its great potential to alleviate food insecurity, malnutrition and poverty, it has remained an underutilised resource in Ghana. For instance, it is uncommon to find sweetpotato served in public places such as local restaurants, canteens and schools. Locally available varieties have very sweet taste (Missah and Kissiedu, 1994; Baafi *et al.*, 2015), and recently introduced orange-fleshed genotypes which possess the precursor to combat vitamin A deficiency at relatively cheaper cost have low dry matter content, limiting their consumption as a staple food. There is a need to develop staple-type high beta-carotene and high dry matter content varieties to increase sweetpotato use as a food security, health and industrial crop commodity in Ghana.

Screening populations with high frequencies of favourable alleles can provide the best parents for breeding (Gasura *et al.*, 2008). Knowledge on the genetic potential of the available germplasm is essential to find source(s) of variation for the end-user traits improvement. Information on the nature and magnitude of variability and

heritability in a population owing to genetic and non-genetic factors is one of the prerequisites in any breeding programme (Kumar *et al.*, 1985). There is knowledge gap on the genetic variability and breeding potential of sweetpotato for the end-user traits improvement in Ghana. The objective of this study was to assess the breeding potential of sweetpotato accessions in Ghana, for the development of non-sweet, high dry matter and high beta-carotene varieties through combination of desired traits into a common genetic background.

MATERIALS AND METHODS

This research was carried out at Fumesua (Forest ecozone) and Pokuase (Coastal Savannah ecozone) in the major and minor cropping seasons in 2011. The sweetpotato accessions used in the study are presented in Table 1. The planting arrangement was one row per ridge, with a distance of 1 m between ridges. The length of a ridge was 3.6 m and within row planting space was 0.3 m, giving a total of 12 plants per ridge. Randomised complete block design (RCBD) was used.

Data collection. Harvesting was done at three and half months after planting. The 10 central plants per row were harvested and were used for storage root yield assessment. Storage root yield data taken were total root yield and marketable root yield. In addition, harvest index was recorded as the ratio of the total root yield to total biomass. Thereafter, one large, one medium and one small storage root, were randomly selected for determination of the physico-chemical traits.

TABLE 1. Description of 115 sweetpotato accessions collected and evaluated in the study

Local accessions	Local improved varieties	Accessions from NARS		Accessions from CIP
CRIWAC 01-10	SANTOMPONA	TAG 03-019	B-REGARD	CIP 442903
CRIWAC 02-10	FAARA	NS 001	FIASO RED	CIP 442291
CRIWAC 03-10	TEKSANTOM	OK 03-015	TAG 03-030	CIP 440069
CRIWAC 04-10	OGYEFO	DOS 03-021	GWERI	CIP 440390
CRIWAC 05-10	OKUMKOM	CARROT C	BD 96-029	CIP 442462
CRIWAC 06-10	OTOO	HUMBERCHERO	FREMA	CIP 442776
CRIWAC 07-10	HISTARCH	B/FASO 002	DOS 03-006	CIP 440062
CRIWAC 08-10	SAUTI	FA10-026	NS 003	CIP 442589
CRIWAC 09-10	APOMUDEN	RESISTO	AAT 03-004	CIP 442145
CRIWAC 10-10		CEMSA	OK 03-021	CIP 442147
CRIWAC 11-10		199062.1	BOT 03-030	CIP 440095
CRIWAC 12-10		OK 03-014	OK 03-017	CIP 441771
CRIWAC 13-10		JONATHAN	KAYIA WHITE	CIP 442901
CRIWAC 14-10		H-ASIATOR	UKEREWE	CIP 443016
CRIWAC 15-10		TANZANIA	OK 03-018	CIP 440071
CRIWAC 16-10		NINGSHU 1		CIP 442896
CRIWAC 17-10		SPK 004		CIP 442162
CRIWAC 18-10		MOHC		CIP 442775
CRIWAC 19-10		BOT 03-028		CIP 443027
CRIWAC 20-10		BOT 03-020		CIP 443129
CRIWAC 21-10		J-ORANGE		CIP 442264
CRIWAC 22-10		BOT 03-027		CIP 442654
CRIWAC 23-10		ADA 001		CIP 443035
CRIWAC 24-10		DOS 03-017		CIP 442913
CRIWAC 25-10		NAV 001		CIP 442237
CRIWAC 26-10		KEMB 37		CIP 443019
CRIWAC 27-10		AAT 03-025		CIP 442850
CRIWAC 28-10		B/FASO 001		
CRIWAC 29-10		ZAMBEZI		
CRIWAC 30-10		BOT 03-020		
CRIWAC 31-10		NASPOT 1		
CRIWAC 32-10		AAT 03-017		

