

Yield stability analysis of *Ipomoea batatas* L. cultivars in diverse environments

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Abstract

Sweet potato is an important food crop in tropical and sub-tropical regions of the world. There is limited published research on yield stability of sweet potato in tropical environments. To identify cultivars with improved agronomic and stable yield characteristics, five elite genotypes obtained from the sweet potato breeding program in Uganda and International Potato Center (CIP) and five land race genotypes were evaluated for yield stability at 12 environments. The Additive Main Effects and Multiplicative Interaction (AMMI) model was used for stability analysis. The analysis of variance of yield data ($t\ ha^{-1}$) for genotypes x locations, genotypes x seasons and locations x seasons was highly significant ($P < 0.01$) showing the variable response of the genotypes across environments and seasons. The average root yield of sweet potato genotypes was significantly ($P < 0.01$) greater at Kachwekano (KARDC) than at Namulonge (NAARI) and Serere (SAARI) locations. Based on AMMI statistical model, Araka Red and Tanzania were the most stable genotypes; while NASPOT 6 and NASPOT 2 had the lowest stability. The model predicted the highest yield from Dimbuca cultivar in 4 of 12 environments and New Kawogo as the cultivar with the lowest yield in 6 of 12 environments. Within each environment and cropping season, the ranking of the genotypes for yield stability was not consistent. Selective deployment of cultivars across environments can improve Sweet potato tuber yield in the lowland and highland tropics.

Key words: Sweet potato; AMMI model; Yield; Cultivars; Genotype × Environment.

Introduction

Sweet potato (*Ipomoea batatas* L.) is an important food crop in many tropical and sub-tropical regions of the world. In Uganda, approximately 2.2 million tonnes of Sweet potato is produced per year making it the second largest producer of the crop in the world (CIP, 1996; Hakiza *et al.*, 2000; FAO, 2002). The utilization of Sweet potato as a food security crop and source of pro-Vitamin A for malnourished children has greatly enhanced the production of the crop in diverse locations (Osiru *et al.*, 2007, Mwangi *et al.*, 2001, 2002). However, current on-farm yields remain low ($4.1\ t\ ha^{-1}$; FAO, 2002) when compared with potential yields in sub-Saharan Africa ($45\ t\ ha^{-1}$;

PRAPACE, 2003). The low yields are attributed to various biotic and edaphic factors such as the *Alternaria* leaf petiole and stem blight disease (Lenne, 1991; Bashaasha *et al.*, 1995; Carey *et al.*, 1997; Skoglund & Smit, 1994).

Although Sweet potato is widely cultivated, agronomic performance and yield stability of crop cultivars under standard management conditions in various agro-ecological conditions are not well documented. The Sweet potato breeding programme in Uganda has registered and released six cultivars (Mwangi *et al.*, 2003) for increased crop production based on dry matter content, tolerance to Sweet

