

1 **Gains in Grain Yield of Extra-early Maize during Three Breeding Periods under Drought and**
2 **Rain-fed Conditions.**

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11

12 **Abstract**

13 Drought is a key maize (*Zea mays* L.) production constraint in sub-Saharan Africa (SSA). Fourteen,
14 15 and 25 extra-early maturing maize cultivars, with varying *Striga* resistance, drought and low soil
15 nitrogen tolerance, were developed from 1995 to 2000 (Period 1), 2001 to 2006 (Period 2) and 2007 to
16 2012 (Period 3), respectively. The objectives of this study were to examine yield gains in the cultivars,
17 investigate inter-trait relationships and yield stability under six drought and 17 rain-fed conditions in
18 West Africa, 2013-2016. Annual rate of yield increase across cultivars was 0.034 Mg ha⁻¹ (3.28 %)
19 and 0.068 Mg ha⁻¹ (2.25 %) while yield gains per period were 0.17 and 0.38 Mg ha⁻¹ under drought and
20 rain-fed environments, respectively. Yield gains under drought and rain-fed environments were related
21 to prolonged flowering period, increased plant and ear heights, improved stalk lodging, ear and plant
22 aspects, whereas delayed leaf senescence and increased number of ears per plant (EPP) accompanied
23 yield improvement under drought only. Ear aspect and EPP were primary contributors to yield and
24 could be used as selection criteria for yield enhancement under drought and rain-fed conditions. High
25 yielding and stable cultivars across all environments based on additive main effects and multiplicative

26 interaction (AMMI) biplot included 2004 TZEE-Y Pop STR C₄, and TZEE-W Pop STR BC₂ C₀ of
27 Period 2 and 2009 TZEE-W STR, TZEE-Y STR 106, TZEE-W STR 107, and TZEE-W DT C₀ STR C₅
28 of Period 3. These cultivars could be commercialized to improve food self-sufficiency in SSA.

29

30 **Abbreviations:** AMMI, additive main effects and multiplicative interaction; EASP, ear aspect; EPP,
31 number of ears per plant; G, cultivar; GGE, genotype main effect plus genotype × environment
32 interaction; E, environment; IITA, International Institute of Tropical Agriculture; IPCA1, interaction
33 principal component axes 1; SSA, sub-Saharan Africa; WAP, weeks after planting; WCA, West and
34 central Africa.

35

36 **MAIZE IS A MAJOR STAPLE CROP** in West and Central Africa (WCA). The development and
37 commercialization of extra-early maize that matures in 80-85 days have made it possible for maize to
38 spread into the savannas of WCA. This has resulted in the expansion of the crop and rapid replacement
39 of the traditional crops, including the indigenous sorghum (*Sorghum bicolor*) and millet (*Pennisetum*
40 *glaucum*), particularly in the savannas of WCA. This is attributable to the fact that extra-early maize
41 cultivars respond better to application of fertilizer, have a shorter growing cycle, and are ready for
42 harvest much earlier than the indigenous sorghum and millet crops. In addition, as a result of the dry
43 spell usually experienced from November of each year to March of the following year, the early and
44 extra-early crops are preferred to reduce the hunger gap in July of each year because of the shorter
45 maturity period of the crop. An important factor constraining maize production in the savanna
46 agroecology is drought, which accounts for huge yield losses annually in sub-Saharan Africa (SSA).
47 Global warming, which is usually associated with irregular rainfall patterns, calls for an urgent and
48 effective genetic intervention to increase grain yield and tolerance to drought stress (Badu-Apraku and
49 Fakorede, 2017).

50 Drought stress and poor soil fertility of tropical soils, especially N, compounds the effects of
51 *Striga hermonthica* on maize because of enhanced secretion of strigolactones, plant hormones that
52 stimulate the germination of *Striga* seeds (Cechin and Press 1993; Mumera and Below 1993; Kim and
53 Adetimirin 1995). Therefore, it is of critical importance to introgress genes for drought tolerance into
54 *Striga*-resistant cultivars in the Guinea and Sudan savannas, which frequently experience intermittent
55 drought stress and low soil fertility. It is therefore not surprising that farmers, who cultivate maize in
56 *Striga*-endemic agro-ecologies of WCA, prefer cultivars with combined *Striga* resistance and drought
57 tolerance. The WCA farmers are reluctant to accept maize cultivars that are susceptible to both drought
58 stress and *Striga* infestation (Badu-Apraku and Fakorede, 2013).

59 To facilitate the development of drought-tolerant cultivars and improved technologies targeted
60 at the different agro-climatic conditions in SSA, particularly drought stress, a program was designed
61 specifically to capitalize on the inherent mechanisms for drought escape and drought tolerance in
62 maize and the prevailing production conditions in WCA. The cultivars possessing drought-escape
63 mechanisms usually complete critical physiological processes of the life cycle before the onset of
64 drought. This is highly desirable in cultivars developed for farmers in agro-ecologies prone to terminal
65 drought stress in WCA. On the other hand, drought tolerance is a physiological mechanism in plants,
66 which is genetically controlled and can enable plants to minimize or withstand the adverse effects of
67 drought. Drought-tolerant cultivars are especially invaluable in environments where the occurrence of
68 drought is unpredictable during crop growth and development in WCA. Two approaches have been
69 adopted since 1995 for developing extra-early maize cultivars with enhanced drought tolerance for
70 drought-prone agro-ecologies of WCA. The first one involves the development of extra-early cultivars
71 that mature before the onset of severe drought. The second strategy involves the development of
72 drought-tolerant cultivars under induced drought stress. Breeding for extra-early-maturing cultivars
73 has been carried out in the savanna agro-ecologies and many cultivars have been developed, released,
74 and commercialized following extensive testing in the diverse agro-climatic conditions of WCA. Since

75 2007, an important strategy of the International Institute of Tropical Agriculture (IITA) maize program
76 has been to evaluate extra-early maize inbred lines from diverse sources for drought tolerance.
77 Selected outstanding drought-tolerant inbred lines are also screened for *Striga* resistance under
78 artificial infestation. The outstanding inbred lines possessing both drought tolerance and *Striga*
79 resistance are used to develop hybrids that are then evaluated for adaptation to drought-prone and
80 *Striga*-endemic locations. The selected lines have served as invaluable sources of drought-tolerance
81 alleles for genetic enhancement of two source populations of extra-early maturity that are being
82 improved using the S₁ family recurrent selection scheme. Genetic enhancement of the extra-early
83 source populations under managed-drought stress using the S₁ recurrent selection method has
84 generated new productive cultivars possessing alleles for both drought-tolerance and *Striga*-resistance.
85 The selection for enhanced resistance to *Striga* and improved grain yield carried out under low N has
86 resulted in extra-early maize with increased tolerance to low N (Badu-Apraku et al., 2009).

87 Studies conducted in temperate countries have been used to document breeding progress by
88 comparing the performance of released cultivars developed during different eras in environments
89 similar to those of the tropical regions (Russell, 1984; Voldeng et al., 1997; Specht et al., 199). For
90 example, Russell (1984) documented genetic gain in grain yield of 0.68% yr⁻¹ for cultivars developed
91 in the USA between 1930s and 1980s. Much higher yield gains of 1.7% yr⁻¹ were reported by
92 Tollenaar (1989) for outstanding maize hybrids developed between the late 1950s and late 1980s and
93 evaluated under drought conditions in Canada. However, only a few reports are available on yield
94 gains for tropical maize evaluated under drought stress. For example, Masuka et al. (2017a)
95 demonstrated annual gains in grain yield of 0.029, 0.085, 0.11, and 0.193 Mg ha⁻¹ for early-maturing
96 open-pollinated varieties (OPVs) under natural drought, low N, optimal conditions, and infestation of
97 the maize streak virus (MSV), respectively, in Eastern and Southern Africa (ESA). Genetic gains
98 under random drought, low N, rain-fed conditions, and MSV for the intermediate-late maturing
99 cultivars were reported to be 0.042, 0.053, 0.079, and 0.109 Mg ha⁻¹ year⁻¹, respectively (Masuka et

100 al., 2017a). However, the authors did not observe any significant gains in grain yield of both early-
101 maturing and late-intermediate-maturing cultivars under managed-drought conditions. Annual genetic
102 gains for grain yield of maize hybrids developed by CIMMYT in ESA during the 2000 - 2010 period
103 and evaluated under managed-drought stress, random drought, low N, optimal conditions, and MSV
104 infestation were estimated to be 0.325, 0.227, 0.209, 0.109, and 0.141 Mg ha⁻¹, respectively (Masuka
105 et al., 2017b). In contrast, studies on genetic gains have been conducted for only OPVs in WCA. For
106 example, Kamara et al. (2004) conducted a study to examine genetic gains from selection of maize
107 cultivars of late-maturity, released between 1970 and 1999, in the savannas of Nigeria; and reported an
108 annual genetic gain in grain yield of 0.41%. The increase was attributed to higher total biomass
109 production and kernel weight, accompanied by reduction in days to flowering and plant height. Bello
110 et al. (2014) conducted a comparative study on the response of six maize hybrids, two each from the
111 1980, 1990, and 2000 eras to under three nitrogen levels (0, 30 and 90 kg N ha⁻¹); the N levels were
112 used as main plots and the six hybrids as sub-plots. Results revealed that mean grain yield increased by
113 48.4 and 62.4 %, as N increased from 0 to 30 kg ha⁻¹ and from 30 to 90 kg ha⁻¹, respectively (Bello et
114 al., 2014). The genetic gains in grain yield of 42% (between 1980 and 2000) and of 9% (between 1990
115 and 2000) were obtained under optimal-N fertilization (90 kg of N ha⁻¹). The two hybrids of the 2000
116 era were outstanding in all the agronomic traits and leaf chlorophyll concentration at all N levels. It
117 was concluded that improving traits associated with fertilizer N response could accelerate rate of
118 genetic gains in maize hybrid yields. In another study conducted by Badu-Apraku et al. (2017a),
119 genetic gains in grain yield of 56 extra-early open-pollinated maize cultivars developed during three
120 breeding eras (1995–2000, 2001–2006, and 2007–2012) were estimated under low N and high soil
121 nitrogen (high N) in Nigeria in 2013 and 2014. They reported genetic gains in grain yield of 0.314 Mg
122 ha⁻¹ era⁻¹ (13.29%) under low N and 0.493 Mg ha⁻¹ era⁻¹ (16.84%) under high N. In a similar study
123 conducted between 1988 and 2010 under induced drought stress and optimal (stress-free) growing
124 conditions, Badu-Apraku et al. (2013a) showed that the annual yield gains for early-maturing OPVs

125 were 0.040 and 0.014 Mg ha⁻¹ under optimal conditions and induced drought, respectively. Genetic
126 gains in yield of the cultivars tested under drought conditions were accompanied by improved plant
127 aspect and husk cover, whereas under optimal conditions, yield gains were associated with improved
128 plant aspect and ear aspect, increased number of ears per plant, increase in plant and ear heights and
129 improved husk cover. Badu-Apraku et al. (2015a) also evaluated maize cultivars of early maturity
130 under low N conditions in WCA and reported an increase in grain yield of 0.165 Mg ha⁻¹ era⁻¹, and a
131 yield range of 2.28 to 2.61 Mg ha⁻¹ for the first era (1955-2000) to the third era (2007-2012) cultivars,
132 respectively. Despite the results of these studies, there is complete lack of information on yield gains
133 and changes in other agronomic traits of extra-early-maturing cultivars of the three breeding periods
134 under drought stress and optimal growing conditions. Furthermore, information on trait association
135 during the different breeding periods is crucial for identifying valuable traits and on different breeding
136 strategies for enhancing progress in improving extra-early maize for stress tolerance (Badu-Apraku et
137 al., 2015b). The current study was therefore conducted to: (a) assess yield gains in extra-early maize
138 cultivars of the three breeding periods (1995-2000 = Period 1; 2001-2006 = Period 2; and 2007-2012 =
139 Period 3) under drought and rain-fed environments; (b) investigate trait associations during the three
140 breeding periods, and (c) assess the performance of the cultivars relative to grain yield and stability
141 across target research environments.