Storage roots selected were approximately 3 cm or more in diameter and without cracks, insect damage or rotten parts (Ekanayake *et al.*, 1990). They were washed, peeled and cut into four equal parts longitudinally. Two opposite quarters of the peeled storage roots were sliced into pieces and 50g fresh sample weighed into a polythene envelope. The fresh samples were frozen using deep-freezer after which it was freeze-dried for 72 hours using freeze-dryer. The dry weights of the freeze-dried samples were recorded. The freeze-dried samples were milled and used for the determination of all the physico-chemical traits except dry matter content using the near-infrared reflectance spectroscopy (NIRS) (Tumwegamire

et al., 2011). The physico-chemical traits measured were beta-carotene, dry matter, fructose, glucose, sucrose, total sugar, starch, protein, iron and zinc content. Dry matter content was calculated after freeze drying as the ratio of the weight of the dry sample expressed as a percentage of the weight of the fresh sample. The laboratory work was carried out at the International Potato Centre (CIP) Fumesua, Ghana and Lima, Peru, respectively.

Data analysis. Data for 102 out of the 115 accessions were analysed because of lack of plants for some accessions. The analysis also excluded minor cropping season data for Pokuase

because the experiment failed due to erratic rainfall. The data were subjected to Analysis of Variance (ANOVA) using GenStat statistical package (Genstat, 2007). To test the efficiency of RCBD for the analysis, relative efficiency (RE) of an Alpha Lattice design over RCBD was determined. The RE was determined as the ratio of the error means square of RCBD to that of the Alpha Lattice and the RE was not significant. RE is significant if the ratio is >1 and vice versa. Hence, RCBD was used to analyse the data employing the method of Steel and Torrie (1980).

Variance components were used to determine genotypic variance (GV) and phenotypic variance (PV) as per Prasad *et al.* (1981) as follows:

Genotype x Location interaction variance

$$(\sigma_{GL}^2) = (MS_{GL} - MS_E)/r$$

$$\text{Genotypic variance } (\sigma_G^2) = (MS_G - MS_{GL})/rl =$$

$$(\sigma_E^2 + r\sigma_{GL}^2 + rl\sigma_G^2 - \sigma_E^2 + r\sigma_{GL}^2)/rl$$

$$\text{Phenotypic variance } (\sigma_p^2) = \sigma_G^2 + \sigma_{GL}^2/l + \sigma_E^2/rl =$$

$$\sigma_G^2 + ((MS_{GL} - MS_E)/r)/l + \sigma_E^2/rl$$

$$\text{Error variance } (\sigma_E^2) = MS_E$$

Where: MS = Means Square for a particular source of variation;

$$\sigma_G^2 = \text{Genotypic variance};$$

$$\sigma_{GL}^2 = \text{Genotype by location interaction variance};$$

$$\sigma_E^2 = \text{error variance};$$

$$r = \text{number of replications/location};$$

$$l = \text{number of locations or environments}$$

Variance components were used to compute broad-sense heritability (h_b^2), Genotypic Coefficient of Variation (GCV), Phenotypic Coefficient of Variation (PCV) and Genetic

Advance (GA) (Burton, 1952; Johnson *et al.*, 1955; Kumar *et al.*, 1985), as presented below:

$$\text{Heritability in broad - sense } (H_b^2) = \sigma_G^2/\sigma_p^2 =$$

$$\sigma_G^2/(\sigma_G^2 + \sigma_{GL}^2/l + \sigma_E^2/rl)$$

$$\text{Genotypic Coefficient of Variation} = (\sigma_G \times 100)/X$$

$$\text{Phenotypic Coefficient of Variation} = (\sigma_p \times 100)/X$$

Where: X is the grand mean of the trait.

$$\text{Genetic Advance (R)} = H_b^2 \cdot k \cdot \sigma_p$$

Where: k (selection differential expressed in phenotypic standard deviation at 5%) = 2.06.

The genetic advance from selection expressed as percent of grand mean was obtained as Genetic Advance ($R_{(X)}$) = $H_b \cdot k \cdot GCV$.

