Variability and Heritability of Morphological Traits in Collection of Cotton Genotypes (*Gossypium hirsutum* L.) and Their Potential Use for the Selection

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Abstract

This study was conducted to estimate variability and heritability of some morphological traits available in 14 introduced cotton genotypes. The experiment was carried out in randomized complete block design with three replications. The experimental site is based in Cotton and Fibres Center for Agricultural Research at Cana located in the south of Republic of Benin.Analysis of variance showed highly significant differences among the 14 cotton genotypes (p<0.001). The results showed high heritability for plant height, plant height to first fruiting branch, number of vegetative branches, length of vegetative branch, length of fruiting branch, number of fruiting branches and number of nodes among 14 varieties. Heritability values vary from 0.720 (NFB) to 0.999 (PH and LVB).Positive correlations were found between plant height and length of fruiting branch and number of fruiting branches and other morphological characters. A highly significant correlation (r = 0.80) was observed between plant height to first fruiting branch and plant height. The relatively high variability and heritability estimates for morphological traits suggested that they could be used in breeding programs.

Keywords: Gossypium hirsutum, fruiting branch, vegetative branch, heritability, variability

Introduction

Cotton (Gossypiumhirsutum L.) is very imperative non-food economy oriented fiber and cash crop of Benin.Cotton network constitutes the main source of national economic increase and constitutes a strong the main source of national economic increase and constitutes a strong strategic tool to fight against the poverty (INRAB, 2013). Benin, through Cotton and Fiber Center for Agricultural Research, has more than 100 genotypes cotton currently in its Genebank. Although there is a big source of variability for breeding and good collections of *G.hirsutum* in Benin cotton Genebank, no information is available on the variability and heritability of their morphological characters. However the selection should be more efficient if morphological as well as physiological characteristics are identified and used as criteria of sifting in the classic process of plants improvement (Hamliat al. 2015). Several characters are reported in the identified and used as criteria of sifting in the classic process of plants improvement (Hamli*et al.*, 2015). Several characters are reported in the literature like having a more or less close connection with the tolerance or the performance of the plants under conditions of stress. Among these characters appear fruiting branches, vegetative branches, plant height to first fruiting branch, main stem of plant and nodes. Thus, according to Khalil *et al.* (2015), plant traits such as plant height play an important role in the sustainable pest management of cotton crop. It is therefore imperious to evaluate the variability and the heritability of these characters within Benin cotton. Genbank in order to assure their stability in the descent cotton Genbank in order to assure their stability in the descent. Cotton Genbank in order to assure their stability in the descent. Morphological characteristics are used by breeders in the development of improved cultivars. The evaluation of genetic variability available is a preparatory to start a program of selection and particularly to choose the parents to be crossed (Hamli *et al.*, 2015). In the same way, the heritability of the characters determines the response to the selection and depends on the genetic material studied as well as experimental device (Sekloka, 2004; Shukla et al., 2006; Atta et al., 2008).

Keeping in view the work of above scientists, the present study was conducted to investigate genetic diversity and heritability of fourteen (14) genotypes of cotton using morphological marker.

Materials and methods Genetic materials

Fourteen genotypes of cotton (Gossypium hirsutum L.) were chosen from the collection in the field for analysis in this experiment (table1) as they presented interesting divers forms and maturenesses. The collection in the field of cotton plants laid within the experiment site of Cotton and Fibres Center for Agricultural Research at Cana (2°5'E, 7°6'N) located in south of Republic of Benin at an altitude of about 89 meters above sea level.

Genotypes	Origins	Characteristics
A24	Tchad (CIRAD/ITRA)	Slender shrub, leaves of lobed form, small boll
CD14	Zambia(CIRAD/MNRWD)	Slender shrub, broad to lobed leaves, medium boll
Chaco 520	Argentina	Compact shrub, leaves of lobed form, big boll
CR 92-498	Costa-Rica (CIRAD)	Slender and bush shrub, broad to lobed form, big boll
CR 92-534	Costa-Rica (CIRAD)	Slender shrub, trend to bend, lobed leave, medium boll
CS 189	Australia (CSIRO)	Bush to compact shrub, broad leave, medium boll
Deltapine 90	USA (Deltapine)	Bush shrub, broad to lobed leave, medium boll
Guazuncho II	Argentina	Bush to compact shrub, broad to lobed leave, medium to
		big boll
H 279-1	Togo (CIRAD/ITRA)	Slender shrub, broad to lobed leaves, medium boll
Irma 772	Cameroun (CIRAD/IRAD)	Slender shrub, broad to lobed leaves, medium boll
Irma Blt-pf	Cameroun (CIRAD/IRAD)	Slender shrub, trend to bend, broad leave, medium to big
		boll
Irma Z 856	Cameroun (CIRAD/IRAD)	Bush to slender shrub, broad leave, medium boll.
Nta 88-6	Mali (CIRAD/IER)	Slender shrub, broad to lobed leaves, medium to big boll
Sicala 34	Australia (CSIRO)	Compact shrub, broad leave, small boll

