

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

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. The objective of this study was to identify soyabean genotypes with high stability for growth and seed weight under CMV and disease-free conditions. 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' N; long 6°30' 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. Additive Main Effects and Multiplicative Interaction (AMMI) analysis revealed that environments' effects (infected and uninfected) were significant ($p < 0.05$) and accounted for 100 % Genotype \times Environment (G \times E) interaction for growth and seed weight. Disease-free soyabean plants produced significantly higher growth and seed weight than the CMV-infected plants. The AMMI and Genotype main effects (G) plus Genotype \times 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. In the meantime, the soyabean genotype TGX 1993-4FN can be exploited for breeding purposes and strategies that will prevent CMV infection in soyabean fields should 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 a mirage for several decades. Inadequate intake of protein-rich food sources further worsens food crisis in the West African subregion. Soyabean (*Glycine max* L. Merrill), as an

27 annual crop is one of the major sources of high quality and inexpensive protein for human
28 consumption. According to FAO [1], the global soyabean output in 2017 was approximately 352.6
29 million tonnes, with about 3.1 million tonnes from Africa. Nigeria with about 0.7 million tons,
30 accounted for 23.3 % of the total for Africa. Being a leguminous crop, soyabean plays an important
31 role in biological nitrogen fixation (BNF) into the soil. The ability of soyabean to increase soil nitrogen
32 is aided by the activity of symbiotic bacteria [2]. Studies have shown that soyabean represents 77 % of
33 the total nitrogen fixed by crop legumes by fixing 16.4TgN per annum. This is a major benefit in
34 African farming systems, where there is a serious problem of soil infertility and application of
35 inorganic fertilizer is constrained by high cost and scarcity of supply. Soyabean can be processed into
36 soya milk, soya meat, bread and oil [3]. Soyabean seeds are also used in formulation of livestock, fish
37 and poultry feeds while its haulms are a good source of fodder in the livestock industry.

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

42 *Cucumber mosaic virus* is a member of the genus *Cucumovirus* in the family *Bromoviridae* [4]. It has a
43 wide host range and causes significant losses in several crops. The virus is transmitted by aphids,
44 infected seeds and through sap inoculation. *Cucumber mosaic virus*, a single stranded RNA (ssRNA)
45 virus, contains about 30 nm icosahedral particles with a tripartite genome encapsidated in three
46 distinctive particles. There are numerous strains of CMV worldwide with variety of symptoms [5].
47 Visible symptoms in vulnerable plants include leaf chlorosis, mosaic, vein necrosis and stunting. The
48 virus can be controlled through application of insecticides to curtail its aphid vectors. Other measures
49 include the use of healthy soyabean seeds but the most ecologically sound and sustainable approach is
50 the cultivation of resistant soyabean varieties.

51 Genotype \times environment (G \times E) interaction can be computed using Additive Main Effects and
52 Multiplicative Interaction (AMMI) model. On the other hand, Genotype main effects (G) plus
53 Genotype \times Environment (GGE) interaction biplots are a modification of the AMMI model [6]. The
54 AMMI analysis is a two-stage process: Analysis of Variance (ANOVA) and Principal Components
55 Analysis of the ANOVA adjusted means. In the PCA, G \times E interaction is partitioned into IPCA (I for
56 interaction) with the first component accounting for the greatest variation. The efficiency of AMMI and
57 GGE is enhanced by the graphical representation of the output expressed as biplots. A biplot gives a
58 better understanding of the genotypes with specific or broad adaptability and environments which elicit
59 strong (or weak) interactive forces. Although interpretation of AMMI biplot is similar to the GGE
60 biplot, the latter provides information on total genetic variation by approximating the joint effects of
61 the genotypes and G \times E interaction. Identification of soyabean genotypes with stable growth and seed
62 weight under CMV endemic and disease-free conditions will be useful for breeding CMV resistant
63 soyabean varieties. Therefore, this study was conducted to identify soyabean genotypes with high
64 stability for growth and seed weight under CMV and disease-free conditions for use in hybridization
65 studies to develop high yielding and CMV resistant soyabean varieties.

66 **2. MATERIALS AND METHODS**

67 68 **2.1 Study Location**

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

75

76 **2.2 Treatments and Experimental Layout**

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

83

84 **2.3 Sowing and Seedling Inoculation**

85 Plastic pots with 30cm diameter and 23cm deep were filled with heat sterilized loamy soil. Soyabean
86 seeds were sown on 23rd August, 2018. An isolate of CMV-infected soyabean leaves obtained from the
87 stock in the Department of Crop Production, Federal University of Technology, Minna was used for
88 inoculation. Virus inoculum was prepared by grinding (1g/mL) the CMV-infected soyabean leaves in
89 inoculation buffer containing 0.1M sodium phosphate dibasic, 0.1M potassium phosphate monobasic,
90 0.01M ethylenediamine tetraacetic acid and 0.001M L-cysteine per litre of distilled water, adjusted to
91 pH 7.2. One μ L of β - mercapto ethanol was then added. At 10 days after sowing (DAS), the upper leaf
92 surface of the soyabean seedlings was dusted with carborundum powder (600-mesh) and the virus
93 extract was rubbed on the dusted leaf surface. Distilled water was applied on the inoculated plants and
94 they were observed for symptom development, growth and seed weights. Uninoculated plants of each
95 soyabean genotype were evaluated in a separate screenhouse to serve as control.

