

Characterization and selection of interspecific hybrids of *Brachiaria decumbens* for seed production in Campo Grande – MS

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Abstract: *The breeding program Brachiaria developed by Embrapa Beef Cattle provides studies to obtain forage with agronomic characters desired by farmers. In this regard, in 2013 and 2014, a study was carried out in order to select the best Brachiaria decumbens intraspecific hybrids, which are superior in relation to production, forage nutritional value, and resistance to spittlebugs. Estimates of genetic parameters and gains with selection were carried out. It was found that there was significant variability between genotypes for all characters. Gain with selection (GS%) ranged from 12 to 324%, and the highest percentage was found for weight of pure seeds (PS) of seed collectors of the second harvest. For reproductive tillers (RT), C001 and R091 hybrids had better performance than the control, and this character may be considered as a parameter to estimate production of pure seeds before flowering starts.*

Key words: *Seed production potential, genetic variability, genotypes, forage improvement.*

INTRODUCTION

At first, selection of forage plants was based only on mass production potential and on forage quality under pasture conditions. However, in recent years, some changes have been observed in strategies of breeding programs of these plants, aiming at obtaining superior cultivars in all aspects. Cultivars development process is long, comprising several stages, and the evaluated characters should be correlated in order to result in crops that have good performance at all stages and variables of study and research. In order to reach the release of a new cultivar, the development process involves several research lines, such as improvement; cytogenetics of the reproductive system; plant nutrition; microbiology; plant health; pasture management and nutritional quality of plants; and seed production and technology, among others (Karia et al. 2006, Valle et al. 2009).

Improvement of tropical forage plants is relatively new compared to other cultures (Karia et al. 2006, Araújo et al. 2008, Valle et al. 2009), and aims to release more productive plants in production and forage quality, and in various agronomic aspects, such as production of good quality seeds and in satisfactory quantity, adaptation to different soil and climate conditions, and especially

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resistance to spittlebug.

Brachiaria plants have been widely used as forage in tropical America, and most species found in Brazil are considered exotic. Recently, there has been a reclassification of the *Brachiaria* genus to *Urochloa*. However, discussions remain among researchers regarding its characteristics and new taxonomic classification. *Brachiaria decumbens*, for its high forage production potential and high adaptability to acid soils and of low fertility, has great importance for national beef cattle. Even being economically very relevant, there is only one cultivar in the market, cv. Basilisk, which was released in the 1960s. One of the limitations of this cultivar is its susceptibility to spittlebugs.

It is extremely important that cultivars from breeding programs solve problems such as susceptibility to spittlebugs. Cv. Marandu is an example, which has been available since the 1980s. A few options of cultivars available to producers have good seed production, so that the ratio production cost x price of seed production is satisfactory. Seed production may vary depending on the location of production, management, and on the form of harvest, and it may reach up to 300 kg ha⁻¹ (Monteiro et al. 2016).

This work aims to characterize and evaluate the potential of seed production by intraspecific hybrids of *Brachiaria decumbens*, developed and pre-selected by the breeding program of Embrapa Beef Cattle, aiming at selecting superior genotypes, which could be candidates for new cultivars or potential parents of sexual propagation to be used in new crosses.

MATERIAL AND METHODS

Genetic materials used in the development of this work were selected based on previous experimental results obtained by the Forage Cultivars Breeding and Production Program of Embrapa Beef Cattle. Promising intraspecific hybrids of *Brachiaria decumbens* (sexual and apomictic), parents and candidates for new cultivars were selected among 324 hybrids, which were obtained by crossing between three plant of sexual reproduction of *B. decumbens* artificial tetraploids (D24/2, D24/27 e D24/45) with cv. *Basilisk* (apomictic). Selection was based on agronomic characteristics of production, forage nutritional value, and resistance to spittlebugs. *B. brizantha* cv. Marandu was used as control. Table 1 shows the genotypes evaluated during the first (2013) and the second (2014) harvest.

Trials were carried out for two consecutive years in Campo Grande - MS, at Embrapa Beef Cattle (lat 20° 25' 03" S, long 54° 42' 20" W, alt 530 m asl). Local climate is classified as Aw tropical rainy savanna, characterized by rainy summer and dry winter. Trials were carried out under normal environmental conditions, in rainy condition. Soil was classified as clayey dystroferic Red Latosol (Oxisol) (53% clay, 38% sand and silt 9%).

