

Genetic variability and advance, heritability, Path coefficient analysis and inter-characters relationships in colocynth (*Citrullus colocynthis* **[L.] Schrad.) collected in Southeastern villages of Benin Republic**

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Abstract

Genetic improvement of crops for important traits requires reliable estimates of genetic variability. heritability and genetic advance of intending parent materials to identify traits useful in planning an efficient breeding program through selection. Effectiveness of genetic improvement of a crop depends on the variability in the agro-morphological traits of individual genotype. Understanding the variation that exists will allow the breeder to determine the breeding strategies to adopt in his breeding program. The objectives of the study were to estimate the magnitude of genetic variability and advance, heritability, inter-characters relationships by simple correlations and path coefficient analysis for yield contributing traits of 40 local colocynth genotypes cultivated in Southeastern Benin. The experiment was carried out in a randomized complete block design with three replications in three locations and during three years. 34 morpho-agronomic traits were observed in each genotype. Analysis of variance revealed that effects of genotype, genotype by year or location and genotype by year by location interactions were significant ($p \le 0.01$ or 0.05) for all the characters. PCV values were relatively greater than GCV for all traits. High magnitude of phenotypic and genotypic coefficient of variations as well as high

heritability along with high genetic advance were recorded particularly for qualitative traits. Correlation and path coefficient analysis were performed on all 34 traits. It was found that all of the yield components except time to emergence, time to tailspins, leaf limb width, fruit number per plant, fruit width, seed length, seed tegument percentage and stem pubescence texture had significant and positive correlations with yield attributes represented by thousand-seeds weight (TSW) and seed number per plant (SNP). Path coefficient analysis indicated that all the characters (except leaf pubescence texture, stem pubescence texture, fruit design produced by secondary color and male flower size) had positive direct effect on TSW and partly SNP. Results suggested that most of the 34 quantitative and qualitative traits studied could be effectively used as selection criteria in the breeding program of *Citrullus colocynthis* varieties with high yielding.

Keywords: *Citrullus colocynthis*, Genetic advance (GA, GAM), Genetic variability, Genotypic coefficients of variation (GCV), Heritability, Phenotypic coefficients of variation (PCV), Path coefficient analysis

Introduction

The origin of colocynth [*Citrullus colocynthis* (L.) Schrad.] is thought to be Africa, where it emerged as an oilseed crop for human consumption. Distribution of the crop around the world was initially based on its use as a food. The crop is a good oil and protein source for segments of the population. It is an ancient crop in West Africa and the region is considered to be the center of diversity for the species (Achigan-Dako et al. 2015, Gama et al. 2013, Simonds 1979, Whitaker and Davis 1962). In Benin, it is annually cultivated traditionally on the family consumption basis. Ecologically, colocynth can be grown in a wide range of environments, and is presently cultivated under diverse agro-climatic conditions in Benin. It can be grown under various rainfalls, temperatures and soil regimes. Depending on the location and maturity period of the cultivar, it is grown during the short rainy growing season between September and February. It is mainly cultivated under a multiple cropping system. Despite the aforementioned importance, its productivity is very low. Documentation of the contribution of plant breeding to a given crop yield improvement and evaluation of the past gains are useful for identifying areas with potential for planning a future breeding program (Waddington et al. 1987). Evans (1993) advocated that an understanding of changes produced by crop breeding on grain yield and its determinants was important to evaluate the efficiency of past improvement work on the advances in genetic yield potential, and to define future selection criteria to facilitate further progress. Genotype, environment and management interact to determine the yield of a crop. However, no method

of estimating long term improvement progress can completely separate genetic effects per se and their interaction effects. Nevertheless, evaluation of popular cultivars from different years in common environments is the most comprehensive and direct method that has been used to estimate progress in yield improvement (Girmay, 2013). Progress made in genetic yield potential and associated changes in morpho-physiological attributes produced by genetic improvement and benefits obtained, thereof, have been documented in different crops in different countries (Girmay 2013, Perry and d'Antuono 1989) by comparing old and modern varieties. Lipids are nutritionally important; they are either fats (solid) or oils (liquid) that consist of fatty acids . A minimum amount of body fat is necessary to provide insulation that prevents heat loss and protects vital organs from shock due to ordinary activities (Anhwange et al. 2010, Boelsma et al. 2001). Deficiency of lipid in the body reflects the type of fatty acids that is lacking in the diet (Anhwange et al. 2010). Therefore, since the common dietary sources of lipids are of animal origin which contained only saturated lipids, it will be of great significance if the essential components of lipids are source from plant products, which had been known to contain them. The usefulness of the essential lipids, which are needed for the proper body functioning and the expensiveness of their processed forms, let us think to provide a veritable channel of harnessing them from local available vegetable sources like *Citrullus colocynthis*.

In low-input farming systems farmers often use a wide range of crop varieties, to provide harvest security, yield stability and the possibility to adapt to changing ecological conditions (Nuijten and van Treuren 2007, Teshome et al. 1999, Hardon and De Boef 1993). Farmers in these situations adopt modern varieties only to a limited extent, or not at all (Nuijten and van Treuren 2007). The main reason for not adopting modern varieties is that they do not meet farmers' requirements. In recent years, various participatory plant breeding initiatives have been set up to develop varieties to meet farmers' requirements better (Nuijten and van Treuren 2007, Almekinders and Elings 2001, Sperling et al. 2001). Some of these programs aim to collaborate with farmers in very early stages of crop development, with the breeder acting more as a facilitator providing the raw genetic material. It is suggested that farmers maintain high levels of crop diversity through developing new varieties and by matching specific varieties to particular conditions (Nuijten and van Treuren 2007). How much genetic diversity these varieties represent, however, has not often been studied and different processes seem to regulate the amount and development of genetic diversity in crops in low-input farming systems. Rather than continuing to assume that the obvious morphological diversity of traditional varieties assures a broad genetic base on-farm, diagnostic surveys of genetic variation and genetic

distance between varieties are urgently needed (Nuijten and van Treuren 2007, Wood and Lenné 1997).

An important step in cultivar development is studying the genetic variability found in genetic resources. The use of genetic resources to create new varieties is important for obtaining higher yields and for the technological transformations required for modernization of agribusiness. It is a dynamic process, but requires continuous enrichment and characterization of the materials maintained in germplasm collections, or by farmers (Gama et al. 2013, Valls 2007, Whitaker and Davis 1962). Attaching a value to genetic resources is a complex task. Describing the kinds of benefits associated with these resources is easier. The simplest benefit arises from the direct use of genetic resources: to produce food and fiber or to help create new varieties of crops and livestock. The ultimate direct-use benefits of crop genetic resources are measured in the increased output, higher quality, better resistance to pests, diseases, and other stress, and other characteristics found in improved crop varieties. These benefits derive not only from the genetic resources contained in precursor wild relatives, but also from the efforts of farmers who domesticated the crop and developed landraces through many years of selection; the work of collectors and gene banks that assembled and preserved genetic material in the form of landraces and wild relatives; and the work of plant breeders who have continued to develop and improve crop varieties.

