



Genotype × Environment Interaction and Stability Analysis for Root Yield in Sweet Potato [*Ipomoea batatas* (L.) Lam]

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Sweet potato breeding in Africa, more especially in Nigeria, has mainly focused on improving productivity on farmers' fields and on fresh root consumption. In order to target the breeding program, the study was conducted to estimate the magnitude of genotype × environment interaction (G × E) and to select stable and high yielding sweet potato genotypes for fresh root yield and root *Cylas* severity in two locations, and to identify the most discriminating and representative test environments in Nigeria. The 41 genotypes were evaluated across two diverse environments using a randomized complete block design (RCBD) with three replications. Data were collected on total number of roots per plant, number and weight of marketable roots per plant, fresh root yield, and root *Cylas* severity. The data were subjected to analysis of variance using the Generalized Linear Model procedure of SAS 9.2 where genotype was treated as a fixed factor and replication treated as a random variable. Stability analysis was conducted using Genotype and Genotype × Environment Interaction (GGE) bi-plot. Environment, genotype, and G × E interaction variances were highly significant ($p < 0.01$) among the assessed agronomic traits. Moreover, the analysis of variance revealed highly significant ($p < 0.01$) differences among genotypes, environments, and G × E interaction effects for all the studied traits. The GGE biplot analyses identified three promising genotypes—G13, G11, and G14—that possess both high mean root yield and high stability, closest to the ideal genotype for root performance and consistency of performance across environments. This study provides valuable information that could be utilized in a breeding program to ameliorate local clones of sweet potato in Nigeria.

Keywords: sweet potato, yield, environment, GGE biplot, genotypes

INTRODUCTION

Sweet potato (*Ipomoea batatas* [L.] Lam), is a hexaploid ($2n = 6x = 90$) and is one of the most important food security crops globally. It belongs to the family *Convolvulaceae*, genus *Ipomoea*, and, according to Vaeasey et al. (2008), the genus has over 600 species, of which *batatas* is the only one with economic value. In many developing countries, sweet potato is reported to be the fifth most important food crop after rice, wheat, maize, and cassava (Aina et al., 2012). Over 110 million metric tons of sweet potato was produced in 2018, with China producing 53.01 million

metric tons representing 65.6% of the world sweet potato production (FAOSTAT, 2018). Africa was responsible for 20.7 million tons which represents about 25.4% of the world production. Nigeria is the second highest sweet potato producer in Africa and the third highest producer in the world, with production of 4.03 million metric tons, which is 5.0% of the world's production (FAOSTAT, 2018). Over the years, sweet potato production has been on the increase in Nigeria. In the last two decades, production has increased 10-fold. However, the increased production can be attributed to the expansion of land under sweet potato cultivation rather than increased yield per unit area, as yield remains abysmally low at an average of <3.0 tons/ha. This low yield is mainly due to the widespread use of obsolete production inputs and methods, chief of which are the use old, unimproved cultivars and the practice of mixed cropping with incompatible crops. Therefore, one important way of mitigating against poor root yield in farmers' fields is to develop and release new sweet potato varieties with stable and high root yield potential into the farming system.

Understanding the differential response of crop genotypes to changing environmental conditions is of key importance in plant breeding. One major step toward the development of improved crop genotypes is the assessment of the nature of interactions that exist between genotypes and the production environment for a particular triat (Sabri et al., 2020). When genotypes are evaluated across a range of different locations and/or years, their yield performances could differ significantly. The existence of large $G \times E$ interaction usually causes serious confounding effects in comparing and recommending good genotypes for wide adaptation (Moussa et al., 2011). Previous $G \times E$ studies on several traits have demonstrated that sweet potato is sensitive to environmental changes. According to Madawal et al. (2015), Gurmu et al. (2017), and Ngailo et al. (2019), changes in environmental conditions have been reported to affect sweet potato storage root yield and yield components. This makes the analysis for $G \times E$ interaction crucial for genotype selection, cultivar release, and identification of suitable production environments for optimum yield. Therefore, having a basic understanding on $G \times E$ interactions, stability parameters, and genetic correlations for root yield and yield components are considered necessary for sweet potato breeders in making an informed choice concerning which locations and input systems should be used in their breeding efforts (Gruneberg et al., 2005).

Statistical tools such as the Additive Main Effect and Multiplicative Interaction (AMMI) (Gauch, 1992) and genotype and genotype-by-environment interaction (GGE) biplot analyses (Yan and Kang, 2003; Yan and Tinker, 2006) have been reported as appropriate for use in GEI analyses. These statistical tools have then been extensively used in several sweet potato improvement programs by authors such as Caliskan et al. (2007) (AMMI model analysis for GEI and stability analysis of sweet potato genotypes across different environments in Turkey) and Laurie and Booyse (2015) (GGE biplots used in and identifying suitable sweet potato genotypes and representative environments in South Africa). The objective of this study, therefore, was to determine the magnitude of GEI for storage root yield, yield-related traits, and sweet potato weevil (*Cylas* spp.) damage among candidate sweet

potato genotypes, as well as to assess the adaptability and stability of 41 improved sweet potato genotypes in two sweet potato representative and contrasting production environments.

MATERIALS AND METHODS

Plant Materials

Forty-one sweet potato genotypes comprising breeding lines, farmer cultivars, and released varieties (checks) were used for the trials. The genotypes were selected based on their high root dry matter contents (RDMC), flesh color (as an indicator of the level of β -carotene content), and fresh root yield.

Research Locations

The field evaluation was conducted in two consecutive seasons at two different locations of Abakaliki (Ebonyi State) and National Root Crops Research Institute (NRCRI) (Iresi Osun State). The two locations differed in ecological characteristics, altitude, rainfall, and atmospheric temperature; the climate could be described as hot humid tropic, with high humidity and adequate rainfall (Table 1).

Field Layout and Experimental Design

The field trial was conducted using a randomized complete block design (RCBD) with three replications. The experimental plots consisted of a three—row plot of three meters long for each genotype. The spacing between rows was 1.0 m and within rows was 0.3 m, giving a total of 10 plants per row and 30 plants per plot (Afuape et al., 2019). The sweet potato was planted in June and harvested in October in 2018 and 2019 in Abakaliki, Ebonyi state (rain forest belt) and Iresi, Osun state (savannah transition ecology) for 2 years. The experiment was conducted under rain-fed conditions. Fertilizer (NPK 15:15:15) was applied at the rate of 400 kg/ha in the test sites, while weeding was done as necessary.

Quality Traits Analysis

The sweet potato genotypes evaluated in Abakaliki were analyzed for chemical quality traits. Dry matter content was determined according to Seruwu (2012), crude protein, fiber, and ash content were determined using the micro Kjeldahl method of AOAC (2010), while fat content was determined according to AOAC (2010) soxhlet extraction method. Carbohydrate was determined using Gravimetric Copper Reduction Method (AOAC, 2010; Okporie et al., 2013). Total carotenoid content of the sweet potato genotypes was determined by the procedure described by Amaya (2001).