142

143 **MATERIALS AND METHODS**

144 *Development of extra-early cultivars possessing mechanisms for drought-escape and tolerance to* 145 *drought, Striga, and maize streak virus*

146 The extra-early populations used for the extraction of inbred lines and cultivars were derived from
147 crosses involving superior accessions, including introduced germplasm selected after extensive testing
148 in WCA (Badu-Apraku and Fakorede 2001; Badu-Apraku et al. 2007). For about two decades, the S₁
149 family selection scheme, artificial *S. hermonthica* field infestation, and screening under managed and

150 random drought have been used by the IITA maize scientists to develop one each of white (TZEE-W
151 Pop STR) and yellow (TZEE-Y Pop STR) source populations of extra-early maturity. Following
152 genetic enhancement of these populations, a large number of cultivars and inbred lines of extra-early
153 maturity, combining drought tolerance and resistance to *S. hermonthica* and MSV, were extracted from
154 each population. Several extra-early inbred lines in the IITA Maize Program possessed drought-escape
155 mechanism(s) and drought-tolerance genes. It was therefore expected that the inbred lines would
156 withstand the drought stress occurring during flowering and grain filling in the savannas of WCA, as
157 had been observed in cultivars of other maturity groups. Thus, a tremendous opportunity existed for
158 improvement of the performance of the cultivars in the program by introgressing genes for improved
159 tolerance to drought and *Striga* resistance. We recognized at the very early stages of the IITA extra-
160 early maize improvement program that several genes governed the expression of drought tolerance in
161 maize. Therefore, a major strategy of the program was to adopt various methods to identify maize
162 inbred lines with tolerance to drought from diverse germplasm sources. Since 2007, various strategies
163 have been employed in the program for the genetic enhancement of the populations for drought
164 tolerance at various testing sites in Nigeria. The focal point of the IITA extra-early-maturing maize
165 program for improving adaptation to drought has been to screen maize inbred lines from diverse
166 genetic backgrounds for tolerance to drought under managed moisture stress at Ikenne (Supplementary
167 Table 1). The soil at the Ikenne experiment station is classified as eutric nitrosol (Soil survey staff,
168 1999) and the research fields are flat and uniform and characterized by high water-holding capacity. A
169 sprinkler irrigation system was used to apply 17 mm of water weekly to the maize crop during the first
170 three weeks of growth in the dry season. The maize plants therefore depended on stored water in the
171 soil for growth and development. This strategy ensured that flowering and grain-filling periods
172 coincided with occurrence of induced drought stress. Under the optimal conditions at Ikenne, the
173 plants were irrigated throughout the growing period using the sprinkler irrigation system, as described
174 by Badu-Apraku et al. (2013a; 2017b). The trials were also evaluated under optimal conditions at

175 Mokwa and Zaria (high-yield environments) in Nigeria to assess the yield of the cultivars. At Bagauda
176 (characterized by terminal drought), the cultivars were exposed to drought stress that occurred from
177 flowering till physiological maturity.

178 Badu-Apraku and Fakorede (2017) have described in detail the strategies adopted to enhance
179 cultivar resistance to *Striga* and tolerance to low N. Briefly, promising drought-tolerant, extra-early
180 inbred lines selected for the development of the cultivars evaluated in the present genetic gain study
181 were also screened for *Striga* resistance under artificial *Striga* infestation at Mokwa and Abuja.
182 Drought-tolerant and *Striga*-resistant inbred lines also possessed tolerance to low N, even though they
183 had not been specifically selected for tolerance to low N (at 30-40 kg N ha⁻¹).

184 By 2007, extra-early inbreds and hybrids that possessed genes for tolerance to drought during
185 flowering and grain-filling periods, and which were also capable of escaping drought (characteristic of
186 extra-early maturing cultivars) and had low-N tolerance genes, had been identified (Badu-Apraku and
187 Fakorede, 2017). A program was therefore commenced in 2011 to generate extra-early cultivars
188 possessing genes for tolerance to drought. Towards this end, tolerance to drought and low N in the
189 extra-early white (TZEE-W Pop STR C₅) and the extra-early yellow (TZEE-Y Pop STR C₅) *Striga*-
190 resistant source populations was improved by introgressing drought and low-N tolerance genes from
191 19 white and 20 yellow extra-early inbred lines with elevated levels of tolerance to drought and/or low
192 N (Badu-Apraku and Fakorede, 2017). Two-hundred testcrosses generated from crosses, which
193 involved each population and outstanding inbreds with enhanced drought tolerance, were evaluated at
194 Ikenne under induced drought stress during the 2011/2012 dry season. The top-performing 25%
195 testcrosses from each source population were selected and recombined to reconstitute each population.
196 This was followed by recombination of the top 10 testcrosses of each population to form experimental
197 cultivars that were designated as 2012 TZEE-W DT STR C₅ and 2012 TZEE-Y DT STR C₅. A total of
198 56 extra-early-maturity maize cultivars from the three breeding periods (1995-2000, 2001-2006, and

199 2007-2012), possessing enhanced drought tolerance and *Striga* resistance, were used for the present
200 study (Supplementary Table 2).

201

202 ***Field evaluation and data collection***

203 The extra-early cultivars were evaluated under six induced or terminal-drought environments in
204 Nigeria and Ghana, and 17 optimal environments in Ghana, Republic of Benin, and Nigeria, from
205 2013 to 2016 (Supplementary Table 3). The drought trials at Ikenne were planted during the dry
206 season and 17 mm of water was supplied to the plots weekly using the sprinkler irrigation system. To
207 create induced drought stress at this location, the drought trials were irrigated for only the first 21 days
208 after planting, causing the maize plants to rely on residual moisture in the soil for growth and
209 development. In contrast, terminal drought was achieved by delaying the planting of the trials such that
210 the occurrence of drought stress coincided with 1-2 weeks before flowering. Optimal environments
211 used in the present study refer to environments where water and nitrogen were adequate for plant
212 growth and development. An 8×7 lattice design, with three replications, was adopted for the trial.
213 Each experimental unit comprised two 4 m long rows, with inter-row spacing of 75 cm and a spacing
214 of 40 cm between plants within rows. Initially, three seeds were planted per hill and two weeks after
215 planting (2 WAP), thinning was done to two seedlings per hill to obtain a final population density of
216 66,666 plants ha^{-1} . Basal fertilizer (60 kg each of N, P and K ha^{-1}) was applied to the managed
217 drought-stress experiments during planting, whereas 60 kg ha^{-1} N was top-dressed at 2 WAP.
218 However, for terminal drought and rain-fed environments, basal fertilizer application rates were 60 kg
219 ha^{-1} each of N, P and K at 2 WAP and 60 kg of N ha^{-1} at 4 WAP. Crop management practices were
220 similar for both drought-stress and rain-fed experiments. Weeds were controlled manually as well as
221 through the use of herbicides, as needed.

222 Data were recorded on the measured traits as described in detail by Badu-Apraku et al.
223 (2015b). Briefly, in the drought-stressed and rain-fed plots, days to 50% anthesis (DA) and days to
224 50% silking (DS) were recorded as the number of days from planting to when 50 % of plants per plot
225 had started shedding pollen or extruding silks, respectively. Anthesis-silking interval (ASI) was
226 computed as the difference between DS and DA. Plant height (PHT) and ear height (EHT) were
227 measured as the length from the base of the plant to the first tassel branch and the upper ear node,
228 respectively. Root lodging (RL) was estimated as the percentage of plants leaning more than 30° from
229 the vertical while stalk lodging (SL) was computed as percentage of plants with broken stalks at or
230 below the highest ear node. Plant aspect (PASP) was rated on a scale of 1 to 9 based on plant type,
231 where 1 = excellent and 9 = poor. Ear aspect (EASP) was scored on a scale of 1 to 9, where 1 = clean,
232 uniform, large, and well-filled ears and 9 = ears with undesirable features. Husk cover (HUSK) was
233 rated on a scale of 1 to 5, where 1 = husks tightly arranged and extended beyond the ear tip and 5 = ear
234 tips exposed. The number of ears per plant (EPP) was determined by dividing the total number of ears
235 harvested by the number of plants in the plot at harvest. In addition, stay green characteristic (STGR)
236 was scored for the drought-stressed plots at 70 days after planting (DAP) on a scale of 1–9, where 1
237 represented plants with almost all leaves green and 9 indicated plants with virtually all leaves dead. .
238 Grain yield for drought trials was adjusted to 150 g kg⁻¹ moisture and estimated from the shelled grain
239 weight. In the rain-fed experiments, grain yield was determined from ear weight using 80% shelling
240 percentage, adjusted to moisture content of 150 g kg⁻¹.

