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# AMMI and GGE Analyses of Soyabean (*Glycine max* L. Merrill) Genotypes Infected and Uninfected with *Cucumber mosaic virus*

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## Authors' contributions

This work was carried out in collaboration among all authors. Author MTS designed the study, performed the statistical analysis, wrote the protocol and wrote the first draft of the manuscript. Authors AAA, CJA and KEO managed the analyses of the study. Author ACW edited and managed the literature searches. All authors read and approved the final manuscript.

## Article Information

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Original Research Article

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# ABSTRACT

Soyabean is an important source of protein for millions of people in developing countries. However, infection by *Cucumber mosaic virus* (CMV) causes devastating losses. Cultivation of resistant varieties has been identified as the best management strategy in many crops. The present study was, therefore, conducted to identify soyabean genotypes with high stability for growth and seed weight under CMV and disease-free conditions. Thus, eight soyabean genotypes were evaluated as CMV-infected and uninfected, using completely randomised design replicated five times and set up in the screenhouse at the School of Agriculture and Agricultural Technology, Federal University of Minna, (lat.9°40 \cdot N;long 6°30 \cdot E at an altitude of 220 m.a.s.l), Nigeria in 2018. Soyabean seedlings were infected with the virus by sap transmission at 10 days after sowing (DAS). Additive Main Effects and Multiplicative (AMMI) analyses of the evaluated parameters for growth and seed weight of the test genotypes showed that environments' effects -infected and uninfected- were

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significant (*p*<0.05). They accounted for 100% Genotype × Environment (G×E) interaction. Disease-free soyabean plants enhanced significantly higher growth and seed weight than the CMV-infected plants. The AMMI and Genotype main effects (G) plus Genotype×Environment (GGE) analyses showed that TGX 1993-4FN was the genotype with the greatest stability for leaf diameter, leaf length, number of leaves per plant, number of days to flowering and seed weight. It is recommended that, the soyabean genotype TGX 1993-4FN, can be exploited for breeding purposes and strategies that will prevent CMV infection in soyabean fields should so be adopted by farmers.

Keywords: AMMI biplot; CMV; GGE biplot; seed weight; soyabean; stability.

# **1. INTRODUCTION**

Food is an important basic need for human survival and in developing countries, ensuring food sufficiency has been difficult for several decades [1]. Inadequate intake of protein-rich food sources further worsens food crisis in the West African sub-region [1]. Soyabean (*Glycine max* L. Merrill) [2], as an annual crop is the most important legume cropped worldwide, with important roles in human and animal nutrition, besides broad industrial applications [3]. It is also one of the major sources of high quality and inexpensive protein for human consumption [4,5].

According to FAO [6], the global soyabean output in 2017 was approximately 352.6 million tonnes, with about 3.1 million tonnes from Africa. Nigeria with about 0.7 million tons, accounted for 23.3% of the total for Africa. Being a leguminous crop, soyabean plays an important role in biological nitrogen fixation (BNF) into the soil [7]. The ability of soyabean to increase soil nitrogen is aided by the activity of symbiotic bacteria [8]. Studies have shown that soyabean represents 77% of the total nitrogen fixed by crop legumes by fixing 16.4TgN per annum [9]. This is a major benefit in African farming systems, where there is a serious problem of soil infertility and application of inorganic fertilizer is constrained by high cost and scarcity of supply. The legume has been rated as the highest contribution of biological nitrogen fixation (BNF) among the grain producers, with reports of rates of up to 450 kgNha<sup>-1</sup> [9].

The United States and South American countries account for most of the world grain production, but cropped areas in "low-income food-deficit countries" are increasing, highlighting an important role for soyabean as a protein source for impoverished populations [10].

Soyabean can be processed into soya milk, soya meat, bread and oil [11]. Soyabean seeds are also used in formulation of livestock, fish and

poultry feeds while its haulms are a good source of fodder in the livestock industry [12,13].

The crop is well adapted to tropical, subtropical and temperate climates. However, its production is threatened by bacterial, fungal and virus diseases [14]. The economically important viruses infecting soyabean include *Cucumber mosaic virus* (CMV), *Cowpea aphid-borne mosaic virus* (CABMV), *Soybean mosaic virus* (SMV), and *Bean yellow mosaic virus* (BYMV).

Cucumber mosaic virus is a member of the genus Cucumovirus in the family Bromoviridae [15], and being one of the most common plant viruses, is known to infect more than 1,300 species across the world [16]. Cucumber mosaic virus is aphid transmitted in a stylet-borne nonpersistent manner [17] and is seed borne in some hosts [18,19]. It has a wide host range and causes significant losses in several crops. Cucumber mosaic virus, a single stranded RNA (ssRNA) virus, contains about 30 nm icosahedral particles with a tripartite genome encapsidated in three distinctive particles. There are numerous strains of CMV worldwide with variety of symptoms [20]. Visible symptoms in vulnerable plants include leaf chlorosis, mosaic, vein necrosis and stunting. The virus can be managed through application of insecticides to curtail its aphid vectors. Other measures employed to manage CMV include the use of healthy soyabean seeds but the most ecologically sound and sustainable approach is the cultivation of resistant soyabean varieties.

