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Estimates of phenotypic diversity and genetic parameters of *Hibiscus* cannabinus L. grown in Burkina Faso

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ABSTRACT

Hibiscus cannabinus is a traditional leafy-vegetable consumed more and more in Burkina Faso. Its leaves and fruits are some good food supplement against nutritional deficiencies. Its fibers are also a source of income for households. Despite of its nutritional an economic potential, this species is underdeveloped in Burkina Faso. The study aims to determine genetic variability and level of heritability among agronomic interest traits of *H. cannabinus* using agro-morphological descriptors. This enable to identify outstanding accessions which could be involved in further breeding programs. A total of 70 accessions of *H. cannabinus* were characterized according randomized incomplete block, using 5 qualitative characters and 15 quantitative characters. The study showed great variability for most of the qualitative and quantitative characteristics studied. Several correlations were observed between characters related to the biomass, the fruits and the fibers production. These correlations are particularly interesting because the improvement of only one character induces improvement of the others. The study, moreover, shows high broad sense heritability (H^2) and weak differences between the phenotypic (PCV) and genotypic (GCV) coefficients of variation. These results indicate that the characters expression is under genetic control. So a direct selection method of *H. cannabinus* improvement is possible for these characters.

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Keywords: Agromorphological, characterization, Hibiscus, selection, Burkina Faso.

INTRODUCTION

Hibiscus cannabinus is a species of multiple uses in Burkina Faso through its organs of interest such as the leaves, the fibers and the seeds. The leaves are cooked as sauce, because they have properties of digestive ballast and intestinal purifier (Millogo-Rasolodimby, 2001). They are used as a medicine to purge, to treat diarrhea, hypertension, dysentery and bilious unrest (Bermudez-Soto et al., 2007). The seeds are

used for the manufacturing of a local condiment, named "*soumbala*" in dioula, "*Bicalga*" in mooré and "*gio-sola*" in Gurunsi-Lélé. This condiment is even considered tastier than the *soumbala* (another traditional condiment) made on the basis of *Vitelaria paradoxa* seeds. The fibers are used in ropes manufacturing, masks manufacturing called "*biiddu*" and in funeral rituals to make the cadavers ready for burial. The aqueous excerpts and methanolic of the seeds are used

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as antioxidants. Regarding its socio-economic importance, H. cannabinus is an indispensable resource for the local populations. However, it neglected by research because few is investigations have been dedicated to it. So, its potential is under-exploited by the population and its genetic varieties are still less known. Some studies of the agromorphological diversity were carried out in East Africa, notably in Kenya, Tanzania and Malawi. Its diversity was studied in West Africa especially in Nigeria, where the species is well-known and cultivated as a plant having commercial fibers (Lakshmana et al., 2009). In Burkina Faso, the studies that have already been done are only focused on the tenacity of its fiber and on the traditional knowledge and expertise about the management of the species (Kabré et al., 2017). For the need of documentation of this species in Burkina Faso, the current study was initiated and aimed at characterizing the genetic variability of H. cannabinus. Its basic objectives are to (i) identify the agro-morphological characters that discriminate H. cannabinus genotypes, (ii) establish the relations which exist between them, and (iii) estimate genetic parameters in a collection of H. cannabinus accessions of Burkina Faso, such as broad sense heritability (H^2) , genotypic (GCV) and phenotypic (PCV) coefficient of variation and genetic advance (GA) expected from selection.

MATERIALS AND METHODS Materials

Genetic material

Seventy (70) accessions of H. cannabinus were randomly collected in fiftysix (56) villages distributed into seven (7) provinces (Table 1) of the three climatic zones of Burkina Faso (Figure 1).

Experimental site

A trial was conducted at "Institut du Développement Rural" (an agronomic research center) located at Gampela (1°21'9, 6''W; 12° 24' 29''N) in Ouagadougou. Gampela is located precisely in the North-Sudanian zone with an annual rainfall ranging from 600 to 900 mm (Thiombiano & Kampmann, 2010). Rainfall patterns recorded during 2017 rainy season was 677.1 mm of water between May and October. The test was implanted on a sand-clay soil with a water pH of 5.20. The analysis of this soil shows that it is composed of total nitrogen (0.046 - 0.071%), phosphate (9.71 - 10.91 ppm) and potassium (84.18 - 101.23 ppm).