96

97 **2.4 Data Collection and Analysis**

98

99 Both the CMV-infected and uninoculated plants were observed for height, leaf diameter, leaf length,
100 number of leaves per plant, number of days to flowering and seed weight per plant. Data were
101 subjected to analysis of variance (ANOVA) at 5 % probability level. Determination of genotype

102 stability was based on AMMI and GGE analyses, using Breeding Management software [7]. In the
103 analyses, infected and uninfected plants were designated as two different environments - diseased and
104 disease-free-. From AMMI biplot, the closest genotype to the axis origin was considered to be the most
105 stable. As for GGE biplot, genotype with the shortest vector projection relative to the biplot origin was
106 rated as the most stable. Wreck's ecovalence method was used for stability coefficients determination.
107 Genotype with the lowest stability coefficient was considered as the most stable.

108 **3. RESULTS**

109 **3.1 Growth and seed weight variability**

110 The plants infected with CMV exhibited leaf chlorosis, mosaic and reduced vigour, whereas uninfected
111 plants were apparently healthy. Apart from number of days to flowering and seed weight, genotypic
112 effects were not significant ($p>0.05$) in all the evaluated parameters. On the other hand, the effects of
113 environments, that is, infected and uninfected were significant ($p<0.05$) Table 1). Combined mean
114 heights for infected and uninfected varied from 27.7 cm for genotype TGX 2025-6E to 33.2 cm for
115 genotype TGX 1448-2A. However, the grand mean height of infected plants of 26.3 cm was
116 significantly ($p<0.05$) lower than the grand mean of uninfected plants of 33.1 cm. Considering the
117 infected plants alone, plant height varied between 22.7 cm for genotype TGX 1993-4FN and 30.7 cm
118 for genotype TGX 1448-2A. The mean heights of genotypes TGX 1448-2A of 30.7 cm, TGX 1951-3F
119 of 28.0 cm, TGX 1987-10F of 26.7 cm and TGX 1994 of 28.7 cm were higher than the grand mean of
120 26.3 cm. In contrast, the heights of uninfected plants ranged between 29.7 cm for TGX 1987-10F and
121 36.7 cm for TGX 1951-3F (Table 2). As observed in TGX 1951-3F with 36.7 cm tall plants, the
122 genotypes TGX 1448-2A with 35.7 cm, TGX 1993-4FN with 33.7 cm and TGX 1994 with 34.7 cm had
123 higher mean heights than the grand mean of 33.1 cm (Table 2).

124 The infected plants produced narrow and deformed leaves contrary to the broad and normal shaped
125 leaves from uninfected plants. Combined leaf diameter means varied between 2.5 cm for genotype

126 TGX 1993-4FN and 4.0 cm for genotype TGX 2017-6E (Table 2). The grand mean of leaf diameter of
127 3.0 cm from infected plants was significantly ($p<0.05$) lower than that of healthy plants with 3.6 cm.
128 From the infected plants, the lowest leaf diameter was observed in genotype TGX 1993-4FN with 2.3
129 cm, whereas genotype TGX 2025-6E recorded the highest leaf diameter of 3.7cm. Moreover, the
130 infected plants of genotypes TGX 1987-10F with 3.3 cm, TGX 2023-1E with 3.3 cm and TGX 2025-
131 6E with 3.7 cm recorded higher leaf diameter than the grand mean with 3.0 cm for the group.
132 Conversely, the leaf diameter of uninfected plants varied between 2.7 cm for TGX 1993-4FN and 5.0
133 cm for TGX 2017-6E. In addition to genotype TGX 2017-6E with 5.0 cm tall plants, the uninfected
134 plants of genotypes TGX 1987-10F with 3.7 cm, TGX 2023-1E with 4.0 cm and TGX 2025-6E with
135 3.7 cm tall plants recorded higher leaf diameter than the grand mean of 3.6 cm (Table 2).

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

143
144 The leaf length of uninfected plants varied between 6.0cm in genotypes TGX 1951-3F, TGX 1987-10F
145 and TGX 2017-6E and 7.7cm in genotype TGX 2025-6E. Besides genotype TGX 2025-6E, uninfected
146 plants of genotypes TGX 1993-4FN with 7.0 cm, TGX 1994 with 7.3 cm and TGX 2023-1E also
147 with 7.3 cm recorded higher leaf lengths than the grand mean of 6.7 cm for the group.