Plant genetic resources, used for various purposes (agronomy, industry, environment, ecology, medicine, etc.), and thus possessing an important economic and social value are essential for humanity survival. To satisfy the future needs in genetic resources, it is imperative to collect and conserve representative stocks of plant genetic diversity (Koffi et al. 2008, Given 1987). Indeed, the chance for fulfilling future demand of genetic resources is better when a high level of genetic diversity is conserved and made available for breeders. This challenge should not be missed, particularly for the crops such as neglected and underutilized by both national and international research programs, so called minor or orphan crops (Koffi et al. 2008, Rasul et al. 2007). The indigenous edible-seeded colocynths are classified into the minor crops. There are several species of cucurbits in tropical Africa and Asia, cultivated mainly for their oleaginous seeds that are important in the social and cultural life of several peoples (Achu et al. 2005, Zoro Bi et al. 2005, Enujiugha and Ayodele-Oni 2003, Das et al. 2002, Badifu, 1993). *Citrullus colocynthis* belonging to this category of crops, is one of the most widely distributed and consumed at both rural and urban levels in Sub-Saharan Africa. In Sub-Saharan Africa, *C. colocynthis* is prized for its oleaginous seeds consumed as thickeners of a traditional soup called "Egusi" soup in Cameroon, Nigeria or Benin and

pistachio soup in Côte d'Ivoire (Koffi et al. 2008, Loukou et al. 2007, Achu et al. 2005, Zoro Bi et al. 2005, Enujiugha and Ayodele-Oni 2003). This cucurbit is reported to be rich in nutrients (Achu et al. 2005, Enujiugha and Ayodele-Oni 2003, Badifu 1993), namely protein (2.17-36%) and fat (4.73- 45.89%). Thus, *C. colocynthis* represents an excellent plant model for which improved cropping systems implementation can insure the economic prosperity of rural women from tropical Africa. In spite of the nutritional and agronomic potentials of *C. colocynthis*, in depth basic investigations on the crop are scant (Koffi et al. 2008, Osuji et al. 2006). For example, to our knowledge, no detailed study has been devoted to genetic diversity, genetic parameters and reproduction biology. However, investigations reported for others species, suggested that cucurbit family is predominantly outcrossing (Koffi et al. 2008, Montes-Hernandez and Eguiarte 2002). Such expectations are based on the fact that indigenous edible-seeded cucurbits are generally monoecious and entomophilous (Gusmini 2003). The first step of such investigations is the documentation and the assemblage of genetic stocks representative of the total genetic diversity displayed by this species (Koffi et al. 2008, Brown and Briggs 1991, Chapman 1989, Kjellqvist 1975). Results from such investigations are useful to improve both their quality and their productivity through selection and breeding, as well as to implement reliable genetic resources collecting and conservation strategy (Koffi et al. 2008).

Understanding of the relationship between the traits, for the selection of the important traits, is the utmost importance. The goal of the path analysis, is that, the acceptable descriptions of the correlation between the traits, based on a model of cause and effect, is presented, and the importance of the affecting traits on a specific one is estimated. In fact, the basic relationships between the traits are expressed by this analysis. So that, the correlation coefficients dividing to the direct and the indirect effects of the set of the independent variables on a dependent variable, and their importance is calculated (Rasaei et al. 2011). Several researches on the relationships among traits effects of plants have been reported (Rasaei et al. 2011, Amaranthath, and Viswantaha 1990).

The main objectives of this research were therefore to analyze extent of genetic variability and advance, broad sense heritability and the correlations existing between seed yield and related traits in *C. colocynthis* by applying sequential path analysis and identifying traits of genotypes, which may be useful in breeding higher-yielding genotypes.

Materials and methods Plant material

It consisted of forty accessions of *Citrullus colocynthis* collected from farmers in Southeastern of Benin Republic. Those materials are named CC 1 to CC 40, and were multiplied in Ko-Anagodo (in Ifangni Commune belonging to the Department of Plateau in Benin during the short rainy growing season from September 2013 to February 2014.

Study sites and experimental design

Experiments were laid out in a random complete block design with three replications at each location and every year. The experiment plots consisted of two rows, with each comprising five plants. The within- and between-rows were 2m respectively. Adjacent plots were also separated by 2m. One seed was sown per hole. Weeds were removed manually during the growing season. No irrigation was applied. Field trials were carried out at three locations and during three years (2014-2017) in Southeastern Benin. The three locations were Ko-Anagodo – $02^{\circ}72$ 'E, $06^{\circ}67N$ - (Ifangni), Késsounou - $02^{\circ}55E$, $06^{\circ}58N$ - (Dangbo commune) and Idiotchè - 02° 64E, 06° 73N - (Sakété Commune), respectively in the Plateau Department for Ko-Anagodo and Idiotchè locations, and the Ouémé Department for Késsounou cited above. The soils at the three stations were well-drained sandy loams at all locations during the three years of experimentations.

Traits recorded

In total, 34 morpho-agronomic characters were recorded. Data were measured on plot basis. Eight (8) plants from the central rows were used to obtain the plot mean. Characters observed, their codes and the measurement procedures are presented in Zanklan (2024, submitted). Among the traits recorded, 20 were quantitative and 14 qualitative. The qualitative characters were treated as quantitative since they showed continuous variation between the genotypes. Quantitative characters were: time to emergence (TE), time to tailspins (TT), time to male flowering (MF), time to female flowering (FF), time to fruit maturity (MT), limb peduncle length (LPL), limb length (LLL), limb width (LLW), number of fruits per plant (FN), plant height (PH), internode length (IL), number of branches per node (NBN), fruit weight (FW), fruit length (FL), fruit width (FWI), seed number per plant (SNP), seed length (SL), seed width (SWI), seed tegument percentage (TP) and thousand-seeds weight (TSW). Qualitative traits consisted of leaf shape (LS), leaf size (LSi), leaf color (LC), leaf pubescence density (LPD), leaf pubescence texture (LPT), internode length (ILq), stem pubescence density (SPD), stem pubescence texture (SPT), plant canopy coverage (PCC), fruit primary skin color (PSC), fruit secondary skin color (SSC), design produced by fruit secondary color (DPSC), female flower size (FFS) and male flower size (MFS).

Statistical analyses

Mean value over all 9 environments (according to years and locations) for each trait evaluated was used to determine the minimum, maximum and the range for different parameters. Data were subjected to analysis of variance (ANOVA) and covariance using JMP 7.0 (SAS Institute 2007) and Minitab 19 (Minitab 2019) statistical packages. The plot yield can be described as:

$$
Y_{ijkl}=\mu_i+g_{ij}+j_{ik}+l_{il}+\epsilon_{ijkl}
$$

where Y_{ijkl} represents the observed value of the ith trait of the jth genotype for the kth location and the lth year. μ is the trial mean of a given trait; g_{ii} , i_{ik} lil are respectively the effects of genotypes, years and locations; ε_{ijkl} is the error, comprising the genotype \times location interactions, the genotype \times year interactions and the plot error.

Variance components were assessed from the mean squares in ANOVA (Becker 2011, Hill et al. 1998, Wricke and Weber 1986, Falconer 1989). Phenotypic variance of the means for genotypes over years, locations and replications σ_{p}^{2} , was calculated as:

$$
\sigma^2{}_{p\, =}\sigma^2{}_{g}+\sigma^2{}_{gl}/Y+\sigma^2{}_{gk}/L+\sigma^2{}_{glk}/YL+\sigma^2{}_{glkr}/RYL
$$

where σ_{g}^{2} represents the variance component due to genotypes, σ_{g}^{2} is the variance components due to genotype by year interactions, $\sigma_{g_k}^2$ represents the variance components due to genotype by location interactions, σ^2_{glk} represents the variance components due to genotype by location within years, σ^2 _{glkr} is the variance components due to genotype by replication within locations and years (pooled the whole plot error), while R, Y and L represent the number of replications, years and locations, respectively.

Phenotypic (PCV) and genotypic (GCV) coefficients of variation were calculated as:

$$
PCV = (\sigma_p / \underline{X}) \times 100
$$

$$
GCV = (\sigma_g / \underline{X}) \times 100
$$

where σ_p , σ_g and X are roots of the phenotypic, genotypic variances, and grand mean across years and locations, respectively.