This expression shows that the expected genetic advance from selection when expressed as a percent of the mean is the product of the selection differential measured in terms of the phenotypic standard deviation, the genetic coefficient of variation, and the square root of the heritability ratio (Johnson *et al.*, 1955). Estimates of the genotypic correlation coefficients were computed according to Miller *et al.* (1958) and (IRRI, 2006) as presented below:

$$\text{Genotypic correlation Coefficient } r_{G(1,2)} =$$

$$\sigma_{1,2}/\sqrt{(\sigma_{G1}^2)(\sigma_{G2}^2)}$$

Where $\sigma_{G(1,2)}$ is the genetic covariance between two traits, σ_{G1}^2 is the genetic variance of the first trait and σ_{G2}^2 is the genetic variance of the second trait.

RESULTS

Genotype by environment interaction (G x E) was significant ($P < 0.05$) for all the traits except dry matter, sucrose, total sugar and starch content

(Table 2). Highly significant differences ($P < 0.01$) were found between the sweetpotato accessions for all the traits except sucrose content which was significant at $P < 0.05$. The range of values for the traits, their grand mean, coefficient of variation (CV), and standard error of mean (SE) are presented in Table 3.

Generally, values for $G \times E$ variance were lower than those of the genotypic variance and the error variance (Table 4). Similarly, the values for the genotypic variance were lower than those of the error variance for all the traits except beta-carotene, fructose, glucose and iron content as well as harvest index, root weight and marketable root weight.

The phenotypic variance was slightly higher than the genotypic variance, except for dry matter and iron content (Table 5). The lowest and the highest values for the genotypic variance were obtained for dry matter (0.001) and beta-carotene content (21.235). A similar trend was found for the phenotypic variance, with values of dry matter and beta-carotene as 0.001 and 23.488, respectively.

The PCV values were higher than GCV values (Table 6). The PCV values ranged from 4.78% for starch content to 63.40% for marketable root weight. The GCV ranged from 4.07% for starch content to 55.35% for marketable root weight.

Broad-sense heritability ranged from 0.30 for sucrose content to 0.90 for beta-carotene content (Table 6). The expected gain from selection and genetic advance (as percent of grand mean) ranged from 0.05% (dry matter content) to 9.03 mg 100g⁻¹DW (beta-carotene content), and 7.13% (starch content) to 105.34% (fructose content), respectively (Table 6).

Dry matter content showed positive and strong association with starch content ($r = 0.71$), but its relationship was strong and negative with beta-carotene ($r = -1.20$), fructose ($r = -0.76$), glucose ($r = -0.78$), total sugar ($r = -0.77$) and harvest index ($r = -0.69$) (Table 7). Beta-carotene content had strong and negative association with starch content ($r = -0.61$), but its association with glucose ($r = 0.62$), sucrose ($r = 0.74$), total sugar ($r = 0.56$), iron ($r = 0.74$) and zinc content ($r = 0.58$) were strong and positive. Fructose and glucose showed strong and negative correlation with starch content but, strong and positive correlation

TABLE 2. Mean squares for the agronomic and physico-chemical traits of the 102 sweetpotato accessions

Source of variation	Df	Beta-carotene	Dry matter	Fructose	Glucose	Sucrose	Total sugar	Starch	Protein	Fe	Zn	HI	Root weight	Marketable root weight
Block	1	0.83	0.032	62.55	116.60	352.64	0.84	296.97	2.71	0.64	0.01	0.054	2.41	3.08
Env.	2	721.91**	0.067**	6.53**	7.36*	66.88*	119.73**	1356.31**	46.32**	2.83**	0.51**	1.358**	74.82**	76.62**
Gen.	101	140.93**	0.008**	8.78**	15.04**	29.35*	75.75**	64.52**	2.91**	0.16**	0.06**	0.146**	12.14**	9.13**
GXE	202	13.52**	0.003 ^{ns}	1.32*	2.16*	20.48 ^{ns}	23.02 ^{ns}	17.80 ^{ns}	1.06*	0.07**	0.02**	0.021**	2.11**	1.65**
Error	299	7.12	0.003	1.06	1.64	20.32	21.38	14.79	0.83	0.05	0.01	0.013	1.12	0.95

*Significant at 5% ($p < 0.05$) level of probability, ** Significant at 1% (0.01) level of probability, ^{ns} = not significant