Data Collection and Analysis

Agronomic data (number and weight of marketable and unmarketable roots, sweet potato root Cylas severity) were collected and subjected to analysis of variance (ANOVA) using the Generalized Linear Model procedure of SAS 9.2 where genotype was treated as a fixed factor and replication treated as a random variable according to the model of Steel and Torrie (1980). Number of marketable (or saleable) roots represents the number of roots that were more than or equal to 100 g (Levette, 1993) or with diameters at the widest point >25 mm

TABLE 1 | Weather information of the research locations.

Location	Latitude (°N)	Longitude (°E)	Altitude (m)	Soil texture	Min and max daily Temp (°C)	Average annual rainfall (mm)
Abakaliki	6° 19' 30N	8° 6' 49E	116	Sandy loam	26.2 – 29.0 °C	1800.3
NRCRI Iresi	7°30'0"N	4°30'0" E	246	Sandy loam	24.7 – 27.8 °C	2024.1

Umudike: Agro-meteorological Unit, National Root Crops Research Institute, Umudike, Abia State.

roots. These were counted and the number recorded per plot. Number of unmarketable roots represents the number of roots that are <100 g or 25 mm at the widest point (Levette, 1993; Stathers et al., 2003). Weight of marketable roots is the weight (kg/plot) of roots suitable for marketing. Weight of unmarketable roots is the weight (kg/plot) of roots not suitable for marketing. Total root yield (t/ha) weight is obtained as the sum of weights of marketable and unmarketable roots converted to tons per hectare. Severity of root *Cylas* infection was taken as the mean damage level of SPVD-diseased plants in a plot on a 1–5 infection scale where 1 = no apparent damage, 2 = mild/very little damage, 3 = moderate damage, 4 = considerable/severe damage, and 5 = severe/very severe damage (Mwanga et al., 2002).

The agronomic data were first analyzed on a location basis since the variances of the two locations were significantly different following a Bartlett's test for homogeneity of the variances of the two locations. The Least Square (LS) means of the genotypes in each location were estimated and separated using standard error of difference (SED). However, the 41 genotypes were then analyzed as a combined trial to increase the resolving power of the estimate of genotype × environment interaction across the two locations and years.

The ANOVA model used for the single-site analysis is as stated below:

$$Y_{ij} = \mu + \alpha_i + \gamma_j + \beta_k + e_{ij}$$

where

y_{ij} = observation on experimental unit in block k assigned treatments i and j ;

μ = overall mean averaged over all treatments and all blocks;

α_i = effect of genotype i ; considered as fixed variable;

γ_j = effect of year j considered as random variable;

β_k = effect of block k considered as random variable;

e_{ijk} = random error associated with experimental units assigned to treatments i and j in block k .

The ANOVA table and expected mean squares for combined sites and population are as shown in **Table 2**.

Genotype × Environment Interaction (GEI) and Stability Analysis

The data generated were analyzed for GEI using GenStat (14th edition) GGE biplot procedure (Yan, 2001, 2002; Yan and Ma, 2006). The model for the GGE biplot based on singular value decomposition (SVD) of t principal components is:

$$Y_{ij} = \mu + \alpha_i + \beta_j + j_{ij}$$

where Y_{ij} is the measure of the i th genotype in the j th environment, μ is the grand mean, α_i is the main effect of the i th genotype, β_j is the main effect of the j th environment, and j_{ij} is the interaction between the i th genotype and j th environment.

RESULTS

The mean squares of the analyses of variance (ANOVA) of the agronomic traits evaluated in two different locations (Abakaliki and Iresi) revealed that there were significant ($p < 0.01$) variations among the genotypes for all the traits in each location and across both locations (**Tables 3–5**). The combined analysis (**Table 6**) also showed that all the agronomic traits varied with respect to genotype, the location, and Genotype-by-Location Interaction. Correlation coefficients for the agronomic traits evaluated in the two locations and 2 years are presented in **Table 7**. Generally, all the traits except root *Cylas* spp. severity exhibited a positive and significant ($p < 0.01$) correlation with root yield. Most of the traits also exhibited significant and positive association among themselves, except root *Cylas* spp. severity at harvest which did not correlate with all other traits.

Genetic Studies

The genotypic (V_g), environmental (V_e), and phenotypic (V_p) variances, as well as the broad sense heritability (H_B) estimates of the agronomic traits at each and combined locations, are presented in **Tables 8–10**. For all the agronomic traits in each location and across locations, V_e was higher than V_g . Heritability estimate for number of marketable roots at Abakaliki was 0.44, while at Iresi it was 0.69. The broad-sense heritability (H_B) estimates for number of unmarketable roots were 0.33 and 0.02 at Abakaliki and Iresi, respectively, and 0.30 and 0.74 for weight of marketable roots at Abakaliki and Iresi, respectively. H_B for root yield at Abakaliki and Iresi, respectively, were 0.48 and 0.81. Understanding the genetic parameters of the traits that impact on farmers and consumers' variety preference is a good guide during the selection of parental lines. **Table 9** shows the genotypic, phenotypic, and environmental variances and the broad sense heritability estimates (H_B) of the agronomic traits in both locations. The result in **Table 9** shows that environmental variances (V_e) were slightly higher than genotypic variance (V_g) for all traits at both locations, depicting environmental influence on the expression of the traits. The heritability (H_B) estimates were either low or high and ranged between 0.02 and 0.52 for all the agronomic traits.

TABLE 2 | Description (background information) of sweet potato genotypes used for the study.

S/No.	Genotypes	Pedigree	Status	Flesh color*
1	OP/87/0087	TIS 87/0087	Breeding line	Light orange
2	F2M1/31	Centennial X TIS 8164	Breeding line	Orange
3	F5M1/3	CIP199034.1 X TIS 8164	Breeding line	Orange
4	MD/23	Mother's Delight	Breeding line	Cream
5	F2M1/14	Centennial X TIS 8164	Breeding line	Very light orange
6	F2M1/18	Centennial X TIS 8164	Breeding line	Light orange
7	F2M1/21	Centennial X TIS 8164	Breeding line	Light orange
8	F2M1/22	Centennial X TIS 8164	Breeding line	Yellow-orange
9	F2M1/35	Centennial X TIS 8164	Breeding line	Light orange
10	Ex-Igbariam/22	Ex-Igbariam	Breeding line	Cream
11	Ex-Igbariam/26	Ex-Igabriam	Breeding line	White
12	Ex-oyunga/17	Ex-Oyunga	Breeding line	White
13	MD/12	Mother's Delight	Breeding line	Light orange
14	Progeny 1	Not available	Breeding line	Orange
15	Progeny 3	Not available	Breeding line	Light orange
16	F1M1/23	Mother's Delight X TIS8164	Breeding line	White
17	F1M1/57	Mother's Delight X TIS 8164	Breeding line	Cream
18	F1M1/64	Mother's Delight X TIS 8164	Breeding line	Light orange
19	F2M1/35	Centennial X TIS 8164	Breeding line	Orange
20	F2M5/13	Centennial X Solo-Abuja	Breeding line	Cream
21	F2M5/5	Centennial X Solo-Abuja	Breeding line	White
22	F2M5/9	Centennial X Solo-Abuja	Breeding line	White
23	F2M6/1	Centennial X Ex-Igbariam	Breeding line	Light orange
24	F2M6/17	Centennial X Ex-Igbariam	Breeding line	Cream
25	F2M6/20	Centennial X Ex-Igbariam	Breeding line	White
26	F2M6/27	Centennial X Ex-Igbariam	Breeding line	Deep orange
27	Solo-1/165	Solo-1	Breeding line	White
28	Solo-1/21	Solo-1	Breeding line	Light orange
29	Solo-1/88	Solo-1	Breeding line	Cream
30	Solo-Abuja/12	Solo-Abuja	Breeding line	Yellow
31	TIS 87/0087/01	TIS 87/0087	Breeding line	White
32	TIS 87/0087/03	TIS 87/0087	Breeding line	Cream
33	TIS 87/0087/07	TIS 87/0087	Breeding line	White
34	TIS 87/0087/23	TIS 87/0087	Breeding line	White
35	TIS 87/0087/25	TIS 87/0087	Breeding line	White
36	UMUSPO-2/02	UMUSPO-2	Breeding line	Cream
37	UMUSPO-2/95	UMUSPO-2	Breeding line	Very light orange
38	MD	Not available	Released variety	Orange
39	TIS 87/0087	Not available	Released variety	Cream
40	KING J	Not available	Released variety	Light orange
41	MD/03	Not available	Released variety	Cream

