Genetic divergence of Heliconiaceae species in the Central West Brazil region

Divergencia genética de las especies de Heliconiaceae en la región Centro Oeste de Brasil

Cintia Graciele da Silva¹, Edneia Zullian Dalbosco¹, Petterson Baptista da Luz¹, Willian Krause¹, Vivian Loges², and Celice Alexandre Silva^{1*}

ABSTRACT

The purpose of this study was to describe morphological traits and estimate genetic divergence and parameters between accessions of the genus *Heliconia* sp. from different municipalities in the state of Mato Grosso, Brazil. A set of 25 traits, 15 quantitative and 10 qualitative were evaluated. The genetic divergence was estimated based on Mahalanobis' distance, with the clustering methods known as Unweighted Pair Group Method using Arithmetic Averages (UPGMA). Genetic variability was observed for all assessed quantitative traits and the accessions were grouped in different classes. The traits with highest relative contribution to variability were longevity of flower stems and inflorescence length. The results indicated the existence of genetic variability among accessions of the *Heliconia* sp. germplasm bank, which can be used in breeding programs.

Key words: morphological description, pre-breeding, tropical ornamentals.

Introduction

The family *Heliconiaceae* consists of 182 neotropical species distributed in Central and South America. Twenty-nine species are found in Brazil, five of which are endemic (Braga, 2014). *Heliconiaceae* are distributed in the Central-West, Northern, Northeastern, and Southeastern regions of the country. Nine species of the genus inhabit the state of Mato Grosso, i.e., *Heliconia psittacorum, H. rostrata, H. episcopalis, H. marginata, H. subulata, H. acuminate, H. hirsuta, H. densiflora,* and *H. stricta;* the last two were first recorded in Mato Grosso (Braga, 2014).

The state of Mato Grosso lies in the so-called Ecological Transition Zone (ETZ) between the biomes savanna (Cerrado) and rainforest (Amazon region). The ETZ is a complex morphoclimatic domain at the North of the Cerrado and Southwest of the Amazon, where savannas and tropical forests coexist under similar environmental conditions (Furley *et al.*, 1992).

RESUMEN

El objetivo de este estudio fue describir los rasgos morfológicos, estimar la divergencia genética y los parámetros entre accesiones del género *Heliconia* sp. en diferentes municipios del estado de Mato Grosso, Brasil. Se evaluó un conjunto de 25 rasgos, 15 cuantitativos y 10 cualitativos. La divergencia genética se estimó sobre la base de la distancia de Mahalanobis, con los métodos de agrupación Unweighted Pair Group Method utilizando promedios aritméticos (UPGMA). Se observó variabilidad genética para todos los rasgos cuantitativos evaluados y las accesiones se agruparon en diferentes clases. Los rasgos con mayor contribución relativa a la variabilidad fueron la longevidad de los tallos de las flores y la longitud de las inflorescencias. Los resultados indicaron la existencia de variabilidad genética entre las accesiones del banco de germoplasma de *Heliconia sp*, que pueden ser explotadas en programas de mejoramiento de la especie.

Palabras clave: descripción morfológica, pre-reproducción, plantas ornamentales tropicales.

Heliconiaceae species have herbaceous plants used in floriculture as ornamental plants, grown under full sun or partial shade, or as cut flowers, because the terminal inflorescences with intense colors of different shapes and sizes are greatly appreciated for event decoration and floral arrangements. Only few and recent studies addressed the agronomic potential of ornamental interest species. In the forest region Zona da Mata of Pernambuco, some studies analyzed genetic parameters of seven genotypes of *H. psit*-*tacorum* (Costa, 2007; Rocha *et al.*, 2010; Araujo *et al.*, 2015).

Due to the large natural variability in *Heliconia* sp. populations and the potential of these species as ornamental, research on breeding, agronomic and genetic characterization should be intensified. A study target of breeding programs is to collect accessions in genebanks to develop genotypes with traits of economic interest that meet the demands of the ornamental market (Rocha *et al.*, 2010). Moreover, the genetic divergence and parameters can be

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¹ Programa de Pos Graduação de Genética e Melhoramento de Plantas, University of Mato Grosso State, Tangará da Serra, MT (Brazil).