TABLE 3. Descriptive statistics for the agronomic and physico-chemical traits of the 102 sweetpotato accessions

Trait	Range	Grand mean	CV (%)	SE
Dry matter	27.00–50.00	38.00	14.1	5.00
Beta-carotene	3.72–33.67	11.44	23.3	2.67
Fructose	0.63–8.05	2.01	51.2	1.03
Glucose	1.74–11.66	3.97	32.3	1.28
Sucrose	6.77–17.15	10.19	44.2	4.51
Total sugars	9.83–30.34	16.46	28.1	4.62
Starch	54.72–75.64	68.56	5.6	3.85
Protein	2.68–6.62	4.47	20.4	0.91
Iron	1.53–2.42	1.99	11.1	0.22
Zinc	0.76–1.28	1.01	11.9	0.12
Harvest index	0.08–0.68	0.40	28.0	0.11
Root weight	0.34–6.43	2.58	41.0	1.06
Marketable root weight	0.10–4.45	2.02	48.3	0.98

TABLE 4. Components of variance for agronomic and physico-chemical traits of the 102 sweetpotato accession

Trait	σ^2_G	$\sigma^2_{G \times E}$	σ^2_E	Ratio $\sigma^2_G : \sigma^2_{G \times E} : \sigma^2_E$
Dry matter	0.001	0.00	0.003	1 : 0.0 : 3.6
Beta-carotene	21.235	3.20	7.120	1 : 0.15 : 0.34
Fructose	1.243	0.13	1.060	1 : 0.10 : 0.85
Glucose	2.147	0.26	1.640	1 : 0.12 : 0.76
Sucrose	1.478	0.08	20.320	1 : 0.05 : 13.75
Total sugars	8.788	0.82	21.380	1 : 0.09 : 2.43
Starch	7.787	1.51	14.790	1 : 0.19 : 1.90
Protein	0.308	0.12	0.830	1 : 0.37 : 2.69
Iron	1.243	0.01	0.050	1 : 0.55 : 0.04
Zinc	0.007	0.01	0.010	1 : 0.75 : 1.43
Harvest index	0.022	0.01	0.010	1 : 0.23 : 0.46
Root weight	1.672	0.50	1.120	1 : 0.30 : 0.67
Marketable root weight	1.247	0.35	0.950	1 : 0.28 : 0.76

with total sugar content. Similarly, negative but very strong relationship existed between total sugar and starch content ($r = -0.99$). Harvest index had strong and positive relationship with the yield traits (Table 7).

DISCUSSION

Significant genetic diversity was observed among the accessions (Table 2) indicating the possibility of developing non-sweet high dry matter and high beta-carotene varieties simultaneously, using this population. G x E

interaction is important in evaluating genotype adaptation, selection of parents and developing genotypes with improved end-product quality (Ames *et al.*, 1999; Zhou *et al.*, 2012). The significant G x E interactions observed may complicate selection for all the traits studied, except dry matter, sucrose, total sugar and starch content. G x E complicates selection by confounding the determination of true genetic values since relative performance of genotypes vary across environments. The existence of G x E indicates that selection should be carried out in a range of environments (Falconer and Mackay,

TABLE 5. Estimate of genotypic and phenotypic variance for the agronomic and physico-chemical traits of the 102 sweetpotato accessions

Trait	Genotypic variation	Phenotypic variation	Ratio $\sigma_G^2 : \sigma_P^2$
Dry matter	0.001	0.001	1 : 1.0
Beta-carotene	21.235	23.488	1 : 1.1
Fructose	1.243	1.463	1 : 1.2
Glucose	2.147	2.507	1 : 1.2
Sucrose	1.478	4.892	1 : 3.3
Total sugars	8.788	12.625	1 : 1.4
Starch	7.787	10.753	1 : 1.4
Protein	0.308	0.485	1 : 1.6
Iron	1.243	0.030	1 : 0.02
Zinc	0.007	0.010	1 : 1.5
Harvest index	0.022	0.025	1 : 1.1
Root weight	1.672	2.023	1 : 1.2
Marketable root weight	1.247	1.522	1 : 1.2

TABLE 6. Genotypic and phenotypic coefficient of variation, heritability and expected genetic advance for the traits