Methods

Experimental design

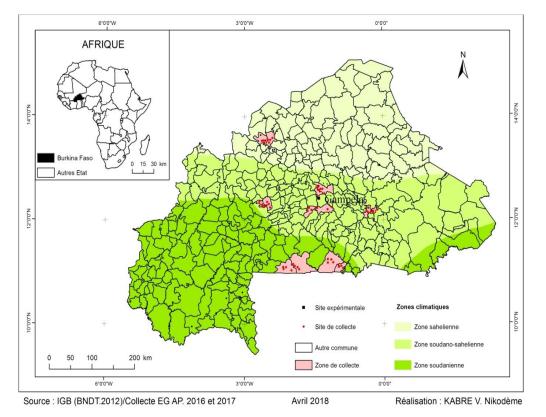
The experimental design consists of a randomized incomplete block with three-repeat. One meter distance separated the blocks. Each block then was subdivided into two (02) elementary blocks that are spaced also by one meter (1 m) distance. As far as sowing is concerned, each seed hole was spaced by a 0.6 m distance on a 4.2 m long line and lines were separated by 0.8 m space. *Observed variables*

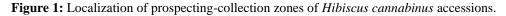
Twenty (20) variables including fifteen (15) quantitative and five (05) qualitative derived from H. cannabinus descriptor of the National Office of Genetic Resources Plantation have been valued. A11 the observations and measurements were made 80 days after sowing date, corresponding to the 50% flowering stage, at four plants per line and per accession. The number of days 50% flowering (DFL) has been valued on the whole line. Qualitative traits include stem color (COT), ribs color (CNE), petal color (CPE), shape of the leaves (FFE) and the pubescence of the stems (PTI). Quantitative traits measured are plant height (HPL), stem diameter (DIT), dry fibers weight (PFI), petiole length (LPE), limb length (LOL), limb width (LAL), leaves biomass (PFF), petal length (LOP), petal width (LAP), Number of sepal (NSE), length and diameter of the fruit respectively (LFR) and (DFR), number of fruits (NFR) and 100 seeds weight (PCG). Data analysis

The descriptive analysis has been made using qualitative data. Thus, some frequencies have been calculated and the diagrams constructed from the frequencies we had. The quantitative data were used for variance (ANOVA) using the GenStat v4.10.3 software to determine those discriminate accessions. Correlations between these traits were studied using Pearson correlation tests (1% level). Principal component analysis (PCA) was performed to know the associations between the studied characters. All the accessions were consecutively grouped from the hierarchical ascending classification (CAH) based on the segregation of Ward method based on using Euclidean distance between individuals. The genotypic and phenotypic variances (VG&VP), the phenotypic and genotypic coefficients of variations (PCV&GCV), the heritability (H^2), the genetic advance (GAx), formulas proposed by Johnson et al. (1955); Assefa et al. (2001) and Hosseini et al. (2012) have been used (Table 2). Excel, XLstat 7.5.2 and Genstat v4.10.3 software, have been used for these analyses.

Table 1: Distribution of	f the producers	according to	province and sex.
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Provinces of collection	Number of producers	Proportions of men (%)	Proportions of women (%)
Kadiogo	9	75	25
Kouritenga	25	78	22
Nahouri	29	60	40
Oubritenga	11	82	18
Sanguié	19	75	25
Sissili	18	50	50
Yatenga	4	68	32





N°	Parameters	Formula used
1	Genotypic Variance (VG)	VG = (MSG - MSE)/r
2	Phenotypic Variance (VP)	VP = VG + (MSE/r) = MSG/r
3	Broad sense heritability (H ²)	H^{2} (%) = (VG/VP)*100
4	Genotypic coefficient of variation (GCV)	GCV (%) = $(\sigma_G/X)*100$
5	Phenotypic coefficient of variation (PCV)	PCV (%) = $(\sigma_P/X)*100$
6	Genetic advance (GA)	$GA = H^2 \sigma_P K$
7	GA (% of trait mean) [GAx (%)]	GAx (%) = (GA/trait mean)*100

Table 2: Formula used for the estimates of the genetic parameters

 \overline{MS}_{G} is the means squared of genotype, \overline{MS}_{E} is the means squared of the residual (error), and r is the number of replication, σ_{G} is the genotypic standard deviation, σ_{P} is the phonotypic standard deviation, X is the trait mean; K is a constant, which at the selection intensity of 5% is 2.06.