MD, Mother's Delight.

*means significant at $P = 5\%$; **means Highly significant at $P = 1\%$; ***means Very highly significant at $P = 0.1\%$.

Genotype and Genotype by Environment (GGE) Biplots

Storage Root Yield

The GGE biplot analyses (which-won-where, mean, stability, and genotype ranking) for storage root yield are presented in **Figures 1A–C**. **Figure 1A** shows the “which-won-where” root yield performance of the sweet potato genotypes evaluated in the

four environments. The two principal components (Axis 1 and 2) revealed about 73% of the total variation observed with Axis 1 accounting for 50% of the total variation, while Axis 2 influenced 23% of the observed variation. Genotype G11 had the highest root yield performance at Iresi 1 and Iresi 2 environments, while G14 was best adapted to the two Abakaliki environments as both were located at the vertices of the polygon in the environments

TABLE 3 | Format of ANOVA Table for combined sites and population.

Source of variation	Df	E(MS)
Block (L)	l(b-1)	$\sigma^2e + gl\sigma^2b$
Location (L)	l-1	$\sigma^2e + b\sigma^2gl + bg\sigma^2l$
Genotypes (G)	g-1	$\sigma^2e + b\sigma^2gl + bl\sigma^2g$
Year (Y)	y-1	$\sigma^2e + b\sigma^2g/l(y) + gb\sigma^2yl + bl\sigma^2g(y) + gbl\sigma^2y$
G(Y)	y(g-1)	$\sigma^2e + b\sigma^2g/l(y) + gb\sigma^2yl + rl\sigma^2g(y)$
G × L	(g-1)(l-1)	$\sigma^2e + b\sigma^2gl$
Y × L	(y-1)(l-1)	$\sigma^2e + b\sigma^2g/l(y) + gb\sigma^2yl$
G(Y) × L	(g-1)y(l-1)	$\sigma^2e + b\sigma^2g/l(y)$
Error	(b-1)(gyl-1)	σ^2e
Total	gylb-1	

b, Blocks; G, Genotypes; Y, Year; e, Error; df, Degree of freedom; MS, Mean squares; E(MS), Expected mean squares.

TABLE 4 | Mean squares of the analyses of variance of agronomic traits of sweet potato genotypes evaluated at Abakaliki across 2 years.

Sources of variation	Mean squares								
	Degrees of freedom	Number of marketable roots	Number of unmarketable roots	Total root number	Wt. of marketable roots	Wt. of unmarketable roots	Total root weight	Root yield	Cylas severity
Rep	2	16.0733	4.6426	11.4314	2.6747	0.0328	3.2441	109.1065	1.9259
Year	1	34.7041 ^{ns}	741.3983 ^{***}	1071.2141 ^{**}	1.7998 ^{ns}	0.1894 [*]	3.4475 ^{ns}	52.1796 ^{ns}	2.5556 [*]
Genotype	40	75.1750 ^{***}	109.0174 ^{***}	280.1644 ^{***}	4.7343 ^{***}	0.0562 [*]	5.0521 ^{***}	83.1419 ^{***}	0.4375 ^{ns}
Gen. × Year	40	41.8796 ^{**}	72.6024 ^{**}	169.7607 ^{**}	3.2940 ^{**}	0.0387 ^{ns}	3.6141 ^{**}	43.1045 ^{ns}	0.5001 ^{ns}
Error	120	23.5595	40.3275	91.2070	1.3839	0.0328	1.5771	30.6534	0.4303

^{*}means significant at P = 5%; ^{**}means Highly significant at P = 1%; ^{***}means Very highly significant at P = 0.1%. Wt., weight; ns, not significant.

(Yan and Tinker, 2006; Ngailo et al., 2019). Genotype G13 was second best in all the environments. The three genotypes were better than the rest in all the environments.

The GGE biplot “Mean vs. Stability” results are as presented in **Figure 1B**. The result showed that G14 had the highest mean root yield across all environments, followed by G11 and G13. The other genotypes had mean root yield around the grand mean value by their relative locations in the biplot. Among the genotypes with high mean root yield, G13 was the most stable, with G11 and G14 exhibiting marked variability in root yield in the various environments. However, there were some genotypes with high stability, but their mean root yields were very poor. According to Yan and Tinker (2006), only stable genotypes with high mean performance are desirable.

An ideal genotype should have both high mean performance and high stability across environments. The center of the concentric circle (**Figure 1C**) is the location for the ideal genotype. Among the test genotypes, the one closest to the point is the best. Though G14 had the highest storage root yield among the 41 genotypes, G13 that possessed both high mean root yield and high stability is closest to the ideal genotype for root yield with consistency of performance across environments.

Root Cylas spp. Severity

The GGE biplot results for the severity of sweet potato root *Cylas* damage are as presented in **Figures 2A–C**. The two principal

component axes used for making the biplots accounted for 62% of the total variation, with Axis 1 and Axis 2 accounting for 38.09 and 24.77% of the total variation, respectively. For root *Cylas* spp. severity, genotypes with high and positive axis 1 scores entailed high susceptibility and those with negative axis 1 were mostly resistant. Sweet potato root *Cylas* spp. severity has been reported to be a major factor that affects the production of sweet potato in Africa (Gibson et al., 1998). Root *Cylas* damage is further enhanced by the cultivation of infested genotypes and a lack of efficient control measures which, in addition, result in low sweet potato yields in many countries. Improved sweet potato genotypes that are resistant to root *Cylas* severity with high yield potential would increase sweet potato production in sub-Saharan Africa (Rukundo et al., 2017; Ngailo et al., 2019).