Genotype × environment (G×E) interaction can be computed using Additive Main Effects and Multiplicative Interaction (AMMI) [21,22,23,24, 25]. On the other hand, Genotype main effects (G) plus Genotype x Environment (GGE) interaction biplotsare a modification of the AMMI model [26]. The AMMI analysis is a two-stage process: Analysis of Variance (ANOVA) and Principal Components Analysis (PCA) of the ANOVA adjusted means. In the PCA, G×E interaction is partitioned into IPCA (I for interaction) with the first component accounting for the greatest variation. The efficiency of AMMI and GGE is enhanced by the graphical representation of the output expressed as biplots. A biplot gives a better understanding of the genotypes with specific or broad adaptability and environments which elicit strong (or weak) interactive forces. Although interpretation of AMMI biplot is similar to the GGE biplot, the latter provides information on total genetic variation by approximating the joint effects of the genotypes and G×E interaction [27]. Identification of soyabean genotypes with stable growth and seed weight under CMV endemic and diseasefree conditions will be useful for breeding CMV resistant soyabean varieties. This is because CMV is difficult to manage, being extremely of broad natural host range, and its ability to be transmitted in a non-persistent manner by more than 60 species of aphids [28]. The only practical solution to these problems is the incorporation of host-plant resistance into soyabean. Therefore, this study was conducted to identify soyabean genotypes with high stability for growth and seed weight under CMV and disease-free conditions for use in hybridization studies to develop high yielding and CMV resistant soyabean varieties.

# 2. MATERIALS AND METHODS

# 2.1 Study Location

The study was conducted at the Teaching and Research Farm, School of Agriculture and Agricultural Technology, Federal University of Technology, Minna, Nigeria(9° 40' N and 6°30' E; 220m.a.s.l). The site is located in the Southern Guinea Savanna with a mean annual rainfall of 1200 mm. The rainy season normally spans between April and October. The major crops cultivated in Minna include soyabean, cowpea, groundnut, rice, maize, sorghum, millet and rice. Soyabean may be grown as a sole crop or intercropped with maize or sorghum.

# 2.2 Treatments and Experimental Layout

Treatments consisted of eight soyabean genotypes viz: TGX 1448-2A, TGX 1951-3F, TGX 1987-10F, TGX 1993-4FN, TGX 1994,TGX 2017-6E, TGX 2023-1E and TGX 2025-6E obtained from the Genetic Resources Unit of the National Cereals Research Institute (NCRI), Badeggi, Niger State, Nigeria. The soyabean genotypes were selected from those designated for screening against biotic and abiotic stresses in the country. The experiment was conducted under screen house conditions using completely randomised design with five replications.

# 2.3 Sowing and Seedling Inoculation

Plastic pots with 30 cm diameter and 23 cm deep were filled with heat sterilized loamy soil. Soyabean seeds were sown on 23rd August, 2018. An isolate of CMV-infected soyabean leaves obtained from the stock in the Department of Crop Production, Federal University of Technology, Minna was used for inoculation. Virus inoculum was prepared by grinding (1g/mL) the CMV-infected soyabean leaves in inoculation buffer containing 0.1M sodium phosphate dibasic, 0.1M potassium phosphate monobasic, 0.01M ethylene diamine tetraacetic acid and 0.001M L-cysteine per litre of distilled water, adjusted to pH 7.2. One  $\mu$ L of  $\beta$ - mercapto ethanol was then added.

At 10 days after sowing (DAS), the upper leaf surface of the soyabean seedlings was dusted with carborundum powder (600-mesh) and the virus extract was rubbed on the dusted leaf surface. Distilled water was applied on the inoculated plants and they were observed for symptom development, growth and seed weights. Uninoculated plants of each soyabean genotype were evaluated in a separate screenhouse to serve as control.

# 2.4 Data Collection and Analysis

Both the CMV-infected and uninoculated plants were monitored and data collected on height, leaf diameter, leaf length, number of leaves per plant, number of days to flowering and seed weight per plant. Plant height was measured with a metre rule from ground level to the highest leaf and the mean heights per pot of the tagged plant were recorded. The leaf lengths of the tagged plants were also measured with a metre rule from the base to the tip of each leaf. The number of leaves per plant was determined using a hand operated tally counter where each mean total count per plant was recorded. Seed weight for each genotype was taken at harvest after threshing. Data were subjected to analysis of variance (ANOVA) at 5% probability level. Determination of genotype stability was based on AMMI and GGE analyses, using Breeding Management software [29]. In the analyses, infected and uninfected plants were designated as two different environments - diseased and disease-free-. From AMMI biplot, the closest genotype to the axis origin was considered to be the most stable. As for GGE biplot, genotype with the shortest vector projection relative to the biplot origin was rated as the most stable. The ecovalence method of [30] was used for stability coefficients determination and the genotype with the lowest stability coefficient was considered as the most stable.