Trait	Genotypic coefficient of variation	Phenotypic coefficient of variation	Heritability (H_p^2)	Expected selection gain (R)	Expected selection gain (R % of mean)
Dry matter	7.60	9.61	0.63	0.05	12.37
Beta-carotene	40.28	42.36	0.90	9.03	78.90
Fructose	55.48	60.18	0.85	2.12	105.34
Glucose	36.91	39.88	0.86	2.79	70.35
Sucrose	11.93	21.70	0.30	1.38	13.51
Total sugars	18.01	21.59	0.70	5.10	30.95
Starch	4.07	4.78	0.72	4.89	7.13
Protein	12.46	15.58	0.64	0.91	20.40
Iron	6.80	8.70	0.61	0.22	10.96
Zinc	8.08	9.90	0.65	0.14	13.60
Harvest index	36.80	39.53	0.84	0.27	68.40
Root weight	50.13	55.13	0.83	2.42	93.83
Marketable root weight	55.35	63.40	0.76	2.01	99.40

1996). This is because progress from selection is realised only when genotypic effects can be separated from environmental effects (Miller *et al.*, 1958). Beta-carotene content may be an exception because the orange-fleshed colour associated with it makes it easy to select.

Differences in genotypic and phenotypic variances (Table 5) may be attributed to environmental effects. It is essential that germplasm testing procedures are designed to

maximise the genetic effects relative to the environmental and interaction effects. Information on the relative magnitudes of the different sources of variance observed in this study (Table 4) provides a guide towards maximising the genetic effects relative to the environmental and interaction effects. The magnitudes of G x E interaction variances were smaller than the error variances (Table 4), implying that the accessions were tested in

TABLE 7. Genotypic correlation coefficient for 10 physico-chemical and three agronomic traits of 102 sweetpotato accessions

Trait	DM	BC	F	G	S	TS	ST	P	Fe	Zn	HI	RWT	MKTRWT
DM													
BC	-1.20												
F	-0.76 0.30												
G	-0.78 0.62 0.98												
S	-0.23 0.74 0.04 0.05												
TS	-0.77 0.56 0.85 0.91 1.09												
ST	0.71 -0.61 -0.61 -0.78 -0.71 -0.99												
P	0.08 0.15 -0.24 -0.19 0.75 0.15 0.21												
Fe	-0.11 0.74 0.02 0.11 1.07 0.60 -0.71 0.75												
Zn	0.18 0.58 -0.22 -0.10 1.15 0.39 0.39 0.75 0.90												
HI	-0.69 -0.41 0.30 0.30 -0.14 0.22 0.57 0.27 -0.08 0.00												
RWT	-0.12 -0.41 0.06 0.06 -0.42 -0.12 -0.19 -0.48 -0.54 -0.51 0.56 1.00												

DM = Dry matter, BC = Beta-carotene, F = Fructose, G = Glucose, S = Sucrose, TS = Total sugar, ST = Starch, P = Protein, Fe = Iron, Zn = Zinc, HI = Harvest index, RWT = Root weight, MKTRWT = Marketable root weight

adequate sample environments. However, a large number of replicates may be required for traits with lower genetic variance than the error variance (root dry matter, sucrose, total sugar, starch, protein and zinc content) to enhance the optimisation of their genetic effects.

The trends for the PVC and the GVC values (Table 6) were in agreement with the results reported for *Solanum anguivi* (Denton and Nwangburuka, 2011), *Corchorus olitorius* (Nwangburuka and Denton, 2012), *Triticale* (Kumar *et al.*, 1985) and oat (Prasad *et al.*, 1981). The observed trend between PVC and GVC could be attributed to environmental effects (Denton and Nwangburuka, 2011). This is because variance of phenotype is due to genotypic and environmental factors combined. High PCV and GCV values which were detected for marketable root weight, root weight, fructose content, beta-carotene content, glucose content, harvest index, and total sugar content suggest that these traits accounted for the highest variation observed in the sweetpotato accessions. This means that the sweetpotato accessions showed high diversity for these traits which could lead to higher selection precision for superior accessions for the traits.

GCV provides a measure for comparing the genetic variability present in various quantitative traits. However, it is not possible to estimate heritable variation with the help of the GCV alone (Prasad *et al.*, 1981). The GCV together with heritability estimates would give the best picture of the amount of advance to be expected from selection (Burton, 1952). Heritability indicates the effectiveness with which selection of genotypes can be based on phenotypic performance (Johnson *et al.*, 1955; Nwangburuka *et al.*, 2012; Mwije *et al.*, 2014). In this study, broad-sense heritability estimates varied from medium to high, with the exception of sucrose content which had low heritability (Table 6).