241 **Statistical analyses**

242 Observations recorded on plot means for grain yield and other agronomic traits were subjected to
243 analysis of variance (ANOVA) for drought stress and optimal environments separately using PROC
244 GLM statement of Statistical Analysis Systems (SAS) 9.3 (SAS Institute, 2011). The environments
245 were regarded as the location-year combinations in the combined ANOVA. The environments,

246 replicates-within environment, and blocks-within-replicates of each experiment were treated as random
247 effects, whereas the entries were considered fixed effects. Means of the 56 cultivars for each variable
248 were regressed on the year when the cultivar was developed to estimate gain year⁻¹ for the respective
249 traits. The means of grain yield and other traits of the maize cultivars were used as dependent
250 variables, and regressed on the year of breeding, as the independent variable to obtain the linear
251 regression coefficient (b-value) under drought stress and rain-fed environments. The relative genetic
252 gain per year was estimated as the b-value divided by the intercept and multiplied by 100 (Badu-
253 Apraku *et al.*, 2009). Similarly, the yield gain per period was computed by regressing mean grain yield
254 of cultivars on the respective periods of development. Annual yield gains for cultivars of each of the
255 breeding period were also computed following a similar procedure. The Excel software in the
256 Microsoft Office suite 2007 was used for the regression analysis as well as for the estimation of the
257 parameters and the graphical display of the regression lines. Correlation coefficients between grain
258 yield and other measured traits of maize cultivars were computed for drought stress and rain-fed
259 growing conditions using SAS version 9.3 (SAS Institute, 2011). To facilitate the estimation of
260 variance components, cultivars were treated as a random factor in this context. Variance components
261 were computed using the restricted maximum likelihood (REML) option in PROC MIXED command
262 (SAS institute, 2011). The estimates of broad-sense heritability (H²) for grain yield were computed for
263 each environment, and all the environments included in the present study revealed an H² value of ≥
264 0.30 (Supplementary Table 2). The H² of grain yield and other traits were estimated as follows:

265

$$H^2 = \frac{\sigma_g^2}{\sigma_g^2 + \frac{\sigma_e^2}{r}}$$

266

267

268

269 where σ_g^2 is the variance attributable to genotypic effects, σ_e^2 is experimental error variance; and $r =$
 270 the number of replicates within each environment (Fehr, 1991).

271

272 Repeatability estimates of the traits (Falconer and Mackay, 1996) across environments were calculated
 273 on a cultivar-mean basis as follows:

274

$$R = \frac{\sigma_g^2}{\sigma_g^2 + \frac{\sigma_{ge}^2}{e} + \frac{\sigma^2}{re}}$$

275

276

277 where e is the number of environments; σ_{ge}^2 is the component of variance attributable to cultivar \times
 278 environment interaction; and σ^2 is the error variance.

279 Step-wise regression analysis and sequential path diagrams were employed to show the cause
 280 and effect relationships among traits in the present study. The Statistical Package for Social Sciences
 281 (SPSS Inc, 2007) was used for the step-wise regression analyses to obtain information on the path
 282 coefficients and the causal relationships required for the path diagrams. Following the method
 283 proposed by Mohammadi et al. (2003), the predictor traits were organized into first, second, and third
 284 order, based on their contributions to the total variation in grain yield, with minimized multicollinearity
 285 (Badu-Apraku et al., 2014; Talabi et al., 2017). To perform the step-wise regression analysis, grain
 286 yield was regressed on measured traits to identify traits with significant contributions to the total
 287 variation in grain yield at $P \leq 0.05$, which were categorized as first order traits. The first-order traits
 288 thereafter were each regressed on other traits that were not in the first order category, to identify traits
 289 with significant contributions to grain yield through the first-order traits. These traits were classified as
 290 second order traits. The same procedure was repeated to identify third order trait(s) and so on. The
 291 path coefficients were obtained from the standardized b values of the stepwise regression analysis
 292 (Badu-Apraku et al., 2014; Talabi et al., 2017). The significance of the path coefficients was tested

293 using the standard errors at 0.05 probability level, with only traits having significant path coefficients
294 retained in each order.

295 A selection index for drought tolerance, which incorporated grain yield of the cultivars under
296 drought, along with the expression of traits such as ASI, PASP, EASP, STGR and EPP, was used to
297 characterize the extra-early maize cultivars as drought tolerant and drought sensitive (Oyekunle and
298 Badu-Apraku, 2012). The effect of different scales was minimized by standardizing each parameter
299 using a mean and standard deviation of zero and one, respectively. Thus, a cultivar characterized by a
300 positive value was considered drought tolerant, whereas the drought sensitive cultivars were those with
301 negative values. The selection index was calculated as follows:

302 Selection index = $[(2 \times \text{Yield}) + \text{EPP} - \text{ASI} - \text{PASP} - \text{EASP} - \text{STGR}]$.

303 Based on the characterization, 35 cultivars (top 25, middle five and worst five genotypes) were
304 selected for stability analysis. The additive main effects and multiplicative interaction (AMMI)
305 analysis was adopted to investigate the relationships among cultivars (G), environments (E), and G \times E
306 interaction (GEI) components of the yield of the selected 35 extra-early cultivars. The AMMI model
307 partitioned G \times E into several interaction principal component axes (IPCAs) through principal
308 component analysis (Zobel et al., 1988; Gauch and Zobel, 1988; Crossa, 1990). The AMMI analysis
309 was performed using the genotype main effect plus G by E interaction (GGE) biplot (Yan, 2001a,
310 2001b) and the AMMI model equation used was that reported by Sadeghi et al. (2011). The AMMI
311 biplot provided information on the performance and stability of the selected cultivars across drought
312 and rain-fed environments.

313

314 **RESULTS**

315 *Analysis of variance for grain yield and other traits and yield gains*

316 Results of combined ANOVA for grain yield and other measured traits under contrasting drought and
317 rain-fed environments (Table 1) revealed significant environment (E), period, cultivar (period),

318 cultivar (period) \times E interaction, and period \times E interaction mean squares for all measured traits,
319 except period mean square for ASI, and E \times period mean square for DS, EHT, RL, and EPP under
320 drought conditions. Grain yield varied from 1.19 to 1.54 Mg ha⁻¹ for cultivars of Period 1 and Mg
321 Period 3, respectively under drought, which corresponded with an overall annual yield gain of 3.28 %
322 (Tables 2 and 3). Annual yield gains of 0.0480, 0.0500 and 0.0002 Mg ha⁻¹ were obtained under
323 drought for cultivars developed during Periods 1, 2 and 3, respectively, whereas the gain in grain yield
324 per period across the 56 cultivars was 0.17 Mg ha⁻¹. Grain yield of cultivars ranged from 3.30 Mg ha⁻¹
325 for Period 1 cultivars to 4.06 Mg ha⁻¹ for Period 3 cultivars under rain-fed conditions, which translated
326 to an annual genetic gain of 2.25%. Under rain-fed environments, cultivars of Period 1 showed an
327 annual yield gain of 0.12 Mg ha⁻¹, whereas the gains in grain yield obtained for Period 2 and Period 3
328 cultivars were 0.022 and 0.014 Mg ha⁻¹, respectively. The yield gain per period of the 56 cultivars was
329 0.38 Mg ha⁻¹ across rain-fed environments. The realized annual increase in grain yield was 0.034 and
330 0.068 Mg ha⁻¹ under drought stress and rain-fed environments, respectively. The significant yield
331 increase from Period 1 to Period 3 under drought stress and rain-fed environments was associated with
332 prolonged flowering period, increase in EHT and PHT, and improvement in SL resistance, EASP, and
333 PASP. Other characters that accompanied the significant yield improvement under drought conditions
334 included prolonged STGR and increased EPP (Table 3).

335 Under drought stress, positive and significant b-values (gain per year) were obtained for grain
336 yield, DA, DS, PHT, EHT, and EPP, whereas significant negative b-values were observed for SL,
337 PASP, EASP, and STGR. The same set of traits showed similar trends under rain-fed environments,
338 except EPP, for which no significant gain was obtained; STGR was not measured under rain-fed
339 environments (Table 3).

340 Regression of mean grain yield of the extra-early maize cultivars tested under drought conditions on
341 mean grain yield under rain-fed environments, and vice versa, clearly separated the maize cultivars
342 into three distinct breeding periods (Figs. 1a and 1b). However, some cultivars from Period 2 produced

343 yields comparable to those of Period 3 extra-early cultivars, whereas one Period 2 cultivar (TZEE-Y
344 SR BC₁ × 9450 STR S₆ F₂) produced yield lower than those of Period 1 cultivars. The extra-early
345 Period 3 cultivars exhibited the most outstanding performance under both drought stress and rain-fed
346 environments. The grain yield of the cultivars under drought stress adequately predicted the yield
347 performance of the cultivars under rain-fed environments and vice versa ($R^2 = 58\%$; Figs.1a and 1b).

348

349 *Interrelationships among traits*

350 Under drought environments, the step-wise regression analysis identified EPP, EASP, RL, and EHT as
351 first order traits; these traits explained about 80 % of the variability in grain yield (Fig. 2). Number of
352 ears per plant had the largest path coefficient, whereas RL had the smallest path coefficient. The
353 second order traits identified under drought included PASP, DS, HUSK, STGR, SL, DA, and PHT;
354 each contributed to the variation in grain yield through one or two first order traits. The highest
355 indirect effect (0.82) was observed for DA through EHT, whereas the lowest indirect effect (-0.15) was
356 obtained for HUSK through EHT. Five out of the seven second order traits made significant
357 contributions to grain yield through EPP, four through EHT, and one each through EASP and RL.
358 Anthesis-silking interval was the only third-order trait identified under drought conditions in this
359 study, which made significant contributions to grain yield through DA.

360 Under rain-fed environments, step-wise regression analysis classified seven traits (EPP, EASP,
361 PASP, RL, SL, PHT, and DA) as the first-order traits (Fig. 3). These traits together contributed about
362 93% to the total variation in grain yield. Five of the traits contributing directly to grain yield showed
363 negative effect, whereas two of the traits had positive effects. The largest direct contribution to grain
364 yield was that of PHT (0.44), whereas the smallest contribution was that of EPP (0.09). Second-order
365 traits identified under rain-fed environments were ASI, EHT, and DS. While EHT made significant
366 contributions to grain yield through six first-order traits, ASI and DS each contributed through only
367 one of the first-order traits.

368

369 ***Performance and stability of extra-early maize cultivars***

370 The AMMI biplot for grain yield clearly depicted the performance of the selected 35 extra-early-
371 maturing maize cultivars of the three breeding periods and stability across drought and rain-fed
372 environments (Fig. 4). The grand mean of grain yield was represented by the vertical dotted line,
373 whereas the interaction principal component axis 1 (IPCA1) value of zero was represented by the
374 horizontal dotted line (y ordinate). The stable cultivars were those placed close to the horizontal line,
375 with little interactions with the environments, whereas the less stable cultivars were those farther from
376 the horizontal line. The high-yielding cultivars were placed to the right of the grand mean line and the
377 farther such cultivars were from the grand mean, the greater their grain yield. Across drought and rain-
378 fed environments, the percentage contributions of E (environment), G (cultivar), and the IPCA1 to the
379 total variation in grain yield sum of squares were 80.78, 9.22, and 2.8, respectively. The 84.6% of the
380 grain yield sum of squares captured by AMMI analysis was a clear indication that the biplot was
381 effective in decomposing the $G \times E$ interaction across drought stress and rain-fed environments (Fig.
382 4). Cultivars 2004 TZEE-Y Pop STR C₄, and TZEE-W Pop STR BC₂ C₀ of Period 2 and 2009 TZEE-
383 W STR, TZEE-Y STR 106, TZEE-W STR 107, and TZEE-W DT C₀ STR C₅ of Period 3 were the
384 most productive ones and stable relative to grain yield across drought and rain-fed environments.
385 Cultivar 2009 TZEE-OR₁ STR yielded more than the mean grain yield but was adapted to high-yield
386 environments. A large number of cultivars, among which TZEE-W STR 108 was outstanding, were
387 high-yielding, with adaptation to low-yield environments.