# 3. RESULTS

## 3.1 Growth and Seed Weight Variability

The plants infected with CMV exhibited leaf chlorosis, mosaic and reduced vigour, whereas uninfected plants were apparently healthy. Apart from number of days to flowering and seed weight, genotypic effects were not significant (p>0.05) in all the evaluated parameters. On the other hand, the effects of environments, that is, infected and uninfected were significant (p < 0.05) Table 1). Combined mean heights forinfected and uninfected varied from 27.7 cm for genotype TGX 2025-6E to 33.2 cm for genotype TGX 1448-2A. However, the grand mean height of infected plants of 26.3 cm was significantly (p<0.05) lower than the grand mean of uninfected plants of 33.1 cm. Considering the infected plants alone, plant height varied between 22.7 cm for genotype TGX 1993-4FN and 30.7 cm for genotype TGX 1448-2A. The mean heights of genotypes TGX 1448-2A of 30.7 cm, TGX 1951-3Fof 28.0 cm, TGX 1987-10F of 26.7 cm and TGX 1994 of 28.7 cm were higher than the grand mean of 26.3 cm. In contrast, the heights of uninfected plants ranged between 29.7 cm for TGX 1987-10F and 36.7 cm for TGX 1951-3F (Table 2). As observed in TGX 1951-3F with 36.7 cm tall plants, the genotypes TGX 1448-2A with 35.7 cm, TGX 1993-4FN with 33.7 cm and TGX 1994 with 34.7 cm had higher mean heights than the grand mean of 33.1 cm (Table 2).

The infected plants produced narrow and deformed leaves contrary to the broad and normal shaped leaves from uninfected plants. Combined leaf diameter means varied between 2.5 cm for genotype TGX 1993-4FN and 4.0 cm for genotype TGX 2017-6E (Table 2). The grand mean of leaf diameter of 3.0 cm from infected plants was significantly (p<0.05) lower than that of healthy plants with 3.6 cm. From the infected plants, the lowest leaf diameter was observed in genotype TGX 1993-4FN with 2.3 cm, whereas genotype TGX 2025-6E had the highest leaf

diameter of 3.7 cm. Moreover, the infected plants of genotypes TGX 1987-10F with 3.3 cm, TGX 2023-1E with 3.3 cm and TGX 2025-6E with 3.7 cm exhibited higher leaf diameter than the grand mean with 3.0 cm for the group. Conversely, the leaf diameter of uninfected plants varied between 2.7 cm for TGX 1993-4FN and 5.0 cm for TGX 2017-6E. In addition to genotype TGX 2017-6E with 5.0 cm tall plants, the uninfected plants of genotypes TGX 1987-10F with 3.7 cm, TGX 2023-1E with 4.0 cm and TGX 2025-6E with 3.7 cm tall plants had wider leaf diameter than the grand mean of 3.6 cm (Table 2).

Infection of the soyabean plants with CMV resulted in reduced leaf length. Combined means of leaf length ranged from 5.5 cm in genotype TGX 1987-10F to 7.2 cm in genotype TGX 2025-6E (Table 2). The grand mean of leaf length from infected plants of 5.8 cm was significantly (p<0.05) lower than that of healthy plants of 6.7 cm length. As for the infected plants, the lowest leaf length was observed in genotype TGX 1987-10F with 5 cm, whereas the highest length came from TGX 2025-6E with 6.7 cm. Genotypes TGX 2025-6E, TGX 1994 and TGX 2017-6E had the same length of 6.0 cm while genotype TGX 2023-1E produced higher leaf length of 6.3 cm than the grand mean of 5.8 cm.

The leaf length of uninfected plants varied between 6.0 cm in genotypes TGX 1951-3F, TGX 1987-10F and TGX 2017-6E, and 7.7 cm in genotype TGX 2025-6E. Besides genotype TGX 2025-6E, uninfected plants of genotypes TGX 1993-4FN with 7.0 cm, TGX 1994 with 7.3 cm and TGX 2023-1E also with 7.3 cm produced higher leaf lengths than the grand mean of 6.7 cm for the group.

Cucumber mosaic virus infection lowered leaf production (Table 3). Combined number of leaves varied from 38 to 47 per plant in TGX 1987-10F and TGX 1951-3F, respectively. The grand mean number of leaves per plant from infected plants of 40 leaves was significantly (p < 0.05) lower than that of uninfected plants with 45 leaves. Considering the infected plants alone, genotype TGX 1987-10F produced the lowest number of leaves per plant of 36 leaves. In contrast, genotypes TGX 1994, TGX 2017-6E and TGX 2025-6E produced the highest number of leaves per plant of 42 leaves. These three genotypes were the only ones with higher number of leaves than the grand mean of 40 leaves for the group (Table 2). With respect to uninfected plants, a range of 40 in genotype TGX 1987-10F to 53 leaves in genotype TGX 1951-3F

was observed per plant. The genotypes which produced higher number of leaves than the grand mean of 45 leaves were TGX 1951-3Fwith 53 leaves, TGX 2017-6Ewith 46 leaves and TGX 2025-6E with 47 leaves.