Plants were transferred to the field in early 2012 in the

Table 1. Genotypes evaluated during the first and the second years of production

Genotypes	1st year	0	2nd year
A001	A036		B006
A002	A038		C001
A003	A041		R025
A004	A042		R033
A005	A043		R041
A007	A044		R044
A008	B005		R071
A009	B009		R078
A011	B010		R087
A012	B026		R091
A013	C001		R101
A015	R184		R107
A017	S044		R110
A018	T038		R120
A019	X121		R124
A020	-		R126
A021	-		R144
A023	-		R181
A024	-		S018
A025	-		S031
A026	-		S036
A027	-		T005
A028	-		T012
A029	-		T026
A030	-		T054
A031	-		X030
A032	-		X072
A033	-		X117
A035	-		Y021



Figure 1. Seed collectors installed in the plot (A). Inflorescence inside the collector (B).

form of seedlings. Each plot contained two seedlings, with useful area of 2 m², spaced 1m between plots. Evaluations were carried out in the first quarter of 2013, with first-year plants, and between December 2013 and the first quarter of 2014, with second-year plants. For the first year of evaluations (2012/2013 crops), it was carried out uniform cutting of plants for seed production in October 2012. For the second year of evaluation, standardization was carried out on October 15, 2013. The experimental design was a complete block with two replications, and two observations per plot. After uniform cuttings (2012 and 2103), fertilization of plants was carried out based on soil chemical analysis. Evaluations were carried out in plots and in seeds collectors. Two collectors per plot were installed and, inside them, it was placed five inflorescences in complete anthesis without threshing (Figure 1).

To determine the typical inflorescences of genotype, ten inflorescences selected at random were collected. When at least six of them were of the same type, it was considered typical inflorescence. Eight typical inflorescences were collected in each plot, and from them, the following characteristics were determined: number of seeds per raceme (NSR), number of racemes (NR), number of seeds per inflorescence (NSI), length of racemes (LR), and length of inflorescences (LI). To determine these characteristics, it was carried counting in increasing order from the inflorescence base to the apex. LR was determined with the aid of a graduated scale. To this end, each raceme was measured from its insertion point in the rachis to the apex. LI was obtained with the aid of the graduated scale, by measuring the distance between the insertion point of the first and last raceme in the inflorescence.

It was also carried out evaluations in the plots at the beginning of the formation of the first reproductive tillers, and before the formation of the first panicle (inflorescence), in 0.25 m² in relation to the number of reproductive tillers (RT). After threshing of inflorescences' seeds, collectors were removed from the plots with their respective seeds and flowers. The collected material was taken to the Seed Laboratory of Embrapa Beef Cattle and evaluated for weight of pure seeds (PS) and weight of empty seeds (ES).

To determine the quality of seeds, it was carried out harvest of the total area of the plot (2 m²). To this end, in the maturation point of each plot, it was carried out plant cutting of the entire plot (except the inflorescences that were inserted in the collectors). Cutting was carried out manually at 20 cm from the soil, with a rice cutter, and was carried out at the beginning of threshing, of 15 to 20% of inflorescences' seeds. The material collected was packed in paper bags, which were closed and placed to dry in the shade. After drying, samples were submitted to manual separation of inflorescences' seeds. Seeds were then subjected to processing by pre-cleaning and cleaning using mesh and air column blower. After seeds processing, which were harvested in 2013, it was carried out the germination standard test (G%), according to RAS (Seed Analysis Rules) (Brasil 2009).

The collected data were submitted to analysis of deviance (ANADEV) using the Selegen-Reml/Blup software (Resende 2006) in order to simultaneously estimate genetic parameters and predict breeding values (BLUP). For the evaluations carried out in 2013, it was used the model of the 20 Selegen-Reml/Blup software, as follows:

$$y = Xr + Zg + e$$

In which y is the data vector; r is the vector of replication effects (fixed) added to the mean; g is the vector of genotypic effects (random); e is the error vector or residues (random); X and Z are the incidence matrices for these effects. For the other characters, evaluated in 2014, it was used the model 2 of the software, as follows:

$$y = Xr + Zg + Wp + e$$

In which y is the data vector; r is the vector of replication effects (fixed), added to the mean; g is the vector of genotypic effects (random); p is the vector of plot effects; e is the vector error or residue (random); X , Z and W are the incidence matrices for these effects.