RESULTS

Agro-morphological variation related to the qualitative traits of *Hibiscus cannabinus*

All accessions of *H. cannabinus* collection have pubescent leaves and fruits. The stems are more pubescent than the leaves. Indeed 27.14% of accessions collected have stems deprived of pubescence compared to 72.86% of stems having epidermal hairs. Meanwhile a high variability has been observed for the other characters (Table 3). This variability is mainly perceptible on the color of the different organs. Thus the stems present 6 colors (Figure 2) such as green (57.14%), purple (22.86%), violet (7.14%), black (5.71%), red (4.29%) and green speckled (2.86%). The leaves are alternate, with a simple serrated or serrated-lanceolate or finger-lobed limb with veins and petioles generally of the same color as the stem.

Agro-morphological variation related to the quantitative traits of *Hibiscus cannabinus*

Analysis of variance reveals a highly significant difference between accessions (p <0.0001) at 1% levels (Table 4). The coefficients of variation of the majority of the studied characters are low (CV<30%). That shows a low heterogeneity of plant material for these characters. However, they are high (CV >30%) for fruits number (CV=31.52%) and leaves biomass (CV=71.25%). That indicates a high heterogeneity of studied accessions for these characters. The accessions of *H. cannabinus* flower between

59 and 96 days after having 2.7m height. Full bloom flowers (5 petals) occur between 7 and 10 days after button floral shaping. Fruits occur after petals fall that is 2 and 3 days after their emergence. The majority of accessions had leaves having long petiole (12.5 to 28.12 cm) and a limb having a less varied length and width (1.85 <ET<1.91cm). A high variability between accessions has been observed regarding the number of fruit per plant (ET=13), the weight of dry fibers and (ET=14.05g) the leaves biomass (ET=397.5g).

Correlation between the studied traits

Table 5 displays many positive and highly significant correlations obtained at $\alpha =$ 1%. The most important ones are those observed between plant height and stem diameter (r = 0.471), dry fibers weight (r =0.310); number of fruits and stem diameter (r = 0.252), dry fibers weight (r = 0.328) and leaves biomass (r = 0.604) as well as between number of days 50% flowering and fruit diameter (r = 0.421); between length of fruit and weight of 100 seeds (r = 0.322). The most significant negative correlations are observed between days 50% flowering and number of fruits per plant (r = -0.235), length of fruit (r =-0.470), 100 seeds weight (r = -0.482) and leaves parameters. There are also negative correlations between leaves biomass and fruits diameter (r = -0.277) and finally between fruit diameter and number of fruits (r = -0.361), length of fruit (r = -0.248).

Associations of morphological traits

The total variability expressed by the first three axes is 55.413% (Table 6). Axis 1 expresses 26.890% of this total variability. Seven characters contribute the most to achievement of axis 1. These are variables number of days 50% flowering (r = 0.667), number of fruits (r = 0.250), petal length (r =0.670), number of sepals (r = 0.194), length of fruit (r = 0.403), fruit diameter (r = 0.386) and 100 seeds weight (r = 0.374). This axis characterises the accessions of H. cannabinus having a long flowering cycle; producing long and heavy fruits and containing numerous and heavy seeds. Axis 2 which expresses 15.749% of this total variability is positively correlated to petiole length (r = 0.315), plant height (r =(0.327) and stem diameter (r = (0.641)). This axis gathers accessions which have an important vegetative development. At last, axis 3 expresses 12.773% of this total variability and is correlated to limb length (r =(0.295) and leaves biomass (r = (0.314)).