Generally, flowering of uninfected plants was earlier than those infected with CMV (Table 3). Combined data revealed that time of flowering varied between 35 days in genotype TGX 1951-3F and 39 days in genotypes TGX 2017-6E and TGX 2025-6E after inoculation. The grand mean time of flowering in uninfected plants of 36 DAS was significantly (p<0.05) lower than that of infected plants of 38 DAS.

Taking the infected plants alone, time of flowering was observed between 36 days in genotype TGX1951-3F and 40 days in genotype TGX 2017-6E after inoculation. With the exception of genotypes TGX 1987-10F, TGX 1994 and TGX 2025-6E which flowered in 39 days and TGX 2017-6E which flowered in 40 days, all other genotypes exhibited lower days to flowering than the grand mean of 38 days for the group.





Note: G1=TGX 1448-2A; G2=TGX 1951-3F; G3=TGX 1987-10F; G4=TGX 1993-4FN; G5=TGX 1994; G6=TGX 2017-6E; G7=TGX 2023-1E; G8=TGX 2025-6E

Table 1. Mean squares of the growth and seed weights from soyabean genotypes infected an	d
uninfected with Cucumber mosaic virus	

Source of variation	DF	Mean square						
		Plant height	Leaf diameter	Leaf length				
Genotypes	7	31.4	1.4	2.1				
Environments	1	553.5*	3.5*	11.0*				
Sensitivities?	7	8.9	0.6	0.4				
Residual	32	44.3	0.6	0.9				
Total	47	48.0	0.8	1.2				
Source of variation	DF	Leaves per plant	Days to flowering	Seed weight per plant				
Genotypes	7	41.3	9.4	3.4*				
Environments	1	374.1*	38.5	8.1*				
Sensitivities	7	16.7	1.6	0.1				
Residual	32	22.8	2.3	0.3				
Total	47	32.1	4.0	0.9				

As for uninfected plants, flowering was earliest at 35 days in genotypes TGX 1951-3F, TGX 1993-4FN and TGX 2023-1E. These three genotypes exhibited lower time of flowering than the grand mean of 36 days for the group. Next were genotypes TGX 1448-2A, TGX 1993-4FN and TGX 2023-1E which flowered at 36 DAS. On the other hand, genotypes TGX 2017-6E and TGX 2025-6E flowered at 37 and 39 DAS, respectively.

Combined seed weights varied between 1.3 g per plant in genotype TGX 1448-2A and 3.5 g per plant. in genotype TGX 1993-4FN (Table 3). The grand mean of seed weight from uninfected plants of 2.3 gper plant was significantly (p<0.05) higher than that of infected plants of 1.5 g per plant. From the infected plants, genotypes TGX 1993-4FN and TGX 2025-6E with 3.2 and 2.3 g per plant, respectively were the only genotypes whose seed weights were higher than the grand mean of 1.3 g per plant. As for uninfected plants, the lowest seed weight was observed in genotype TGX 1994 with 1.8 g per plant, whereas genotype TGX 1993-4FN with an average of 3.7 g per plant was the highest. Besides genotype TGX 1993-4FN, the seed weights of TGX 2017-6E of 2.4 g and genotype TGX 2025-6E of 2.9 g were also higher than the grand mean for the group of 2.3 g.

# 3.2 Growth and Seed Weight Stability

None of the genotypes exhibited consistent stability for the entire set of parameters. Generally, the first axis (IPCA) accounted for 100% variation in all the parameters (Table 4). Additionally, the two environments (infected and uninfected) were far away from the axis origin. For plant height, AMMI analysis showed that genotype TGX 2025-6E was the closest to biplot origin, followed bygenotypes TGX 1994 and TGX 2017-6E, whereas the remaining genotypes were far away (Fig. 1a). From GGE biplot, uninfected plants or disease-free environment elicited a longer vector along the axis. The genotype TGX 2025-6E exhibited the shortest vector projection to the biplot origin, followed by TGX 1994 and TGX 2017-6E (Fig. 1b). Wricke's stability analysis indicated that genotype TGX 2025-6E had the lowest stability coefficient of 0.147, followed by TGX 1994 and TGX 2017-6E which gave stability coefficient of 0.313 and 0.383, respectively (Table 5). With respect to leaf diameter, genotype TGX 1448-2A was the closest to the AMMI biplot origin, followed by TGX 2023-1E (Fig. 2a). In GGE analysis,

diseased environment or infected plants, gave longer vector projection relative to the biplot origin. In all, genotype TGX 1448-2A exhibited the shortest vector projection, followed by TGX 2023-1E (Fig. 2b). Moreover, both genotypes had the lowest stability coefficient of 0.008. Next to them were genotypes TGX 1987-10F, TGX 1993-4FN and TGX 1994 with uniform stability coefficient of 0.022 (Table 5).