² Federal University of Pernambuco, Recife, PE (Brazil).

^{*} Corresponding author: celice@unemat.br



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analyzed to estimate genetic gains and determine the most appropriate breeding method (Cruz *et al.*, 2012).

This study addressed the morphological description and the estimation of genetic parameters and divergence of *H. psittacorum* and *H. densiflora* accessions, with a view to evaluate the genetic variability in the state of Mato Grosso, identifying possible parents to breed future hybrids with ornamental potential.

Material and methods

The germplasm collection of *Heliconias* (BAG) of the State University of Mato Grosso (UNEMAT) was established in March 2014, in an experimental field in the municipality of Tangará da Serra, MT (14°39' S and 57°25' W; elevation 321 m a.s.l.). The climate is tropical, with well-defined dry and rainy seasons and annual means varying from 1,300 to 2,000 mm rainfall and 16-36°C temperature range (Martins *et al.*, 2010). The soil, with a flat to slightly wavy relief, is classified as Latossolo vermelho distrófico (Embrapa, 2006).

The area for planting of seedlings was harrowed and limed according to soil analysis. Fertilization at planting consisted of 50 g monoamonic phosphate (MAP) per planting hole and topdressing of monthly applications of 50 g urea and 20 g potassium chloride per hole and of 50g MAP every 6 months.

Four accessions of *H. densiflora* and 14 of *H. psittacorum*, from 13 municipalities in the state of Mato Grosso, were described (Tab. 1). The clumps were divided and four rhizomes planted, at a spacing of 3.0 m between rows and 1.5 m between plants in full sun. When needed, irrigation was applied three times a week. The experiment was conducted as described by Costa *et al.* (2007), in a randomized block design with 18 treatments (accessions), four blocks (replications) with one rhizome per plot.

Qualitative and quantitative morphological traits were evaluated around 400 d after planting, when the plants were fully established. Twenty-five descriptors were used, of which 15 were quantitative and 10 qualitative. The quantitative descriptors were: LL (cm) leaflength; LW (cm) leaf width; NLS (n) number of leaves on the flower stem; NIC (n) number of inflorescences per clump; SW (g) shoot weight (leaves and floral stem); FSW (g) flower stem weight without leaves, flower peduncle and inflorescence; FSL (cm) flower stem length; FSD (cm) flower stem diameter, measured 20 cm below the inflorescence; IL (cm) inflorescence length; LI (cm) inflorescence width; NFI (n) number of flowers per inflorescence; NBI (n) number of bracts per inflorescence; BL (cm) bract length; BD (cm) bract depth; and SL (days) longevity of the flower stem. The qualitative

Accession	Species	Municipality	Latitude	Longitude	Altitude m a.s.l.
1	H. densiflora	Alta Floresta	9°51'05"	56°12'31"	281
2	H. densiflora	Alta Floresta	9°51'47"	56°12'04"	271
3	H. densiflora	Alta Floresta	9°52'43"	56°9'22"	281
4	H. densiflora	Carlinda	10°10'8"	55°48'53"	299
5	H. psittacorum	Nova Canaã	10°36'44"	55°42'05"	265
6	H. psittacorum	Colíder	10°46'55"	55°27'00"	310
7	H. psittacorum	Matupá	10°12'26"	54°57'39"	260
8	H. psittacorum	Guarantã Norte	9°46'02"	54°53'55"	348
9	H. psittacorum	Guarantã Norte	9°44'26"	54°53'16"	336
10	H. psittacorum	Peixoto Azevedo	10°16'9"	55°01'15"	324
11	H. psittacorum	Terra Nova do Norte	10°44'5"	55°08'43"	295
12	H. psittacorum	Santo Afonso	14°35'9"	57°10'56"	494
13	H. psittacorum	Nova Marilândia	14°21'5"	57°02'01"	355
14	H. psittacorum	Tangará da Serra	14°42'2"	57°47'31"	204
15	H. psittacorum	Barra do Bugres	15°07'6"	57°04'34"	156
16	H. psittacorum	Porto Estrela	15°18'1"	57°10'11"	168
17	H. psittacorum	Porto Estrela	15°24'2"	57°11'51"	148

15°35'37"

57°11'51"

TABLE 1. Accessions of Heliconia densiflora and H. psittacorum collected in different municipalities in the state of Mato Grosso, Brazil.