388 **DISCUSSION**

389 The significant cultivar means squares for all traits measured under drought and rain-fed environments
390 suggested that the cultivars were genetically distinct in the expression of these traits, which should
391 facilitate the identification and selection of superior cultivars under the research conditions, i.e.,
392 drought and rain-fed environments. Similarly, significant mean squares for environments for all

393 measured traits under drought and optimal environments were an indication that the environments
394 were unique in their ability to discriminate among the cultivars under drought and optimal
395 environments. These findings corroborate results reported by Badu-Apraku et al. (2013a), who
396 compared 50 early-maturing maize cultivars developed during three breeding eras under drought stress
397 and optimal environments in WA. The significant cultivar \times environment interactions detected for all
398 the measured traits under drought and optimal conditions suggested that the environments influenced
399 the performance of the cultivars differentially and that multi-environment testing was desirable.
400 However, this is inconsistent with the results of Badu-Apraku et al. (2013a), who observed lack of
401 significant $E \times$ era and $E \times$ cultivar (era) effects for all the measured traits of early-maturing genotypes
402 evaluated under drought conditions. The observed differences between the findings of Badu-Apraku et
403 al. (2013a) and the results of the present study might have resulted from the fewer drought testing sites
404 used in the former study.

405 An important objective of the present study was to investigate yield gains of 56 extra-early-
406 maturing cultivars developed during three breeding periods under drought and rain-fed environments.
407 The extra-early cultivars showed an annual genetic gain of 3.28%, with a realized yield increase of
408 $0.034 \text{ Mg ha}^{-1} \text{ yr}^{-1}$ under drought conditions, and 2.25% annual yield gain corresponding to an annual
409 increase of 0.068 Mg ha^{-1} under rain-fed conditions, which are greater than the yield gains obtained for
410 the early-maturing cultivars reported by Badu-Apraku et al. (2013a), who reported an annual yield gain
411 of 1.1% (0.014 Mg ha^{-1}) and 1.3% (0.040 Mg ha^{-1}) under drought and well-watered conditions,
412 respectively. The annual yield gain obtained for this set of extra-early maize cultivars under drought
413 was also higher compared with the annual percentage yield gains of 2.56 reported under artificial
414 *Striga* infestation (Badu-Apraku et al., 2016), 2.14 under low soil nitrogen (Badu-Apraku et al.,
415 2017a), and 2.72 under multiple-stress environments (Badu-Apraku et al., 2017b) for the same set of
416 extra-early cultivars. Furthermore, the annual yield gain of 0.034 Mg ha^{-1} achieved under drought in
417 the present study was greater than the 0.029 Mg ha^{-1} gain obtained for CIMMYT's ESA early-

418 maturing OPVs (Masuka et al., 2017a) and was comparable with the annual yield gain of 0.042 Mg
419 ha⁻¹ reported for the CIMMYT's ESA intermediate-late OPVs under random drought stress. The
420 implications of the results obtained in the present study are that the extra-early OPVs had better
421 responses to selection for improved grain yield and drought tolerance than the early and intermediate-
422 late varieties tested under drought stress. Furthermore, the relative annual yield gain of 3.28 %
423 obtained for the extra-early cultivars under drought conditions was higher than the 2.25 % achieved
424 under rain-fed environments in the present study. A plausible reason for this was that the emphasis of
425 the breeding program was more on improvement in drought tolerance rather than performance of
426 cultivars under rain-fed environments. With the recent advances in molecular breeding techniques,
427 marker-assisted selection (MAS) and genomic selection (GS) schemes are presently being employed to
428 fast-track breeding processes and accelerate yield gains in our program. In addition to the MAS and
429 GS, several other strategies outlined by Masuka et al. (2017a) for increasing genetic gains in Eastern
430 and Southern Africa breeding pipeline are being used in WCA under the DTMA/STMA Project for
431 accelerating genetic gains. These include, among others, increase in the size of the IITA maize
432 breeding program to facilitate the use of higher selection intensity and increase in the precision of
433 selection to achieve higher heritability.

434 Meseka et al. (2006) indicated that drought-tolerant genotypes might be characterized using a
435 selection index combining superior grain yield with desirable expression of PASP, EASP, and STGR,
436 reduced ASI, and increased EPP under drought as well as high grain yield under optimal conditions. In
437 the present study, the increased grain yields under drought and rain-fed environments were associated
438 with prolonged DA and DS, increase in EHT and PHT, and improvement in SL resistance, EASP and
439 PASP. In contrast, improved STGR and EPP accounted for yield gains only under drought
440 environments. Gains achieved in grain yield associated with delayed leaf senescence during the
441 breeding periods under drought may be attributed to longer grain filling duration period. The results of
442 this study showed that the traits included in the selection index for characterizing drought tolerance

443 were indeed effective in the development of superior cultivars under this stress factor. However, it was
444 not effective in keeping constant the EHT and PHT and the flowering dates of the cultivars. Selection
445 for improved tolerance to drought is usually conducted under drought stress, whereas field evaluations
446 for drought tolerance are conducted under drought and optimal environments. However, results of our
447 studies have demonstrated repeatedly that outstanding cultivars identified under stress usually
448 displayed outstanding performance under stress-free conditions (Badu-Apraku et al., 2013a). In this
449 study, cultivar grain yield under drought adequately predicted yield performance of the cultivars under
450 optimal environments ($R^2 = 58\%$). The implication of this result is that the performance of the
451 cultivars relative to grain yield under drought is a reliable indicator of expected yield performance of
452 the cultivars under optimal environments, and vice versa. Therefore, cultivars with outstanding
453 performance under drought stress also display superior grain yield under optimal conditions, and vice
454 versa. Similar results were obtained by Badu-Apraku et al. (2013a) when early-maturing maize
455 cultivars were evaluated under drought and optimal conditions.

456 Badu-Apraku et al. (2014) and Talabi et al. (2017) used the path coefficient analysis (Wright,
457 1921; Dewey and Lu, 1959) to quantify the contributions of various agronomic traits to the variation in
458 grain yield. Of particular interest was the sequential path analysis, which allowed for categorization of
459 traits into orders corresponding to the relative importance of the traits in explaining the variation in
460 grain yield (Mohammadi et al., 2003). Under drought conditions, the identification of EPP, EASP, RL,
461 and EHT as first order traits implied that these traits could be useful for index selection for genetic
462 enhancement of grain yield under drought stress. Of four first-order traits, only EPP and EASP were
463 among the traits included in the selection index (i.e., EASP, PASP, EPP, STGR, and ASI) along with
464 grain yield for improvement of drought tolerance, emphasizing the importance of these traits when
465 cultivars are subjected to drought stress. The categorization of PASP and STGR among the second-
466 order traits was also an indication that these traits had potential value in selecting for drought
467 tolerance. However, identification of ASI as a third-order trait in this study suggested that not only was

468 this trait of least importance but also that it did not play a prominent role in justifying its use in the
469 drought-tolerance selection index in maize. The results of this study are inconsistent with the findings
470 of Talabi et al. (2017), who identified ASI, EASP, PASP, STGR, and EPP as the primary traits directly
471 responsible for the variability in grain yield of early-maturing full-sib progenies under drought stress.
472 The difference in the findings may be explained by the differences in the genetic materials used for the
473 present study; the cultivars evaluated in the present study were extra-early-maturing, whereas Talabi et
474 al. (2017) evaluated early-maturing full-sib progenies. This suggested that specific selection indices
475 may be needed for the different types of genetic materials as well as maturity groups. Under rain-fed
476 environments, the identification of EASP, PASP, EPP, RL, SL, PHT, and DA as first-order traits
477 implied that these traits were key in determining the variation observed in grain yield. Again, ASI was
478 not in the first order traits, as observed under drought but was among the second-order traits under
479 optimal conditions. The consistent identification of EASP and EPP as first-order traits under the
480 contrasting environments confirmed their reliability for selection to improve grain yield across diverse
481 environments. It is striking that Badu-Apraku et al. (2017) placed EASP and PASP among the first-
482 order traits under high- and low-N conditions for the same set of cultivars as used in this study. An
483 important observation from the findings of several researchers (Badu-Apraku et al., 2011b; 2014;
484 2017; Talabi et al., 2017) is that EASP is a key trait accounting for the variation observed in grain
485 yield under diverse stress conditions. Hence, EASP should be accorded the desired emphasis in
486 selection programs designed to improve grain yield under contrasting environments to achieve
487 concomitant improvement in tolerance to diverse stress environments.

488 Development of outstanding maize hybrids for adoption by small-scale farmers in SSA remains
489 the most sustainable approach for increasing food security, alleviating poverty, and improving
490 livelihoods in the sub-region. The AMMI biplot identified the following cultivars from Period 2: 2004
491 TZEE-Y Pop STR C₄ and TZEE-W Pop STR BC₂ C₀ as well as 2009 TZEE-W STR, TZEE-Y STR
492 106, TZEE-W STR 107, and TZEE-W DT C₀ STR C₅ from Period 3 as highly productive and stable

493 genotypes across drought and optimal environments. These outstanding cultivars should be extensively
494 tested in on-farm trials and commercialized for improving food self-sufficiency and farmers' incomes
495 in SSA. The cultivar 2009 TZEE-OR₁ STR, which was high yielding but adapted to high-yield
496 environments and TZEE-W STR 108, which was promising relative to grain yield but was adapted to
497 low-yield environments should be further tested for commercialization in the specific environments in
498 which they displayed outstanding performance.