RESULTS AND DISCUSSION

For germination percentage (G%) of the first harvest, there was genotypic difference. Block effect, which was considered as fixed effect, was tested by the Snedecor F test. When genotype effects are significant, respective components of variance are significantly different from zero, and so are their coefficients of determination (Resende and Duarte 2007). Heritability between genotypes means (h^2mc) exceeded in 50%, and it can be considered of medium magnitude, showing

that more than half of the observed phenotypic variation, on average, was due to genetic causes (Table 2).

Of the seeds trashed in the collectors, it was observed accuracy (Acc) of 84 and 97% (Table 2). Thus, it was within the established standards, and was classified as high (Resende and Duarte 2007), which means good reliability for the prediction of genotypic values of treatments. Both for weight of pure seeds (PS) and for weight of empty seeds (ES), it was found significant differences by the Chi-square test (LRT) for the genotype effect, and their heritability estimates (h^2mc) presented percentage above 0.50 (Table 2).

Table 2. Deviance Analysis (ANAEV), variation components of genetic parameters for germination (G%) in the first harvest, weight of pure seeds (PS), weight empty seeds (ES), number of reproductive tillers (RT), number of seeds per inflorescence (NSi), number of seeds per racemes (NSR), number of racemes (NR), length of raceme (LR) and length of inflorescences (LI) in the second harvest

	ANAEV			Variation Components			Genetic Parameters				
	Genotypes	Plot	Blocks	Genotypes	Plot	Blocks	h^2g	h^2mc	Acc	General Mean	CVe (%)
¹ G%	4.52*	-	1.63	11.93	-	-	0.42 +- 0.24	0.59	0.77	10.32	39.06
² PS (g m ⁻²)	26.42**	0.01	0.00	0.30	8.10 ⁴	0.06	0.83 +- 0.30	0.95	0.97	0.25	70.98
² ES (g m ⁻²)	7.14**	0.00	0.00	0.05	7.10 ⁴	0.08	0.37 +- 0.20	0.7	0.84	0.83	24.08
³ RT	11.86**	0.00	157.55**	871.77	0.50	115.13	0.88 +- 0.31	0.97	0.98	38.61	19.73
³ NSi	33.24**	22.98**	0.05	1070.14	388.28	973.40	0.44 +- 0.11	0.77	0.88	30.77	18.61
³ NSR	43.43**	0.21	0.10	33.18	1.34	51.74	0.38 +- 0.10	0.82	0.91	30.77	12.27
³ NR	235.46**	2667.90**	250000**	0.54	0.48	10 ⁻⁶	0.53 +- 0.12	0.69	0.83	4.42	15.68
³ LR	64.85**	5.08*	224.64**	5.32	0.47	3.17	0.59 +- 0.13	0.89	0.95	6.23	18.05
³ LI	32.02**	132.55**	366.99**	8.16	4.44	2.33	0.55 +- 0.12	0.76	0.87	11.34	19.77

¹Evaluations carried out in beds during the first harvest. ²Evaluations carried out in seed collectors during the second harvest. ³Evaluations carried out in beds during the second harvest. * Significant at the likelihood ratio test, considering 5% probability by the X² test. ** Significant at the likelihood ratio test, considering 1% probability by the X² test.

Table 3. Predicted genotypic values (BLUP) of 44 hybrids (Hib) and gain with selection (GS% at 30, 20 and 10% selection intensity) for the germination (G%) of *B. decumbens* hybrids in the first harvest