Broad sense heritability (h^2) was estimated on genotypic mean basis as described by Hill et al. (1998), Allard (1999) and Falconer (1989) as:

$$
h^2 = \sigma_{g}^2 / \sigma_{p}^2
$$

Expected genetic advance (GA), and GA as percent of the mean assuming selection of the superior 5% of the genotypes were estimated in accordance to Falconer (1989), Fehr (1987) and Johnson et al. (1955) as:

$$
GA = k \times \sigma_p \times h^2
$$

GA (in % of the mean) = $(GA / \underline{X}) \times 100$

where k is the selection differential (with 5% selection intensity, $k = 2.063$), σ_p is the phenotypic standard deviation, h^2 is the heritability, and X refers to the grand mean of a given trait.

Genetic correlation coefficient was estimated from the analysis of covariance as suggested by Miller et al. (1958) and Robinson et al. (1951). Estimations of genetic correlation coefficient (r_g) for any two traits x and y are defined as:

$$
r_g\equiv \sigma_{g1/2}\,/\,\sigma_{g1}\,x\;\sigma_{g2}
$$

where $\sigma_{g1/2}$ is the genetic covariance between two traits, and σ_{g1} , σ_{g2} are the genotypic standard deviations of the first and second characters, respectively. In similar manner, the phenotypic correlation (r_p) can be obtained (Falconer 1989).

Path analysis partitions the total correlation coefficients into direct and indirect effects of various characters, and then helps to quantify the relationships among variables based on *a priori* model. Direct and indirect path coefficients were calculated as initially proposed by Dewey and Fu (1959) and earlier Wright (1921) as:

$$
r_{yi} = P_{yi} + \sum_{i=1}^{n} \cdots r_{ii} P_{yi} \cdots
$$
 for $i \neq i$, and $i' \neq 1$

where r_{yi} is the simple correlation coefficient between the i-th causal variable (x) and effect variable (y) , r_{ii} is the simple correlation coefficient between the i-th and i'-th causal variables, P_{yi} is the path coefficient (direct effect) for the i-th causal variable (x), $r_{ii}P_{yii}$ ^[1] is the indirect effect of the i-th causal variable via the i'-th causal variable. To determine P_{vi} values, square

matrices of the correlation coefficient between independent traits in all possible pairs were inverted and then multiplied by the correlation coefficients among the independent and dependent traits.

Traits closely related to seed yield, namely seed number per plant (SNP) and thousand-seeds weight (TSW) were used as dependent variables for path coefficient analysis in the present study.

Results

Variability, (phenotypic [PCV] and genotypic [GCV] coefficient of variation). heritability (h²) and genetic advance (GA. GAM)

Descriptive statistics for the 20 quantitative and 14 qualitative characters in 40 *Citrullus colocynthis* collections including the minimum, maximum, mean and their standard deviations for data averaged over three locations in three years are summarized in Tables 1 and 2. In general, all the traits exhibited wide range of variation. Significant variation existed then in all traits.

Table 1. Descriptive statistics (range, mean and standard deviation SD) for 20 quantitative traits in 40 tested genotypes over three years and three locations in each year

| Traits | Range | Mean | SD | Traits | Range | Mean | SD | | |
|---------------|----------------|----------------|--------|---------------|--------------|----------------|----------------|--------|--------|
| | Minimum | Maximum | | | | Minimum | Maximum | | |
| TE | 0.00 | 9.00 | 2.66 | 1.49 | IL | 0.80 | 104.00 | 71.84 | 13.21 |
| TT | 1.00 | 9.00 | 2.38 | 1.35 | NBN | 0.40 | 3.12 | 1.48 | 0.43 |
| MF | 19.00 | 63.00 | 39.02 | 7.49 | FW | 50.00 | 231.00 | 158.28 | 35.92 |
| FF | 22.00 | 63.00 | 38.45 | 7.85 | FL | 10.50 | 40.80 | 24.14 | 6.01 |
| MT | 85.00 | 186.00 | 117.45 | 25.69 | FWI | 8.00 | 35.00 | 22.41 | 5.57 |
| LPL | 23.20 | 708.00 | 44.15 | 61.52 | SNP | 222.00 | 1571.00 | 710.99 | 241.18 |
| LLL | 1.50 | 203.00 | 117.62 | 25.02 | SL | 1.30 | 3.00 | 1.77 | 0.27 |
| LLW | 0.95 | 137.00 | 81.32 | 14.85 | SWI | 0.80 | 1.60 | 1.05 | 0.13 |
| FN | 2.00 | 24.00 | 8.94 | 3.47 | TP | 1.44 | 18.20 | 9.81 | 2.79 |
| PН | 1.40 | 16.40 | 8.52 | 2.21 | TSW | 4.40 | 96.6 | 60.03 | 2.83 |

Table 2. Descriptive statistics (range, mean and standard deviation SD) for 14 qualitative traits in 40 tested genotypes over three years and three locations in each year

Combined analysis of variance showed that there were significant differences for all the parameters considered among the colocynth genotypes collected (Tables 3 and 4). Based on the results obtained, statistical analysis revealed that the genotypes included in the study had highly significant variation ($p \leq 0.01$) for all traits under study. It could be concluded that

differences between colocynth collections may be due to genetical differences between cultivars and indicating considerable amount of variation present in these material and revealing a high level of diversity among the genotypes for these traits, excepted for number of branches per node (NBN), seed length (SL), seed tegument percent (TP) and most all the qualitative characters. Effects of the factors yield (Y) and location (L) were quite non-significant for all the quantitative traits compared to qualitative ones. Furthermore, genotypic mean variance significance was lower for qualitative traits than for quantitative characters (Tables 3 and 4). Interactions between genotypes and the factors years and location were highly significant ($p \le 0.05$ or 0.01) for all traits observed.

The magnitude of phenotypic variation does not reveal the relative amount of genotypic and non-genetic components of variation. Moreover, it is difficult to compare the variances between various traits because they are not unit free. Thus, variance components, estimates of phenotypic coefficient of variation (PCV), genotypic coefficient of variation (GCV), broad sense heritability, genetic advance (GA) and genetic advance as a percentage of mean (GAM) were calculated and compared (Tables 5 and 6). In general, higher PCV values than GCV were obtained for all the characters. For quantitative traits PCV were very high and ranged from 76.251 (leaf limb width - LLW) to 839.988 % (thousand-seeds weight - TSW). It ranged from 74.275 (leaf size - LSi) to 284.484 % (design produced by fruit secondary skin color - DPSC) for qualitative characters. Quantitative characters were observed to have the highest PCV. The genotypic coefficient of variation (GCV) measures the genotypic variance relative to the mean of a given character and is independent from the units of measurements. These values allow direct comparison of various characters. GCV showed similar trends as PCV and ranged from 15.906 (time from sowing to maturity - MT) to 291.046 % (limb peduncle length) concerning quantitative characters. For qualitative traits, GCV ranged from 44.830 (leaf size - LSi) to 213.666 % (DPSC).