Structuring the diversity of studied accessions

The dendrogram resulting from the hierarchical ascending classification (CAH) of Figure 3 reveals a distribution of the 70 accessions in 3 groups. This variability depends more on genotypes than the origin of accessions. An examination of the coefficient of determination and Fisher's F statistic values indicates that five characters such as cycle of flowering ($R^2 = 3.5\%$), number of fruits ($R^2 =$ 54.1%), fruit diameter ($R^2 = 4.4\%$), 100 seeds weight ($R^2 = 4.7\%$) and leaves biomass ($R^2 =$ 97.5%) with a p-value <0.0001 are more discriminating of the 3 agromorphological groups obtained. That shows that they are very distinct. These results are confirmed by the Newmann Keuls test (Table 7). So, group 1 consists of 30 accessions (43% of all collected accessions). These accessions come from 7 provinces. This group is characterized by accessions having long flowering cycle (80.52 days). They produce high diameter fruit (1.59cm) and low biomass leave (577.02g). Group 2 includes 26 accessions (37% of accessions collected) come from 5 provinces belonging to the Sudanian and Sudano-Sahelian zone. This group is

characterized by accessions having precocious cycle (76.36 days) and producing many fruits (83.21) having mean diameter (1.50 cm) but 100 seeds weight most elevated. At last, group 3 includes 14 accessions (20% of collected coming from 4 provinces accessions) distributed in three climatic zones of the country as group 1. This group is characterized accessions by having intermediate cycle (78.85 days) between group 1 and group 2 producing as group 2, a lot of fruits (85.33), heavy seeds (2.97g). It produces the most high leaves biomass (3946.34 g).

Estimates of Genetic the Parameters of *Hibiscus cannabinus*

The heritability is high $(H^2 > 80\%)$ for the number of days 50% flowering, number of fruits, 100 seeds weight, dry fibers weight and leaves biomass. It is moderate for the other characters except petal width where it is low. Phenotypic variances are higher than genotypic variances for all studied characters (Table 8). Phenotypic variances varied from 0.01 to 1277949.33 and genotypic variances from 0 to 1275258. The leaves biomass, the number of fruit, the dry fibers weight and the number of days 50% flowering have very high genotypic and phenotypic variance values. But it is very low (<01) for other studied characters. The phenotypic coefficients of variation are higher than the genotypic coefficients of variation for all studied characters. The maximal gap (11.03%) is obtained for the character "fruit length" (PCV = 14.71% and GCV=3.68%). The characters, leaves biomass, number of fruits and dry fiber weights had at a time elevated PCV and GCV (>20%). Characters such as DFL, LPE, NSE, DFR, had low (<11%). All the other characters have moderate PCV (>20). The genetic advance (GAx) varied from 0.11% (petal width) to 146.43% (leaves biomass). The characters of interest such as leaves biomass, number of fruits, dry fibers weight and 100 seeds weight record the most elevated genetic advance compared to the other characters. The flowering cycle, the length and the width of the limb had genetic moderate advance (18.34%, 14.67%, and 12.45% respectively).