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149 *Cucumber mosaic virus* infection lowered leaf production (Table 3). Combined number of leaves
150 varied from 38 to 47 per plant in TGX 1987-10F and TGX 1951-3F, respectively. The grand mean

151 number of leaves per plant from infected plants of 40 leaves was significantly ($p<0.05$) lower than that
152 of uninfected plants with 45 leaves. Considering the infected plants alone, genotype TGX 1987-10F
153 produced the lowest number of leaves per plant of 36 leaves. In contrast, genotypes TGX 1994, TGX
154 2017-6E and TGX 2025-6E produced the highest number of leaves per plant of 42 leaves. These three
155 genotypes were the only ones with higher number of leaves than the grand mean of 40 leaves for the
156 group (Table 2). With respect to uninfected plants, a range of 40 in genotype TGX 1987-10F to 53
157 leaves in genotype TGX 1951-3F was observed per plant. The genotypes which produced higher
158 number of leaves than the grand mean of 45 leaves were TGX 1951-3F with 53 leaves, TGX 2017-6E
159 with 46 leaves and TGX 2025-6E with 47 leaves.

160

161 Generally, flowering of uninfected plants was earlier than those infected with CMV (Table 3).
162 Combined data revealed that time of flowering varied between 35 days in genotype TGX 1951-3F and
163 39 days in genotypes TGX 2017-6E and TGX 2025-6E after inoculation. The grand mean time of
164 flowering in uninfected plants of 36 DAS was significantly ($p<0.05$) lower than that of infected plants
165 of 38 DAS.

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167 Taking the infected plants alone, time of flowering was observed between 36 days in genotype
168 TGX1951-3F and 40 days in genotype TGX 2017-6E after inoculation. With the exception of
169 genotypes TGX 1987-10F, TGX 1994 and TGX 2025-6E which flowered in 39 days and TGX 2017-
170 6E which flowered in 40 days respectively, all other genotypes recorded lower days to flowering than
171 the grand mean of 38 days for the group.

172

173 As for uninfected plants, flowering was earliest at 35 days in genotypes TGX 1951-3F, TGX 1993-4FN
174 and TGX 2023-1E. These three genotypes recorded lower time of flowering than the grand mean of 36

175 days for the group. Next were genotypes TGX 1448-2A, TGX 1993-4FN and TGX 2023-1E which
176 flowered at 36 DAS. On the other hand, genotypes TGX 2017-6E and TGX 2025-6E flowered at 37
177 and 39 DAS respectively.

178
179 Combined seed weights varied between 1.3 g per plant.in genotype TGX 1448-2A and 35.g per plant.
180 in genotype TGX 1993-4FN (Table 3). The grand mean of seed weight from uninfected plants of.3 g
181 per plant was significantly ($p<0.05$) higher than that of infected plants of 1.3 g per plant. From the
182 infected plants, genotypes TGX 1993-4FN and TGX 2025-6E with 3.2 and 2.3 g per plant, respectively
183 were the only genotypes whose seed weights were higher than the grand mean of 1.3 g per plant. As for
184 uninfected plants, the lowest seed weight was observed in genotype TGX 1994 with 1.8 g per plant,
185 whereas genotype TGX 1993-4FN with an average of 3.7 g per plant was the highest. Besides genotype
186 TGX 1993-4FN, the seed weights of TGX 2017-6E of 2.4 g and genotype TGX 2025-6E of 2.9 g were
187 also higher than the grand mean for the group of 2.3 g.

188 189 **3.2 Growth and seed weight stability**

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

200 1448-2A was the closest to the AMMI biplot origin, followed by TGX 2023-1E (Fig. 2a). In GGE
201 analysis, diseased environment or infected plants, gave longer vector projection relative to the biplot
202 origin. In all, genotype TGX 1448-2A exhibited the shortest vector projection, followed by TGX 2023-
203 1E (Fig. 2b). Moreover, both genotypes had the lowest stability coefficient of 0.008. Next to them were
204 genotypes TGX 1987-10F, TGX 1993-4FN and TGX 1994 with uniform stability coefficient of 0.022
205 (Table 5).

206 For the leaf length, 50 % of the evaluated genotypes made up of genotypes TGX 1448-2A, TGX 1951-
207 3F, TGX 1987-10F and TGX 1993-4FN were the closest to AMMI biplot origin (Fig. 3a). GGE
208 analysis revealed that uninfected plants or disease free environment produced longer vector projection
209 along the axis. Genotypes TGX 1448-2A, TGX 1951-3F, TGX 1987-10F and TGX 1993-4FN
210 exhibited relatively shorter vector projections compared to the remaining genotypes (Fig. 3b), with an
211 equal stability coefficient of 0.001 (Table 5).