| | three locations within each year in Southeastern of Benin Republic | | | | | | | | | | | |
|---------------|--|------------|-------------|-----------------------------|--------------|-----------------------|--------|--|--|--|--|--|
| Traits | | | | Sources of variation | | | | | | | | |
| | Year | Location | Genotype | $G \times Y$ | $G \times L$ | $G \times Y \times L$ | Pooled | | | | | |
| | (Y) | (L) | (G) | | | | error | | | | | |
| TE | 65.878 | 54.269 | $5,778$ * | $2.883***$ | $3.044**$ | $2.104**$ | 1.879 | | | | | |
| TT | 65.389 | 0.889 | $5.510*$ | 2.556^{**} | $2.922**$ | $1.624**$ | 1.567 | | | | | |
| MF | 6166.80 | 482.70 | 94.56** | 58.470** | 130.034** | 60.874** | 42.14 | | | | | |
| FF | 11840.9 | 1263.5 | $63.9**$ | $76.505**$ | $19.220**$ | $60.463**$ | 36.6 | | | | | |
| MT | 249905 | 12983 | 349^{**} | 220.385** | 89.643** | 284.309** | 167 | | | | | |
| LPL | $63.9**$ | 61916.3 | $16513.0**$ | 6594.196** | 772.358** | 6974.364** | 3206.9 | | | | | |
| LLL | 5087.68 | 460.08 | 2224.70** | 1683.827** | 1290.289** | 1224.217** | 558.49 | | | | | |
| LLW | 376.88 | 1181.35 | 587.43** | 556.858** | 535.366** | $402.374**$ | 204.84 | | | | | |
| FN | 62.381 | 5.458 | 35.580** | 16.808** | $17.594**$ | $9.541***$ | 11.115 | | | | | |
| PH | $1.684**$ | $7.374***$ | $14.537**$ | $12.408**$ | $5.963**$ | $5.320**$ | 4.570 | | | | | |
| \mathbf{L} | 9743.7 | 328.1 | $560.7***$ | 228.053** | 470.018** | 183.130** | 141.3 | | | | | |
| NBN | 0.571 | 0.177 | 0.675 | $0.354**$ | $0.280**$ | $0.230**$ | 0.171 | | | | | |
| FW | 9743.7 | 562.4 | 5681.9** | 3481.922** | 2587.484** | 1200.447** | 1104.9 | | | | | |
| FL | $16.015***$ | $2.842**$ | 178.942** | 85.309** | $56.615**$ | 42.714** | 31.001 | | | | | |
| FWI | 530.219 | 170.553 | 131.834** | 93.386** | $41.457**$ | 28.417** | 26.094 | | | | | |
| SNP | 146689 | 33892 | 129509** | 105449.743** | 72934.410** | 109837.211** | 55456 | | | | | |
| SL | 15.682 | 2.150 | 0.086 | $0.093**$ | $0.063**$ | $0.055***$ | 0.040 | | | | | |
| SWI | 3.808 | 0.761 | 0.012 | $0.008***$ | $0.010**$ | $0.013***$ | 0.008 | | | | | |
| TP | 1587.00 | 242.95 | 7.23 | $11.578***$ | 3.806^{**} | $5.168***$ | 4.34 | | | | | |
| TSW | 2167.2 | 420.6^* | 834.0** | 93487.179** | 74548.564** | 82046.698** | 801.4 | | | | | |

Table 3. Mean squares from combined analysis of variance for 40 *Citrullus colocynthis* genotypes evaluated by 20 quantitative morpho-agronomic characters across three years and three locations within each year in Southeastern of Benin Republic

*and **: significant at 0.05 and 0.01. respectively

Table 4. Mean squares from combined analysis of variance for 40 *Citrullus colocynthis* genotypes evaluated by 14 qualitative characters across three years and three locations within each year in Southeastern of Benin Republic

| Traits | Sources of variation | | | | | | | | | | | |
|---------------|-----------------------------|-----------------------|--------------------------------|---|--------------|-----------------------|--------------|--|--|--|--|--|
| | Year (Y) | Location (L) | Genotype (G) | $G \times Y$ | $G \times L$ | $G \times Y \times L$ | Pooled error | | | | | |
| LS | 0.000^{**} | $0.900**$ | $30.146*$ | 0.000^{**} | $25.207**$ | $0.000**$ | 0.000 | | | | | |
| LSi | $0.000***$ | 1.200 | 7.684 | $0.000**$ | $11.353**$ | $0.000**$ | 0.856 | | | | | |
| LC | 0.000^{**} | $0.300**$ | 34.923** | $0.000**$ | 19.992** | $0.000**$ | 1.508 | | | | | |
| LPD | $0.000**$ | $0.300**$ | 23.353** | $0.000**$ | $21.530**$ | $0.000**$ | 1.624 | | | | | |
| LPT | $0.000**$ | $0.000**$ | 18.800 | $0.000**$ | $20.923**$ | $0.000**$ | 1.578 | | | | | |
| ILq | $0.000***$ | $0.300**$ | 9.418 | $0.000**$ | $7.513**$ | $0.000**$ | 0.659 | | | | | |
| SPD | $0.000***$ | $0.000**$ | 37.507** | $0.000**$ | 13.230** | $0.000**$ | 0.998 | | | | | |
| SPT | $0.000**$ | $0.300**$ | 36.892 | $0.000**$ | 15.684** | $0.000**$ | 1.183 | | | | | |
| PCC | $0.000**$ | $0.000**$ | 15.069 | $0.000**$ | 18.461** | $0.000**$ | 1.392 | | | | | |
| PSC | $0.000***$ | $0.900**$ | 77.038 | $0.000**$ | $23.361**$ | $0.000**$ | 1.762 | | | | | |
| SSC | 0.000^{**} | $0.300**$ | 34.700 | $0.000**$ | $11.069**$ | $0.000**$ | 0.835 | | | | | |
| DPSC | $0.000**$ | 1.200 | 28.915 | $0.000**$ | 19.661** | $0.000**$ | 1.483 | | | | | |
| FFS | $0.000**$ | $0.000**$ | 22.646 | $0.000^{\ast\ast}$ | $14.153***$ | $0.000**$ | 1.067 | | | | | |
| MFS | $0.000**$ | $0.000**$ | 23.107 未来 \cdot \sim | $0.000**$ \sim \sim \sim \sim \sim | $13.230**$ | $0.000**$ | 0.998 | | | | | |

: significant at 0.01

Broad sense heritability (h^2) has also to be considered to predict the amount of progress from farmer selection in the present study. Broad sense heritability was generally low and showed values under 50 % for all 20 quantitative traits (Table 5). In general, values estimated for qualitative characters were higher than for quantitative traits. It ranged from 0.132 (Time to maturity - MT) to 43.281 % (fruit length - FL). For qualitative

traits, heritability ranged from 36.429 (leaf size - LSi) to 95.859 % (male flower size - MFS). Obtained results showed that qualitative characters are highly heritable, and almost traits presented high heritability around 50 % (Table 6). This fact is maybe interesting as it shall be possible to select for them limiting costs involved in research activities. Heritability estimates along with genetic advance is more helpful in foresee the genetic gain under selection than heritability estimates alone. High genetic advance (GA) and genetic advance related to trait mean (GAM) joined with heritability was observed for all quantitative traits, particularly leaf peduncle length (LPL), leaf limb length (LLL), leaf limb width (LLW), fruit weight (FW), fruit length and width (FL and FWI) and seed number per plant (SNP). Therefore, selection of genotypes based on these indicated traits will be effective. Nevertheless, GA and GAM were relatively low for thousand-seeds weight (TSW) in spite its importance in enhancing seed yield in colocynth (Table 5). For qualitative characters, GA was relatively low. However, GAM was higher, and sometimes above 100 % (Table 6).