Characters	Modalities	Frequencies (%)		
	Black	5,71		
	Purple	22,86		
СОТ	Red	4,29		
	Green	57,14		
	Green speckled	2,86		
	Violet	7,14		
	Black	5,71		
~~~	Purple	25,71		
CPE	Red	7,14		
	Green	57,15		
	Violet	4,29		
	Black	5,71		
	Purple	25,71		
CNE	Red	7,14		
CIVE	Green	57,15		
	Violet	4,29		
	Simple jagged	88,57		
FFE	Jagged lanceolate	4,29		
	palmate polylobe	7,14		
	No pubescent	27,14		
PTI	Pubescent	72,86		

Table 3: Variation of the qualitative characters in the collection of *H. cannabinus* 

Stem color (COT), petiole color (CPE), rib color (CNE), leaves shape (FFE), stem pubescence (PTI).









a. green stem; b. purple stem; c. violet stem; d. black stem; e. red stem; f. speckled green stem

Figure 2: H. cannabinus stem colors.

**Table 4:** Performance of the 70 accessions for the different quantitative traits and F Fisher observed as well as their.

Variables	Minimum	Maximum	Mean	ЕТ	CV(%)	R ² (%)	F value	<b>Pr</b> > <b>F</b>
DFL (day)	59	96	78.65	10.03	12.75	37.86	118.64**	< 0.0001
NFR	21	93.5	52.11	12.94	31.52	14.65	33.67**	< 0.0001
LPE (cm)	12.5	28.12	19.75	2.78	14.10	11.68	71.92**	< 0.0001
LOL (cm)	2.87	18.12	14.14	1.85	13.09	9.42	76.84**	< 0.0001
LAL (cm)	2.3	15.62	10.98	1.91	17.43	15.83	59.78**	< 0.0001
HPL (m)	1.37	8.3	2.70	0.57	21.34	20.39	46.55**	< 0.0001
DIT (cm)	1.7	6.12	3.14	0.68	21.64	9.59	38.70**	< 0.0001
LOP (cm)	3.5	9.37	6.98	1.14	16.32	21.70	68.38**	< 0.0001
LAP (cm)	2	8.25	3.81	0.85	22.39	20.39	42.66**	< 0.0001
NSE	3	9.25	7.47	0.70	9.420	10.24	10.13**	< 0.0001
LFR (cm)	1.35	7.07	2.22	0.55	24.99	10.76	30.43**	< 0.0001
DFR (cm)	0.9	2.17	1.54	0.20	12.98	9.70	80.91**	< 0.0001
PCG (g)	1.4	3.80	2.86	0.41	14.62	51.73	118.22**	< 0.0001
PFI (g)	25.6	125.7	61.59	14.05	22.81	60.68	81.67**	< 0.0001
PFF (g)	122.5	9175	1209.49	797.58	71.25	64.05	16.62**	< 0.0001

Number of days 50% flowering (DFL); number of fruits per plant (NFR); Petiole length (LPE); Limb width (LAL), Leaf length (LOL); Plant height (HPL); Stem Diameter (DIT); Petal length (LOP); Petal width (LAP); Number of sepal (NSE); Length of fruit (LFR); Diameter of fruit (DFR); Weight of 100 seeds (PCG); Weight of dry fibers (PFI); Leaves biomass (PFF); Standard deviation (ET); CV: coefficient of variation; F: F of Fisher; R²: coefficient of determination; ** Significant at 1% probability levels.

CHAR.	DFL	NFR	LPE	LOL	LAL	HPL	DIT	LOP	LAP	NSE	LFR	DFR	PCG	PFI	PFF
DFL	1														
NFR	-0,235	1													
LPE	0,060	0,109	1												
LOL	-0,402	0,000	0,136	1											
LAL	-0,253	0,053	0,328	0,812	1										
HPL	-0,015	0,093	0,113	0,231	-0,033	1									
DIT	0,101	0,252	0,373	0,090	0,033	0,471	1								
LOP	-0,673	0,262	-0,063	0,341	0,251	0,028	-0,074	1							
LAP	-0,496	0,236	-0,035	0,248	0,215	0,041	-0,013	0,818	1						
NSE	0,402	-0,136	0,224	-0,230	-0,122	-0,070	0,072	-0,179	-0,183	1					
LFR	-0,470	0,082	0,057	0,338	0,255	0,095	0,061	0,610	0,593	-0,093	1				
DFR	0,421	-0,361	0,024	-0,084	-0,076	-0,039	0,027	-0,344	-0,171	0,345	-0,248	1			
PCG	-0,482	0,146	-0,108	0,214	0,027	0,169	0,112	0,508	0,473	-0,133	0,322	-0,218	1		
PFI	-0,047	0,328	-0,035	0,087	-0,018	0,310	0,259	0,166	0,076	-0,097	0,043	-0,270	0,027	1	
PFF	-0,113	0,604	-0,017	-0,012	-0,015	0,157	0,152	0,191	0,090	-0,092	-0,007	-0,277	0,084	0,824	1

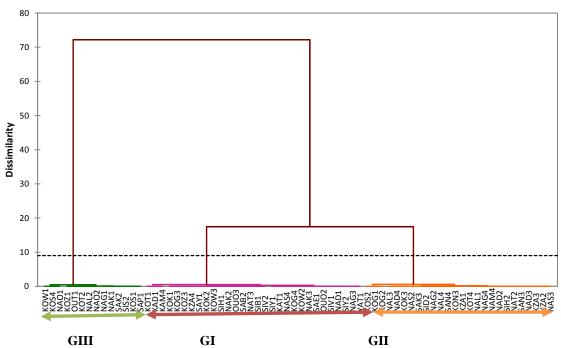
 Table 5: Correlation among some of the agromorphological traits significant at 1%.

Number of days 50% flowering (DFL); number of fruits per plant (NFR); Petiole length (LPE); Limb width (LAL), Leaf length (LOL); Plant height (HPL); Stem Diameter (DIT); Petal length (LOP); Petal width (LAP); Number of sepal (NSE); Length of fruit (LFR); Diameter of fruit (DFR); Weight of 100 seeds (PCG); Weight of dry fibers (PFI); Leaves biomass (PFF).

Characters	<b>F</b> 1	F2	F3
DFL	0.667	0.069	0.008
NFR	0.250	0.233	0.184
LPE	0.000	0.315	0.291
LOL	0.100	0.002	0.295
HPL	0.030	0.327	0.004
DIT	0.007	0.641	0.034