499 For more than two decades, early and extra-early maize cultivars have been developed for the
500 savannas of SSA and extensively evaluated by IITA scientists in the sub-region. Based on the results
501 of studies conducted under *Striga*-infested and *Striga*-free conditions, as well as those obtained from
502 studies involving 50 early-maturing cultivars evaluated under drought, *Striga*-infestation, and optimal
503 conditions (Badu-Apraku et al., 2013b; 2014, and 2017), the conclusion was that early and extra-early
504 maize responded favorably to selection under biotic and abiotic stresses encountered in SSA. Selection
505 for drought and/or *Striga* tolerance/resistance has inadvertently led to improvement in the level of
506 tolerance to low N but not as much as the response to direct selection for low-N tolerance.
507 Furthermore, selection under stress conditions results in improved performance of extra-early maize
508 cultivars under stress-free environments. In addition, efforts at genetically enhancing maize for
509 tolerance to drought in WCA has led to several conclusions that should guide breeders in SSA. The
510 products of the research efforts include drought-tolerant early and extra-early populations, OPVs,
511 inbred lines, and hybrids. Our experience has demonstrated unambiguously that the early and extra-
512 early materials are capable of escaping drought and also possess genes for drought tolerance and can
513 withstand drought stress that occurs randomly during the cropping season. Based on information on
514 the DA and DS used as maturity indices, we have clearly established that there is tremendous genetic
515 variability for the flowering traits in each maturity group. These flowering traits have been shown to
516 have high heritability and significant negative phenotypic and genetic correlations with grain yield
517 (Badu-Apraku and Fakorede, 2017). Therefore, early and extra-early maturities are under genetic

518 control and are amenable to genetic enhancement and many maize improvement methods such as
519 recurrent selection, pedigree selection, backcross breeding, double haploid, marker-assisted selection
520 and genomic selection.

521 522 **CONCLUSIONS**

523 Based on the average annual rate of increase in grain yield under drought conditions (0.034 Mg ha^{-1})
524 and under optimal conditions (0.068 Mg ha^{-1}), it can be concluded that considerable progress has been
525 made during the last three decades in the genetic enhancement of extra-early maturing maize cultivars
526 for drought tolerance in WCA. The availability of these extra-early cultivars is expected to contribute
527 to improved food self-sufficiency, farmers' incomes, and farmers' livelihoods in SSA. The significant
528 improvements in grain yield under drought and optimal conditions were associated with prolonged DA
529 and DS, increased EHT and PHT, and improvement in SL resistance, EASP, and PASP. In addition,
530 delayed senescence and increased EPP accompanied significant improvement in productivity under
531 drought. The EASP and EPP were consistently identified as highly reliable indirect selection criteria
532 for improving grain yield through index selection under drought and rain-fed environments. High
533 yielding and stable cultivars across all environments based on additive main effects and multiplicative
534 interaction (AMMI) biplot included 2004 TZEE-Y Pop STR C₄, and TZEE-W Pop STR BC₂ C₀ of
535 Period 2 and 2009 TZEE-W STR, TZEE-Y STR 106, TZEE-W STR 107, and TZEE-W DT C₀ STR C₅
536 of Period 3. These cultivars could be commercialized to improve food self-sufficiency in SSA.
537 Considerable improvement has been achieved in development and commercialization of drought-
538 tolerant maize cultivars in the extra-early maturity group for the sub-region.

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- 665

666 Fig 1a and b. Regression of grain yield of extra-early maize cultivars of three breeding periods under
667 drought on yield performance under rain-fed environments and vice versa.

668

669 Fig. 2. Path analysis model diagram showing causal relationships of measured traits of extra-early
670 maize cultivars of three breeding periods, evaluated under drought stress at six environments in WA,
671 2013-2016. Bold value is the residual effect; values in parenthesis are direct path coefficients while
672 other values are correlation coefficients. R1 is residual effects; ASI, anthesis-silking interval; DA,
673 days to 50 % anthesis; DS, days to 50 % silking; EASP, ear aspect; EHT, ear height; EPP, ears per
674 plant; HUSK, husk cover; PASP, plant aspect; PHT, plant height; RL, root lodging; SL, stalk lodging;
675 STGR, stay green characteristics; and YD, grain yield.

676

677

678 Fig. 3. Path analysis model diagram showing causal relationships of measured traits of extra-early
679 maize cultivars of three breeding periods, evaluated under rain-fed conditions at 17 environments in
680 WA, 2013-2014. Bold value is the residual effect; values in parenthesis are direct path coefficients
681 while other values are correlation coefficients. R1 is residual effects; ASI, anthesis-silking interval;
682 DA, days to 50 % anthesis; DS, days to 50 % silking; EASP, ear aspect; EHT, ear height; EPP, ears
683 per plant; HUSK, husk cover; PASP, plant aspect; PHT, plant height; RL, root lodging; SL, stalk
684 lodging; STGR, stay green characteristics; and YD, grain yield.

685 Fig. 4. Mean performance and stability of selected 35 extra-early maturing maize cultivars of three
686 breeding periods in terms of grain yield as measured by principal components across 23 drought and
687 rain-fed environments in West Africa between 2013 and 2016. E1 = Ikenne, drought, 2013; E2 =
688 Bagauda, drought, 2013; E3 = Dusu, drought, 2013; E4 = Kpeve, drought, 2014; E5 = Ikenne, drought,
689 2014; E6 = Ikenne, drought, 2015; E7 = Ikenne, rain-fed, 2013; E8 = Ife, high-N, 2013; E9 = Zaria,
690 rain-fed, 2013; E10 = Mokwa, high-N, 2013; E11 = Ina, rain-fed, 2013; E12 = Angaradebou rain-fed,
691 2013; E13 = Maini-Hari, rain-fed, 2013; E14 = Nyankpala, rain-fed, 2013; E15 = Ikenne, rain-fed,
692 2014; E16 = Ife high-N, 2014; E17 = Mokwa, high-N, 2014; E18 = Zaria, rain-fed, 2014; E19 =
693 Bagauda, rain-fed, 2014; E20 = Ina, rain-fed, 2014; E21 = Angaradebou, rain-fed, 2014; E22 = Manga,
694 rain-fed, 2014; and E23 = Fumesua, rain-fed, 2014.

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Table 1. Mean squares for grain yield and other agronomic traits of extra-early maize cultivars of three breeding periods evaluated under drought stress in six environments and under rain-fed conditions in 17 environments in Nigeria, Benin, and Ghana, 2013 – 2016.

| Entry | DF | Grain yield (Mg ha ⁻¹) | Days to anthesis | Days to silk | Anthesis silking interval (days) | Plant height (cm) | Ear height (cm) | Root lodging (%) | Stalk lodging (%) | Husk cover [†] | Plant aspect [‡] | Ear aspect [¥] | Ear rot (%) | Ears/p lant | Stay green charact eristic [§] |
|-----------------------|------|---------------------------------------|---------------------|-----------------|---|-------------------------|--------------------|------------------------|-------------------------|----------------------------|------------------------------|----------------------------|----------------|----------------|--|
| Drought environments | | | | | | | | | | | | | | | |
| Environment (E) | 5 | 122796029** | 4015.6** | 3839.1** | 100.0** | 53250.1** | 12167.6** | 2555.2** | 1057.7** | 479.9** | 775.5** | 505.9** | 4258.1** | 8.35** | 594.1** |
| Block (E × Rep) | 108 | 582011** | 5.1** | 9.1** | 2.4** | 650.4** | 266.2** | 13.1** | 21.1** | 0.8** | 0.9** | 0.9** | 6.6** | 0.03** | 1.3** |
| Rep(E) | 12 | 798912** | 26.4** | 31.8** | 1.5 | 1247.4** | 426.4** | 39.2** | 19.7** | 1.4** | 1.8** | 2.8** | 14.3** | 0.02 | 2.8** |
| Era | 2 | 10230293** | 163.5** | 128.6** | 2.1 | 4712.2** | 1604.7** | 54.0** | 129.8** | 3.1** | 20.7** | 16.9** | 27.2** | 0.19** | 7.5** |
| Cultivar (Period) | 53 | 787274** | 26.2** | 31.1** | 2.3** | 801.2** | 363.5** | 12.9** | 31.1** | 0.8** | 1.8** | 1.3** | 19.5** | 0.03** | 1.2** |
| E × Cultivar (Period) | 265 | 250476** | 5.3** | 6.1** | 2.1** | 268.9** | 138.7** | 11.1** | 16.0** | 0.7** | 0.9** | 0.7** | 14.2** | 0.02** | 0.9** |
| E × Period | 10 | 574222** | 7.4** | 5.7 | 3.4* | 631.0** | 129.4 | 10.4 | 20.7** | 3.1** | 1.2** | 1.4** | 33.1** | 0.01 | 2.6** |
| Error | 548 | 131885 | 2.3 | 3.3 | 1.4 | 190.9 | 113.2 | 7.8 | 8.7 | 0.2 | 0.4 | 0.3 | 5.2 | 0.01 | 0.4** |
| Repeatability | | 0.78 | 0.84 | 0.84 | 0.09 | 0.74 | 0.71 | 0.26 | 0.54 | 0.14 | 0.67 | 0.63 | 0.28 | 0.42 | 0.3** |
| Rain-fed environments | | | | | | | | | | | | | | | |
| Environment (E) | 16 | 12755930** | 1962.8** | 2574.8** | 134.8** | 97075.2** | 54264.3** | 6919.2** | 33292.6** | 46.6** | 75.5** | 57.7** | 657.5** | 2.00** | - |
| Block (E × Rep) | 306 | 830616** | 4.5** | 5.3** | 1.1** | 286.1** | 216.1** | 34.2** | 84.3** | 0.2** | 0.6* | 0.4** | 2.7** | 0.01** | - |
| Rep(E) | 34 | 4759476** | 22.8** | 23.0** | 1.1 | 1118.4** | 673.4** | 200.1** | 497.5** | 0.4** | 0.8* | 0.8** | 4.6** | 0.01** | - |
| Era | 2 | 124088801** | 743.1** | 608.2** | 5.4** | 14815.0** | 10611.3** | 100.8** | 499.6** | 3.0** | 13.5** | 31.2** | 6.8** | 0.10** | - |
| Cultivar (Period) | 53 | 8815082** | 79.9** | 91.9** | 1.3** | 1309.9** | 974.6** | 59.7** | 242.1** | 0.7** | 2.1** | 2.1** | 4.4** | 0.01** | - |
| E × Cultivar (Period) | 848 | 698835** | 3.7** | 4.1** | 0.9** | 220.0** | 154.5** | 27.5** | 80.9** | 0.2** | 0.6** | 0.2** | 1.9** | 0.01* | - |
| E × Period | 32 | 1142530** | 7.1** | 8.0** | 2.0** | 585.8** | 330.6** | 35.6** | 133.3** | 0.2* | 0.8** | 0.5** | 3.1** | 0.01* | - |
| Error | 1564 | 360244 | 1.9 | 2.2 | 0.8 | 144.4 | 127.2 | 19.6 | 38.7 | 0.2 | 0.5 | 0.2 | 1.4 | 0.01 | - |
| Repeatability | | 0.95 | 0.97 | 0.96 | 0.39 | 0.89 | 0.89 | 0.58 | 0.69 | 0.76 | 0.78 | 0.93 | 0.56 | 0.44 | - |

700 *, ** Significant at 0.05 and 0.01 probability level, respectively.