Table 5. Variability (phenotypic [PCV] and genotypic [**G**CV] coefficient of variation). heritability (h²) and genetic advance (GA. GAM) for 40 farmers' genotypes of *Citrullus*

| | coroo <i>jimino</i> tootou | | | | |
|---------------|----------------------------|---------|-----------|---------|---------|
| Traits | PCV $(\frac{9}{6})$ | GCV(%) | $h^2(\%)$ | GA | GAM |
| TE | 437.055 | 90.140 | 4.253 | 1.022 | 38.353 |
| TT | 376.500 | 98.528 | 6.848 | 1.267 | 53.192 |
| MF | 214.934 | 24.917 | 1.344 | 2.325 | 5.959 |
| FF | 300.581 | 20.786 | 0.478 | 1.140 | 2.965 |
| MT | 437.483 | 15.906 | 0.132 | 1.401 | 1.193 |
| LPL | 701.903 | 291.046 | 17.193 | 109.925 | 248.969 |
| LLL. | 95.167 | 40.101 | 17.756 | 41.002 | 34.860 |
| LLW | 76.251 | 29.803 | 15277 | 19.543 | 24.032 |
| FN | 140.721 | 66.677 | 22451 | 5.830 | 65.177 |
| PН | 84.489 | 44.734 | 28.033 | 4.164 | 48.862 |
| П. | 150.268 | 32.959 | 4.810 | 10.714 | 14.913 |
| NBN | 105.820 | 55.453 | 27.461 | 0.888 | 59.950 |
| FW | 98.613 | 47.623 | 23.322 | 75.098 | 47.446 |
| FL | 84.213 | 55.402 | 43.281 | 18.155 | 75.193 |
| FWI | 142.644 | 51.233 | 12900 | 8.507 | 37.961 |
| SNP | 113.722 | 50.615 | 19.809 | 330.436 | 46.475 |
| SL | 240.097 | 16.518 | 0.473 | 0.041 | 2.344 |
| SWI | 204.310 | 10.412 | 0.259 | 0.011 | 1.094 |
| TP | 439.848 | 27.407 | 0.388 | 0.345 | 3.523 |
| TSW | 839.988 | 48.103 | 0.327 | 3.411 | 5.683 |

colocynthis tested by 20 quantitative traits

| | conceptures wowe or чиш uu ve uu u | | | | | | | | | | |
|---------------|---|---------|-----------|--------|------------|--|--|--|--|--|--|
| Traits | PCV $(\frac{9}{6})$ | GCV(%) | h^2 (%) | GA | GAM | | | | | | |
| LS | 112.784 | 82.564 | 53.590 | 8.291 | 124.690 | | | | | | |
| LSi | 74.275 | 44.830 | 36.429 | 3.451 | 55.820 | | | | | | |
| LC | 145.770 | 114.378 | 61.567 | 9.566 | 185.148 | | | | | | |
| LPD | 127.482 | 90.046 | 49.892 | 7.041 | 131.214 | | | | | | |
| LPT | 131.154 | 88.487 | 45.519 | 6.034 | 123.163 | | | | | | |
| IL q | 101.511 | 73.653 | 52.643 | 4.593 | 110.246 | | | | | | |
| SPD | 153.036 | 130.304 | 72.498 | 10.757 | 228.887 | | | | | | |
| SPT | 148.036 | 122.292 | 68.243 | 10.351 | 208.416 | | | | | | |
| PCC | 104.592 | 68.705 | 43.150 | 5.260 | 93.107 | | | | | | |
| PSC | 162.430 | 140.434 | 74.749 | 15.655 | 250.482 | | | | | | |
| SSC | 156.242 | 134.387 | 73.980 | 10.452 | 238.462 | | | | | | |
| DPSC | 284.484 | 213.666 | 56.409 | 8.331 | 331.063 | | | | | | |
| FFS | 109.884 | 84.978 | 59.805 | 7.592 | 135.574 | | | | | | |
| MFS | 119.748 | 117.243 | 95.859 | 9.709 | 236.812 | | | | | | |

Table 6. Variability (phenotypic [PCV] and genotypic [**G**CV] coefficient of variation). heritability (h²) and genetic advance (GA. GAM) for 40 farmers' genotypes of *Citrullus colocynthis* tested by 14 qualitative traits

Correlations between characters

Correlation analysis figures out the intensity of relationship between two traits. For the present breeding material genotypic correlations were computed among yield contributing components and are presented in Table 7.

The genotypic correlation coefficients among 34 morpho-agronomic quantitative and qualitative characters in colocynth in nine environments (three years and three locations per year) are presented in Table 7. Seed number per plant (SNP) and thousand-seeds weight (TSW) are important components of seed yield. SNP presented significant positive genotypic correlations with all traits studied, except time to emergence (TE), time to tailspins appearance (TT), leaf limb width (LLW), fruit number per plant (FN), fruit width (FWI), seed length (SL), seed tegument percentage (TP) and stem pubescence density (SPD). Apart from TE, TT, MF, FF, MT, FL, SL, LPT, SPD and MFS, TSW exhibited positive associations with the remaining 24 characters (Table 7). From Table 7, the differences in both magnitude and direction of correlation coefficients were observed considering the nine environments taken in the present study for some of the characters. Positive significant correlation coefficients were observed for most traits except for those cited above.

Table 7. Associations of 20 morpho-agronomic quantitative and 14 qualitative characters in 40 genotypes of *Citrullus colocynthis* evaluated across three years (2014-2017) and three locations each year in Southeastern Benin

Path coefficient analysis

Path coefficients are the separation of correlation coefficients into components of direct and indirect effects. The use of methods needs cause and effect situation among the variables and direction in the casual system is assigned, based upon experimental evidences. The Tables 8 and 9 present estimates of direct and indirect contributions of 33 traits to seed yield contributing components represented in this study by seed number per plant (SNP) and thousand-seeds weight (TSW). Simple correlation coefficients indicate only the general association between any two traits without possible causes of such association. Path coefficient analysis presents a better understanding of cause-and-effect relationships among different characters and plays therefore an important role in determining the degree of relationship between yield and its components. Thus, the path coefficient analysis was performed to partition the correlation coefficient into direct and indirect effect of various characters on yield. Results pertaining to path coefficient analysis are presented in Tables 8 and 9. Path coefficient analysis based on SNP and TSW as dependent variables was then studied. It revealed positive direct effect on SNP for most of the traits except for TE, MF, FF, LLW, FN, PH, FWI and SL (Table 8). Moreover, indirect effects of SNP via the remaining 33 independent characters were positive or negative and often very high and then considerable. Because of those indirect effects, this trait should be considered along with direct effects on seed yield. TSW as dependent variable, showed high positive direct effect with all traits apart from LPT, SPT, DPSC and MFS. Its indirect effects via all remaining 33 characters were very various and positive or negative, and need carefully analyses for the use of those traits as selection criteria.

In general, results of the direct and indirect path coefficients that estimated the extent of the relationships between seed yield (number) per plant or thousand-seeds weight and 33 yield related characters using the genotypic correlation values are presented here. All traits evaluated in this study were genotypically significantly correlated with seed yield in the two directions. Path analysis partitioned these correlation coefficients into their components, thus giving an insight to which traits can be selected for improvement in seed yield.

Discussions

Genetic variability

Significant differences were observed among the genotypes for all the traits. These differences indicated the presence of high variability and

opportunity for improvement. A wide range of variation for all traits under observation in colocynth was noted under different environmental conditions. Such results had been reported by Jamshidian et al. (2013), Cervenski et al. (2011), Singh et al. (2011), Sharma (2010), Kaygısız (2009), Meena et al. (2009) and Balkaya et al. (2005) in cabbage., In *Citrullus colocynthis*, studies in that item are scarce. Highly significant means squares for genotypic differences were found in the combined analysis of variance for all the traits observed, confirming the possibility of improving these traits through selection. Also, genotypes responded differently to changes in years and locations, as genotype \times year, genotype \times location and genotype \times year \times location interactions mean squares were highly significant (p≤0.05 or p≤0.01) for all the characters. This indicates that ranking between genotypes was affected significantly by differences in environments even if those genotypes were tested at three various locations during three different years. **Table 8.** Path coefficient analysis (direct and indirect effects) of component traits on seed