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

218 Regarding to number of days to flowering, AMMI analysis showed that genotype TGX 2023-1E was
219 the nearest to the biplot origin. Also close to the biplot origin were genotypes TGX 1951-3F and TGX
220 1993-4FN (Fig. 5a). In GGE analysis, genotypes TGX 2023-1E, TGX 1951-3F and TGX 1993-4FN
221 exhibited relatively shorter vector projections to the biplot origin. Infected plants or diseased
222 environment exhibited longer vector projection along the axis. (Fig.5b). Wreck's analysis revealed that

223 TGX 2023-1E had the lowest stability coefficient of 0.022, whereas genotypes TGX 2023-1E, TGX
224 1951-3F and TGX 1993-4FN gave a uniform stability coefficient of 0.105 (Table 5).

225 As for seed weight per plant, AMMI analysis indicated that genotypes TGX 1951-3F and TGX 1993-
226 4FN were closest to the axis origin (Fig. 5a). Additionally, GGE biplot showed that CMV infection or
227 diseased environment caused longer vector projection relative to the axis origin (Fig. 5b). The
228 soyabean genotype TGX 1951-3F exhibited relatively shorter vector projections relative to the biplot
229 origin, followed by genotype TGX 1993-4FN. Similarly, genotype TGX 1951-3F gave the lowest
230 stability coefficient of 0.001, which was closely followed by genotype TGX 1993-4FN with 0.002.
231 Other genotypes with relatively low stability coefficients were TGX 2017-6E and TGX 2023-1E with
232 0.003 and TGX 1994 with 0.004 (Table 5).

233 **4. DISCUSSION**

234 *Cucumber mosaic virus* is a threat to several crops of economic importance around the globe. The
235 observation that there were no significant effects of genotypes in AMMI analysis was an indication of
236 genetic similarities among the evaluated soyabean genotypes. However, the significant effects of
237 environments underscored the need for adequate measures to prevent infection and adoption of
238 resistant varieties by farmers. Plant height, leaf diameter, leaf length, number of leaves per plant,
239 number of days to flowering are yield components because of their direct relationship with seed
240 production. All these yield contributing factors were affected by CMV, indicating the pathogenicity of
241 the virus on the vulnerable soyabean genotypes. The fact that all the genotypes inoculated elicited
242 disease symptoms indicated absence of immunity. This corroborates the findings of Adamu [8] who
243 obtained similar result from soyabean lines that were inoculated with CMV.

244 Immune varieties are desirable as a preventive measure against plant pathogenic viruses but are not
245 usually available. This is a condition that necessitates adoption of tolerant cultivars. Therefore, the

246 soyabean genotypes studied here can be described as being tolerant to CMV. The infected genotypes
247 did not attain maximum potentials particularly seed weight owing to impairment of the growth
248 structures. This agrees with the findings of Anuradha [9] who reported that various biochemical and
249 physiological processes were compromised in *Bunchy top virus*-banana host-pathosystem. Viruses are
250 obligate parasites that utilise their host resources including ribosome and mitochondrion for self-
251 replication and establishment. The deleterious impacts of CMV infection as observed in this study
252 arose from its systemic movement within the cells and tissues of the host plants. Studies have shown
253 that systemic movement of a virulent virus is facilitated by intercellular translocation of virus particles
254 within a host plant. This is a phenomenon that triggers host – virus interaction and the outcome is
255 defined by their compatibility [10].

256 It was observed that the two environments, infected and uninfected genotypes were far away from the
257 axis origin, indicating that they elicited strong interactive forces. This arose from the differences in
258 genotypes' performance with respect to the parameters studied. Apart from plant height, the
259 observation that diseased environment elicited longer vector projection along the axis revealed that it
260 was the main factor responsible for G×E interaction. Moreover, the observed differences in stability of
261 genotypes were the consequences of their genetic variability. The genotypes that were close to the axis
262 origin can be described as being stable across diseased and disease-free environments. Similarly,
263 genotypes with short vector projections on the biplots exhibited high stability. In addition, the
264 genotypes with low stability coefficients can be described as being stable for the investigated
265 characters. This means that they maintained a uniform performance under diseased and disease-free
266 conditions. However, genotype TGX 1993-4FN which was consistently the closest to the AMMI biplot
267 origin, with the shortest vector projection on the GGE biplot, and with the lowest stability coefficients
268 can be described as the genotype with the greatest stability.

269 Most genotypes were not stable for the entire growth and yield traits probably because the genes
270 controlling these traits are quantitatively inherited, indicating that they are under the influence of
271 several genes. Although polygenic or quantitative traits are desirable in plant disease management, the
272 genes involved may not interact synergistically. Although genotype TGX 1951-3F recorded the lowest
273 stability coefficient for seed weight, it was low-yielding. This will affect its acceptability to the farmers.
274 The same explanation holds for genotypes TGX 2017-6E, TGX 2023-1E and TGX 1994 which
275 exhibited relatively low stability coefficients but were low in seed weight and cannot be given to
276 farmers for planting. The soyabean genotype TGX 1993-4FN with the highest seed weight per plant,
277 combined with the highest stability for most of the quantitative traits evaluated including seed weight
278 can be described as the most promising and which can be exploited in hybridization studies for the
279 development of high yielding CMV resistant soyabean varieties for farmers. Nevertheless, the
280 observation that not all the genotypes were stable for growth and seed weight shows that there is room
281 for improvement [11].