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H psittacorum

Porto Estrela

descriptors evaluated in the inflorescences and leaves were: inflorescence growth, hairiness, waxiness, bract and flower color; arrangement of bracts (flat or spiral); hairiness and waxiness of the flower stem (pseudostem + floral peduncle + inflorescence) and leaves; firmness of the flower stem (bracts + rachis), (Costa *et al.* 2007).

Flower stems with two to three open bracts were cut between 7:00 and 8:00 h, twice a week, for 1 month. The stalks were stored in water recipients and transported in buckets to the postharvest laboratory. The inflorescences were cleaned (removing the flowers from within the bracts), washed, and cut to a standardized stem length of 80 cm). Quantitative and qualitative traits of five flower stems per clump were assessed, with four replications.

The flower stalks were placed in containers with water, which was exchanged every 2 d and maintained at 19°C (cold room). The postharvest shelf-life (days) was evaluated every 2 d, for 21 d and the stems discarded when the bracts darkened on the inflorescences.

For the quantitative descriptors, analysis of variance was performed, the Scott Knott grouping test was applied and genetic parameters associated with effects of genetic and environmental nature, were estimated according to Cruz *et al.* (2012). The values of the genotypic determination coefficient (H^2) were expressed as the proportion of phenotypic variance caused by the genetic variability among treatment means (Cruz, 2006), and it can be used when the effect of genotypes is fixed, as in the case of this study.

To quantify the genetic divergence among accessions, Mahalanobis' generalized distance (D^2) was used for quantitative variables, while the simple match distance was used for the qualitative descriptors. The relative importance of variables was also evaluated by the methodology of Singh (1981). For the simultaneous analysis of qualitative and quantitative variables, the Gower's General Coefficient of Similarity was performed (1971).

The statistical analysis was performed using GENES software (Cruz, 2014). The accessions were grouped by the hierarchical method Unweighted Pair-Group Method Using Arithmetic Averages (UPGMA) and validated by the cophenetic correlation coefficient using the software MEGA, version 5 (Kumar *et al.*, 2009). The Pearson correlation between the distance matrices was also performed and the significance determined by the Mantel test (10,000 permutations). All Mantel tests performed here were conducted using the R packages *vegan* (Oksanen *et al.*, 2012).

Results and Discussion

All H² values exceeded 87%, except for NIC (54.28%) and NFI (67.55%), which were considered low (Tab. 2). For the H² values, the variation index (I_v) was considered. According to Vencovsky and Barriga (1992), an I_v higher than one (1.0) indicates good conditions for selection gains by simple breeding methods, such as mass selection.

TABLE 2. Estimates of the genetic parameters of four accessions of *Heliconia densiflora* and 14 accessions of *H. psittacorum*.

T	Parameters									
Trait	σ_{f}^{2}	σ_{e}^{z}	$\widehat{\varphi}_{\mathfrak{g}}$	H² (%)	CV _g (%)	I _v				
NIC	10.14	4.63	5.50	54.28	22.02	0.54				
FSL	0.06	0.003	0.05	94.54	19.40	2.08				
SW	2,068.08	178.20	1,889.88	91.38	35.30	1.62				
FSW	997.75	86.61	911.13	91.31	34.04	1.62				
FSD	0.49	0.06	0.43	87.74	13.88	1.33				
NBI	1.29	0.08	1.21	93.80	21.32	1.94				
BD	5.56	0.60	4.96	89.14	22.31	1.43				
BL	10.81	0.39	10.41	96.31	26.71	2.55				
IL	12.42	0.48	11.93	96.07	25.11	2.47				
IW	6.75	0.70	6.04	89.53	26.13	1.46				
NFI	26.30	8.53	17.77	67.55	20.58	0.72				
NLS	4.75	0.10	4.64	97.73	27.91	3.28				
LW	2.51	0.24	2.26	90.23	14.38	1.51				
LL	50.78	1.82	48.96	96.41	20.94	2.59				
SL	4.41	0.023	4.39	99.47	26.19	6.88				