701 †Husk cover scored on a scale of 1-9, where 1 = husks tightly arranged and extended beyond the ear tip and 9 = ear tips exposed.; ‡Plant aspect recorded on a scale of 1-9

702 based on plant type, where 1 = excellent and 9 = poor; ¥Ear aspect rated on a scale of 1 – 9, where 1 = clean, uniform, large, and well-filled ears and 9 = ears with undesirable

703 features; §Stay green characteristic scored on a scale of 1 – 9, where 1 represented plants with almost all leaves green and 9 indicated plants with virtually all leaves dead.

Table 2. Means \pm SE for grain yield and other agronomic traits of extra-early maize cultivars of three breeding periods evaluated under drought stress in six environments and under rain-fed growing conditions in 17 environments in Nigeria, Benin, and Ghana, 2013 to 2016.

| Trait | Period | Number of cultivars | Drought conditions | Rain-fed conditions |
|--|-----------|---------------------|--------------------|---------------------|
| Grain yield (Mg ha ⁻¹) | 1995-2000 | 14 | 1.190 \pm 0.0512 | 3.296 \pm 0.1214 |
| | 2001-2006 | 17 | 1.353 \pm 0.0733 | 3.674 \pm 0.1000 |
| | 2007-2012 | 25 | 1.538 \pm 0.0361 | 4.056 \pm 0.1218 |
| Days to anthesis | 1995-2000 | 14 | 53 \pm 0.33 | 51 \pm 0.43 |
| | 2001-2006 | 17 | 54 \pm 0.39 | 53 \pm 0.35 |
| | 2007-2012 | 25 | 54 \pm 0.21 | 53 \pm 0.27 |
| Days to silking | 1995-2000 | 14 | 55 \pm 0.38 | 53 \pm 0.45 |
| | 2001-2006 | 17 | 57 \pm 0.44 | 54 \pm 0.37 |
| | 2007-2012 | 25 | 56 \pm 0.21 | 54 \pm 0.28 |
| Anthesis silking interval (days) | 1995-2000 | 14 | 3 \pm 0.12 | 2 \pm 0.06 |
| | 2001-2006 | 17 | 3 \pm 0.11 | 2 \pm 0.04 |
| | 2007-2012 | 25 | 3 \pm 0.06 | 2 \pm 0.03 |
| Plant height (cm) | 1995-2000 | 14 | 147 \pm 1.49 | 167 \pm 1.58 |
| | 2001-2006 | 17 | 152 \pm 1.95 | 173 \pm 1.57 |
| | 2007-2012 | 25 | 154 \pm 1.53 | 175 \pm 1.22 |
| Ear height (cm) | 1995-2000 | 14 | 66 \pm 1.13 | 79 \pm 1.54 |
| | 2001-2006 | 17 | 69 \pm 1.39 | 84 \pm 1.16 |
| | 2007-2012 | 25 | 70 \pm 0.92 | 85 \pm 1.03 |
| Root lodging (%) | 1995-2000 | 14 | 3.6 \pm 0.23 | 5.5 \pm 0.32 |
| | 2001-2006 | 17 | 4.1 \pm 0.27 | 5.4 \pm 0.20 |
| | 2007-2012 | 25 | 3.3 \pm 0.17 | 4.8 \pm 0.28 |
| Stalk lodging (%) | 1995-2000 | 14 | 5.8 \pm 0.42 | 11.6 \pm 0.63 |
| | 2001-2006 | 17 | 6.1 \pm 0.29 | 11.2 \pm 0.54 |
| | 2007-2012 | 25 | 4.8 \pm 0.29 | 9.8 \pm 0.53 |
| Husk cover [†] | 1995-2000 | 14 | 3.0 \pm 0.05 | 2.1 \pm 0.03 |
| | 2001-2006 | 17 | 2.8 \pm 0.06 | 2.0 \pm 0.04 |
| | 2007-2012 | 25 | 2.8 \pm 0.05 | 2.0 \pm 0.02 |
| Plant aspect [‡] | 1995-2000 | 14 | 4.4 \pm 0.10 | 2.5 \pm 0.05 |
| | 2001-2006 | 17 | 4.0 \pm 0.08 | 2.4 \pm 0.04 |
| | 2007-2012 | 25 | 3.9 \pm 0.07 | 2.3 \pm 0.06 |
| Ear aspect [¥] | 1995-2000 | 14 | 3.7 \pm 0.07 | 2.8 \pm 0.07 |
| | 2001-2006 | 17 | 3.5 \pm 0.09 | 2.6 \pm 0.05 |
| | 2007-2012 | 25 | 3.3 \pm 0.05 | 2.4 \pm 0.06 |
| Ear rot (%) | 1995-2000 | 14 | 4.5 \pm 0.21 | 1.8 \pm 0.09 |
| | 2001-2006 | 17 | 3.9 \pm 0.34 | 1.7 \pm 0.07 |
| | 2007-2012 | 25 | 4.5 \pm 0.26 | 1.7 \pm 0.06 |
| Stay green characteristic [§] | 1995-2000 | 14 | 4.2 \pm 0.08 | - |
| | 2001-2006 | 17 | 3.9 \pm 0.08 | - |
| | 2007-2012 | 25 | 3.9 \pm 0.07 | - |
| Ears per plant | 1995-2000 | 14 | 0.7 \pm 0.011 | 0.9 \pm 0.0044 |
| | 2001-2006 | 17 | 0.8 \pm 0.013 | 0.9 \pm 0.0038 |
| | 2007-2012 | 25 | 0.8 \pm 0.008 | 0.9 \pm 0.0039 |

704 [†]Husk cover scored on a scale of 1-9, where 1 = husks tightly arranged and extended beyond the ear tip and 9 = ear tips
705 exposed.; [‡]Plant aspect recorded on a scale of 1-9 based on plant type, where 1 = excellent and 9 = poor; [¥]Ear aspect rated on
706 a scale of 1 – 9, where 1 = clean, uniform, large, and well-filled ears and 9 = ears with undesirable features; [§]Stay green
707 characteristic scored on a scale of 1 – 9, where 1 represented plants with almost all leaves green and 9 indicated plants with
708 virtually all leaves dead.

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Table 3. Relative genetic gains, in grain yield and other agronomic traits of extra-early maize cultivars of three breeding periods across six drought and 17 rain-fed research environments in Nigeria, Benin, and Ghana, 2013 to 2016.

| Trait | Relative gain (% per year) | R ² | a (intercept) | b (linear regression coefficient) |
|--|-------------------------------|----------------|---------------|---|
| Drought stress environments | | | | |
| Grain yield (Mg ha ⁻¹) | 3.28 | 0.358 | 1.03 | 0.034** |
| Days to anthesis | 0.17 | 0.086 | 52.6 | 0.089* |
| Days to silk | 0.16 | 0.076 | 55.3 | 0.087* |
| Anthesis silking interval (days) | -0.52 | 0.022 | 2.9 | -0.015 |
| Plant height (cm) | 0.49 | 0.185 | 144 | 0.710** |
| Ear height (cm) | 0.65 | 0.152 | 64 | 0.416** |
| Root lodging (%) | -1.05 | 0.045 | 4.1 | -0.043 |
| Stalk lodging (%) | -1.65 | 0.102 | 6.5 | -0.107** |
| Husk cover [†] | -0.17 | 0.010 | 3.0 | -0.005 |
| Plant aspect [‡] | -0.91 | 0.171 | 4.5 | -0.041** |
| Ear aspect [§] | -1.51 | 0.315 | 4.0 | -0.061** |
| Ears rot (%) | -0.23 | 0.001 | 4.4 | -0.010 |
| Stay green characteristic [§] | -0.72 | 0.121 | 4.3 | -0.031** |
| Ears/plant | 0.40 | 0.101 | 0.8 | 0.003* |
| Rain-fed environments | | | | |
| Grain yield (Mg ha ⁻¹) | 2.25 | 0.361 | 3.017 | 0.068** |
| Days to anthesis | 0.30 | 0.199 | 50.7 | 0.150** |
| Days to silk | 0.20 | 0.108 | 52.8 | 0.107** |
| Anthesis silking interval (days) | -0.45 | 0.025 | 2.0 | -0.009 |
| Plant height (cm) | 0.44 | 0.289 | 164.5 | 0.727** |
| Ear height (cm) | 0.77 | 0.264 | 76.8 | 0.591** |
| Root lodging (%) | -1.09 | 0.065 | 5.9 | -0.064 |
| Stalk lodging (%) | -1.17 | 0.077 | 12.2 | -0.143* |
| Husk cover [†] | -0.32 | 0.061 | 2.1 | -0.007 |
| Plant aspect [‡] | -1.52 | 0.217 | 2.6 | -0.040** |
| Ear aspect [§] | -1.75 | 0.284 | 3.2 | -0.055** |
| Ears rot (%) | -0.91 | 0.046 | 2.0 | -0.018 |
| Ears/plant | 2.25 | 0.059 | 0.9 | 0.020 |

712 *, ** Significant at 0.05 and 0.01 probability level, respectively.

713 [†]Husk cover scored on a scale of 1-9, where 1 = husks tightly arranged and extended beyond the ear tip and 9 = ear tips
 714 exposed.; [‡]Plant aspect recorded on a scale of 1-9 based on plant type, where 1 = excellent and 9 = poor; [§]Ear aspect rated on
 715 a scale of 1 – 9, where 1 = clean, uniform, large, and well-filled ears and 9 = ears with undesirable features; [§]Stay green
 716 characteristic scored on a scale of 1 – 9, where 1 represented plants with almost all leaves green and 9 indicated plants with
 717 virtually all leaves dead.

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719 Supplementary Table 1: Description of test locations used for the evaluation of the cultivars of three breeding periods under drought and rain-fed environments, 2013 to 2016.

| Location | Agro ecological zone† | Latitude | Longitude | Altitude (m ASL) | Annual rainfall during growing season (mm) |
|----------|-----------------------|----------|-----------|------------------|--|
| Ikenne | RF | 6°87'N | 3°7'E | 60 | 1500 |
| Kadawa | SS | 11°45'N | 8°45'E | 468.5 | 884 |
| Bagauda | SS | 12°00'N | 8°22'E | 580 | 884 |
| Mokwa | SGS | 9° 18'N | 5° 4'E | 457 | 1100 |
| Zaria | NGS | 11°11'N | 7°38'E | 640 | 1200 |

†NGS, Northern Guinea Savanna; RF, Rain forest zone; SGS, Southern Guinea savanna; SS, Sudan savanna.