282 **5. CONCLUSION AND RECOMMENDATIONS**

283 This study revealed the pathogenicity of CMV on the evaluated soyabean genotypes. Disease-free
284 soyabean plants produced significantly higher growth and seed weight than the CMV-infected plants.
285 The AMMI analysis revealed that environments' effects represented by infected and uninfected
286 genotypes were significant ($p < 0.05$) and accounted for 100 % Genotype \times Environment (G \times E)
287 interaction for growth and seed weight. The AMMI and GGE analyses showed that genotype TGX
288 1993-4FN was the genotype with the greatest stability for leaf diameter, leaf length, number of leaves
289 per plant, number of days to flowering and seed weight. Therefore, the soyabean genotype TGX 1993-
290 4FN can be exploited for breeding purposes. Pending the arrival of such resistant varieties from
291 soyabean breeders, strategies that will prevent CMV infection in soyabean fields should be adopted by
292 farmers.

293 Table 1: Mean squares of the growth and seed weights from soyabean genotypes infected and
 294 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

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

Genotype	Plant height (cm)			Leaf diameter (cm)			Leaf length (cm)		
	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*	

296 *Significant at $p \leq 0.05$

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306 Table 3: Number of leaves per plant, days to fruiting and seed weight per plant in soyabean genotypes infected and uninfected with
 307 *Cucumber mosaic virus*
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Genotypes	Number of leaves per plant			Days to flowering			Seed weight per plant (g)		
	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*	

309 *Significant at $p \leq 0.05$

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317 Table 4: Additive main effects and multiplicative interaction (AMMI) of the soyabean genotypes
 318 infected and uninfected with *Cucumber mosaic virus*
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Source of variation	DF	Sum of square		
		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

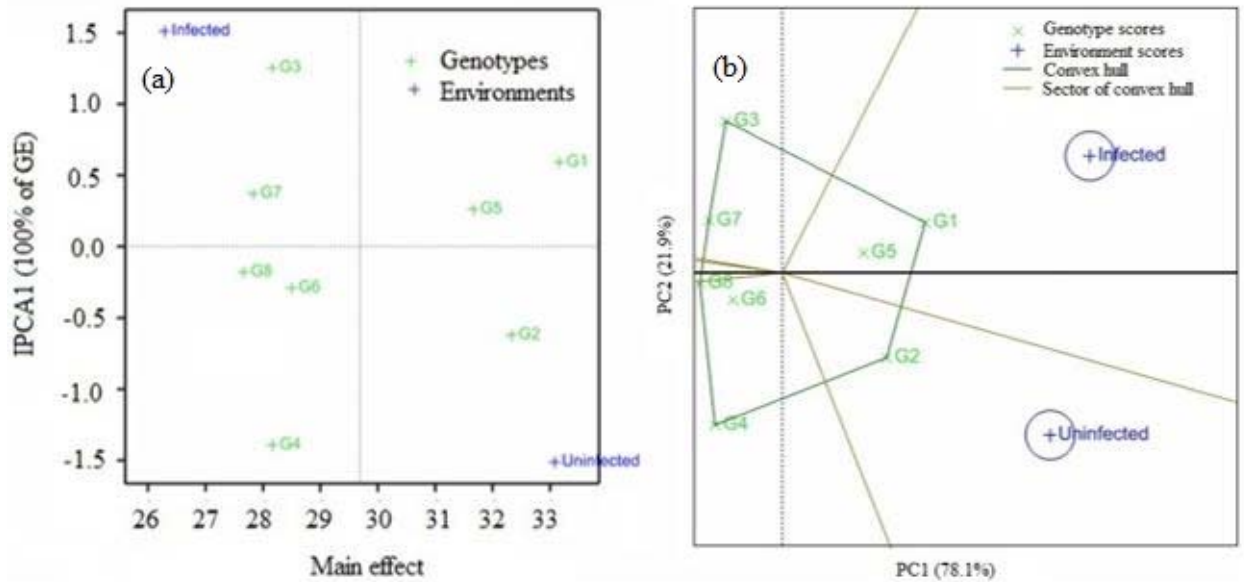
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345 Table 5: Stability coefficients of the growth and yield attributes in soyabean genotypes infected
 346 and uninfected with *Cucumber mosaic virus*

Genotype	Stability coefficient					
	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	0.003
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