Table1: Qualitative description of the 14 cotton genotypes studied

Morphological characteristics

Seven morphological traits were measured in the field experiments on the all 14 genotypes. Eight plants were selected randomly from each genotype and morphological characters were described during three years 2009, 2010 and 2011. Measurements were made on plant height to first fruiting branch (PHFFB); number of vegetative branches (NVB); plant height (PH); length of vegetative branch (LVB); length of fruiting branch (LFB); number of fruiting branches (NFB) and number of nodes (NN).

Field evaluation

Plots were single rows, 10 m in length and 1 m apart with 0.50 m plant spacing. The seeds were sown at the end of June with one genotype per row. The seedlings were thinned to 1 plant per hill 3 weeks after sowing. The N.P.K. fertilizer was applied at thinning with the rate of 200kg/ha 21 days after emergence and N fertilizer was applied with the rate of 50 kg/ha 40 days after sowing. Insect pest was controlled using ten fortnightly sprays of binaries accaricid and binaries aphicid pesticides.

Data analysis

Cotton genotypes were planted in a randomized complete block design (RCBD) or bloc of Fisher with three replications.. Statistical analyzes of the results were performed with the STATISTICA software, version 6 (www.statsoft.com) and R (Version 3.1). Canonical discriminant analysis step (stepwise canonic analysis) was performed on the morphological variables prior to selection of the most discriminating variables. It was

followed by a second canonical discriminant analysis to describe, through a system of axes, genotypes based on variables selected by the canonical discriminant analysis step. Then, analysis of variance followed by tests comparing averages by the test and Newman Keuls were used to characterize the genotypes from the average of their different morphological characters and compare the observed variability of a genotype to another. Heritability genotypic (h^2_{sb}) and genetic advance (GA) of genotypes morphological characters were also estimated in order to assess the proportion of the phenotypic variability of genetic origin which is heritable and fixable in whole or in part.

Cluster analysis using Unweighted Paired Group Mean Average (UPGMA) was also performed to create a dendrogram and group the morphologically similar genotypes. Principal component analysis (PCA) based on morphological characters was also conducted to investigate the relationship between the different genotypes of cotton.

Results

Discrimination of the cotton genotypes based on morphological characteristics

characteristics Univariate analysis showed a significant variability (p>0, 05) among the genotypes for all morphological traits (Table 2).The canonical analysis revealed that all seven morphological variables best discriminated the different genotypes. The results of inferential tests (table 3) confirmed the discriminating power of these variables because they showed that, for all of them, there are very highly significant differences (p / Wilk's Lambda <0.0001) between genotypes.The canonical discriminant analysis showed that the axis 1 (or first canonical discriminant axis) was strongly correlated with variables PHFFB, NVB, LVB, LFB, NFB and NN, while axis 2 best discriminated genotypes on the basis of the variable PH (table 4). It would therefore be possible to make a fairly accurate description of 14 genotypes from these morphological variables.

Variables	SC	MC	\mathbf{F}	Р
PHFFB	449,19	34,55	28,97	0,000***
NVB	10,2279	0,7868	5,453	0,000***
PH	13585,1	1045,0	443,6	0,000***
LVB	4994,5	384,2	251,9	0,000***
LFB	2918,97	224,54	206,13	0,000***
NFB	47,83	3,68	3,42	0,003**
NN	75,27	5,79	2,77	0,011*

Table 2: Univariate tests performed on morphological variables according to different genotypes of cotton

PHFFB=plant height to first fruiting branch; NVB= number of vegetative branches; PH=plant height; LVB=length of vegetative branch; LFB= length of fruiting branch; NFB= number of fruiting branches; NN= number of nodes

Variables	F value	Partial (Lambda)	Wilk's Lambda	p< Lambda
PHFFB	10,54459	0,138295	0,000000	0,000***
NVB	4,55407	0,270926	0,000003	0,000***
PH	63,89030	0,025804	0,000605	0,000***
LVB	54,25153	0,030250	0,017038	0,000***
LFB	37,31082	0,043389	0,124563	0,000***
NFB	3,31340	0,338075	0,477968	0,006**
NN	4,74513	0,262885	0,822341	0,000***

Table 3: Canonical analysis on morphological variables to determine the most discriminating variables to compare the different genotypes.