For the leaf length, 50% of the evaluated genotypes, made up of genotypes TGX 1448-2A, TGX 1951-3F, TGX 1987-10F and TGX 1993-4FN- were the closest to AMMI biplot origin (Fig. 3a). GGE analysis revealed that uninfected plants or disease-free environment produced longer vector projection along the axis. Genotypes TGX 1448-2A, TGX 1951-3F, TGX 1987-10F and TGX 1993-4FN exhibited relatively shorter vector projections compared to the remaining genotypes (Fig. 3b), with an equal stability coefficient of 0.001 (Table 5).

With respect to leaf production, the location of genotype TGX 1993-4FN was exactly on the AMMI biplot origin, whereas genotype TGX 2025-6E was the closest to it (Fig. 4a). From the GGE biplot, CMV infection or diseased environment encouraged longer vector projection relative to the axis origin. Genotypes TGX 1993-4FN and TGX 2025-6E exhibited relatively shorter vector projections to the biplot origin (Fig. 4b). These two genotypes TGX 1993-4FN and TGX 2025-6E gave stability coefficient of 0.003 and 0.170 respectively (Table 5).

Regarding number of days to flowering, AMMI analysis showed that genotype TGX 2023-1E was the nearest to the biplot origin. Also close to the biplot origin were genotypes TGX 1951-3F and TGX 1993-4FN (Fig. 5a). In GGE analysis, genotypes TGX 2023-1E, TGX 1951-3F and TGX 1993-4FN exhibited relatively shorter vector projections to the biplot origin. Infected plants or diseased environment exhibited longer vector projection along the axis (Fig. 5b). Wricke's analysis revealed that TGX 2023-1E had the lowest stability coefficient of 0.022, whereas genotypes TGX 2023-1E, TGX 1951-3F and TGX 1993-4FN gave a uniform stability coefficient of 0.105 (Table 5).

As for seed weight per plant, AMMI analysis indicated that genotypes TGX 1951-3F and TGX 1993-4FN were the closest to axis origin (Fig. 5a). Additionally, GGE biplot showed that CMV infection or diseased environment caused longer vector projection relative to the axis origin (Fig. 5b). The soyabean genotype TGX 1951-3F exhibited relatively shorter vector projections relative to the biplot origin, followed by genotype TGX 1993-4FN. Similarly, genotype TGX 1951-3F gave the lowest stability coefficient of 0.001,

which was closely followed by genotype TGX 1993-4FN with 0.002. Other genotypes with relatively low stability coefficients were TGX 2017-6E and TGX 2023-1E with 0.003 and TGX 1994 with 0.004 (Table 5).



Fig. 2. AMMI plot of genotype and environment means against the first IPCA scores (a) and GGE biplot (b) of the leaf diameter in soyabean genotypes infected and uninfected with *Cucumber mosaic virus* 







Note: G1=TGX 1448-2A; G2=TGx 1951-3F; G3=TGx 1987-10F; G4=TGX 1993-4FN; G5=TGX 1994; G6=TGX 2017-6E; G7=TGX 2023-1E; G8=TGX 2025-6E

Plant height (cm)			Leaf diameter (cm)			Leaf length (cm)			
Genotype	Infected	Uninfected	Combined	Infected	Uninfected	Combined	Infected	Uninfected	Combined
TGX 1448-2A	30.7	35.7	33.2	2.7	3.3	3.0	5.3	6.3	5.8
TGx 1951-3F	28.0	36.7	32.3	3.0	3.0	3.0	5.3	6.0	5.7
TGx 1987-10F	26.7	29.7	28.2	3.3	3.7	3.5	5.0	6.0	5.5
TGX 1993-4FN	22.7	33.7	28.2	2.3	2.7	2.5	5.3	7.0	6.2
TGX 1994	28.7	34.7	31.7	3.0	3.3	3.2	6.0	7.3	6.7
TGX 2017-6E	24.7	32.3	28.5	3.0	5.0	4.0	6.0	6.0	6.0
TGX 2023-1E	25.0	30.7	27.8	3.3	4.0	3.7	6.3	7.3	6.8
TGX 2025-6E	24.0	31.3	27.7	3.7	3.7	3.7	6.7	7.7	7.2
Grand mean	26.3	33.1*		3.0	3.6*		5.8	6.7*	

Table 2. Plant height, leaf diameter and leaf length from soyabean genotypes infected and uninfected with Cucumber mosaic virus

\*Significant at p≤0.05

# Table 3. Number of leaves per plant, days to fruiting and seed weight per plant in soyabean genotypes infected and uninfected with Cucumber mosaic virus