NIC: number of inflorescences per plant; FSL: flower stem length; SW: flower stem weight with leaf; FSW: fresh stem weight without leaf; FSD: diameter of the flower stem; NBI: number of bracts per inflorescence; BD: bract depth; BL: bract length; IL: inflorescence length; IW: inflorescence width; NFI: number of flowers per inflorescence; NLS: number of leaves on the flower stem; LW: leaf width; LL: leaf length (LL); SL: stem longevity.

The selection gain for the traits NIC and NFI may be minimized, since the H² value is low and I_v <1.0. The results also suggest the existence of genetic variability among accessions of *Heliconia* sp. for the studied traits, indicating favorable genetic values for breeding programs. For the 15 evaluated quantitative traits, several morphological groups were identified (Tab. 3)

Analyzing agronomic traits of *H. psittacorum* genotypes under full sun and partial shade, Costa *et al.* (2007) found H^2 varying from 82.25 to 97.33%, except for the trait days to inflorescence cut (20.02%). Similar results were reported by Rocha *et al.* (2010), in a study with *H. psittacorum* cultivars and interspecific hybrids, where I_v values between 0.21 and 1.85 were recorded for seven traits (days until inflorescence sprouting, period until stem harvest, cycle, stem weight without leaves, stem diameter, inflorescence length, number of open bracts per inflorescence). **TABLE 3.** Relative contribution of 15 quantitative traits to genetic variability detection, based on Mahalanobis' generalized distance between 18 accessions of *Heliconia densiflora* and *H. psittacorum*.

Evaluated traits	Diversity (Sj)	Contribution (%)
Number of inflorescences per plant (NIC)	201.06	0.29
Flower stem length (FSL)	3,933.14	5.85
Flower stem weight (FSW)	820.57	1.22
Flower stem weight with leaf (SW)	1,502.58	2.23
Number of bracts per inflorescence (NBI)	3,376.23	5.02
Bract depth (BD)	1,080.07	1.60
Diameter of the flower stem (FSD)	707.27	1.05
Bract length (BL)	3,908.13	5.81
Inflorescence length (IL)	15,402.00	22.93
Inflorescence width (IW)	1,974.05	2.93
Number of flowers per inflorescence (NFI)	90.21	0.13
Number of leaves on the flower stem (NLS)	4,906.55	7.30
Leaf width (LW)	3,555.69	5.29
Leaf length (LL)	6,376.53	9.49
Stem longevity (SL)	19,335.58	28.78

The inflorescence traits such as length, stem length, fresh weight, and durability are of great commercial interest, due to promising ornamental potential. Inflorescences of *Heliconiaceae* can be classified in small (up to 10 cm), medium (10.1 to 30 cm), large (30.1 to 50 cm), and very large (>50 cm) (Castro *et al.*, 2007). Of the 18 accessions, 16 were classified as medium and two (accessions 5 and 16) as small (Tab. 4). The small to medium inflorescences are less difficult to handle and transport (Castro *et al.*, 2007). Genetic diversity studies of *H. psittacorum* and interspecific hybrids in the state of Pernambuco registered inflorescence lengths from 12.1 to 23.3 cm (Rocha *et al.*, 2010).