| | Table 8. Continued | | | | | | | | | | | | |
|--------------|---------------------------|--------------------------|--------------------------|----------|------------|-----------|--------------------------|---------------------|------------|--------------------------|----------|--------------------------|------------|
| Trait | Direct effect | | | | | | | Indirect effect via | | | | | |
| | | NBN | FW | FL | FWI | SL | SWI | TP | TSW | LS | LSi | LC | LPD |
| TE | -0.040 | 2.451 | 1.700 | -1.306 | 0.781 | 0.310 | 0.394 | -0.093 | -1.619 | 1.666 | -2.291 | 0.959 | 1,666 |
| TT | 0.000 | 1.047 | 2.695 | -0.492 | -1.000 | 0.094 | 0.243 | 0.098 | -1.800 | -2.021 | -2.689 | 0.405 | 0,064 |
| MF | -0.170 | 2.500 | 0.459 | -2.233 | -0.108 | -0.065 | 0.253 | 2.666 | 0.515 | -0.191 | 0.000 | 2.214 | -0.096 |
| FF | -0.005 | 2.478 | -0.048 | -0.437 | -0.021 | 0.024 | 0.116 | -0.307 | 2.500 | -0.632 | 1.083 | 2.285 | 1,750 |
| MT | 0.003 | -0.480 | 0.133 | 1.440 | -0.625 | -0.083 | -0.046 | 0.151 | 0.627 | -2.500 | 0.588 | 2.625 | 1,666 |
| LPL | 0.061 | 0.040 | -0.518 | -0.174 | -0.625 | 2.777 | -0.363 | 2.222 | 0.538 | -2.631 | 1.060 | 1.866 | 2,500 |
| LLL | 0.021 | -0.053 | 0.370 | -0.127 | -0.381 | -1.818 | -0.016 | 2.523 | -0.166 | 0.632 | -1.904 | -0.370 | -0.579 |
| LLW | -0.061 | -0.900 | 0.539 | -0.313 | 0.130 | 0.231 | 2.800 | 0.042 | 2.400 | 2.861 | -1.239 | 0.471 | 0,200 |
| FN | -0.047 | -0.410 | 2.133 | -0.355 | -0.024 | 0.432 | 0.769 | 0.054 | -2.473 | 2.000 | -1.369 | -0.553 | $-3,000$ |
| PH | -0.011 | 0.000 | 0.395 | 0.028 | 0.693 | -0.818 | -0.120 | 1.888 | 0.000 | 1.781 | 2.100 | 0.396 | 0,745 |
| \mathbf{L} | 0.028 | -0.958 | 0.130 | 0.248 | 0.831 | -0.212 | -0.122 | 0.372 | 1.107 | 0.500 | -0.333 | -7.428 | 2,833 |
| NBN | 0.029 | $\overline{}$ | 2.200 | -0.115 | -0.271 | -0.509 | -0.227 | 1.404 | -0.888 | -3.062 | 0.465 | 0.362 | 2,538 |
| FW | 0.060 | -1.200 | $\overline{}$ | 0.595 | 1.295 | -0.650 | -0.582 | 0.714 | -1.960 | 2.000 | -0.140 | 1.272 | 2,558 |
| FL | 0.023 | 2.400 | -0.595 | \sim | -0.127 | 0.666 | -1.166 | 1.750 | -2.000 | 0.535 | 1.916 | 0.566 | $-2,250$ |
| FWI | -0.086 | 1.200 | -1.033 | 0.127 | ω | 1.233 | 1.333 | 0.747 | 1.500 | 2.722 | 2.428 | 2.476 | 1,369 |
| SL | -0.017 | 2.200 | -0.752 | 0.045 | 0.806 | \sim | 0.037 | 0.057 | -0.439 | 2.750 | -3.157 | -0.050 | 0,645 |
| SWI | 0.016 | 1.000 | -0.516 | -0.022 | 1.483 | -0.037 | $\overline{}$ | 0.101 | 0.076 | -1.058 | -2.166 | 1.600 | $-2,928$ |
| TP | 0.000 | 1.800 | -1.123 | 0.153 | -0.258 | 0.058 | -0.101 | \sim | 0.944 | 0.000 | -2.214 | -2.500 | 0,764 |
| TSW | 0.038 | 1.600 | -0.550 | -0.013 | 1.387 | -0.031 | -0.005 | -0.944 | | 2.500 | 2.625 | -0.512 | $-1,225$ |
| LS | 0.088 | -2.800 | 0.089 | -0.199 | 3.225 | -0.132 | 0.101 | -2.000 | -2.500 | $\overline{}$ | -1.875 | 0.461 | 1,217 |
| LSi | 0.083 | -2.800 | -0.078 | -0.150 | 2.741 | -0.106 | 0.073 | -1.722 | -2.000 | 1.875 | | 0.552 | $-1,951$ |
| LC | 0.058 | -2.800 | -0.078 | -0.150 | 2.741 | -0.106 | 0.073 | -1.722 | -2.000 | 1.875 | 0.000 | $\overline{}$ | $-2,857$ |
| LPD | 0.011 | 2.200 | -0.977 | 0.111 | 0.161 | 0.035 | -0.077 | -0.240 | 2.333 | 1.875 | -0.701 | 0.607 | |
| LPT | 0.081 | -2.200 | -0.337 | -0.075 | 2.000 | -0.065 | 0.030 | -1.296 | -2.166 | 2.750 | -0.201 | -1.428 | -0.950 |
| ILq | 0.050 | -1.800 | -0.359 | -0.068 | 1.935 | -0.061 | 0.026 | -1.259 | -2.833 | 2.000 | -0.219 | -1.357 | -0.916 |
| SPD | 0.038 | -1.800 | -0.359 | -0.068 | 1.935 | -0.061 | 0.026 | -1.259 | -2.833 | 2.000 | -0.219 | -1.357 | $-0,916$ |
| SPT | 0.116 | -2.200 | 0.224 | -0.238 | 3.612 | -0.153 | 0.124 | -2.222 | -1.500 | -1.500 | 0.236 | -2.214 | $-1,783$ |
| PCC | 0.054 | -1.400 | -0.382 | -0.062 | 1.870 | -0.058 | 0.022 | -1.222 | -2.500 | 2.250 | -0.236 | -1.285 | -0.883 |
| PSC | 0.069 | -1.400 | -0.382 | -0.062 | 1.870 | -0.058 | 0.022 | -1.222 | -2.500 | 2.250 | -0.236 | -1.285 | -0.883 |
| SSC | 0.039 | 0.400 | -0.483 | -0.032 | 1.580 | -0.042 | 0.005 | -1.055 | -1.000 | 2.375 | -0.315 | -0.964 | $-0,733$ |
| DPSC | 0.030 | 1.800 | -0.561 | -0.009 | 1.354 | -0.030 | -0.007 | -0.925 | 0.166 | 3.250 | -0.377 | -0.714 | $-0,616$ |
| FFS | 0.020 | 2.600 | -0.606 | 0.003 | 1.225 | -0.022 | -0.015 | -0.851 | 0.833 | 2.750 | -0.412 | -0.571 | -0.550 |
| MFS | 0.061 | 2.000 | 0.179 | 0.120 | -2.451 | 0.090 | -0.056 | 1.555 | 2.500 | -3.000 | 0.078 | 1.928 | 1,183 |

Table 8. Continued

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Table 9. Path coefficient analysis (direct and indirect effects) of component traits on seed yield parameter represented by thousand-seeds weight (TSW) for data recorded across three different years and three locations in each year.