Traits with medium to high heritability are influenced by additive gene effects (Denton and Nwangburuka, 2011). This suggests that selection based on phenotype will be effective. However, the heritability value in itself does not provide an indication of the amount of genetic progress that would result from selecting the best individuals (Johnson *et al.*, 1955). The utility of estimates of

heritability is increased when they are used in conjunction with the selection differential (the amount that the mean of the selected lines exceeds the mean of the entire population), since genetic advance is commonly predicted as the product of the heritability and the selection differential.

Prediction of the response of an individual to selection is more reliable when GCV, broad-sense heritability and genetic advance are combined (Ghandi *et al.*, 1964; Ibrahim and Hussein, 2006). The predicted improvement over the means for this population is above 10% and up to 105% for all the traits, except starch content (7.13%). Miller *et al.* (1958), observed 13 – 15% response to selection for lint yield of upland cotton, and noted that it was particularly encouraging. This suggests that sufficient useful genetic variation is present in this population that could be used to provide substantial improvement through the selection of superior accessions. The type of gene action operating is also critical since the expected amount of superiority will be realised in subsequent generations only if all of the genetic effects are additive (Miller *et al.*, 1958; Falconer and Mackay, 1996). Conversely, non-additive effects (epistasis, dominance, and interactions) may decrease the amount of genotypic superiority passed on. It is, therefore, important to seek additional information concerning the nature of gene action in order to buttress this point in sweetpotato.

Genetic relationships between traits (Table 7) may result from pleiotropic gene effects, linkage of two genes, linkage disequilibrium and epistatic effects of different genes or environmental influences (Falconer and Mackay, 1996). Except for the very high (>1) values observed for dry matter and beta-carotene content (-1.2), sucrose and total sugar content (1.09), sucrose and iron content (1.07), and sucrose and zinc content (1.15), the values for the genetic correlation coefficient were in agreement with those reported by Grüneberg *et al.* (2009). Denton and Nwangburuka (2011) also reported genotypic correlation coefficients greater than one in a study on *Solanum anguivi*. A similar observation was made by Nwangburuka and Denton (2012) for character association on leaf *Corchorous olitorius*.

The correlations observed are thus population specific because the inter-relationships might be quite different in populations in which different gene associations exist in the parental lines (Miller *et al.*, 1958). The differences observed could also be attributed to the variations in the mean values of the characters under study which are at levels different from the other studies. The strong positive association observed between dry matter content and starch content, and the strong negative relationship found for sugar content and dry matter and starch content indicates that it is possible to develop non-sweet high dry matter sweetpotato varieties which are the preferred sweetpotatoes for most Ghanaians (Baafi *et al.*, 2015; Baafi *et al.*, 2016). A similar observation was made by Grüneberg *et al.* (2009), who reported that development of non-sweet sweetpotato varieties should not be too difficult. However, developing non-sweet, high dry matter and high beta-carotene sweetpotato varieties could be challenging due to the strong negative association between dry matter content and beta-carotene content, and the positive association between beta-carotene content and sugar content. Breeding for such cultivars may require many cycles of selection and hybridisation to break genetic linkages associated with the traits. However, beta-carotene is controlled by a limited number of genes (Oduro, 2013; Baafi, 2014) and should be easy to manipulate. The results also showed that dry matter content, sugar content and most of the physico-chemical traits (except protein) may indirectly be selected using beta-carotene content (the orange-fleshed colour).

CONCLUSION

Significant genetic diversity was established between the accessions studied for dry matter, beta-carotene and sugar content. The relative magnitude of the G x E variance to the other variances indicates high potential for progress from selection since the superior accessions in the population can be readily identified. This means that sufficient useful genetic variation is present in the population which may be exploited to provide for substantial amount of improvement through selection of superior accessions. High

heritability coupled with the high gains expected indicate high breeding value and mostly additive genetic effects. This suggests that selection can be effective for these traits. The strong negative association between dry matter content and sugar content indicates that it is feasible to develop non-sweet high dry matter sweetpotato cultivars. However, developing non-sweet, high dry matter varieties with a high beta-carotene content may require many cycles of selection.

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