PHFFB=plant height to first fruiting branch; NVB= number of vegetative branches; PH=plant height; LVB=length of vegetativebranch; LFB= length of fruiting branch; NFB= number of fruiting branches; NN= number of nodes

Table 4: Canonical discriminant analysis on morphological variables according to different

genotypes						
	Racine1 (68,42%)	Racine2 (27,26%)				
PHFFB	0,8264	-0,7830				
NVB	0,6126	0,4463				
PH	-0,4728	-1,2071				
LVB	-1,0997	0,6744				
LFB	-1,0620	0,6687				
NFB	-0,5706	0,3070				
NN	1,1051	-0,3023				

PHFFB=plant height to first fruiting branch; NVB= number of vegetative branches; PH=plant height; LVB=length of vegetative branch; LFB= length of fruiting branch; NFB= number of fruiting branches; NN= number of nodes

Morphological descriptors

Statistical analysis of variables of table 5 revealed highly significant variability among the 14 genotypes. The plant height varied from 92,1±0,06 cm (Chaco 520) to 147,0±1,15 cm (Irma Blt-pf), plant height to first fruiting branch varied from19,0±0,28cm (Sicala 34) to29,5±0,28 (Nta 88-6), the longest of vegetative branches from 50 cm (Chaco 520) to 95 cm (A 24), the longest fruiting branches from 33,5±0,87cm (Chaco 520) to 70,3±0,68 cm number of fruiting branches from 14,8±0,86 (Chaco 520) to (A 24), vegetative branches from 1,4±0,12 18,6±0,34 (H279-1), number of (Guazuncho II) to 3,4±0,28 (A 24), number of node on main-stem from 20,1±0,63 (Chaco 520) to 25,3±0,17 (Irma Z 856). The all variables are highly significantly heritable and vary from 0.720 (NFB) to 0.999 (PH and LVB).

variables studied according to different genotypes of cotton								
Génotypes	PHFFB	NVB	PH	LVB	LFB	NFB	NN	
A 24	21,2±1,04	3,4±0,28	129,7±0,33	96,9±0,38	70,3±0,68	16,3±0,28	23,4±0,90	
CD 14	23,7±0,40	$2,2\pm0,17$	124,4±0,75	68,0±1,15	40,5±0,58	16,3±0,34	$22,8\pm0,80$	
Chaco 520	19,5±1,15	$1,8\pm0,29$	92,1±0,06	50,9±0,81	33,5±0,87	14,8±0,86	20,1±0,63	
CR 92-498	21,3±0,75	1,9±0,23	$106,6\pm0,81$	62,7±0,63	40,3±0,17	16,2±0,69	22,3±1,15	
CR 92-534	24,9±1,10	$2,2\pm0,17$	123,9±1,27	67,2±0,69	43,1±0,57	15,9±0,05	21,9±0,11	
CS 189	19,3±0,17	1,9±0,23	94,4±0,23	65,0±0,86	41,0±0,57	15,9±0,17	21,9±1,04	
Dp 90	20,3±0,17	$2,1\pm0,06$	102,6±1,15	66,7±0,75	39,4±0,34	16,3±0,17	22,0±0,57	
Guazuncho II	17,2±0,69	$1,4\pm0,12$	96,9±0,98	55,3±0,17	40,1±0,57	16,8±1,39	22,1±1,79	
H 279-1	21,8±0,46	1,9±0,23	124,2±1,27	73,5±0,75	51,9±1,10	18,6±0,34	21,9±0,92	
Irma 772	22,2±0,05	$1,6\pm0,12$	140,1±0,64	77,0±1,15	46,8±0,11	18,3±0,98	24,5±0,28	
Irma Blt-pf	25,9±0,52	$2,1\pm0,1$	147,0±1,15	76,1±0,05	50,0±0,57	18,5±0,28	24,4±0,23	
Irma Z 856	26,3±0,17	2,6±0,2	123,1±1,21	72,8±0,46	42,7±0,17	17,6±0,23	25,3±0,17	
Nta 88-6	29,5±0,28	$2,6\pm0,29$	$143,4\pm0,81$	81,0±0,28	46,1±0,63	17,0±0,57	$23,8\pm0,75$	
Sicala 34	19,0±0,28	$1,6\pm0,17$	102,1±0,06	64,3±0,75	46,1±0,63	16,4±0,23	21,8±0,57	
Moyenne	22,3±0,53	2,09±01	117,9±2,8	69,8±1,71	45,1±1,31	16,7±0,21	22,7±0,27	
EcTyp.	3,431	0,589	18,28	11,08	8,48	1,379	1,807	
h_{bs}^2	0,984***	0.963***	0.999***	0.999***	0.997***	0.720***	0.880***	
GG%	6,95	1,168	37,619	22,80	17,416	2,045	3,275	
P>F	0,000***	0,000***	0,000***	0,000***	0,000***	0,000***	0,000***	

Table 5: Analysis of variance and genotypic heritability performed on morphological variables studied according to different genotypes of cotton