The traits FSL and FSD are important to ensure the quality and success of *Heliconia* species on the market. As standard length of floral stems of *Heliconiaceae*, Loges *et al.* (2005) suggested 80 cm. On the other hand, the higher the FSL, the greater the risk of stem breaking. Consequently, accessions with a FSL equal to or greater than 80 cm are recommended for breeding programs (Tab. 4).

TABLE 4. Mean values of 15 quantitative traits of Heliconia densiflora (accessions 1-4) and H. psittacorum (accessions 5-18).

	Means of evaluated traits														
Accession	IL	FSL	FSW	SL	SW	FSD	NIC	NBI	BD	BL	IW	LW	LL	NLS	NFI
	cn	n	G	days	g	Mm	un		mm		cm			un	
							H. densi	flora							
1	15.6 c	88 e	60.3 c	6.5 d	82.8 c	5.1 b	8.7 b	3.9 d	12.3 a	14.2 b	4.7 c	9.1 c	33.1 c	6.0 d	17.4 b
2	23.2 a	124 c	108.5 b	6.5 d	146.7 b	5.4 b	13.2 a	4.6 c	8.5 c	21.1 a	6.4 c	11.2 b	45.3 a	5.9 d	17.5 b
3	17.0 b	93 e	56.25 c	6.6 d	73.8 c	5.2 b	8.2 b	3.9 d	8.7c	15.3 b	6.4 c	11.0 b	37.7 b	5.9 d	14.3 b
4	18.1 b	107 d	61.46 c	6.4 d	76.9 c	5.4 b	11.7a	3.9 d	12.6 a	15.1 b	4.7 c	12.1 a	31.0 d	4.8 d	16.0 b
							H. psittac	orum							
5	8.3 e	135 b	87.96 c	6.5 d	127.2 b	6.3 a	7.0 b	4.3 c	5.3 d	7.3 d	9.3 b	12.3 a	13.3 f	11.3 a	10.3 b
6	12.7 d	180 a	155.0 a	6.6 d	203.5 a	4.5 c	8.25 b	5.6 b	9.0 c	11.1 c	12.8 a	11.8 a	37.9 b	10.7 a	18.1 b
7	12.7 d	141 b	112.5 b	6.6 d	169.4 b	5.0 b	13.0 a	6.3 a	6.9 d	10.0 c	9.3 b	12.3 a	34.7 c	11.6 a	26.0 a
8	12.6 d	127 c	94.4 b	8.5 b	136.3 b	4.6 c	9.0 b	6.2 a	6.9 d	10.5 c	10.4 b	11.1 b	33.5 c	8.6 b	16.0 b
9	12.1 d	144 b	111.6 b	7.7 c	148.2 b	5.4 b	11.0 a	4.4 c	10.4 d	9.6 d	9.1 b	9.5 c	44.2 a	6.1 d	25.8 a
10	12.0 d	143 b	115.8 b	7.7 c	162.8 b	4.7 c	8.75 b	6.8 a	8.9 c	10.6 c	10.2 b	11.2 b	34.6 c	9.1 b	21.1 a
11	11.3 d	148 b	143.3 a	7.8 c	211.8 a	4.6 c	7.25 b	5.7 b	10.7 b	10.3 c	12.5 a	12.8 a	38.1b	9.6 b	16.9 b
12	11.8 d	93 e	57.5 c	8.6 b	86.5 c	4.1 d	8.25 b	6.4 a	11.6 b	10.7 c	11.1 b	9.3 c	31.2 d	7.6 c	28.75
13	16.2 c	99 e	70.8 c	7.7 c	89.2 c	4.6 c	5.25 b	3.3 d	12.7 a	13.9 b	14.0 a	8.8 c	24.1 e	5.3 d	23.3 a
14	10.7 e	117 d	72.5 c	7.7 c	98.5 c	3.8 d	15.0 a	5.6 b	9.1 c	9.1 d	10.3 b	9.7 c	32.3 d	8.0 c	21.2 a
15	14.8 c	93 e	39.8 c	13.6 a	57.8 c	3.8 d	13.0 a	4.8 c	8.5 c	13.2 b	8.0 c	7.4 d	30.2 d	5.6 d	21.5 a
16	9.7 e	122 c	65.4 c	7.5 c	91.2 c	3.5 d	16.0 a	6.6 b	10.2 b	9.0 d	9.2 b	9.0 c	28.9 d	8.2 c	28.5 a
17	11.9 d	141 b	101.4 b	7.8 c	150.4 b	4.0 d	15.0 a	6.7a	13.5 a	10.9 c	11.1b	10.6 b	35.7 c	8.7 b	25.9 a
18	16.0 c	122 c	80.8 c	13.3 a	102.8 c	4.9 b	13.0 a	3.8 d	13.1 a	14.9 b	9.0 b	8.1 d	35.1 c	5.4 d	18.8 b