| | | | | | | Table 9. Continued | | | | | | | |
|---------------|----------------------|--------------------------|----------|--------------------------|------------|---------------------------|--------------------------|---------------------|----------------|----------|----------|--------------------------|------------|
| Traits | Direct effect | | | | | | | Indirect effect via | | | | | |
| | | FW | FL | FWI | SNP | SL | SWI | TP | LS | LSi | LC | LPD | LPT |
| TE | 0.034 | -2.000 | 0.816 | -1.375 | 0.964 | -0.006 | -0.016 | 0.527 | -1.600 | 1.208 | -1.693 | -2.055 | -0.880 |
| TT | 0.066 | -1.956 | 0.276 | -3.142 | 1.333 | 0.109 | 0.118 | 0.486 | 0.565 | 1.241 | -1.648 | 1.645 | -0.789 |
| MF | 0.124 | -3.400 | 0.114 | -0.366 | -0.583 | 0.082 | 0.093 | 0.212 | 0.326 | 0.862 | -1.351 | 1.290 | $-0,210$ |
| FF | 0.112 | -2.217 | 0.289 | -0.350 | -2.500 | 0.032 | 0.028 | 0.217 | -2.461 | 0.531 | 1.583 | 2.142 | $-2,250$ |
| MT | 0.066 | -1.304 | 0.590 | -0.662 | -2.750 | -0.021 | -0.026 | 0.285 | 2.700 | -0.941 | 1.150 | 2.466 | $-1,538$ |
| LPL | 0.038 | 0.469 | -0.174 | -0.187 | -1.315 | -1.111 | 2.363 | 1.722 | 1.000 | -0.878 | 1.600 | 7.333 | $-0,186$ |
| LLL | 0.045 | -0.102 | 0.297 | 0.084 | 0.000 | 1.818 | -0.836 | 2.000 | 0.068 | 0.190 | -0.467 | -0.275 | 0,195 |
| LLW | 0.046 | -0.113 | -0.052 | 0.130 | 0.205 | 0.380 | 2.800 | 0.418 | -0.027 | 0.154 | -0.679 | -0.650 | 2,000 |
| FN | 0.019 | -0.977 | 0.140 | 0.129 | 0.885 | 0.328 | 0.307 | 1.000 | -0.961 | 0.538 | 1.276 | 2.666 | -0.325 |
| PH | 0.068 | 0.008 | -0.201 | -0.387 | 1.166 | 3.045 | -2.600 | 1.037 | 0.625 | 2.000 | 0.283 | 0.098 | $-1,937$ |
| \mathbf{L} | 0.035 | -0.384 | 0.184 | 0.337 | -0.930 | 0.056 | 0.048 | 0.372 | -0.653 | 1.333 | -1.857 | 1.277 | $-2,300$ |
| NBN | 0.058 | -2.200 | 0.105 | 0.037 | -0.150 | 0.980 | 2.181 | 1.071 | -0.187 | -0.095 | 0.400 | 1.692 | -0.560 |
| FW | 0.079 | $\overline{}$ | -0.303 | -0.323 | 0.213 | 0.640 | 0.810 | 0.207 | 1.750 | 0.180 | 0.290 | 0.176 | $-1,304$ |
| FL | 0.028 | 0.303 | ω | -0.013 | -1.750 | -1.857 | -1.166 | 1.400 | -0.070 | -0.750 | 1.433 | -1.125 | 0,038 |
| FWI | 0.046 | 0.258 | 0.013 | $\overline{}$ | 0.322 | 0.323 | 0.440 | 0.422 | -0.222 | -2.000 | -1.857 | -0.345 | 0,159 |
| SNP | 0.052 | 0.146 | 0.045 | -0.322 | \sim | -2.833 | 1.400 | 1.076 | 0.086 | -0.074 | 1.222 | 0.730 | 0,548 |
| SL | 0.056 | 0.741 | -0.127 | 1.387 | 1.833 | $\overline{}$ | -0.003 | 0.165 | -1.750 | 1.500 | -1.366 | 2.322 | -0.679 |
| SWI | 0.056 | 0.719 | -0.120 | 1.322 | 2.500 | 0.003 | $\overline{}$ | 0.174 | 0.882 | 3.055 | 2.333 | -2.000 | 1,619 |
| TP | 0.106 | -0.325 | 0.183 | -1.677 | -2.000 | 0.167 | -0.174 | \sim | 0.000 | -0.904 | -1.083 | 1.352 | 1,363 |
| ${\bf L S}$ | 0.066 | 0.213 | 0.026 | -0.129 | 1.000 | 0.083 | -0.084 | 0.000 | \overline{a} | -1.250 | 0.207 | 0.320 | 0,026 |
| LSi | 0.025 | -0.180 | 0.750 | 2.000 | 0.074 | -2.000 | -3.055 | 0.904 | 1.250 | | 0.219 | -0.365 | 1,750 |
| LC | 0.051 | -0.290 | -1.433 | 1.857 | -3.222 | 1.366 | -5.333 | 1.083 | 1.375 | -0.219 | | 0.357 | 0,958 |
| LPD | 0.042 | -0.176 | 2.125 | 0.345 | -0.730 | -2.322 | 2.000 | -1.352 | 3.125 | -0.131 | -0.357 | $\overline{}$ | 0,600 |
| LPT | -0.377 | -0.529 | -1.370 | -0.583 | 1.384 | 1.870 | 1.142 | -0.941 | -2.375 | 0.149 | -1.892 | -1.083 | |
| ILq | 0.021 | 0.157 | 0.360 | 0.560 | 0.137 | 1.900 | -1.410 | -2.500 | 0.222 | 0.000 | -0.196 | 0.087 | 0,141 |
| SPD | 0.000 | 1.135 | -0.272 | -0.655 | 0.743 | -0.480 | -0.314 | -0.825 | -0.100 | -0.434 | -1.757 | -0.786 | -0.157 |
| SPT | -0.017 | -0.641 | 1.166 | 0.192 | -0.259 | -0.800 | 0.508 | 1.032 | 0.073 | -0.568 | 0.217 | 1.111 | 0,013 |
| PCC | 0.033 | -0.433 | 2.000 | -1.636 | 0.962 | -2.583 | -1.100 | -1.777 | 0.090 | 0.241 | -0.023 | -0.062 | 0,328 |
| PSC | 0.042 | 0.838 | 0.929 | 2.900 | 1.392 | -1.066 | -1.000 | 0.069 | -1.625 | -1.666 | -0.060 | 0.869 | -0.179 |
| SSC | 0.013 | -1.571 | 1.134 | 0.670 | 2.500 | -1.195 | -1.384 | -0.073 | 1.285 | -0.166 | -0.059 | -0.541 | 3,444 |
| DPSC | -0.004 | 1.280 | 1.950 | 0.820 | 2.090 | -2.512 | 1.600 | -0.071 | -0.451 | -1.108 | -0.074 | -1.130 | 0,953 |
| FFS | 0.010 | -0.130 | 1.178 | 0.453 | 1.714 | 0.972 | 1.000 | 0.547 | -0.117 | 0.073 | -0.024 | 0.000 | 0,157 |
| MFS | -0.052 | 2.565 | 1.200 | -1.500 | -1.022 | -1.000 | -0.185 | -1.571 | -0.188 | -0.714 | 0.283 | -0.340 | -0.112 |

Table 9. Continued

The magnitude of phenotypic variation does not reveal the relative amount of genotypic and non-genotypic components of variation (Falconer 1989). Furthermore, it is difficult to compare variations for various characters because they are not unit free. Since most of the economic characters (grain yield) are complex in inheritance and are greatly influenced by several genes interacting with various environmental conditions, study of phenotypic coefficient of variation (PCV) and genotypic coefficient of variation (GCV) is not only useful for comparing the relative amount of phenotypic and genotypic variations among different traits but also very useful to estimate the scope for improvement by selection (Bello et al. 2012). The reliability of a parameter to be selected for breeding programs among other factors is dependent on the magnitude of its coefficient of variations, especially the GCV. However, the differences between genotypic and phenotypic coefficient of variability indicate the environmental influence (Bello et al. 2012). While a lower value of CV generally depicts low variability among the tested samples, a high proportion GCV to the PCV is desirable in breeding works. Results given in Tables 5 and 6 depicted that phenotypic variances and PCVs were slightly higher than genetic variances and GCVs for all the characters, suggesting the least influence of environment in the expression of these characters Similar results have also been reported by Bello et al. (2012) and Nelson et al. (1992). Estimates of genotype \times year, genotype \times location and genotype \times year \times location interaction variances for the traits in most instances were low considering qualitative characters. This result tends to support the notion that greater heterozygosity confers a buffering effect or stability over a wide range of environments, whereas inbreeding leads to increased homozygosity and less buffering capacity (Falconer 1989, Nigussie and Saleh 2007). High genetic variability for grain yield in the genotypes over years recorded in the test materials suggested that it could be further exploited through improvement and selection programs (Wannows et al. 2010, Najeeb et al. 2009, Kashiani et al. 2008, Aziz et al. 1998).