	Number of leaves per plant			Days to flowering			Seed weight per plant (g)		
Genotypes	Infected	Uninfected	Combined	Infected	Uninfected	Combined	Infected	Uninfected	Combined
TGX 1448-2A	38	43	41	37	36	36	0.7	1.9	1.3
TGx 1951-3F	40	53	47	36	35	35	1.2	2.0	1.6
TGx 1987-10F	36	40	38	39	36	38	0.9	1.9	1.4
TGX 1993-4FN	39	45	42	37	35	36	3.2	3.7	3.5
TGX 1994	42	43	43	39	36	38	1.1	1.8	1.5
TGX 2017-6E	42	46	44	40	37	39	1.5	2.4	2.0
TGX 2023-1E	38	44	41	37	35	36	1.0	1.9	1.5
TGX 2025-6E	42	47	45	39	39	39	2.3	2.9	2.6
Grand mean	40	45*		38*	36		1.5	2.3*	

\*Significant at p≤0.05

	Sum of square					
Source of variation	DF	Plant height	Leaf diameter	Leaf length		
Genotypes	7	73.2	3.3	5.0		
Environments	1	184.5	1.2	3.7		
Interactions	7	20.9	1.4	0.8		
IPCA 1	7	20.9	1.4	0.8		
Residuals	0	0.0	0.0	0.0		
Source of variation	DF	Leaves per plant	Days to flowering	Seed weight per plant		
Genotypes	7	96.3	21.9	8.0		
Environments	1	124.7	12.8	2.7		
Interactions	7	39.0	3.8	0.2		
IPCA 1	7	39.0	3.8	0.2		
Residuals	0	0.0	0.0	0.0		

Table 4. Additive main effects and multiplicative interaction (AMMI) of the soyabean genotype	es
infected and uninfected with Cucumber mosaic virus	











Note: G1=TGX 1448-2A; G2=TGx 1951-3F; G3=TGx 1987-10F; G4=TGX 1993-4FN; G5=TGX 1994; G6=TGX 2017-6E; G7=TGX 2023-1E; G8=TGX 2025-6E

	Stability coefficient						
Genotype	Plant height	Leaf diameter	Leaf length	Number of leaves	Days to flowering	Seed weight	
TGX 1448-2A	1.605	0.008	0.001	0.420	0.633	0.099	
TGx 1951-3F	1.758	0.147	0.001	27.503	0.105	0.001	
TGx 1987-10F	7.188	0.022	0.001	0.781	0.383	0.011	
TGX 1993-4FN	8.855	0.022	0.001	0.003	0.105	0.002	
TGX 1994	0.313	0.022	0.043	9.031	0.730	0.004	
TGX 2017-6E	0.383	1.063	0.070	0.781	0.730	0003	
TGX 2023-1E	0.633	0.008	0.251	0.281	0.022	0.003	
TGX 2025-6E	0.147	0.147	0.459	0.170	1.063	0.041	

Table 5. Stability coefficients of the growth and yield attributes in soyabean genotypes infected and uninfected with Cucumber mosaic virus



Fig. 6. AMMI plot of genotype and environment means against the first IPCA scores (a) and GGE biplot (b) of the seed weightin soyabean genotypes infected and uninfected with *Cucumber mosaic virus* 

Note: G1=TGX 1448-2A; G2=TGx 1951-3F; G3=TGx 1987-10F; G4=TGX 1993-4FN; G5=TGX 1994; G6=TGX 2017-6E; G7=TGX 2023-1E; G8=TGX 2025-6E

#### 4. DISCUSSION

Cucumber mosaic virus is a threat to several crops of economic importance around the globe [31]. The observation that there were no significant effects of genotypes in AMMI analysis was an indication of genetic similarities among the evaluated soyabean genotypes. However, the significant effects of environments underscored the need for adequate measures to prevent infection and adoption of resistant varieties by farmers. Plant height, leaf diameter, leaf length, number of leaves per plant, number of days to flowering are yield components because of their direct relationship with seed production. All these yield contributing factors were affected by CMV, in the present study, indicating the strength of the virulence of the virus on the vulnerable soyabean genotypes. The fact that all the genotypes when inoculated elicited disease symptoms indicated absence of immunity. This corroborates the findings of [32] who obtained similar result from soyabean lines that were inoculated with CMV.

Immune varieties are desirable as a preventive measure against plant pathogenic viruses but are not usually available. This is a condition that necessitates adoption of tolerant cultivars. Therefore, the soyabean genotypes studied here can be described as being tolerant to CMV. The infected genotypes did not attain maximum potentials particularly seed weight owing to impairment of the growth structures. This agrees with the findings of [33] who reported that various biochemical and physiological processes were compromised in Bunchy top virus-banana hostpathosystem. Viruses are obligate parasites that utilise their host resources including ribosomes and mitochondria for self-replication and establishment. The deleterious impacts of CMV infection as observed in this study arose from its systemic movement within the cells and tissues of the host plants. Studies have shown that systemic movement of a virulent virus is facilitated by intercellular translocation of virus particles within a host plant. This is a phenomenon that triggers host - virus interaction and the outcome is defined by their compatibility [34].