IL: inflorescence length; FSL: flower stem length; FSW: fresh stem weight without leaf; SL: stem longevity; SW: flower stem weight with leaf; FSD: diameter of the flower stem; NIC: number of inflorescences per plant; NBI: number of bracts per inflorescence; BD: bract depth; BL: bract length; IW: inflorescence width; LW: leaf width; LL: leaf length; NLS: number of leaves on the flower stem; NFI: number of flowers per inflorescence.

Means followed by the same letter in the column do not differ statistically by the Scott-Knott test at 5% probability.

The fresh weight of flower stems without leaf (FSW) ranged from 39.8 to 155.0 g (Tab. 4). This trait directly affects the stages management, preparation, packaging, and transport. The success of cut flowers on the market depends on the selection of accessions with a stem durability of more than 10 d. Long-lived stems can reach markets that are more distant by extending the shelf-life and maintaining the commercial quality (Castro *et al.*, 2007).

In the assessment of genetic diversity based on the 15 quantitative traits, three groups were determined using 255 as a cut distance (Fig. 1). Group I comprised the accessions 1, 2, 3, 4, 6, 8, 9, 10, 11, 12, 13, 14, 16, and 17, Group II accession 5 and Group III accessions 15 and 18. The isolation of the acession 5 of *H. psittacorum* in Group II was based on the traits of FSD and LL (Tab.4). The FSD expresses greater resistance of the flower stem against breaking by wind in the field, to transportation from the field to the location of cleaning, and in the subsequent selection stages (Albuquerque *et al.*, 2010).

The cophenetic correlation for the data was 0.87, confirming the reliability of the conclusions based on the visual



FIGURE 1. Dendrogram of genetic dissimilarity between *Heliconia densiflora* (accessions 1-4) and *H. psittacorum* (accessions 5-18), obtained by the UPGMA method, based on 15 quantitative traits, using Mahalanobis' generalized. Cutting distance 255 (Cophenetic correlation: 0.87; Distortion: 8.3%; Stress: 28.9%).

assessment of the dendrogram, since values above 0.7 indicate a good adjustment of the dissimilarity matrix and the graphical representation of the distances (Sokal and Rohlfe, 1962). The accessions of the species *H. densiflora* (accessions 1, 2, 3, and 4) were grouped in the same group (G I), while the accessions of the species *H. psittacorum* were separated in two groups.

The highest contributions to the determination of the divergence of the studied accessions (Tab. 4) were related to the traits longevity of the flower stem (SL) with 28.78%, and inflorescence length (IL) with 22.93%, both accounting for 51.71% of the variation among accessions. Stem durability was an important trait for the genetic divergence among accessions, since the accessions 15 and 18 with highest SL formed one group (Group III). On the other hand, the flower stem diameter (FSD) and leaf length (LL) forcing the isolation of accession 5, are traits with no significant contribution to the genetic divergence among accessions (Tab. 4; Fig. 1)

To ensure the success in parental selection for breeding programs, less relevant traits should not be considered in characterization and evaluation studies and the other traits with high discriminatory potential maintained (Cruz *et al.*, 2012). The traits that contributed least to divergence were number of flower inflorescences (NFI) (relative contribution to variation 0.13%) and number of inflorescences per clump (contribution 0.29%) (Tab. 4), which were therefore discarded.