PHFFB=plant height to first fruiting branch; NVB= number of vegetative branches; PH=plant height; LVB=length of vegetative branch; LFB= length of fruiting branch; NFB= number of fruiting branches; NN= number of nodes;

Classification of cotton genotypes based on morphological characters

UPGMA cluster of the 14 genotypes revealed interesting associations based on morphological characters. Cotton genotypes are grouped into two main clusters (figure 1). The first cluster contained 9 genotypes and the second 5. Similarity index of the cluster was <5%, indicating the reliability of the clusters to the original data. Principal component analysis (PCA) based on morphological characters showed that the first two axes make up 80.88% of the total variance (figure 2). On the first axis, all the morphological characters had the highest variance. Moreover, it is interesting to note that principal component analysis classified alsocotton genotypes into two groups. However, PCA had not shown significant difference among genotypes. The first group contained 13 genotypes and the second 1 (A 24). Genotype A 24 was placed in more distant and so can be used in selection or in hybridization programs to produce heterosis with interesting characters.

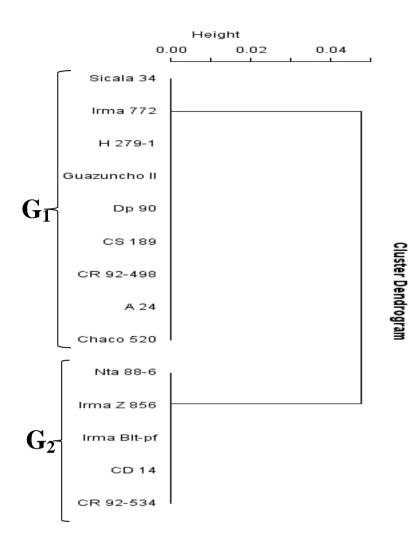


Figure 1: UPGMA cluster of genotypes based on morphological characters

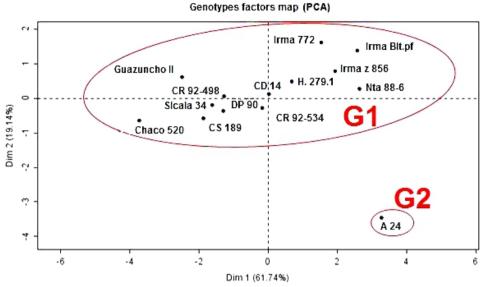


Figure 2: PCA based on morphological characters

Correlation determined for morphological characters

coefficient of correlation determined for morphological The characters showed a high significant positive correlation between plant height to first fruiting branch and plant height with number of nodes, number of vegetative branches with length of vegetative branch and length of fruiting branch, plant height and length of vegetative branch and number of fruiting branches with number of nodes, number of fruiting branches and number of nodes. However, the correlations were no significant between the plant height and number of vegetative branches, plant height to first fruiting branch and number of vegetative branches (table 6).

Table 6: Correlation of Pearson between morphological characters								
	PHFFB	NVB	PH	LVB	LFB	NFB	NN	
PHFFB	1,00							
NVB	0,51ns	1,00						
PH	0,80**	0,45ns	1,00					
LVB	0,49ns	0,78**	0,75**	1,00				
LFB	0,15ns	0,65*	0,53ns	0,88***	1,00			
NFB	0,38ns	-0,04ns	0,68**	0,45ns	0,36ns	1,00		
NN	0,64*	0,42ns	0,76**	0,65*	0,39ns	0,68**	1,00	

PHFFB=plant height to first fruiting branch; NVB= number of vegetative branches; PH=plant height; LVB=length of vegetativebranch; LFB= length of fruiting branch; NFB= number of fruiting branches; NN= number of nodes; ***,**, * = Significant at 0.1; 1 and 5% probability levels, respectivelyns = Non-significant

Discussion

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efficiency (Ribot *et al.*, 2012).This conventional method of improvement assisted by morphological characters may also be complemented by molecular markers at different stages, to analyze the genetic diversity, select progenitors and identify varieties. Since the characters recommended in this study are easy to measure and have high heritability, the use of molecular markers would not be necessary as the efficient use of molecular markers is obtained when selected characters are highly influenced by the environment. However, molecular markers could be useful if one wishes to accumulate various characters in the same genotype.Since there are molecular techniques which allow selection to be made in early generations, an improvement program could be designed to combine the three selection methods (selection assisted by morphological characters, selection assisted by molecular markers and conventional selection).

Conclusion

The study revealed high significant variability and heritability in morphological traits among the 14 cotton genotypes. These results are suggesting that the morphological variability observed is heritable and fixable. This conventional method of selection assisted by morphological characters may be complemented and confirmed by molecular markers to analyze the genetic diversity in collection.

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