720 Supplementary Table 2: Extra-early maize cultivars used in the study, their year of
 721 release/development and reactions to biotic and abiotic stresses.

| Code | Cultivars | Year of development | Reactions to stresses | | |
|------|--|---------------------|-----------------------|---------------------------|-------------|
| | | | Drought | <i>Striga hermonthica</i> | Low N |
| 1 | 95 TZEE-Y | 1995 | Susceptible | Susceptible | Susceptible |
| 3 | 97 TZEE-Y 2-C ₁ | 1997 | Susceptible | Susceptible | Susceptible |
| 5 | CSP SR × TZEE-Y STR | 1997 | Susceptible | Susceptible | Susceptible |
| 6 | TZEE-W ST × GUA 314 BC ₁ | 1997 | Susceptible | Susceptible | Susceptible |
| 7 | TZEE-W-SR BC ₅ (RE) | 1997 | Susceptible | Susceptible | Susceptible |
| 8 | 98 SYN EE-W | 1998 | Susceptible | Susceptible | Tolerant |
| 9 | 98 TZEE-W STR | 1998 | Tolerant | Susceptible | Susceptible |
| 10 | 99 TZEE-Y STR C ₀ | 1999 | Susceptible | Susceptible | Susceptible |
| 11 | 99 TZEF-Y Pop STR QPM C ₀ | 1999 | Susceptible | Susceptible | Susceptible |
| 12 | EV 99 QPM | 1999 | Susceptible | Susceptible | Susceptible |
| 35 | 99 TZEF-Y STR C ₀ | 1999 | Susceptible | Susceptible | Susceptible |
| 36 | TZEE-Y Pop STR C ₀ | 1999 | Susceptible | Susceptible | Susceptible |
| 13 | 2000 SYN EE-W STR | 2000 | Susceptible | Susceptible | Tolerant |
| 14 | 2000 SYN EE-W STR QPM | 2000 | Tolerant | Susceptible | Susceptible |
| 15 | FERKE TZEE-W STR | 2001 | Susceptible | Resistant | Tolerant |
| 16 | SINE TZEE-W STR | 2001 | Susceptible | Susceptible | Susceptible |
| 18 | TZEE-Y Pop STR C ₃ | 2001 | Tolerant | Susceptible | Susceptible |
| 19 | TZEE-W Pop STR C ₃ | 2002 | Tolerant | Resistant | Tolerant |
| 20 | TZEE-Y Pop STR C ₄ | 2002 | Tolerant | Susceptible | Tolerant |
| 21 | 2004 TZEE-W Pop STR C ₄ | 2004 | Tolerant | Tolerant | Tolerant |
| 22 | 2004 TZEE-Y Pop STR C ₄ | 2004 | Tolerant | Tolerant | Tolerant |
| 23 | TZEE-W Pop STR QPM C ₀ | 2004 | Tolerant | Tolerant | Tolerant |
| 24 | TZEE-W Pop STR BC ₂ C ₀ | 2004 | Tolerant | Tolerant | Tolerant |
| 27 | TZEE-W Pop × LD S ₆ (SET 1) | 2004 | Tolerant | Tolerant | Tolerant |
| 28 | TZEE-W Pop × LD S ₆ (SET2) | 2004 | Tolerant | Tolerant | Susceptible |
| 29 | TZEE-W Pop × LD S ₆ (SETA1) | 2004 | Susceptible | Resistant | Tolerant |
| 30 | TZEE-W Pop × LD S ₆ F ₂ (SET A2) | 2004 | Susceptible | Susceptible | Tolerant |
| 31 | TZEE-Y Pop STR QPM C ₀ | 2004 | Susceptible | Susceptible | Susceptible |
| 32 | TZEE-Y SR BC ₁ × 9450 STR S ₆ F ₂ | 2004 | Susceptible | Susceptible | Susceptible |
| 33 | TZEE-Y Pop STR QPM C ₁ | 2005 | Susceptible | Tolerant | Tolerant |
| 34 | TZEE-W Pop STR C ₄ | 2006 | Tolerant | Tolerant | Tolerant |
| 37 | 2008 SYN EE-W DT STR | 2008 | Tolerant | Susceptible | Tolerant |
| 38 | 2008 SYN EE-Y DT STR | 2008 | Susceptible | Susceptible | Susceptible |
| 39 | 2008 TZEE-W STR | 2008 | Tolerant | Tolerant | Tolerant |
| 40 | 2008 TZEE-Y STR | 2008 | Susceptible | Susceptible | Susceptible |
| 41 | TZEE-W Pop STR C ₅ | 2008 | Tolerant | Resistant | Tolerant |
| 42 | TZEE-Y Pop STR C ₅ | 2008 | Tolerant | Susceptible | Susceptible |
| 43 | 2009 TZEE-OR ₁ STR | 2009 | Tolerant | Tolerant | Tolerant |
| 44 | 2009 TZEE-OR ₁ STR QPM | 2009 | Susceptible | Tolerant | Tolerant |
| 45 | 2009 TZEE-OR ₂ STR | 2009 | Tolerant | Resistant | Tolerant |
| 46 | 2009 TZEE-OR ₂ STR QPM | 2009 | Susceptible | Tolerant | Tolerant |
| 47 | 2009 TZEE-W STR | 2009 | Susceptible | Tolerant | Susceptible |
| 48 | TZEE-W STR 104 | 2009 | Tolerant | Resistant | Tolerant |
| 49 | TZEE-W STR 105 | 2009 | Tolerant | Resistant | Tolerant |
| 50 | TZEE-Y STR 106 | 2009 | Susceptible | Tolerant | Susceptible |
| 51 | TZEE-W STR 107 | 2009 | Tolerant | Resistant | Tolerant |
| 52 | TZEE-W STR 108 | 2009 | Tolerant | Resistant | Tolerant |
| 2 | TZEE-W STR 104 BC ₁ | 2010 | Susceptible | Resistant | Tolerant |
| 4 | TZEE-Y STR 106 BC ₁ | 2010 | Tolerant | Tolerant | Tolerant |
| 17 | TZEE-W STR 105 BC ₁ | 2010 | Tolerant | Resistant | Tolerant |
| 25 | TZEE-W STR 107 BC ₁ | 2010 | Tolerant | Tolerant | Tolerant |
| 26 | TZEE-W STR 108 BC ₁ | 2010 | Tolerant | Resistant | Tolerant |
| 53 | 2012 TZEE-W DT STR C ₅ | 2012 | Tolerant | Resistant | Tolerant |
| 54 | 2012 TZEE-Y DT STR C ₅ | 2012 | Susceptible | Tolerant | Susceptible |
| 55 | TZEE-W DT C ₀ STR C ₅ | 2012 | Tolerant | Resistant | Tolerant |
| 56 | TZEE-Y DT C ₀ STR C ₅ | 2012 | Tolerant | Susceptible | Tolerant |

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723 Supplementary Table 3. Environments, locations, research conditions and years of evaluation of extra-
 724 early maturing maize cultivars under drought-stress and rain-fed growing environments in West Africa.
 725

| Environment | Country | Location | Management | Year | Grain yield (Mg ha ⁻¹) | Heritability |
|-------------|----------------|--------------------|-------------------------|------------------|---------------------------------------|--------------|
| 1 | <u>Nigeria</u> | <u>Ikenne</u> | <u>Managed drought</u> | <u>2013/2014</u> | 2.438 | 0.67 |
| 2 | <u>Nigeria</u> | <u>Ikenne</u> | <u>Managed drought</u> | <u>2014/2015</u> | 0.760 | 0.71 |
| 3 | <u>Nigeria</u> | <u>Ikenne</u> | <u>Managed drought</u> | <u>2015/2016</u> | 1.046 | 0.73 |
| 4 | <u>Nigeria</u> | <u>Bagauda</u> | <u>Terminal drought</u> | <u>2013</u> | 2.237 | 0.57 |
| 5 | <u>Nigeria</u> | <u>Dusu</u> | <u>Terminal drought</u> | <u>2013</u> | 0.284 | 0.78 |
| 6 | <u>Ghana</u> | <u>Kpeve</u> | <u>Terminal drought</u> | <u>2014</u> | 1.649 | 0.65 |
| 7 | <u>Nigeria</u> | <u>Bagauda</u> | <u>Rain-fed</u> | <u>2014</u> | 4.421 | 0.76 |
| 8 | <u>Ghana</u> | <u>Fumesua</u> | <u>Rain-fed</u> | <u>2014</u> | 2.493 | 0.54 |
| 9 | <u>Benin</u> | <u>Ina</u> | <u>Rain-fed</u> | <u>2013</u> | 2.858 | 0.67 |
| 10 | <u>Benin</u> | <u>Ina</u> | <u>Rain-fed</u> | <u>2014</u> | 3.053 | 0.72 |
| 11 | <u>Nigeria</u> | <u>Ikenne</u> | <u>Rain-fed</u> | <u>2013</u> | 2.964 | 0.51 |
| 12 | <u>Nigeria</u> | <u>Ikenne</u> | <u>Rain-fed</u> | <u>2014</u> | 3.367 | 0.86 |
| 13 | <u>Nigeria</u> | <u>Mania</u> | <u>Rain-fed</u> | <u>2013</u> | 2.210 | 1.00 |
| 14 | <u>Ghana</u> | <u>Manga</u> | <u>Rain-fed</u> | <u>2013</u> | 3.447 | 0.53 |
| 15 | <u>Ghana</u> | <u>Nyankpala</u> | <u>Rain-fed</u> | <u>2013</u> | 2.975 | 0.34 |
| 16 | <u>Ghana</u> | <u>Nyankpala</u> | <u>Rain-fed</u> | <u>2014</u> | 3.441 | 0.47 |
| 17 | <u>Nigeria</u> | <u>Zaria</u> | <u>Rain-fed</u> | <u>2013</u> | 4.841 | 0.82 |
| 18 | <u>Nigeria</u> | <u>Zaria</u> | <u>Rain-fed</u> | <u>2014</u> | 5.670 | 0.91 |
| 19 | <u>Benin</u> | <u>Angaradebou</u> | <u>Rain-fed</u> | <u>2013</u> | 3.234 | 0.37 |
| 20 | <u>Benin</u> | <u>Angaradebou</u> | <u>Rain-fed</u> | <u>2014</u> | 3.793 | 0.61 |