Among the 10 qualitative traits evaluated, six indicated no polymorphism among accessions, e.g.: upright inflorescence position, absence of hairiness on flower stem and leaves, and firmness and flat arrangement of the bracts. Moreover, the accessions differed in terms of flower and inflorescence colors and presence of waxiness on the inflorescence, leaves and stem (Tab. 5).

Tropical flowers differ from other more traditional plants on the market in their diversity of shapes, resistance to transport and durability (Loges *et al.*, 2005), aside from attractive visual aspects, being widely used for floral arrangements, event and party decor, cultivation in gardens, and projects of ornamental garden design. In studies of *Heliconia* sp species focused on cut flowers, Castro *et al.* (2007) reports that the high contrast of bract colors is one of the traits responsible for the appeal of this ornamental plant on the market. In the accessions, eight colors were identified in the inflorescences and two in the flowers (Tab. 5). Thus, the existence of variability for the traits in

TABLE 5. Qualitative traits evaluated in 18 Heliconia sp. accessions, collected in municipalities of the state of Mato Grosso	, Brazil.
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Accesions		Inflores	cence		В	racts	Flor	St	em	Leaf	
ACESSIONS -	Growth	Color	Hairiness	Hairiness Waxiness Firmness Arrangement		Color	Hairiness	Waxiness	Waxiness		
	H. densiflora										
1	Upright	Orange-red	Absent	Absent	Resistant	Flat	Orange	Absent	Absent	Absent	
2	Upright	Orange-red	Absent	Absent	Resistant	Flat	Orange	Absent	Absent	Absent	
3	Upright	Orange-red	Absent	Absent	Resistant	Flat	Orange	Absent	Absent	Absent	
4	Upright	Orange-red	Absent	Absent	Resistant	Flat	Orange	Absent	Absent	Absent	
					H. psittaco	rum					
5	Upright	Red	Absent	Absent	Resistant	Flat	Yellow	Absent	Present	Absent	
6	Upright	Red	Absent	Absent	Resistant	Flat	Yellow	Absent	Present	Absent	
7	Upright	Red	Absent	Absent	Resistant	Flat	Yellow	Absent	Present	Absent	
8	Upright	Red	Absent	Absent	Resistant	Flat	Yellow	Absent	Present	Absent	
9	Upright	Light pink	Absent	Present	Resistant	Flat	Orange	Absent	Present	Present	
10	Upright	Red	Absent	Absent	Resistant	Flat	Yellow	Absent	Present	Absent	
11	Upright	Red	Absent	Absent	Resistant	Flat	Yellow	Absent	Present	Absent	
12	Upright	Yellow	Absent	Absent	Resistant	Flat	Yellow	Absent	Present	Absent	
13	Upright	Dark red	Absent	Absent	Resistant	Flat	Yellow	Absent	Absent	Absent	
14	Upright	Dark orange	Absent	Absent	Resistant	Flat	Yellow	Absent	Present	Absent	
15	Upright	Dark pink	Absent	Present	Resistant	Flat	Orange	Absent	Present	Present	
16	Upright	Dark orange	Absent	Absent	Resistant	Flat	Yellow	Absent	Present	Absent	
17	Upright	Orange	Absent	Absent	Resistant	Flat	Yellow	Absent	Present	Absent	
18	Upright	Dark pink	Absent	Present	Resistant	Flat	Orange	Absent	Present	Present	

flower and inflorescence color is fundamental for successful breeding.

Waxiness on inflorescences and leaves was detected in 21.43% of the accessions and 92.85% of the floral stems of *H. psittacorum* accessions (Tab. 5). Waxiness is an undesirable trait, since the visual aspect can be affected by handling (Loges *et al.*, 2005). This trait was not observed in *H. densiflora* accessions (Tab. 5).