It was observed that the two environments, infected and uninfected genotypes were far away from the axis origin, indicating that they elicited strong interactive forces. This arose from the differences in genotypes' performance with respect to the parameters studied. Apart from plant height, the observation that diseased environment elicited longer vector projection along the axis revealed that it was the main factor responsible for G×E interaction. Moreover, the observed differences in stability of genotypes were the consequences of their genetic variability. The genotypes that were close to the axis origin can be described as being stable across diseased and disease-free environments. This finding agrees with report by [22] who observed similar result in their adaptability and stability study with rice genotypes in India.

Similarly, genotypes with short vector projections on the biplots exhibited high stability. In addition, the genotypes with low stability coefficients can be described as being stable for the investigated characters. This means that they maintained a uniform performance under diseased and disease-free conditions. This is also similar to reports from several other studies [22,23,24,25]. In the present study, genotype TGX 1993-4FN which was consistently the closest to the AMMI biplot origin, with the shortest vector projection on the GGE biplot, and with the lowest stability coefficients can be described as having the greatest stability.

Most genotypes were not stable for the entire growth and yield traits, probably because the genes controlling these traits are quantitatively inherited. Although polygenic or quantitative traits are desirable in plant disease management, the genes involved may not interact synergistically. Though, genotype TGX 1951-3F exhibited the lowest stability coefficient for seed weight, it was low-yielding. This will affect its acceptability to the famers. The same explanation holds for genotypes TGX 2017-6E, TGX 2023-1E and TGX 1994 which had relatively low stability coefficients but were low in seed weight and cannot be given to farmers for planting. The soyabean genotype TGX 1993-4FN with the highest seed weight per plant, combined with the highest stability for most of the quantitative traits evaluated including seed weight can be described as the most promising and which can be exploited in hybridization studies for the development of high yielding CMV resistant soyabean varieties for farmers. Nevertheless, the observation that not all the genotypes were stable for growth and seed weight shows that there is room for improvement [35].

# 5. CONCLUSION AND RECOMMENDA-TION

This study revealed the pathogenicity of CMV on the evaluated soyabean genotypes. Disease-free soyabean plants produced significantly higher growth and seed weight than the CMV-infected plants. The AMMI analysis revealed that environments' effects represented by infected and uninfected genotypes were significant (p<0.05) and accounted for 100% Genotype × Environment (G×E) interaction for growth and seed weight. The AMMI and GGE analyses showed that genotype TGX 1993-4FN was the genotype with the greatest stability for leaf diameter, leaf length, number of leaves per plant, number of days to flowering and seed weight. Therefore, the soyabean genotype TGX 1993-4FN can be exploited for breeding purposes. Pending the arrival of such resistant varieties from soyabean breeders, strategies that will prevent CMV infection in soyabean fields should be adopted by farmers.

## **COMPETING INTERESTS**

Authors have declared that no competing interests exist.

## REFERENCES

- Matemilola S, Elegbede I. The challenges of food security in Nigeria. Open Access Lib. J. 2017;4:e4185.
- Salaudeen MT, Adama CJ, Ogunsola KE, Ishaq MN, Abdullahi AA, Abdulkadir A. Resistance of soybean lines infected with *blackeye cowpea mosaic virus* under controlled conditions. Int. J. Agric. Rural Dev. 2016;19:2720-2730.
- Riaz MN. Soy applications in food. CRC Press. 2005;304. Available:https://doi.org/10.1201/97814200 37951
- Vesper H, Schmelz EM, Nikolova-Karakashian MN, Dillehay DL, Lynch DV, Merrill AH. Jr. Sphingolipids in food and the emerging importance of sphingolipids to nutrition. J. Nutr. 1999;129:1239-1250.
- Lokuruka MNI. Soybean nutritional properties: The good and the bad about soy foods consumption-a review. Afr J Food Agric Nutrition Dev. 2010;10:2439-2459.
- FAO (Food and Agriculture Organization). Soyabean; 2017. Available:http://www.fao.org/faostat/en/#da ta/QC
- Deacon J. The nitrogen cycle and nitrogen fixation. Institute of Cell and Molecular Biology. The University of Edinburg. Blackwell Scientific Publication; 1998.