The GGE biplot that analyzed for genotype performance in a specific environment (which-won-where) showed that G24 had the least *Cylas* spp. damage in environments IRE 1 and ABA 2, while G37 and G31 exhibited least root damage in IRE 2 and ABA 1, respectively (**Figure 2A**). Genotypes G22 and G30 had the most severe *Cylas* spp. damage across environments. The differential expression of tolerance by the genotypes in different environments is unexpected as tolerance to *Cylas* spp. is known to be largely influenced by strong environmental control.

Across all environments, G37 also had the least mean *Cylas* spp. damage, followed by G23, G14, and G15 in that order (**Figure 2B**). The four genotypes also exhibited high *Cylas*

TABLE 5 | Mean squares of the analyses of variance of agronomic traits of sweet potato genotypes evaluated at Iresi, Osun State across 2 years.

Sources of variation	Degrees of freedom	Mean squares							
		Number of marketable roots	Number of unmarketable roots	Total root number	Wt. of marketable roots	Wt. of unmarketable roots	Total root wt	Root yield	Cylas severity
Rep	1	152.5814	11.2558	218.8140	1.0880	0.0126	0.8526	35.4207	0.5931
Year	1	857.9391***	1404.2250***	4563.5641***	5.2490 ^{ns}	0.02116 ^{ns}	5.9753 ^{ns}	118.8241 ^{ns}	4.0960**
Genotype	40	56.4123**	24.3874 ^{ns}	110.5934**	7.5395**	0.02176 ^{ns}	7.7524***	110.1608**	0.4432 ^{ns}
Gen. × Year	40	20.948 ^{ns}	23.6844 ^{ns}	61.61.2434 ^{ns}	3.1535 ^{ns}	0.02191 ^{ns}	3.3073 ^{ns}	44.2613 ^{ns}	0.4154 ^{ns}
Error	120	28.4289	20.4409	51.8588	3.0167	0.02081	3.0543	59.1476	0.4389

*means significant at $P = 5\%$; **means Highly significant at $P = 1\%$; ***means Very highly significant at $P = 0.1\%$.
^{ns}, not significant.

TABLE 6 | Mean squares of the analysis of variance of sweet potato genotypes evaluated for root yield and yield components at two locations in 2 years.

Sources of variation	Degrees of freedom	Mean squares							
		Total number of roots	Total root weight	Root yield (t/ha)	Number of marketable roots	Number of unmarketable roots	Weight of marketable roots	Weight of unmarketable roots	Cylas spp. severity
Block (L)	4	49.713	1.173	23.020	60.108	6.935 ^{ns}	1.058	0.034	2.108
Year	1	575.474**	0.157 ^{ns}	165.325*	267.022**	46.374 ^{ns}	0.431 ^{ns}	0.043 ^{ns}	6.526**
Location	1	559.649**	27.502***	636.851**	3.584 ^{ns}	530.327***	32.222***	0.256**	1.252 ^{ns}
Location*Year	1	4986.105***	9.161*	5.794 ^{ns}	608.960***	2085.430***	6.514 ^{ns}	0.170*	0.074 ^{ns}
Genotype	40	249.549***	9.252***	128.506***	92.225***	76.166***	8.859***	0.046*	0.562 ^{ns}
Genotype*Location	40	127.882**	3.782**	66.594*	38.144*	50.887*	3.650**	0.029 ^{ns}	0.297 ^{ns}
Genotype (Y)	80	108.439***	2.592 ^{ns}	28.666 ^{ns}	28.768 ^{ns}	44.813 ^{ns}	2.319 ^{ns}	0.034 ^{ns}	0.519 ^{ns}
Genotype (Year)	80	113.280*	4.384**	58.908 ^{ns}	33.275 ^{ns}	46.569*	4.190**	0.025 ^{ns}	0.405 ^{ns}
Location									
Error	326	74.774	2.219	43.572	25.818	31.739	2.090	0.028	0.433
Total	491								

*means significant at $P = 5\%$; **means Highly significant at $P = 1\%$; ***means Very highly significant at $P = 0.1\%$.
^{ns}, not significant.

spp. tolerance stability, thereby enhancing the probability of identifying highly tolerant genotypes that could be deployed across many environments. These genotypes can form an elite genetic resource pool in the global efforts toward the genetic control of *Cylas* spp. damage in sweet potato.

Overall, G37 can be ranked as the most tolerant genotype to *Cylas* spp. damage due to its position very close to the center of the concentric circle that represents the “ideal genotype” (Yan and Tinker, 2006). The ranking of the leading genotypes according to their tolerance to *Cylas* spp. damage performance can be given as: G37 > G23 > G14 > G15. **Figures 3, 4** describe the severity of root *Cylas* spp. in Abakaliki and Iresi Osun State, respectively.

DISCUSSION

Genetic variability is essential for selection (Sarif et al., 2020). Except for *Cylas* weevil damage severity (in all environments) and weight of unmarketable roots (in some environments), significant variation exists among the genotypes evaluated for all the traits, an occurrence that has also been reported by

Vimala et al. (2011), Madawal et al. (2015), and Afuape et al. (2019). In this study, year as a factor exerted a reducing influence on the agronomic traits. *Cylas* weevil damage severity is largely controlled by the environment as genetic resistance to *Cylas* weevil is still unavailable. By its epidemiology, *Cylas* spp. incidence and severity occurs in the dry season when soil moisture level is very low (Stathers et al., 2003). This explains the observed large environmental variance (0.43) compared to the genotypic variance (0.00), and the extremely low broad-sense heritability estimate (0.02) recorded in this study for the trait. The very low genotypic variance (0.00) and heritability estimate ($H_B = 0.00$) for weight of unmarketable roots (roots < 100 g) also portend that environment is largely responsible for the formation of small sweet potato roots. This is true as plant density, drought spell, and poor weed management among other factors could lead to the formation of small storage roots.

The genetic study reveals that environmental variance (V_e) was higher than genotypic variance (V_g), leading to moderate broad sense heritability (H_B) estimates in both locations and in the combined form. As H_B estimate is a reflection of the genetic component of an observed phenotype expression of a

TABLE 7 | Combined Pearson correlation coefficients (*r*) of agronomic traits evaluated in two locations and 2 years.

Traits	Number of marketable roots	Total number of roots	Weight of marketable roots	Total root weight	Root yield	Root <i>Cylas</i> spp. severity
Number of marketable roots	1.00	0.828***	0.721***	0.736***	0.611***	−0.091 ^{ns}
Total number of roots		–	0.509***	0.561***	0.438***	−0.073 ^{ns}
Weight of marketable roots			–	0.995***	0.861***	−0.034 ^{ns}
Total root weight				–	0.861***	−0.031 ^{ns}
Root yield					–	−0.042 ^{ns}

***means Very highly significant at $P = 0.1\%$.
ns, not significant.

TABLE 8 | Variance components and broad-sense heritability estimates of agronomic traits of sweet potato genotypes evaluated at Abakaliki, Ebonyi State across 2 years.