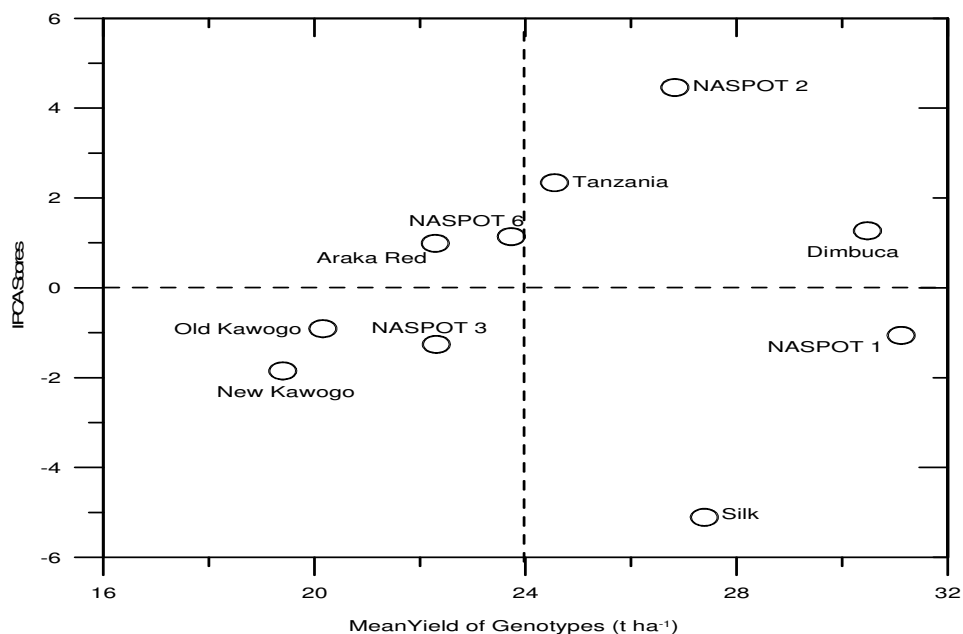


Fig 1. Plot of mean tuber yield (t ha⁻¹) and AMMI interaction (IPCA) scores for 10 Sweet potato genotypes evaluated for yield stability in 12 environments. The IPCA scores are on the Y-axis and root yield are on the X-axis. The genotype Tanzania is a resistant check and NASPOT1 is susceptible check. NASPOT 2, NASPOT 3, and NASPOT 6 are registered cultivars, while Araka Red, Old Kawogo, New Kawogo, Silk and Dimbuca are cultivars selected from diverse regions. The genotypes to the right of mid-point along the X-axis are classified as high yield potential and those to the left side as low yield potential.

potato viruses and high tuber yield. Knowledge of genotype performance and yield adaptation in diverse agro-ecological zones would be highly beneficial for cultivar deployment. Previous research has shown that other crop genotypes can exhibit differences in traits such as yield and disease resistance when grown in diverse environments and may often have various response because of genotype x environment (G x E) interaction (Eberhart & Russell, 1996; Cooper et al., 1996; Crossa et al; 1990; Mulema *et al.*, 2008). Further, differences in yield adaptability among Sweet potato cultivars were noted in other locations (David *et al.*, 1998).

Due to the widespread cultivation of Sweet potato in tropical and sub-tropical regions, assessment of cultivar adaptability is crucial for improved yield. In this research, a multivariate technique known as the additive main effects and multiplicative interaction (AMMI) analysis (Gauch, 1993; Gauch & Zobel, 1990) was used for evaluation of agronomic performance (Sweet potato yield) and genotype adaptation in diverse environments. This manuscript complements a publication on stability of sweet potato cultivars to *Alternaria* leaf and stem blight diseases (Osiru *et al.*, 2009), using the same materials and experimental approach. The objective of this research was to determine the yield performance and

assess yield stability of ten Sweet potato genotypes across a range of environments (seasons and locations) in Uganda using the AMMI statistical model.

Materials and Methods

Site characterisation

The experiments were conducted at three distinct locations of Namulonge Agricultural and Animal Production Research Institute (NAARI), Serere Agricultural and Animal Production Research Institute (SAARI), and Kachwekano Agricultural Research and Development Center (KARDC). These locations represented the main agro-ecologies for Sweet potato production in Uganda as previously described (Osiru *et al.*, 2009).

Plot establishment and experimental design

At each location, 10 Sweet potato genotypes were evaluated for yield. Five cultivars obtained from the Sweet potato breeding programme (Mwanga *et al.*, 2001; 2002; 2003) and five Sweet potato landraces (farmer's varieties) selected from various parts of the country (Osiru *et al.*, 2007).