Considering only the qualitative traits, two groups were determined using 0.148 as a cut distance. Group II contained the accessions 9, 15 and 18 and Group I the remaining accessions (Fig. 2). The cophenetic correlation was 0.94, also considered high. No polymorphism was detected for accessions of the species *H. densiflora* (accessions 1, 2, 3, and 4). Once again, the accessions of species *H. psittacorum* were classified in two distinct groups, with polymorphism within the species.

The correlations between dissimilarity matrices quantitative, estimated by Mahalanobis (0.74), qualitative Simple Match (0.62) and combined Gower distance (0.84) the whole set of traits, were all significant and high.

In the combined analysis (Gower General Coefficient of Similarity), showing correlation between the matrices of original distances and the cluster matrix. Using 3.6 as cut



FIGURE 2. Dendrogram of genetic dissimilarity between *Heliconia densiflora* (accessions 1 to 4) and *H. psittacorum* (accessions 5 to 18), based on the UPGMA method, using 11 qualitative traits and the Simple Match. Cutting distance 0.148 (Cophenetic correlation: 0.94. Distortion: 2.7%. Stress: 16.4%).

distance three groups were determined: Group I contained the accessions 5, 6, 7, 8, 10, 11, 12, 13, 14, 16, and 17, Group II accessions 1, 2, 3, and 4, and Group III accessions 9, 15 and 18 (Fig. 3). The *H. psittacorum* accessions were assigned to Group I and III. The *H. densiflora* accessions were clustered in Group II, showing the low variability of these accessions, probably due to the low number of accessions.



FIGURE 3. Dendrogram of genetic dissimilarity between *Heliconia densiflora* (accessions 1 to 4) and *H. psittacorum* (accessions 5 to 18), obtained by the method UPGMA, based on 15 quantitative and 11 qualitative traits, using the Gower distance. (Cophenetic correlation: 0.84. Distortion: 2.9%. Stress: 17.0%). Cutting distance 3.6.

In Group I, accession 13 stood out with a lower FSL and FSW, along with the absence of waxiness on the inflorescence, stem and leaf. In addition, the inflorescences are dark red and the flowers yellow.

In Group II, the accessions 1, 2 and 4 had low FSL and FSW values and no waxiness on the inflorescence, stem and leaf, as well as highest FSD and orange-red inflorescences and orange flowers.

In Group III, accessions 15 and 18 stood out for good postharvest shelf-life. In breeding programs, parents with high genetic divergence are recommended, to promote the occurrence of superior segregating plants in subsequent generations. However, the choice of individuals with superior plant, flower and fruit traits is also important, and finally, performance analysis for later recommendation of use *per se*. Thus, crosses should be performed between different genotypes with desirable agronomic traits (Cruz *et al.*, 2012).

To this end, we suggest to cross accessions 15 and 18 (Group III) with accessions 1, 2 and 4 of Group II and/or accession 13 of Group I, to breed superior genotypes.

Studies of Gomes *et al.* (2016) pointed out that among *Heliconias* species, *H. psittacorum* cultivars and hybrids have a number of interesting traits, such as year-round production, terminal and upright inflorescences, varied bract number, and different types of flower colors, as observed in this study.

In this sense, based on the traits evaluated, the studied accessions are of interest for ornamental use, enabling a diversification of the ornamentals market. Traits such as bract length are relevant for playing an important role in the composition of floral arrangements and to raise the interest of consumers, as was expressed by Albuquerque *et al.* (2010).

The diameter and length of the flower stem are also key traits to characterize *Heliconia* sp. accessions. They make the flower stem more resistant to winds, to the transport from the field to the location of cleaning and the selection steps, and extend the postharvest shelf-life.

Conclusions

Morphological characterization of *Heliconia* species allowed the recognition of those descriptors which contribute to the detection of genetic divergence and provided key knowledge of the patterns of variation phenotypic of the target species.

The genetic variability among accessions of germplasm collection of *Heliconia* sp. uncovered in the present study can be explored in the breeding programs.

The characteristics that contributed most to the detection of genetic divergence were floral stem durability and inflorescence length.

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