	G (%)					
	Treatments	BLUP (LL-UL)		Treatments	BLUP (LL-UL)	
1	A044	19.95 (15.55 - 24.35)		23	A015	9.95 (4.76 - 15.15)
2	A029	13.70 (9.30 - 18.10)		24	A032	9.95 (4.76 - 15.15)
3	A030	13.70 (9.30 - 18.10)		25	T038	9.95 (4.76 - 15.15)
4	A004	13.34 (8.15 - 18.53)		26	C001	9.83 (5.43 - 14.23)
5	A025	13.10 (8.70 - 17.50)		27	A009	9.57 (4.35 - 14.79)
6	A023	12.49 (7.30 - 17.69)		28	B026	9.57 (4.35 - 14.79)
7	A031	12.49 (7.30 - 17.69)		29	R184	9.57 (4.35 - 14.79)
8	A042	12.49 (7.30 - 17.69)		30	A033	9.53 (4.33 - 4.72)
9	A036	11.69 (6.47 - 16.90)		31	B009	9.53 (4.33 - 14.72)
10	X121	11.69 (6.47 - 16.90)		32	A035	9.24 (4.84 - 13.64)
11	A020	11.65 (6.45 - 16.84)		33	A012	9.15 (3.93 - 14.36)
12	A008	11.62 (7.22 - 16.02)		34	B005	9.10 (3.91 - 14.30)
13	A026	11.32 (6.92 - 15.72)		35	A017	8.34 (3.94 - 12.74)
14	S044	10.84 (5.62 - 16.06)		36	A024	8.30 (3.08 - 13.52)
15	A013	10.80 (5.60 - 15.99)		37	A041	8.30 (3.08 - 13.52)
16	A019	10.80 (5.60 - 15.99)		38	A007	8.26 (3.06 - 13.45)
17	A027	10.43 (6.03 - 14.83)		39	A002	7.75 (3.35 - 12.15)
18	A028	10.37 (5.18 - 15.57)		40	A005	7.41 (2.22 - 12.60)
19	B010	10.37 (5.18 - 15.57)		41	A011	7.41 (2.22 - 12.60)
20	A021	10.13 (5.73 - 14.53)		42	A018	7.41 (2.22 - 12.60)
21	A038	09.99 (4.78 - 15.21)		43	A043	6.86 (2.46 - 11.26)
22	A003	09.95 (4.76 - 15.15)		44	A001	6.14 (0.95 - 11.33)
GS%	30%	20%	10%			
	26	32	47			

LL: Lower limit of the withdrawal period; UL: Upper limit of the withdrawal period.

Heritability of individual plots (h^2g) takes into account the existence of only one genotype replication, while among genotypes means (h^2mc), the presence of replications assists in reducing the environmental effect (Resende and Duarte 2007), contributing to higher h^2mc estimates. Coefficient of variation (CVe) was considered as low for the characters, except for weight of pure seeds (PS), which was 70.98%

For evaluations carried out in beds, accuracy of the morphological characterization of the inflorescence ranged from 83 to 91%. It is considered satisfactory and is classified of high magnitude (Resende and Duarte 2007). Heritability of individual plots (h^2g) for number of seeds in the inflorescences (NSI) and number of seeds per racemes (NSR) did not exceed 0.50. Broad sense heritabilities (h^2mc) were 0.77 and 0.82, respectively.

For the number of reproductive tillers (RT), number of racemes (NR), length of raceme (LR) and length of inflorescences (LI), heritability of individual plots (h^2g) and broad sense heritability (h^2mc) were greater than 0, 50. Genotypic and plot effects were significant for all characters, except for number of seeds per raceme (NSR), which showed no difference in the plot effect. Estimates of heritability between genotypes means (h^2mc) showed high magnitude, exceeding 0.50 (Table 2). From the predicted breeding values (BLUP), it is possible to obtain the ranking of hybrids for each character, and to reliably identify hybrids with superior genotypic values (Resende 2006).

For evaluations during the first year, hybrid A044 was ranked in the first position for G%, differing from the other hybrids, when the confidence interval is observed. The range of BLUP values was high, from 19, 95 for the first ranked, to 6.14, for the last ranked; thus, variation was of 70%. Gain with selection (GS%), at 30%, 20% and 10% of selection intensity, in relation to the population mean, ranged from 26 to 47%. It should be noted that the higher the gain with selection in relation to the population mean, the greater the progress in the selection of superior genotypes (Table 3).

For evaluations carried out in seeds collectors during the second harvest, both for weight of pure seeds (PS) and for weight of empty seeds (ES), control cv. Marandu presented the highest genotypic values (BLUP). Low production of pure seed of the genotypes may be inherited by the next generation, since both the narrow-sense heritability (h^2g)

Table 4. Predicted genotypic values (BLUP), (GS% at 30, 20 and 10% selection intensity) regarding weight of pure seeds (PS) and weight of empty seeds (ES) in relation to cv. Marandu (Mar) for *B. decumbens* hybrids evaluated in seed collectors in the second harvest