Table 1. Total tuber yield (Kg ha⁻¹) of ten sweet potato genotypes evaluated for root yield at three locations during four cropping seasons^w

Genotype ^x	2001A ^y			2001B			2002A			2002B		
	NAARI	SAARI	KARDC	NAARI	SAARI	KARDC	NAARI	SAARI	KARDC	NAARI	SAARI	KARDC
Dimbuca	30.98	27.95	20.97	23.45	21.27	62.65	9.29	11.87	62.47	23.69	20.04	63.34
Tanzania	24.42	34.83	29.74	18.83	3.78	48.42	7.91	7.66	40.74	18.72	4.04	47.73
NASPOT 1	33.93	35.09	30.41	35.86	8.48	68.81	7.89	9.43	35.81	34.37	8.28	65.08
NASPOT 2	31.04	58.68	21.88	18.87	23.35	40.43	6.56	8.57	28.65	16.58	23.83	38.99
Silk	26.74	34.58	19.82	19.45	10.48	90.99	3.73	3.56	9.61	18.50	9.58	84.37
New Kawogo	23.90	13.21	11.80	11.20	2.83	59.79	1.75	4.66	22.65	12.01	3.59	61.19
NASPOT 3	25.51	20.53	30.55	12.18	3.46	61.37	4.28	9.24	24.05	12.29	4.07	60.74
NASPOT 6	15.57	32.84	23.81	12.75	4.74	59.79	3.97	5.26	54.26	13.21	5.68	59.22
Araka Red	6.67	34.85	28.90	9.33	17.47	54.20	2.95	3.43	20.19	9.54	17.08	52.28
Old Kawogo	31.51	24.37	13.76	21.72	8.43	52.69	11.78	0.55	8.31	21.48	8.42	53.88
Means	21.92	31.69	23.16	18.36	10.43	59.91	5.99	6.42	30.67	18.04	10.46	58.68
LSD (0.05)	9.82	11.27	12.21	15.15	12.81	23.05	6.43	6.37	20.95	13.98	12.30	18.98

^wExperimental locations consist of Namulonge Agric. Res. Station (NAARI), Serere Animal Agric Res. Station (SAARI) and Kachwekano Agricultural Research & Development Center (KARDC),.

^xGenotypes evaluated for yield stability in diverse environments, ^ySeasons consist of 2001A , 2001B, 2002a and 2002B.