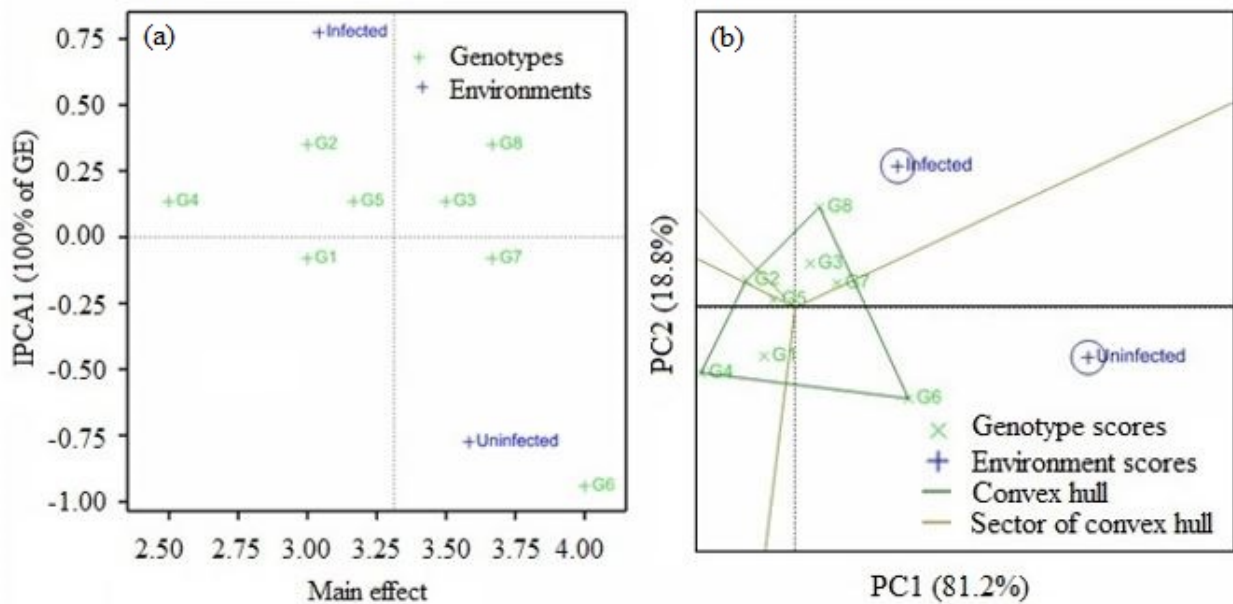
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UNDER PEER REVIEW



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 361 Fig. 1: AMMI plot of genotype and environment means against the first IPCA scores (a) and
 362 GGE biplot (b) of the plant height in soyabean genotypes infected and uninfected with *Cucumber*
 363 *mosaic virus*

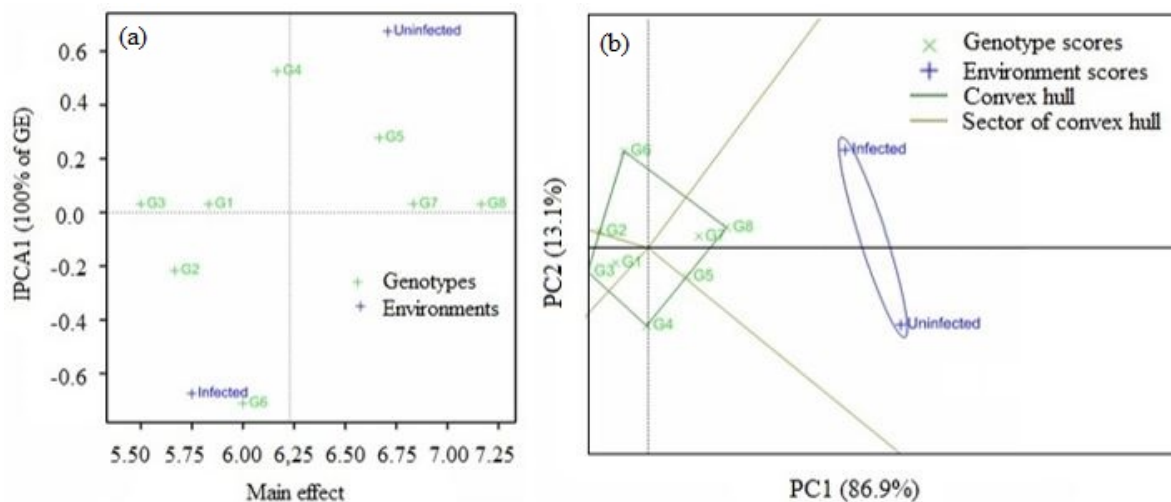
364 **Note:** G1=TGX 1448-2A; G2=TGX 1951-3F; G3=TGx 1987-10F; G4=TGX 1993-4FN;
 365 G5=TGX 1994; G6=TGX 2017-6E; G7=TGX 2023-1E; G8=TGX 2025-6E
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 369 Fig. 2: AMMI plot of genotype and environment means against the first IPCA scores (a) and
 370 GGE biplot (b) of the leaf diameter in soyabean genotypes infected and uninfected with
 371 *Cucumber mosaic virus*