Traits	V _g	V _{gy}	V _e	H _B
Number of marketable roots	8.32	9.16	23.56	0.56
Number of unmarketable roots	9.10	16.14	40.33	0.53
Total root number	27.60	39.28	91.21	0.50
Wt. of marketable roots	0.36	0.96	1.38	0.38
Wt. of unmarketable roots	0.00	0.00	0.03	0.00
Total root wt	0.36	1.02	1.58	0.36
Root yield	10.01	6.23	30.65	0.64
<i>Cylas</i> severity	−0.02	0.03	0.43	0.00

trait, the moderate H_B in the combined genotype-location-year interaction analysis depicts that genetic gain could be achieved in such important traits as total root weight (H_B = 0.5), weight of marketable roots (0.52), root yield (0.46), and number of marketable roots (0.34). Moderate heritability is often deemed acceptable for quantitative traits (Tumwegamire et al., 2011) and has been variously reported for most root yield and yield components by Tumwegamire et al. (2011) and Afuape et al. (2019). The observed differences in the H_B estimates for each trait in both locations shows that location affects trait heritability estimates, with poor environment lowering the value by confounding the true genotypic value with non-experiment wise errors which could lead to selection error in the breeding cycle.

Correlation coefficient is a measure of the extent and direction of the relationship between any two traits (variables). The positive and strong relationships between root yield and number of marketable roots ($r = 0.786$), total number of roots ($r = 0.438$), marketable roots ($r = 0.861$), and total root weight ($r = 0.861$) suggests that these traits are important root yield components, and that their simultaneous selection will be a good approach to increasing root yield. This same relationship had been observed by Afuape et al. (2011) and Yahaya et al. (2015). As the severity of *Cylas* spp. is a yield-reducing affect, the negative correlation observed between it and root yield and yield components is expected as severely infested fields often exhibit reduced plant density at harvest, severe reduction in root weight, significant reduction in good roots, and impaired plant physiological processes due to severed food translocation organs, as infested plants often have the vine cortex and pith eaten up by the *Cylas* weevil larvae developing inside the plant.

The stability analysis aims at helping the breeder to identify which genotypes have specific and/or general adaptability to

various production environments. It also helps in the analyses of the test environments for prudent decision making for future evaluations. As expected, the two locations (Iresi and Abakaliki) clustered in different quadrants, depicting that both locations are truly different. The Iresi environments (IRE 1 and IRE 2) were higher performing environments for root yield and *Cylas* damage suppression compared to the Abakaliki environments (ABA 1 and ABA 2). The differences between these locations were so large that year effect could not nullify it, meaning that testing data from one location cannot represent the performance of same materials in the other location, irrespective of the number of years used in the testing process. As different genotypes performed differently in each of these locations, we are able to identify genotypes that are specifically adapted to each environment, knowledge that will help breeders to adequately advise farmers on what cultivar to use where, provided the various cultivars possess acceptable end-user quality preferences. Conducting a stability analyses has thus helped in identifying specific genotypes for both locations, as well as a stable genotype that can be cultivated across all the locations tested and locations that share similar attributes to the test locations.

Dry matter content according to Mok et al. (1997) and Cervantes-Flores et al. (2011) is among the main attributes that further improve the quality and acceptability of sweet potato for consumption and processing in sub-Saharan Africa. The dry matter content ranged from 43.85% (F2M1/35) to 24.07% (F2M6/17) and varied significantly ($p < 0.0001$) between the sweet potato genotypes. Findings from this study are in accord with records reported by Okunade et al. (2019) in Osun (21.35–42.10%) and Kanu et al. (2018) in Umudike, Abia State (27.30–41.56%), but disagrees with the result (19.4–22.6%) reported by Laurie et al. (2013) for some OFSPs. The low dry matter content recorded in F2M6/17 could be a result of the presence of

TABLE 9 | Variance components and broad-sense heritability estimates of agronomic traits of sweet potato genotypes evaluated at Iresi, Osun State across 2 years.

Traits	Vg	Vgy	Ve	Vp	H _B
Number of marketable roots	8.87	-3.74	23.56	12.89	0.69
Number of unmarketable roots	0.18	1.62	40.33	11.07	0.02
Total root number	12.34	4.69	91.21	37.49	0.33
Wt. of marketable roots	1.10	0.07	1.38	1.48	0.74
Wt. of unmarketable roots	0.00	0.00	0.03	0.01	0.00
Total root wt	1.11	0.13	1.58	1.57	0.71
Root yield	16.47	-7.44	30.65	20.42	0.81
Cylas severity	0.01	-0.01	0.43	0.11	0.08

TABLE 10 | Variance components and broad-sense heritability estimates of agronomic traits of sweet potato genotypes evaluated across two locations and years.

Traits	Vg	Vgl	Vgy	Vgly	Ve	H _B
Total number of roots	10.95	2.43	-1.61	12.84	74.77	0.22
Total root weight	0.76	-0.10	-0.60	0.72	2.22	0.50
Root yield (t/ha)	10.20	1.28	-10.08	5.11	43.57	0.48
Number of marketable roots	5.26	0.81	-1.50	2.49	25.82	0.34
Number of unmarketable roots	2.40	0.72	-0.59	4.94	31.74	0.13
Weight of marketable roots	0.75	-0.09	-0.62	0.70	2.09	0.52
Weight of unmarketable roots	0.00	0.00	0.00	0.00	0.03	0.00
Cylas spp. severity	0.00	-0.02	0.04	-0.01	0.43	0.02

$$H_B = [\sigma^2(G)] / [\sigma^2(G) + \sigma^2_{error/replication} + \sigma^2(GY)/year + \sigma^2(GL)/location + \sigma^2(GY)/year]$$

Vg, Ve, Vp = Genotypic, Environmental and Phenotypic Variance, respectively.

Vgy, Vgl, and Vgly = genotype-by-year, genotype-by-location and genotype-by-location-by-year interaction variances, respectively.

H_B = Broad sense heritability.

high moisture in the tuber. Factors such as variety, crop season, age of the plant, and location could affect dry matter content in sweet potato. According to Eleazu and Ironua (2013), high dry matter content improves storability, texture, and product yield. It also has the potential of being utilized for industrial purposes and for flour production in confectioneries. Proteins are essential nutrients for structural and functional performers of different biomolecules in the human body, and they provide the essential amino acids required for metabolism. The protein content recorded in this study varied significantly ($p < 0.0001$) across the forty sweet potato genotypes. Results from this study revealed that protein was highest in TIS 87/0087/01 (2.67%), followed by UMUSP 2/02 (2.53%), and lowest in SOLO-1/88 (1.54%). This result correlated well with reports credited to Rakesh et al. (2017) in some sweet potato cultivars. Studies show that protein content in sweet potato, especially orange fresh sweet potato (OFSP), is comparable to cassava (1.6%) and yam (1.5) (Woolfe, 2008).