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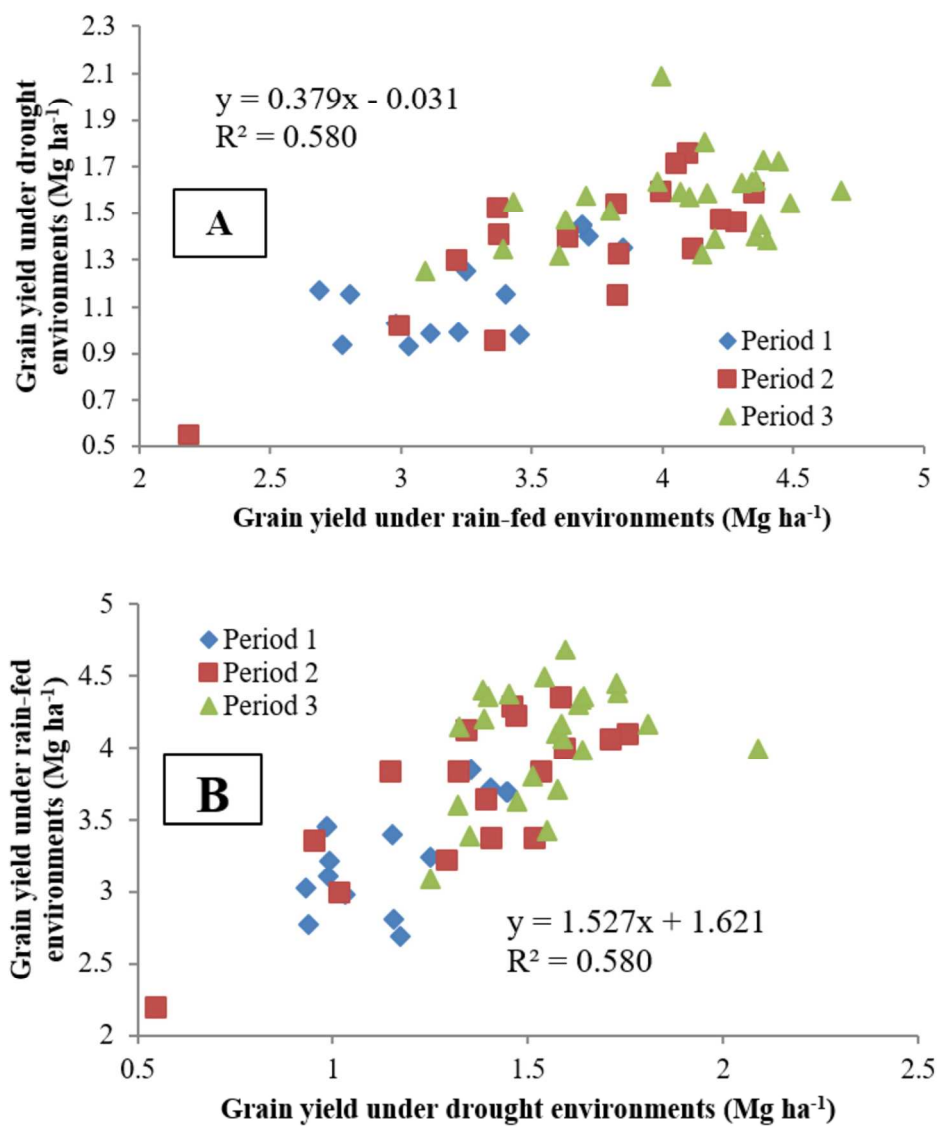


Fig 1a and b

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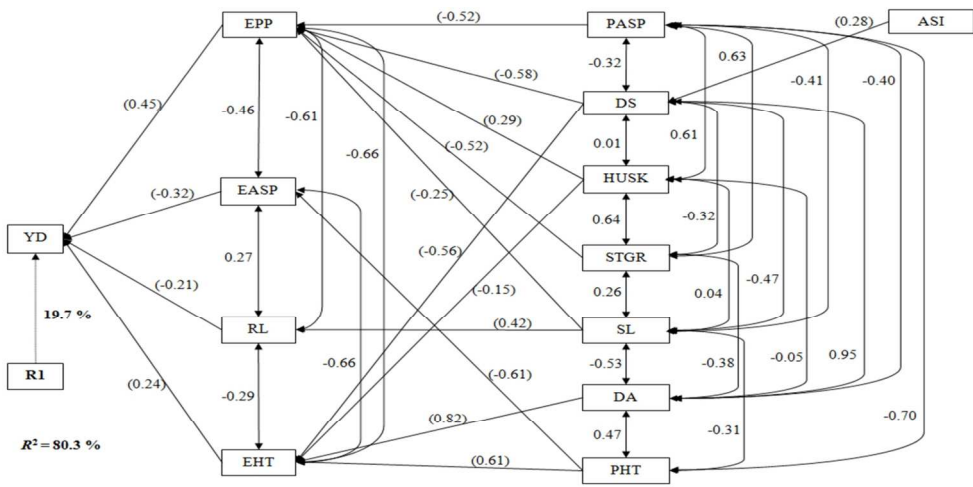


Fig. 2.

127x65mm (300 x 300 DPI)

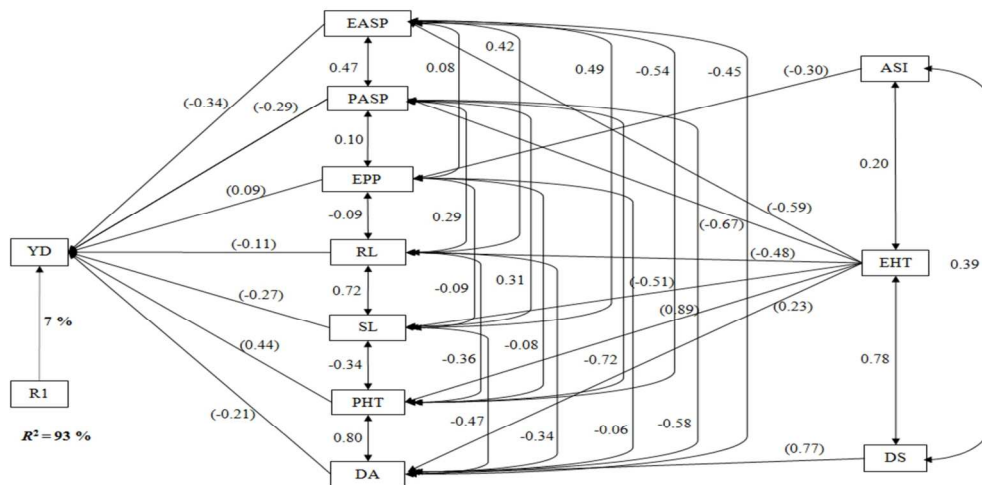
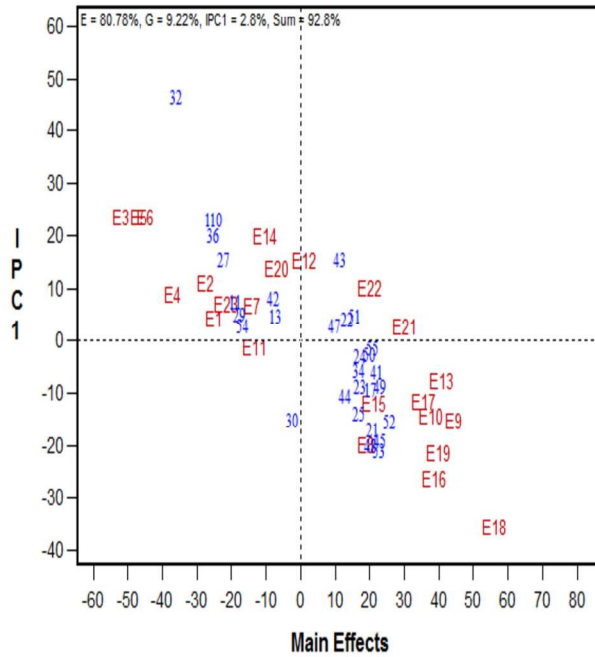


Fig. 3.

130x69mm (300 x 300 DPI)



| Entry | Variety |
|-------|--------------------------------|
| 1 | 95 TZEE-Y |
| 2 | TZEE-W STR 104 BC1 |
| 3 | 97 TZEE-Y 2-C1 |
| 4 | TZEE-Y STR 106 BC1 |
| 5 | CSP SR X TZEE-Y STR |
| 6 | TZEE W ST X GUA 314 BC1 |
| 7 | TZEE-W-SR BC5 (RE) |
| 8 | 98 SYN EE-W |
| 9 | 98 TZEE-W STR |
| 10 | 99 TZEE-Y STR C0 |
| 11 | 99 TZEF-Y POP STR QPM C0 |
| 12 | EV 99 QPM |
| 13 | 2000 SYN EE-W STR |
| 14 | 2000 SYNEE-W STR QPM |
| 15 | FERKE TZEE-W STR |
| 16 | SINE TZEE-W STR |
| 17 | TZEE-W STR 105 BC1 |
| 18 | TZEE-Y POP STR C3 |
| 19 | TZEE-W POP STR C3 |
| 20 | TZEE-Y POP STR C4 |
| 21 | 2004 TZEE-W POP STR C4 |
| 22 | 2004 TZEE-Y POP STR C4 |
| 23 | TZEE-W POP STR QPM C0 |
| 24 | TZEE-W POP STR BC2 C0 |
| 25 | TZEE-W STR 107 BC1 |
| 26 | TZEE-W STR 108 BC1 |
| 27 | TZEE-W POP X LD S6 (SET 1) |
| 28 | TZEE-W POP X LD S6 (SET2) |
| 29 | TZEE-W POP X LD S6 (SETA1) |
| 30 | TZEE-W POP X LD S6 F2 (SET A2) |
| 31 | TZEE-Y Pop STR QPM C0 |
| 32 | TZEE-Y SR BC1 x 9450 STR S6 F2 |
| 33 | TZEE-Y POP STR QPM C1 |
| 34 | TZEE-W POP STR C4 |
| 35 | 99TZEF-Y STR C0 |
| 36 | TZEE-Y Pop STR C0 |
| 37 | 2008 SYN EE-W DT STR |
| 38 | 2008 SYNEE-Y DT STR |
| 39 | 2008 TZEE-W STR |
| 40 | 2008 TZEE-Y STR |
| 41 | TZEE-W POP STR C5 |
| 42 | TZEE-Y POP STR C5 |
| 43 | 2009 TZEE-OR1 STR |
| 44 | 2009 TZEE-OR1 STR QPM |
| 45 | 2009 TZEE-OR2 STR |
| 46 | 2009 TZEE-OR2 STR QPM |
| 47 | 2009 TZEE-W STR |
| 48 | TZEE-W STR 104 |
| 49 | TZEE-W STR 105 |
| 50 | TZEE-Y STR 106 |
| 51 | TZEE-W STR 107 |
| 52 | TZEE-W STR 108 |
| 53 | 2012 TZEE-W DT STR C5 |
| 54 | 2012 TZEE-Y DT STR C5 |
| 55 | TZEE-W DT C0 STR C5 |
| 56 | TZEE-Y DT C0 STR C5 |

Fig. 4.

211x237mm (300 x 300 DPI)