- Thuita M, Vanlauwe B, Mutegi E, Masso C. Reducing spatial variability of soybean response to rhizobia inoculants in farms of variable soil fertility in Siaya County of Western Kenya. Agric. Ecosystems Environ. 2018;261:153-160.
- 9. Herridge DF, Peoples MB, Boddey RM. Global inputs of biological nitrogen fixation in agricultural systems. Plant Soil. 2008;311:1-18.
- Hungria M, Mendes IC. Nitrogen fixation with soybean: The perfect symbiosis? In: Biological nitrogen fixation, Chapter 99, Publisher: John Wiley and Sons Inc, (Ed: Francs J. de Bruijn). 2015;1005-1019. DOI: 10.1002/9781119053095ch99
- Agarwal DA, Billore SD, Sharma AN, Dupare BU, Srivastava SK. Soybean: Introduction, improvement, and utilization in India-problems and prospects. Agric Res. 2013;2:293-300.
- 12. Zehnder GW, Murphy JF, Sikora EJ, Kloepper JW. Application of rhizobacteria for induced resistance. Eur. J. Plant Pathol. 2001;107:39-50.
- Heuze V, Tran G, Noziere P, Leissive M, Lebas F. Soyabean hulls Feepedia, a Programme by INRA, CIRAD, AFZ and FAO; 2017. Available:https://www.feedpedia.org/node/ 42
- 14. Graham PH, Vance CP. Legumes: Importance and constraints to greater use. Plant Physiol. 2003;131:872-877.
- de Breuil S, Giolitti EJ, Bejerman N, Lenardon SL. Effects of Cucumber mosaic virus on the yield and yield components of peanut. J. Plant. 2012;94:669-673.
- García-Arenal F, Palukaitis P. Cucumber mosaic virus. In: García-Arenal, F, Palukaitis P, Mahy BWJ and Van Regenmortel MHV, Editors. Encyclopedia of Virology. Oxford: Academic Press. 2008;614-619.
- 17. Gildow FE, Shah DA, Sackett WM, Butzler T, Nault BA, Fleischer SJ. Transmission efficiency of *Cucumber mosaic virus* by aphids associated with virus epidemics in snapbean. Phytopathol. 2008;98:1233-1241.
- Tomlinson JA, Karter AL. Studies on the seed transmission of *Cucumber mosaic virus* in chickweed (*Stellaria media*) in relation to the ecology of the virus. Ann Appl Biol. 1970;66:381:386.
- 19. Yang Y, Kim KS, Anderson EJ. Seed transmission of *Cucumber mosaic virus*

in spinach. Phytopathol. 1997;87:924-931.

- Agrios GN. Plant pathology. Fifth Edition. Elsevier Academic Publishers Amsterdam; 2005.
- 21. Azizi A, Shams-Baksh M. Impact of *Cucumber mosaic virus* infection on the varietal traits of common bean cultivars in Iran. Virusdis. 2014;25:447-454.
- 22. Das S, Misra RC, Patnaik MC, Das SR. G
  × E interaction, adaptability and yield stability of mid-early rice genotypes. Indian J. Agric. Res. 2010;44:104-111.
- 23. Suryanarayana L, Sekhar MR, Babu DR, Ramana AV, Rao VS. Cluster and principal component analysis in maize. Int. J. Curr. Microbiol. Appl. Sci. 2017;6:354-359.
- 24. Mehrnaz T, Elham B, Ali RA, Ali RAK. Genetic diversity of 13 Maize (*Zea mays* L.) hybrids based on multivariate analysis methods. Int. J. Farming Allied Sci. 2014;3(5):467-470.
- 25. Muhammad RW, Qayyum A, Hamza A, Ahmad MQ, Naseer NS, Liaqat S, Ahmad B, Malik W, Noor E. Analysis of genetic traits for drought tolerance in maize. Gen. Mol. Res. 2015;28:13545-13565.
- Subramanian A, Subbaraman N. Hierarchical cluster analysis of genetic diversity in maize germplasm. Electronic J. Plant Breeding. 2010;1:431-436.
- Dia M, Wehner TC, Arellano C. Analysis of genotype × environment interaction (G×E) using SAS programming. Agron. J. 2016;108:1838-1852.
- 28. BMS (Breeding Management System) Breeding management system. Version

3.0.8. Integrated Breeding Platform (IBP), Mexico; 2015.

- 29. Nault BA, Shah D, Taylor AG. Viruses and aphids everywhere in New York snap bean fields in 2005. In Proceedings of the 2006 Empire State Fruit and Vegetable EXPO. Cornell Coop. Exten., Syracuse, NY. 2006;74–76.
- Wricke G. Evaluation method for recording ecological differences in field trials. Z. Pflanzenzuecht. 1962;47:92-96.
- 31. Zitter TA, Murphy JF. Cucumber mosaic. The Plant Health Instructor; 2009. DOI: 10.1094/PH-1-2009-0518-01
- Adamu AS, Salaudeen MT, Gana AS, Ishaq MN. Response of selected soybean (*Glycine max* [L.] Merr.) lines to *Cucumber mosaic virus* disease in Minna, Niger State. Nig. J. Agric. Food Environ. 2015;11:45-51.
- Anuradha C, Selvarajan R, Vasantha S, Suresha GS. Biochemical characterization of compatible plant virus interaction: A case study with *Bunchy top virus*-banana host-pathosystem. Plant Pathol. J. 2015;14:212-222.
- Pallas V, Garci´a, JA. How do plant viruses induce disease? Interactions and interference with host components. J. Gen. Virol. 2011;92:2691-2705.
- Dia M, Wehner TC, Elmstrom GW, Gabert A, Motes JE, Staub JE, Tolla GE, Widders IE. Genotype × environment interaction for yield of pickling cucumber in 24 U.S. environments. Open Agric. 2018;3:1-16.

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