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

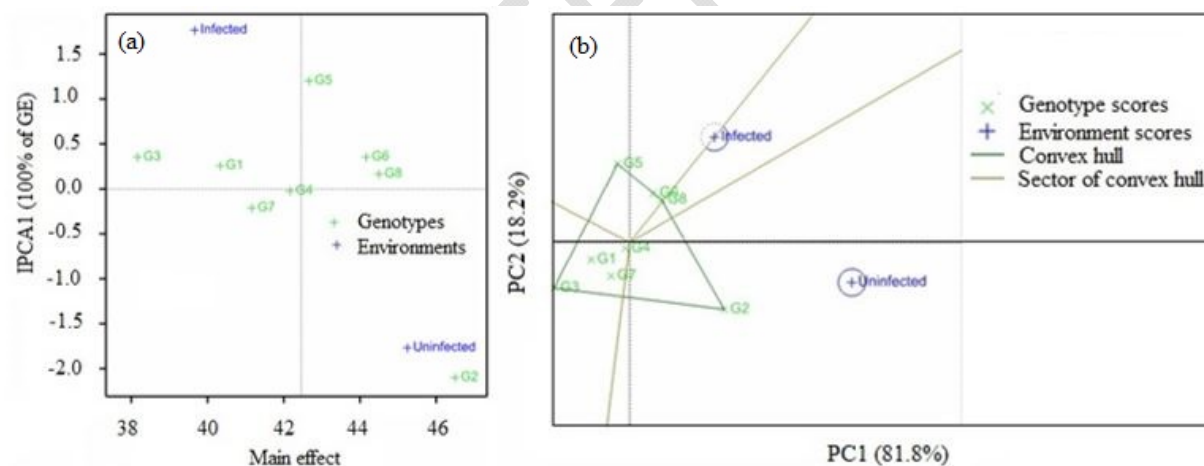
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377 Fig. 3: AMMI plot of genotype and environment means against the first IPCA scores (a) and
378 GGE biplot (b) of the leaf length in soyabean genotypes infected and uninfected with *Cucumber*
379 *mosaic virus*

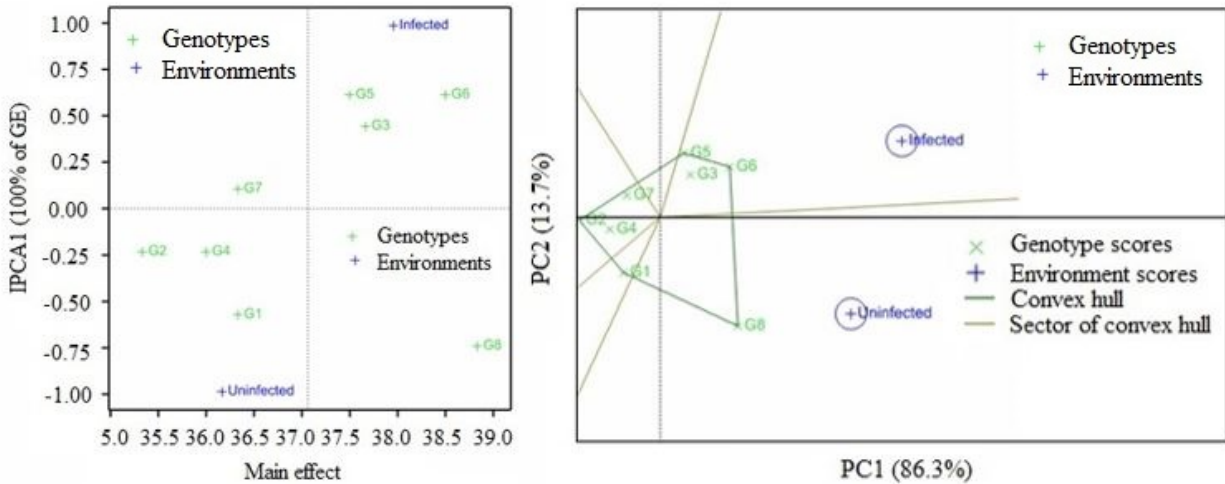
380 **Note:** G1=TGX 1448-2A; G2=TGx 1951-3F; G3=TGx 1987-10F; G4=TGX 1993-4FN;
381 G5=TGX 1994; G6=TGX 2017-6E; G7=TGX 2023-1E; G8=TGX 2025-6E
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384 Fig. 4: AMMI plot of genotype and environment means against the first IPCA scores (a) and
385 GGE biplot (b) of the number of leaves per plant in soyabean genotypes infected and uninfected
386 with *Cucumber mosaic virus*