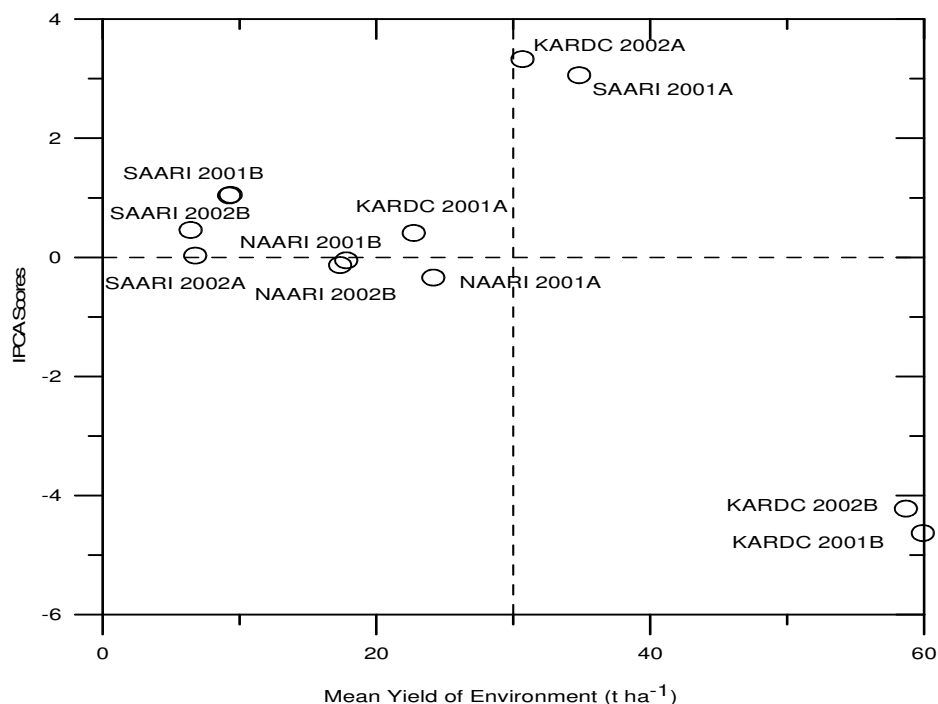


Fig 2. Plot of mean tuber yield ($t\ ha^{-1}$) averaged across genotypes and AMMI interaction (IPCA1) scores for 12 environments in which sweet potato was evaluated. The IPCA1 scores are on the Y-axis and root yield for each environment are on the X-axis. NAARI= Namulonge; SAARI= Serere; KARDC= Kachwekano; 2001A and 2002A correspond to first rains (March to July) of 2001 and 2002, respectively; 2001B and 2002B refer to second rains (September – December) of 2001 and 2002, respectively. The environment to the right of the mid-point along the x-axis is classified high yield potential and those to the left as low yield potential.

The experiments were established in a randomized complete block design (RCBD) with four replications. Each plot consisted of four rows (ridges of 40 cm high) which were 6 m in length. Stem cuttings (30 cm in length with 6 nodes) were planted at a spacing of 25 cm apart within rows and 90 cm between rows. A total of 20 cuttings were planted in each row. Field plots were subjected to normal agronomic and cultural practices. At all locations, the cropping seasons were as described (Osiru *et al.*, 2009).

Yield evaluation

At maturity, fresh foliar weight /10 plants, weight and number of roots per plant; non-marketable, marketable, and total root yield ($t\ ha^{-1}$) were recorded after sizing. Yield data were obtained by harvesting ten plants from the two middle rows of each plot (Kg). Total root yield was then converted to mean tuber yield per hectare ($t\ ha^{-1}$). The marketable tubers consisted of large clean roots (>45 mm diameter) and medium roots were 25-45 mm, while the unmarketable portion consisted of small roots (<25 mm diameter).

Data analysis

Root yield for each cultivar and location were subjected to analysis of variance (ANOVA) using the General Linear Model procedure of the Statistical Analysis System (SAS, 1995). The means, standard errors (SE), and least significant differences, coefficients of variation were computed (Steel *et al.*, 1997; Gomez & Gomez, 1984). In the GLM model, genotypes were designated fixed effects, while locations, cropping seasons and replications were designated random effects. Yield data were additionally subjected to the Additive Main Effects and Multiplicative Interaction (AMMI) statistical model (Gauch, 1993; IRRISTAT- IRRI, 2005). The AMMI model was calculated as previously described (Osiru *et al.*, 2009).

Results and discussion

Site characterisation

Rainfall patterns and annual total rainfall (mm), average relative humidity (%), and average temperatures ($^{\circ}C$) during the cropping cycle were previously described in a previously published

Table 2. Combined additive, multiplicative interaction and the analysis of variance for root yield of 10 sweet potato genotypes grown in 12 environments

Source of variation	Df	F-value	P>F
Treatments	119	10.29	0.004**
Genotypes	9	65.55	0.008**
Environments	11	934.18	0.01**
Replications	24	5.75	0.394
G x E	99	2.79	0.052*
AMMI 1	19	32.37	0.043*
AMMI 2	17	26.35	0.085
AMMI 3	15	16.8	0.149
Residuals	63	2.05	-

** Significant at $P < 0.01$, * significant at $P < 0.05$, ns = non significant

manuscript. Similarly, the morphological characteristics of Sweet potato genotypes as well as agronomic, tuber yield and disease reaction were previously described (Osiru *et al.*, 2009)