The fat content among the forty sweet potato varieties ranged between 0.17 to 0.31% and showed significant ($p < 0.0001$) differences among the genotypes. The fat content was high in solo-1/21 (0.31%) while fat was least in the MD/12 genotype with 0.17%. The results of fat content from this study were in agreement with Ishida et al. (2000) and Mohammad et al. (2016), who reported 0.17–0.30% and 0.2–0.33%, respectively. Sweet potato, like other known roots and tubers, is well-known for

its low fat content. The fat concentration of some sweet potato, especially the orange fresh sweet potato, are still little better than other roots and tubers such as cassava (0.28%) and yam (0.17%) (USDA, 2018). Crude fiber, which is also an important indicator of a healthy food material that plays a major role in reducing the incidences of colon cancer, diabetes, heart disease, and certain digestive diseases, were observed in an appreciable amount. The crude fiber content of the sweet potato ranged from 0.33% (in SOLO-1/21) to 0.54% (in TIS 87/0087/01) and was significantly different among the genotypes. This result is in line with 0.35% reported by Endrias et al. (2016) for different varieties of OFSP.

Ash is an inorganic residue in any food substance which directly denotes the mineral content. The sweet potato has a reasonable amount of ash content, an indication of rich mineral constituents. The ash value varied from 1.14% (in F2M6/1 and SOLO-1/21) to 1.32% (in F2M1/31). The ash values obtained from this study were in line with the results reported by Mohammad et al. (2016), who recorded 1.17–1.31% ash content. Carbohydrate was among the most abundant nutritive constituent of the sweet potato. This is consistent with previous reports credited to Adepoju and Adejumo (2015), Amha and Baruch (2016), and Endrias et al. (2016). The carbohydrate content obtained from this study ranged from 19.95% in F2M6/17 to 39.75% in F5M1/3 and varied significantly among the genotypes, indicating that they are a good source of energy. This study is within FAO Corporate Document Repository

TABLE 11 | Analyses of some chemical quality traits of the Sweet potato genotypes evaluated at Abakaliki across 2 years.

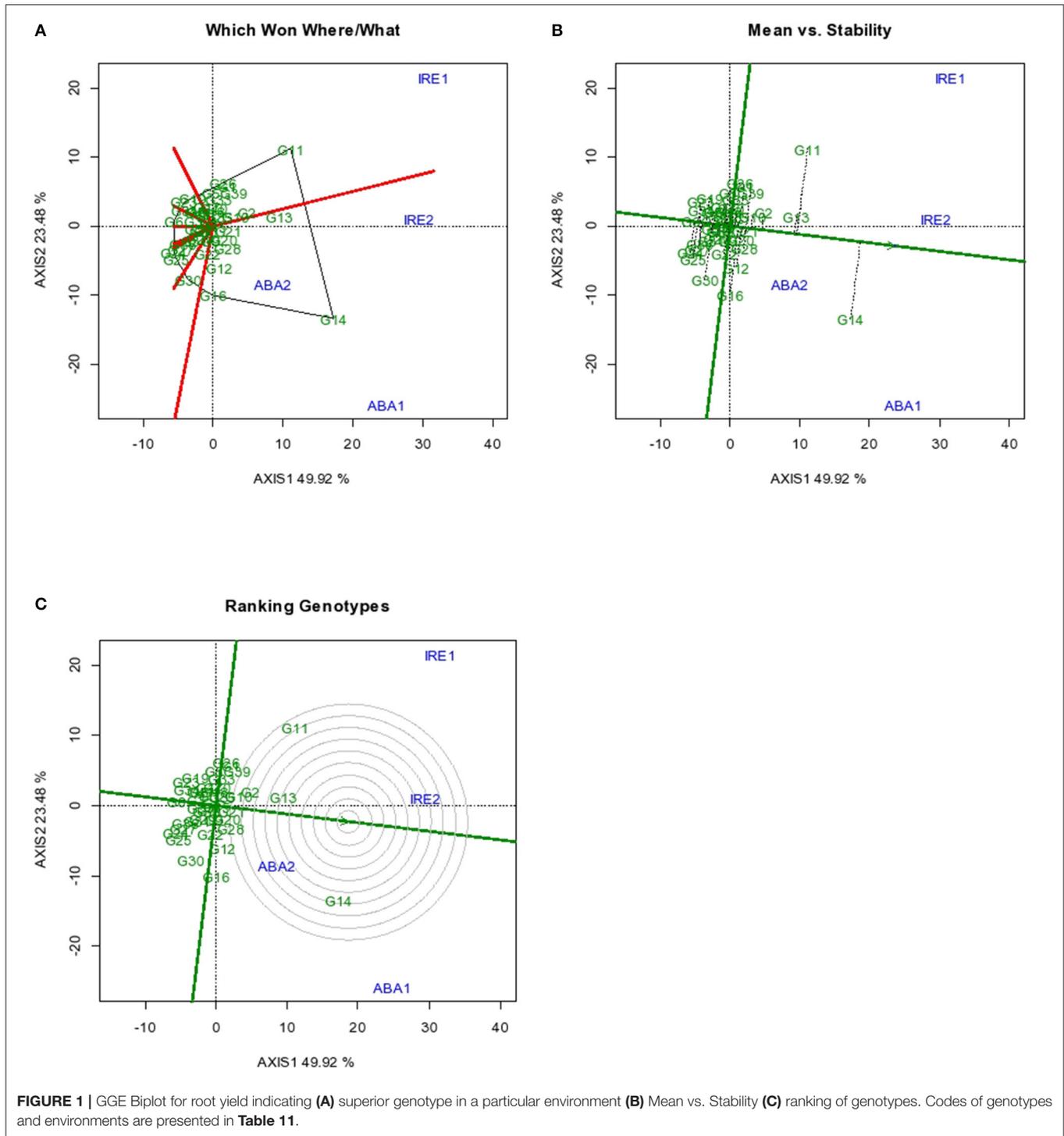
Genotype	DM	Crude protein	Fat	Crude fiber	Ash	Carbohydrate	TCC (mg/100 g)
MOTHER DELIGHT	26.40	1.90	0.27	0.39	1.20	22.64	5.55
F2M6/17	24.07	2.17	0.18	0.51	1.26	19.95	5.31
KING J	27.63	2.29	0.19	0.52	1.28	23.35	4.30
F1M1/64	26.80	2.33	0.18	0.50	1.26	29.53	3.28
F2M1/31	27.49	1.70	0.24	0.50	1.32	31.72	3.23
F2M6/1	25.60	1.86	0.28	0.43	1.14	37.89	3.11
PROGENY 3	38.94	1.94	0.18	0.47	1.28	35.08	2.94
PYT/F2M5/5	41.87	1.64	0.19	0.53	1.29	38.21	2.78
UMUSPO2/95	30.52	2.12	0.27	0.42	1.18	26.54	2.47
F5M1/3	37.84	1.90	0.20	0.52	1.26	33.96	2.37
MD/12	30.84	2.16	0.17	0.49	1.25	26.77	2.30
UMUSP 2/02	42.01	2.53	0.21	0.46	1.27	37.55	2.24
F2M5/13	37.42	2.14	0.24	0.48	1.21	36.36	2.21
F2M1/18	30.53	1.60	0.25	0.38	1.24	30.06	2.09
OP /87/0087	25.97	2.05	0.18	0.51	1.31	24.92	1.98
F1M1/23	41.60	2.16	0.19	0.51	1.27	34.06	1.67
F5M1/3	43.43	1.77	0.28	0.42	1.21	39.75	1.64
F2M1/22	40.75	1.93	0.30	0.42	1.21	36.89	1.60
MD/03	34.61	1.86	0.27	0.36	1.23	30.90	1.53
MD/23	34.42	2.15	0.21	0.50	1.25	30.31	1.45
F2M6/20	35.90	1.75	0.25	0.39	1.27	32.24	1.37
EX IGBARIAM/26	29.81	1.87	0.21	0.49	1.19	26.05	1.24
F2M1/14	36.39	1.77	0.18	0.53	1.28	36.63	1.23
TIS 87/0087/01	36.98	2.67	0.23	0.54	1.30	32.25	1.14
CL2/F2M1/35	43.85	2.47	0.31	0.40	1.22	39.45	0.94
TIS 87/0087/25	40.87	2.50	0.26	0.37	1.19	36.55	0.93
F2M6/27	35.72	2.11	0.22	0.47	1.25	29.67	0.91
F2M5/9	31.54	1.94	0.27	0.38	1.17	27.78	0.89
PYT/F2M1/35	43.74	2.33	0.27	0.37	1.22	39.55	0.79
F2M1/21	40.72	2.13	0.23	0.47	1.26	36.64	0.77
SOLO-1/21	37.72	2.14	0.31	0.33	1.14	33.80	0.71
EX-IGBARIAM/22	35.28	2.47	0.25	0.53	1.27	30.76	0.70
SOLO-ABUJA/12	39.84	1.81	0.26	0.42	1.16	26.20	0.68
F1M1/57	38.53	2.09	0.25	0.43	1.17	34.60	0.68
TIS 87/0087/23	39.49	1.84	0.21	0.39	1.18	35.88	0.62
TIS 87/0087/07	41.04	1.54	0.28	0.40	1.27	37.55	0.61
TIS 87/0087/03	36.09	1.95	0.26	0.37	1.23	32.28	0.53
EX-OYUNGA/17	40.30	1.90	0.22	0.48	1.24	36.47	0.48
SOLO-1/165	35.95	2.05	0.25	0.47	1.16	32.02	0.47
SOLO-1/88	42.52	1.54	0.29	0.41	1.20	38.08	0.18
Mean ± SD	35.78 ± 5.71	2.03 ± 0.28	0.24 ± 0.04	0.45 ± 0.06	1.23 ± 0.05	32.52 ± 5.06	1.75 ± 1.27
F-LSD _{0.05}	0.29	0.05	0.03	0.03	0.03	0.28	0.02