387 **Note:** G1=TGX 1448-2A; G2=TGx 1951-3F; G3=TGx 1987-10F; G4=TGX 1993-4FN;
388 G5=TGX 1994; G6=TGX 2017-6E; G7=TGX 2023-1E; G8=TGX 2025-6E
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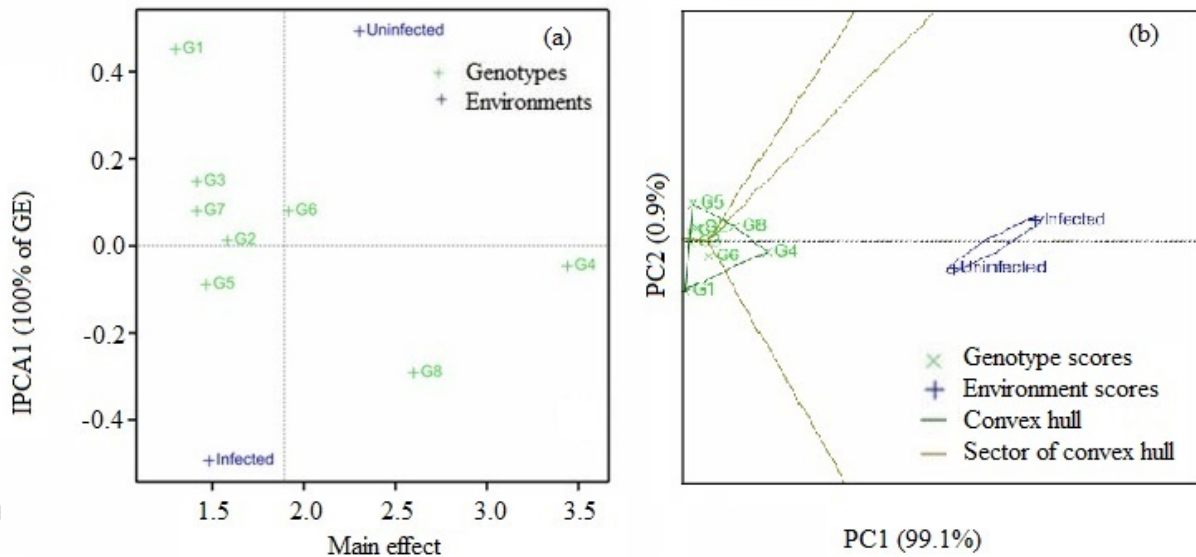
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391 Fig. 5: AMMI plot of genotype and environment means against the first IPCA scores (a) and
 392 GGE biplot (b) of the number of days to flowering in soyabean genotypes infected and
 393 uninfected with *Cucumber mosaic virus*

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

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

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

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406 **REFERENCES**

- 407
- 408 1. FAO (Food and Agriculture Organization) (2017). Soyabean.
409 <http://www.fao.org/faostat/en/#data/QC>
410
- 411 2. Thuita, M., Vanlauwe, B., Mutege, E. and Masso, C. (2018). Reducing spatial variability
412 of soybean response to rhizobia inoculants in farms of variable soil fertility in Siaya
413 County of western Kenya. *Agric. Ecosystems Environ.*, 261:153–160.
414
- 415 3. Agarwal, D. A, Billore, S. D., Sharma, A. N., Dupare, B. U., Srivastava, S. K. (2013).
416 Soybean: Introduction, improvement, and utilization in India—Problems and Prospects.
417 *Agric Res.*, 2:293–300.
418
- 419 4. de Breuil, S., Giolitti, E. J., Bejerman, N, and Lenardon, S. L. (2012). Effects of
420 *Cucumber mosaic virus* on the yield and yield components of peanut. *J. Plant Pathol.*,
421 94:669 – 673.
422
- 423 5. Agrios, G. N. (2005). Plant Pathology. Fifth Edition. Elsevier Academic Publishers,
424 Amsterdam
425
- 426 6. Dia, M., Wehner, T. C. and Arellano, C. (2016). Analysis of Genotype × Environment
427 Interaction (G×E) Using SAS Programming. *Agron. J.*, 108:1838 – 1852.
428
- 429 7. BMS (Breeding Management System) (2015). Breeding management system. Version
430 3.0.8. Integrated Breeding Platform (IBP), Mexico.
431
- 432 8. Adamu, A. S.,Salaudeen, M. T., Gana, A. S. and Ishaq, M. N. (2015): Response of
433 selected soybean (*Glycinemax* [L.] Merr.) lines to cucumber mosaic virus disease in
434 Minna, Niger State. *Nig. J. Agric. Food Environ.*, 11: 45 – 51.
435
- 436 9. Anuradha, C., Selvarajan, R., Vasantha, S. and Suresha, G. S. (2015). Biochemical
437 characterization of compatible plant virus interaction: A case study with *Bunchy top*
438 *virus*-banana host-pathosystem. *Plant Pathol. J.*, 14: 212 – 222.
439
- 440 10. Pallas, V. and Garcí'a, J. A. (2011). How do plant viruses induce disease? Interactions
441 and interference with host components. *J. Gen. Virol.*, 92: 2691 – 2705.
442
443
- 444 11. Dia, M., Wehner, T. C., Elmstrom, G. W., Gabert, A., Motes, J. E., Staub, J. E., Tolla, G.
445 E. and Widders, I. E. (2018). Genotype ×environment interaction for yield of pickling
446 cucumber in 24 U.S. environments. *Open Agric.*, 3: 1 – 16.
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