Tuber yield

Variation in root yield was recorded among genotypes, locations and cropping seasons. Averaged root yield was significantly greater ($P < 0.05$) at KARDC (41.10 t ha⁻¹) than at NAARI (16.08 t ha⁻¹) or SAARI (14.75 t ha⁻¹) across genotypes and seasons (Table 1). The highest mean root yield was recorded at NAARI and SAARI during the 2001A, followed by the 2001B seasons (Table 1). At KARDC, the highest average yield (59.91 t ha⁻¹) was recorded in 2001B season. Overall, Sweet potato yield varied among genotypes. The variation in yield among locations may be attributed to weather or climatic factors, and the duration of growing periods (Osiru *et al.*, 2009). The maturity period or duration of Sweet potato cultivation at KARDC location (2200 m.a.s.l) is considerably longer than at NAARI or SAARI (lower altitude locations), contributing to vigorous physiological growth, and perhaps more dry matter accumulation. Previous research has shown that greater physiological growth and dry matter accumulation could be expected at high altitude locations in the tropics (Mcharo *et al.*, 2001). The variation in yield may also be attributed to genotype response to the environment at specific locations. Our results are similar to the findings of yield differences among Sweet potato genotypes previously reported in other environments (Ngeve, 1993; Nawale & Salvi, 1983).

The seasonal variation in root yield may be due to climatic or soil factor differences among locations, especially during the root bulking period at SAARI

and NAARI. Water stress during critical periods of root bulking have been shown to result in low root yield and quality defects in Sweet potato (Carey *et al.*, 1997; Ekanayake *et al.*, 1988,).

Yield adaptation across environments

The analyses of variance for root yield across environments and seasons resulted in significant differences ($P < 0.052$) in the interactions of genotypes x environments (Table 2). The significant interactions of genotypes x environments suggest that root yield of Sweet potato genotypes varied across environments. Previous research has shown similar results of differences in yield of Sweet potato cultivars among locations (Naskar & Singh, 1992; Mcharo *et al.*, 2001).

Although this research was conducted with the intention of assessing Sweet potato genotype interactions with *Alternaria* leaf spot disease, the disease level was relatively low (data not shown) and did not significantly impact yield. At KARDC, where *Alternaria* disease was slightly greater than other locations (Osiru *et al.*, 2009), disease incidence was not significantly correlated to total yield. We hypothesize that the disease effects was manifested at 5 months after planting at KARDC, and occurred after the critical root bulking period, resulting in minimum effects on Sweet potato yield. It is also possible that the relatively longer duration of crop growth at KARDC may have contributed to yield compensation at this location. It is possible that under intense *Alternaria* disease pressure, differences in yield of Sweet potato may be detected. There is little published data on yield loss relationship between *Alternaria* disease and yield of Sweet potato genotypes. Similarly, pathogen effects on yield were not significant in other studies conducted on the

Table 3. Ranking of genotypes based on AMMI estimates and unadjusted means (in parenthesis) for total root yield of 10 Sweet potato genotypes grown in 12 environments (location by season combinations)

Genotypes	NAARI ¹		NAARI		NAARI		NAARI		SAARI		SAARI		SAARI		SAARI		KARDC		KARDC		KARDC		KARDC			
	2001A ²		2001B		2002A		2002B		2001A		2001B		2002A		2001A		2001B		2002A		2002B		2002A		2002B	
Dimbuca	9	(10)	5	(10)	1	(9)	3	(10)	4	(3)	1	(3)	4	(9)	1	(3)	4	(4)	6	(7)	1	(8)	6	(8)		
Tanzania	1	(4)	9	(2)	3	(2)	9	(2)	6	(7)	5	(2)	6	(1)	5	(2)	3	(7)	4	(3)	4	(1)	5	(3)		
NASPOT 1	8	(1)	2	(1)	4	(4)	2	(1)	1	(2)	2	(5)	1	(2)	2	(6)	5	(2)	3	(2)	2	(4)	3	(2)		
NASPOT 2	7	(3)	10	(5)	5	(5)	10	(6)	3	(1)	6	(1)	2	(4)	6	(1)	1	(6)	1	(10)	3	(5)	1	(10)		
Silk	2	(6)	1	(8)	10	(6)	1	(8)	2	(9)	3	(9)	3	(3)	3	(8)	9	(1)	5	(4)	5	(6)	4	(5)		
New Kawogo	4	(9)	3	(7)	7	(7)	4	(7)	10	(6)	9	(7)	10	(6)	8	(7)	10	(5)	10	(6)	10	(2)	10	(6)		
NASPOT 3	6	(8)	4	(9)	6	(10)	5	(9)	7	(10)	7	(10)	7	(7)	7	(10)	8	(10)	8	(5)	8	(7)	8	(4)		
NASPOT 6	5	(2)	7	(3)	2	(1)	7	(3)	9	(8)	4	(6)	9	(10)	4	(5)	7	(9)	9	(8)	6	(10)	9	(7)		
Araka Red	10	(5)	8	(4)	8	(8)	8	(5)	5	(5)	8	(4)	5	(8)	9	(4)	2	(8)	2	(1)	7	(9)	2	(1)		
Old Kawogo	3	(7)	6	(6)	9	(3)	6	(4)	8	(4)	10	(8)	8	(5)	10	(9)	6	(3)	7	(9)	9	(3)	7	(9)		