LOP	0.670	0.038	0.030
NSE	0.194	0.046	0.084
LFR	0.403	0.013	0.192
DFR	0.386	0.000	0.087
PCG	0.374	0.004	0.008
PFF	0.144	0.203	0.314
Proper value	3.227	1.890	1.533
Variability (%)	26.890	15.749	12.773
% cumuli	26.890	42.639	55.413

Table 6: Squared Cosines of 12 Characters on three factors and total variability explained of 12.

Number of days 50% flowering (DFL); number of fruits per plant (NFR); Petiole length (LPE); Leaf length (LOL); Plant height (HPL); Stem Diameter (DIT); Petal length (LOP); Number of sepal (NSE); Length of fruit (LFR); Diameter of fruit (DFR); Weight of 100 seeds (PCG); Leaves biomass (PFF).



Dendrogramm

**Figure 3:** Dendrogram of Dissimilarity among 70 accessions of *H. cannabinus* using Ward's Minimum Variance Method of Cluster Analysis.

	Groups					
VAR.	Group 1	Group 2	Group 3	<b>R²%</b>	F.Fisher	Pr>F
DFL	80.52A	76.35B	78.85AB	3.5	3.69	0.027
NFR	51.34B	83.21A	85.33A	54.1	121.40	< 0.0001
LPE	19.65A	19.91A	19.66A	0.2	0.19	0.824
LOL	13.96A	13.94A	13.80A	0.1	0.081	0.922
HPL	2.67A	2.72A	2.76A	0.3	0.35	0.706
DIT	3.06A	3.25A	3.13A	1.6	1.66	0.192
LOP	6.76A	7.16A	7.13A	2.8	3.00	0.052
NSE	7.50A	7.47A	7.42A	0.2	0.18	0.834
LFR	2.24A	2.23A	2.20A	0.1	0.06	0.933
DFR	1.59A	1.50B	1.52AB	4.4	4.77	0.009
PCG	2.77B	2.97A	2.84AB	4.7	5.095	0.007
PFF	577.02C	1681.68B	3577.66A	97.5	3946.34	< 0.0001

Table 7: Evaluation of the average performance of the 4 groups by the Newmann Keuls test.

Number of days 50% flowering (DFL); number of fruits per plant (NFR); Petiole length (LPE); Leaf length (LOL); Plant height (HPL); Stem Diameter (DIT); Petal length (LOP); Number of sepal (NSE); Length of fruit (LFR); Diameter of fruit (DFR); 100 seeds Weight (PCG); Leaves biomass (PFF). F: F of Fisher; R²: coefficient of determination.

Variables	SMG	SME	VG	VP	H ² (%)	√VG	√VP	GCV(%)	PCV(%)	GA	GAx(%)
DFL	221.31	40.89	60.14	73.77	81.52	7.75	8.59	9.86	10.92	14.42	18.34
NFR	1157.6	149.1	336.17	385.87	87.12	18.33	19.64	26.23	28.10	35.25	50.43
LPE	9.76	5.7	1.35	3.25	41.60	1.16	1.80	5.89	9.13	1.55	7.83
LOL	7.75	2.97	1.59	2.58	61.68	1.26	1.61	9.07	11.55	2.04	14.67
LAL	6.03	3.24	0.93	2.01	46.27	0.96	1.42	8.89	13.07	1.35	12.45
HPL	0.42	0.27	0.05	0.14	35.71	0.22	0.37	8.28	13.86	0.28	10.20
DIT	0.4	0.33	0.02	0.13	17.50	0.15	0.37	4.86	11.63	0.13	4.19
LOP	2.14	0.8	0.45	0.71	62.62	0.67	0.84	9.57	12.10	1.09	15.61
LAP	0.96	0.58	0.13	0.52	0.01	0.36	62.63	9.34	16.72	0.00	0.11
NSE	0.6	0.44	0.05	0.20	26.67	0.23	0.45	3.09	5.99	0.25	3.29
LFR	0.32	0.3	0.01	0.11	6.25	0.08	0.33	3.68	14.71	0.04	1.89
DFR	0.04	0.03	0.00	0.01	25.00	0.06	0.12	3.75	7.50	0.06	3.86
PCG	0.52	0.01	0.17	0.19	98.08	0.41	0.42	14.42	14.56	0.84	29.41
PFI	891.2	13.99	292.40	297.07	98.43	17.10	17.24	25.08	25.28	34.95	51.25
PFF	3833848	8074	1 275 258.00	1 277 949.33	99.79	1 129.27	1 130.46	71.16	71.23	2 323.85	146.43

Table 8: Evaluation of the genetic parameters of the accessions of *H. cannabinus* of Burkina Faso.

Number of days 50% flowering (DFL); number of fruits per plant (NFR); Petiole length (LPE); Limb width (LAL), Leaf length (LOL); Plant height (HPL); Stem Diameter (DIT); Petal length (LOP); Petal width (LAP); Number of sepal (NSE); Length of fruit (LFR); Diameter of fruit (DFR); Weight of 100 seeds (PCG); Weight of dry fibers (PFI); Leaves biomass (PFF); VG: Genotypic variance, VP: Phenotypic variance H²: Heritability, GCV: Genotypic coefficient of variation, PCV: Phenotypic coefficient of variation, GAx: Genetic advance expected.