DM, dry matter content; TCC, Total carotenoid content.

carbohydrate range but appreciably higher than 21 and 25% for nine orange-fleshed sweet potato varieties grown in Bangladesh by Mohammad et al. (2016).

There was high variation among the genotypes with respect to total carotene content. The orange fleshed sweet potato recorded among the genotypes had the highest amount of total carotene content. The amount of total carotenoid content of

the roots ranged from 0.18 to 5.55 mg. Highest total carotene content was observed in Mother Delight with 5.55 mg/100 g (fresh weight basis), followed by F2M6/17 (5.31 mg/100 g) which are orange fleshed, and was lower in genotype Ex- Oyunga/17 (0.48 mg/100 g), Solo-1/165 (0.47 mg/100 g), and Solo-1/88 (0.18 mg/100 g). This result agrees with the findings of Adepoju and Adejumo (2015), Mohammad et al. (2016), and Islam et al. (2016)



but inconsistent with Donado-Pestana et al. (2012) who reported 0.390 to 8.823 mg/100 g.

CONCLUSION

The current study determined the magnitude of genotype-by-environment interaction and stability for storage root yield

and root *Cylas* severity. Since most sweet potato breeding programs are often tailored toward the development of high yielding, biotic and abiotic resistance and/or tolerance, this work has identified G14 as a high yielding and *Cylas* spp. tolerant genotype that is stable for both traits. These combined attributes will make it possible for its deployment across many environments. Genotypes such as G11 and G13 could