High values of PCV and GCV observed in most of the traits show that the selection can be effective for these traits but also indicated the existence of substantial variability, ensuring ample scope for their improvement through selection. These observations are in confirmation with the findings of Bishwas and Singh (2024), Mitra et al. (2023) and Bello et al. (2012).

High magnitude of broad sense heritability estimated in all qualitative characters (Table 6). This implied the possibility of effective selection for genetic improvement of these traits. Values of genetic advance as percentage of mean (GAM) ranged from 1.094 for seed width (SWI) to 248.969 % for leaf peduncle length (LPL) considering quantitative traits. For qualitative

characters, GAM ranged from 55.820 in leaf size (LSi) to 331.063 % in fruit design produced by secondary skin color (DPSC). High heritability estimates coupled with high estimates of genetic advance expected in the next generation in several traits indicate the preponderance of additive gene action for the expression of these traits which is fixable in subsequent generations. This also provides the evidence that larger proportion of phenotypic variance has been attributed to genotypic variance, and reliable selection could be made for these traits on the basis of phenotypic expression (Bello et al. 2012). These results find support from the earlier studies by Akbar et al. (2008) and Kashiani et al. (2008) even though it was on another crop. Those authors suggested that concerned parameters were under the control of additive genetic effects. Sumathi et al. (2005) suggested also that these parameters could be manipulated according to requirements, and worthwhile improvement could be achieved through selection. Those authors concluded that the selection at an early segregating generation will prove beneficial for selecting superior varieties of maize. Such conclusions should also be made in the study presented in the current research. To sum up, it can be concluded that heritability estimates along with genetic advance is more helpful in predicting the genetic gain under selection than heritability estimates alone as mentioned in Akbar et al. (2008) in maize.

Correlations among traits

In plant breeding, correlation coefficient analysis measures the mutual relationships between various plant characters and determines the component characters on which selection can be based for genetic improvement in yield (Cousin et al. 1985). Genotypic correlation coefficient is the heritable association between two variables Hussain et al. (2011). However, phenotypic correlation includes both phenotypic and environmental effect. Hence, significant phenotypic correlation without significant genotypic correlation has no value (Bello et al. 2012, Falconer 1989). In the present study correlation showed positive and significant association between traits contributing to yield (TSW and SNP) and most of the other characters at genotypic level (Table 7).

Phenotypic correlation is a composite of genotypic and environmental correlations. In this study, it was observed that the genotypic correlation coefficients were, in most cases, lower than their corresponding phenotypic correlation coefficients indicating that the inherent association between the characters is governed largely by environmental causes. Non similar finding were earlier observed by Adekoya et al. (2014) and Scod et al. (1995) in okra. Characters which are phenotypically correlated but not genotypically correlated will not produce repeatable estimates of intercharacter associations and any selection based on the relationships is likely to be unreliable.

Path coefficient analysis

Correlation measures mutual association with no regard to causation, whereas path analysis specifies causes and measures their relative importance (Dewey and Lu, 1959). Time to emergence (TE), time to male flowering (MF), time to female flowering (FF), leaf limb width (LLW), fruit number per plant (FN), plant height (PH), fruit width (FWI), seed length (SL) and thousand-seeds weight (TSW) had negative direct effect on seed number per plant (SNP). However, apart from leaf pubescence texture (LPT), stem pubescence texture (SPT), design produced by secondary skin color (DPSC) and male flower size (MFS), all other traits showed positive direct effect on TSW. These observations suggested the inefficiency of selection based on correlation alone. In conclusion, the above findings illustrated that the performance of the genotypes varied from location and year to another (Ogunniyan et al. 2015, Adekoya et al. 2014).

Understanding of the relationships between the traits, for the selection of the important ones, is of crucial importance. The goal of the path analysis, is that, the acceptable descriptions of the correlation between the traits, based on a model of cause and effect, is presented, and the importance of the affecting traits on a specific trait is estimated. In fact, the basic relationships between the traits are expressed by this analysis (Becker 2011, Falconer 1989). So that, the correlation coefficients dividing to the direct and the indirect effects of the set of the independent variables on a dependent variable, and their relative importance are calculated. Several researches of the relationships and trait effect of plants have been reported. In the path analysis of some quantitative traits of maize (Ogunniyan et al. 2015, Bello et al. 2012) and soybean were identified. In soybean, the number of pods per plant, the weight of 100-grains and the number of seeds per plant compared to other traits, had the most direct effect on the yield as reported in Edache (1996). In an experiment that performed on the peas plant, it was cleared that, there was positive and significant correlation between the number of pods per plant, the weight of 100-grains and the number of lateral branches with grain yield. Also, indicated that, the number of pods per plant, the pod length and vine length had most direct effect on the grain yield of the peas (Rasaei et al. 2011, Ghobary 2010, Amaranthath and Viswaantaha 1990). From these reports, pea grain yield had a positive correlation with the plant height, the number of pod, the number of grains per pod and the yield of the pod, and pod yield showed the high direct effect in this plant (Rasaei et al. 2011).

Moreover, the use of simple correlation analysis cannot fully explain the relationships among characters. Therefore, path coefficient analysis has been used by many researchers for a more complete determination of the impact of an independent variable on a dependent one (Bello et al. 2012, Udensi and Ikpeme 2012, Majid et al. 2011, Karademir et al. 2009, Zhao et al. 2008, Lal 2007, Akinyele and Osekita, 2006) in okra, fennel, wheat, cotton, potato and *Cajanus cajan*, respectively). Path coefficient analysis helps the breeder to explain direct and indirect effects which have been extensively used in breeding works of different crop species by various researchers (Adekoya et al. 2014). The knowledge of inter-character relationships is very important in plant breeding for indirect selection for characters that are not easily measured and for those that exhibit low heritability. A path coefficient analysis simultaneously captures the effects of intricate relationships among various traits under investigation (Falconer 1989). Information obtained from correlation coefficients can be enhanced by partitioning them into direct and indirect effects for a set of a priori cause–effects interrelationships, as has been demonstrated in various crops (Adekoya et al. 2014). However, there has been relatively little information on some of the newly collected genotypes of colocynth in Africa. A careful study of relationships among quantitative characters is necessary in order to ascertain the magnitude and direction of changes to be expected during selection in *Citrullus colocynthis*. Knowledge of relationships between yield and its components is essential as this may help in constructing suitable selection criteria for yield. In order to determine the relationships between yield and the other examined traits, correlation coefficients were calculated. Simple correlation coefficients calculated among the examined traits are shown in Table 7. Results suggested that any positive increase in these traits will accelerate the yield potential of colocynth in Africa. So, these traits should be paid attention in breeding programs. Positive and significant correlations between yield and yield components in colocynth were also reported.

Conclusions

The study presented here revealed that information about the extent of variation, estimates of heritability and expected genetic advance in respect of the performance of colocynth and particularly its yield contributing characters represents the basic requirement for a crop improvement program. Broad sense heritability is useful for measuring the relative importance of additive portion of genetic variance that can be transmitted to the offspring. The preponderance of additive gene effects controlling a trait usually resulted to both high heritability and genetic advance, while those governed by non-additive gene actions could give high heritability with low genetic

advance. However, in the present research, expected genetic advance values were based on broad sense heritability, which integrates additive portion of the total phenotypic variance. Effective selection for superior genotypes is possible considering some of the quantitative and especially the qualitative characters investigated. Those traits could be sources of alleles that can be manipulated with other promising cultivars from farmers' origin in Benin and West Africa.

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