#### DISCUSSION

This study revealed the existence of a high morphological variability within accessions of Hibiscus cannabinus grown in Burkina Faso. The higher coefficients of variation values of studied characters give an idea of this variability. The most variable characters are the number of fruits (CV=31.52%) biomass and leaves (CV=71.25%). According to Aljane and Ferchichi (2007), a high value of the coefficient of variation indicates the heterogeneity of the studied genetic material. So the studied accessions are heterogeneous for the variables, number of fruits and the leaves biomass. Six morphotypes have been identified on the basis of the stem color compared to three morphotypes identified according to farmers's nomenclature during ethnobotanical investigation. This difference could be resulted from the interpretation of the morphotypes coloration by the farmers. In fact, farmers don't distinguish the red color and violet, the black color and purple. The prevalence of the green morphotype that is observed within the cultivars may be explained by peasants' farming method which is more oriented by the preference of the green color. Indeed, the leaves of the green morphotype are more valued for its food value and its fibers are more suitable for the (Kabré stringing et al., 2017). The performances of the limb length (13.9 cm), limb width (10.85 cm), petal length (6.98 cm) and 50% flowering days (78 days), are close to those found by Faruq et al. (2013) on Malaysian morphotypes. However the diameter of the stem (1.4 cm) and the height of the plant (2.1 m), measured by Faruq et al. (2013) were, however, smaller than those observed in our study. Indeed the cumbersome span of the plant vegetative part varies from a country to another and could be influenced by photosensitivity and climate type according to Mahieu et al. (2003). These authors argue that H. cannabinus has a better growth in the tropical and subtropical zones, an altitude of no less than 500 m above sea level. This is due to the fact that it adjusts itself to both hot and humid climate.

The study also revealed interesting correlations between several characters related to the plant height (HPL) and the stem diameter (DTI) on the one hand and between the plant height and the dry fibers weight on the other hand. These correlations show that plants of large size have a well-developed stem to assure its stability and also produce much fiber. Accessions organization in three distinct groups without referring to their origin could indicate the existence of diversity within accessions of the original population. This is due to the perpetual plant material exchanges between the farmers of different or neighboring localities, revealed during ethnobotanical investigation (Kabré et al., 2017). However, the morphological diversity between groups showed that there are possibilities of a genetic improvement of the species; either by direct selection within every group or by hybridization between groups. For a possible selection of Hibiscus cannabinus producing more leaves, seeds and fibers, group 3 is more reliable because it allows genitors' choice for new creation of varieties meeting producers' needs. Moreover, the high distance between groups (820-049; 1117-717 and 1937-766 for between 1-3; 1-2 and 2-3 respectively) would facilitate the transgressive segregation, because the greater the genetic distance, the more heterosis in the offspring can be observed. The testing of the genotypic coefficient of variation (GCV) and heritability (H²) provides interesting information on the choice of the parents to be hybridized in order to get the desired traits (Ibrahim et al., 2006). Thus, the identification of characters to be improved genetically is based on the variability and heritability of the characters. A good knowledge of the genetic parameters is thus necessary for the choice of the optimal selection criteria (Merour et al., 2008). In order to understand whether the observed variation was due to genetic factors, the genetic parameters were estimated. For the studied characters, phenotypic coefficient of variation (PCV) is higher than genotypic coefficient of variation (GCV) for all the characters. But the differences between them are low, indicating a significant genetic