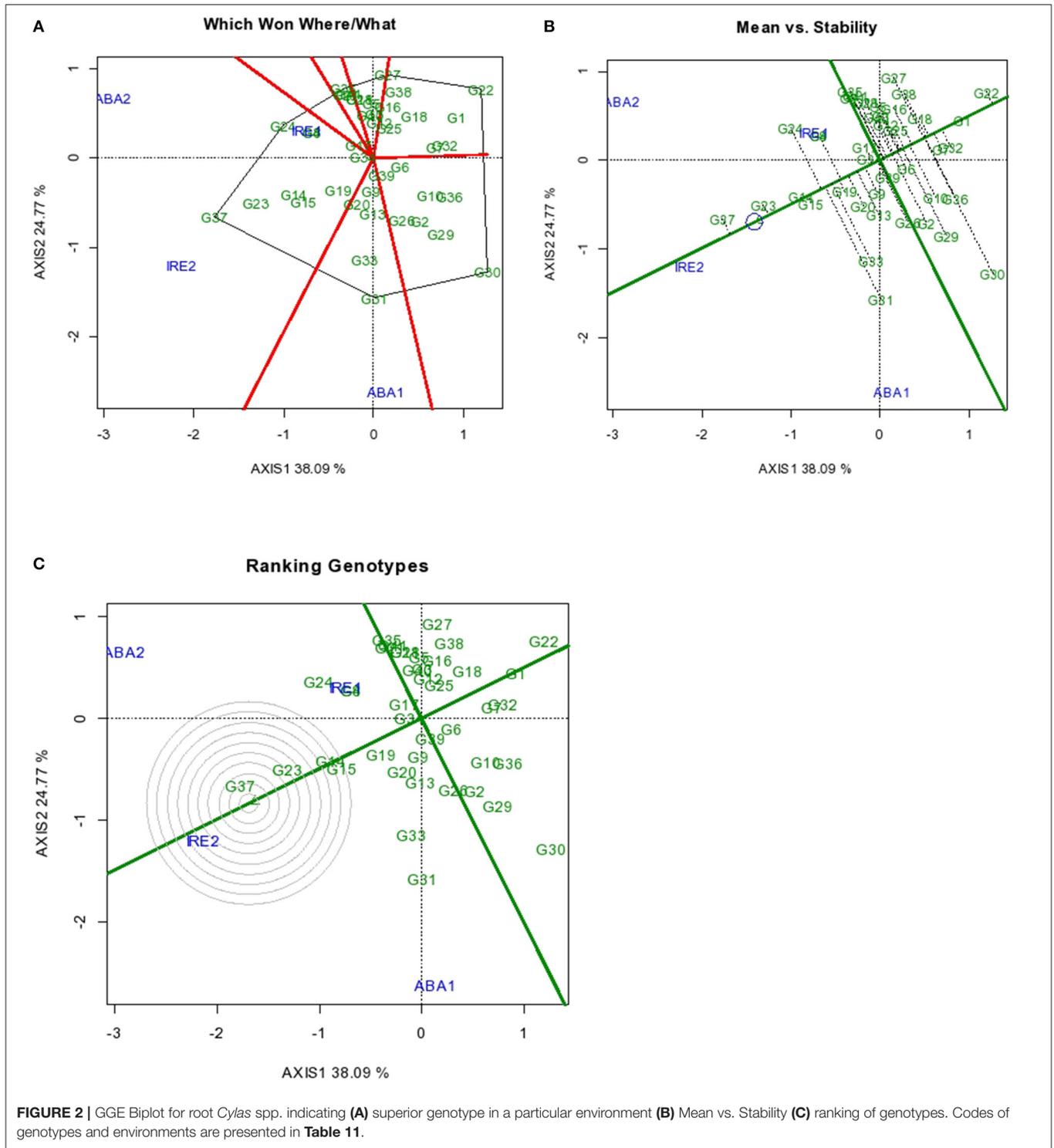
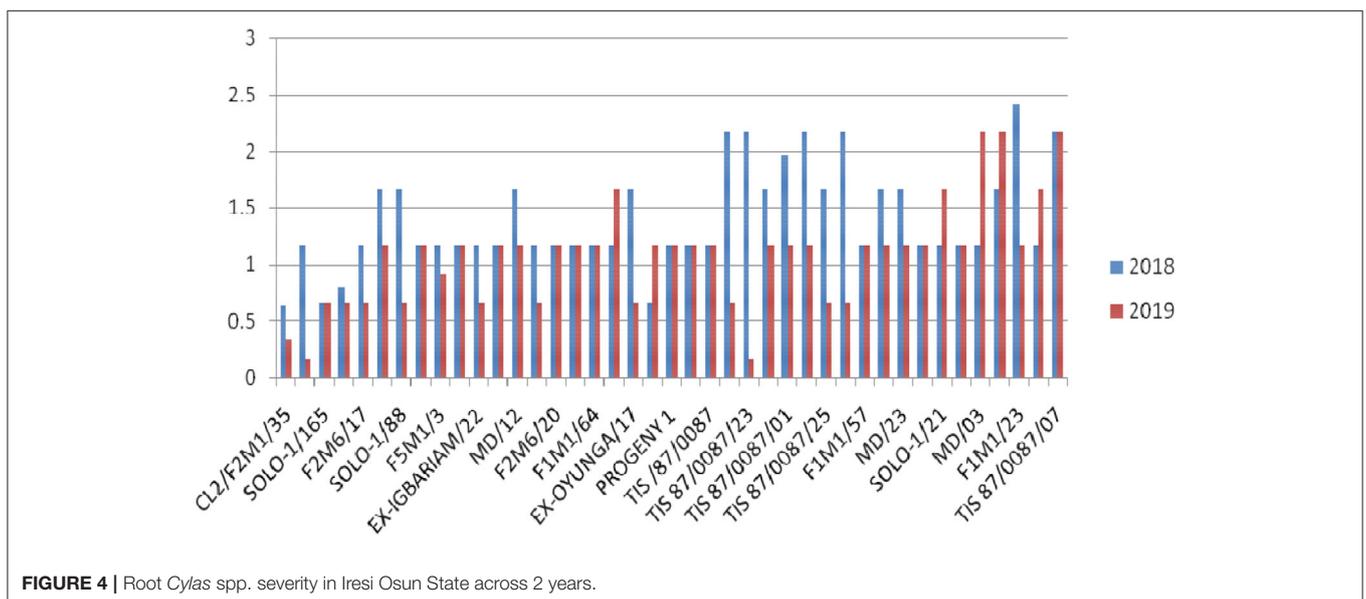
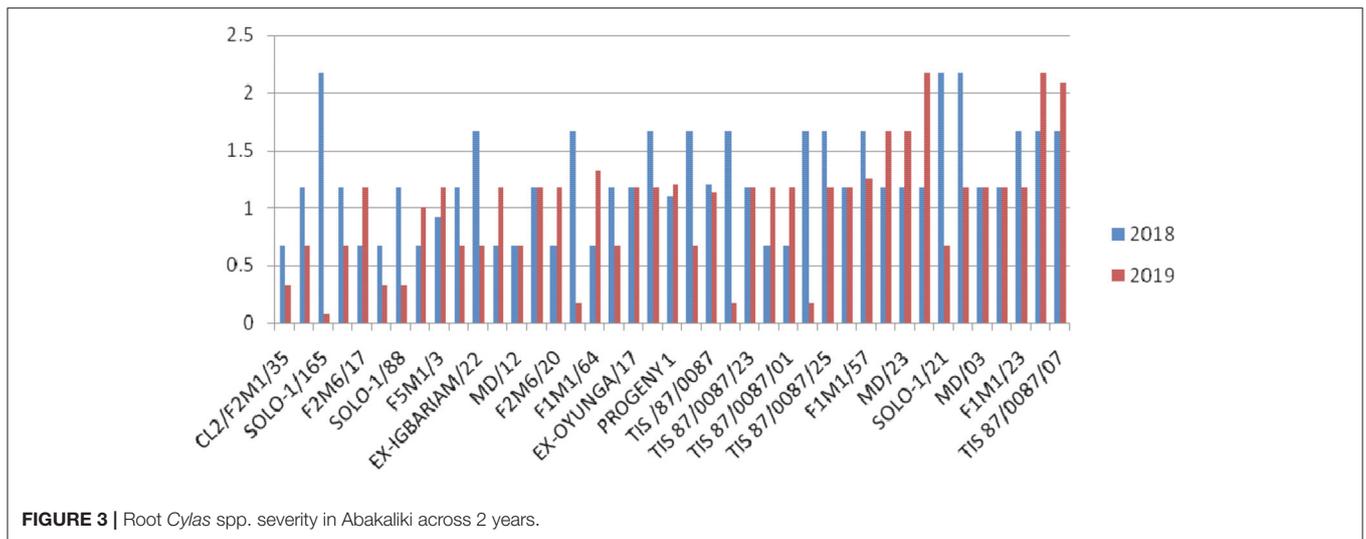


FIGURE 2 | GGE Biplot for root *Cylas* spp. indicating (A) superior genotype in a particular environment (B) Mean vs. Stability (C) ranking of genotypes. Codes of genotypes and environments are presented in Table 11.

be cultivated in environments with low *Cylas* spp. pressure as they are highly productive but exhibit low *Cylas* spp. tolerance. Genotypes G24, G37, and G31, which had average yield but high and stable tolerance to *Cylas* damage, could be used as elite germplasm toward the development of new varieties that combine high yield and *Cylas* tolerance.

In general, the knowledge and extent of the genotype-by-environment interaction (GEI) provided by this study would assist breeders in deploying limited resources in the right varietal development cycle. This study suggests that deploying resources into conducting genotype testing in many locations and years would be a worthwhile investment.



DATA AVAILABILITY STATEMENT

The original contributions presented in the study are included in the article/supplementary material, further inquiries can be directed to the corresponding author/s.

AUTHOR CONTRIBUTIONS

All authors listed have made a substantial, direct and intellectual contribution to the work, and approved it for publication.

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Conflict of Interest: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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