¹NAARI= Namulonge; SAARI= Serere; KARDC= Kachwekano; ²2001A and 2002A correspond to first rains (March to July) of 2001 and 2002, respectively; 2001B and 2002B to second rains (September – December) of 2001 and 2002, respectively. 1-10 indicate ranking of genotype with respect to the ten genotypes under evaluation at each environment.

Alternaria-potato pathosystem and yield relationship (Shtienberg & Fry, 1990; Rotem, 1994).

A comparison of the ranking of root yield of Sweet potato genotypes predicted by AMMI model resulted in variations in genotype performance across environments (Table 3). Prediction of genotypes with high root yield also varied within environments. The genotype Dimbuca was predicted as the best yielding cultivar in four environments by AMMI model while New Kawogo was predicted as having the lowest yield in six out of twelve environments (Table 3). The low accuracy of prediction implies that AMMI model may not be a good predictor of Sweet potato yield at the tested locations. AMMI prediction of environments where the best root yield could be attained was often accurate suggesting that the model can consistently identify environments (KARDC 2001B, KARDC 2002B) where highest root yield could be attained. In previous research, genotype x environment interaction, and yield have been quantified in some Sweet potato genotypes based on regression analysis (Ngeve, 1993; David *et al.*, 1998). In contrast to other research findings which indicate that AMMI analysis increased the accuracy of yield predictions in diverse crop genotypes (Gauch and Zobel, 1990), our experimental results did not show increased accuracy of predictions on Sweet potato.

A plot of total root yield of Sweet potato genotypes versus the principal component scores (IPCA) showed that the genotypes Dimbuca, Tanzania, NASPOT 2, NASPOT 6, Araka Red, had positive principal component scores (PCA1), while genotypes Old Kawogo, NASPOT 1, NASPOT 3, Silk and New Kawogo had negative IPCA scores (Fig. 1). The genotypes Araka Red, Old Kawogo and NASPOT 1 showed PCA scores in close proximity to 0, implying a small interaction with the environment. The genotypes or environments with large PCA scores (negative or positive) indicate high interactions with the environment even though the accuracy is low. Mean yield at environments showed differences at KARDC 2002B and KARDC 2001B relative to the rest of the testing locations (Fig. 2).

We conclude that high and positive PCA scores derived from AMMI statistical model indicate that Sweet potato genotypes can be expected to perform better in an environment. Similarly, high and negative PCA scores indicate that genotypes may have lower yield potential in an environment, therefore, poorly adapted to the environment. The AMMI statistical model can be utilized to estimate yield performance and adaptation of sweet potato in various environments even though with a lower degree of accuracy. Yield performance of Sweet potato cultivars with similar genotypic characteristics (dry matter content, growth characteristics, reaction to

disease or agronomic conditions) to the genotypes described in this experiment can be estimated if environmental parameters are known. Similarly, the root yield of Sweet potato genotypes as impacted by other biological stress agents such as Sweet potato virus disease, Sweet potato chlorotic stunt virus and other fungal agents could be investigated using a similar methodology. Selective deployment of cultivars across environments can greatly improve Sweet potato root yield in the lowland and highland tropics.

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