control on the expression of the studied characters (Lakshmana et al., 2009). Similar results were reported on Burkina Faso sweet grains sorghum (Sawadogo et al., 2014), on millet (Drabo et al., 2013; Govindaraj et al., 2011; Hosseini et al., 2012) and on rice. According to the classification of Sumathi et al. (2010), the genotypic and phenotypic coefficients of variation are low when they are below 10%, moderate when they are between 11 and 20%; and higher when they are above 20%. So, the number of sepals, petiole length and fruit diameter have lower PCV and GCV. These coefficients of variation revealed the extent of the variability but the heritability allows knowing the proportion of variability transferable to the progeny. About studied characters expressed all high heritability (H²>20%). Danquah and Ofori (2012) got the same result with Ghana eggplants, Vetriventhan and Nirmalakumari (2007) on millet, Ali et al. (2012) and Nebié (2014) on sorghum. This high heritability of characters confirms the low influence of environmental factors characters on expression. Thus, the phenotype allows a good prediction of genotype accessions (Visscher et al., 2008). That means that the contribution of genotypes to the expression of characters is important. Heritability alone doesn't permit to predict if the selection brings a substantial improvement. Nevertheless, both heritability (H²) and genetic progress (GAx) evaluation can bring more reliable information (Govindaraj et al., 2011; Drabo et al., 2013). So heritability and the huge genetic progress (GAx) that have been observed, the number of days 50% flowering, the number of fruits, the limb length and width, the plant height, the petal length, the 100 seeds weight, the dry fibers weight and the leaves biomass, show large genetic effects of additive type in these characters determination (Kashif et al., 2003). Thus a direct selection method for the enhancement of H. cannabinus is possible for these traits.

#### Conclusion

This study revealed important agromorphological diversity within accessions of H. cannabinus in Burkina Faso. This variability is perceptible on all the studied characters and much more on the number of fruits, stem diameter, plant height, leaves biomass and dry fibers weight. These characters presented interesting relationships between them. indicating that the improvement of a single character would cause that of the others. The study also revealed a structuring of H. cannabinus accessions into three (3) agro-morphological groups without reference to their origin whose parameters such as number of fruits, dry fibers weight and leaves biomass are those that differentiate them. Morphological group 3 accessions have the best agronomic performances in relation to those of groups 1 and 2. The study, besides, shows more important genetic control on characters expression proved by high heritability and low extent difference between phenotypical and genotypical coefficients of variation. Besides heritability and genetic progress values suggests that most quantitative characters of H. cannabinus would be controlled by additive genes. Thus a genetic improvement can be considered from the variability highlighted in this study.

### **COMPETING INTERESTS**

Authors have declared that no competing interests exist.

### **AUTHORS'S CONTRIBUTION**

This work was carried out in collaboration between all the authors. NVK wrote the draft of the manuscript, performed the statistical analysis and the literature searches. BS, MK and ZK read and amend the manuscript and also managed the analyses of the study. RKN gave practical advice for the activities designed. PB-K wrote the research protocol, gave practical advice for the activities designed, read and enhanced the quality of the final manuscript and financially supported the study. All the authors read and approved of the final manuscript.

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