	Treatments	PS (g m ⁻²)			Treatments	ES (g m ⁻²)		
		BLUP (LL-UL)				BLUP (LL-UL)		
1	Mar	2.38 (2.04 - 2.72)			Mar	1.17 (0.93 - 1.41)		
2	R087	0.76 (0.35 - 1.16)			R181	1.15 (0.90 - 1.39)		
3	R120	0.41 (0.08 - 0.75)			R091	1.02 (0.77 - 1.260)		
4	C001	0.30 (-0.03 - 0.64)			S036	1.00 (0.75 - 1.24)		
5	R126	0.30 (-0.04 - 0.64)			R120	0.94 (0.70 - 1.19)		
6	X030	0.20 (-0.13 - 0.54)			S031	0.93 (0.68 - 1.17)		
7	R124	0.09 (-0.25 - 0.43)			X030	0.92 (0.68 - 1.17)		
8	Y021	0.07 (-0.27 - 0.40)			X072	0.82 (0.58 - 1.07)		
9	R091	0.06 (-0.27 - 0.40)			X117	0.82 (0.58 - 1.07)		
10	B006	0.05 (-0.31 - 0.41)			R110	0.81 (0.56 - 1.05)		
11	R041	0.05 (-0.29 - 0.39)			B006	0.80 (0.53 - 1.06)		
12	T012	0.05 (-0.29 - 0.38)			R041	0.79 (0.55 - 1.04)		
13	X117	0.05 (-0.29 - 0.38)			R126	0.77 (0.53 - 1.02)		
14	R181	0.04 (-0.36 - 0.44)			T012	0.77 (0.52 - 1.01)		
15	S031	0.04 (-0.30 - 0.38)			T026	0.77 (0.52 - 1.01)		
16	S036	0.04 (-0.36 - 0.44)			C001	0.70 (0.46 - 0.94)		
17	R110	0.03 (-0.30 - 0.37)			R124	0.69 (0.45 - 0.93)		
18	T054	0.03 (-0.31 - 0.37)			Y021	0.58 (0.34 - 0.83)		
19	X072	0.03 (-0.31 - 0.36)			R087	0.56 (0.27 - 0.85)		
20	T026	0.02 (-0.31 - 0.36)			T054	0.50 (0.26 - 0.75)		
	GS (%)	30%	20%	10%	GS (%)	30%	20%	10%
		149	221	324		23	27	34

LL: Lower limit of the withdrawal period; UL: Upper limit of the withdrawal period.

Table 5. Predicted genotypic values (BLUP) and gain with selection (GS% in 30, 20 and 10% selection intensity) in relation to the number of reproductive tillers (RT), number of seeds per inflorescence (NSI), number of seeds per raceme (NSR) and number of racemes (NR) compared to the treatments (Treat) cv. Marandu (Mar) and *B. decumbens* hybrids evaluated in the beds in the second harvest

		RT			NSI			NSR			NR					
	Treat	BLUP (LL-UL)		Treat	BLUP (LL-UL)		Treat	BLUP (LL-UL)		Treat	BLUP (LL-UL)					
1	C001	140.84 (124.15 - 157.54)		R120	201.50 (174.53 - 228.48)		Mar	47.61 (43.28 - 51.94)		R120	6.12 (5.75 - 6.50)					
2	R091	71.41 (54.71 - 88.11)		Mar	189.63 (162.66 - 216.61)		T012	36.83 (32.50 - 41.17)		R126	5.30 (4.93 - 5.68)					
3	Y021	68.99 (52.29 - 85.69)		R181	160.27 (133.30 - 187.25)		S031	35.57 (31.24 - 39.90)		R181	5.30 (4.93 - 5.68)					
4	R124	49.63 (32.93 - 66.33)		R091	151.07 (124.10 - 178.05)		S036	33.50 (29.17 - 37.84)		R091	4.89 (4.52 - 5.27)					
5	Mar	36.57 (19.87 - 53.27)		T012	150.36 (123.39 - 177.34)		C001	32.75 (28.41 - 37.08)		T026	4.89 (4.52 - 5.27)					
6	B006	36.08 (19.39 - 52.78)		S036	148.95 (121.97 - 175.92)		R120	31.78 (27.45 - 36.11)		T054	4.89 (4.52 - 5.27)					
7	S031	33.91 (17.21 - 50.60)		R126	147.69 (120.72 - 174.67)		R091	30.70 (26.36 - 35.03)		R110	4.49 (4.11 - 4.86)					
8	S036	30.52 (13.82 - 47.22)		T026	140.34 (113.37 - 167.32)		R041	30.21 (25.88 - 34.54)		R041	4.49 (4.11 - 4.86)					
9	R041	29.79 (13.10 - 46.49)		R110	134.30 (107.32 - 161.27)		R181	29.93 (25.60 - 34.26)		S036	4.49 (4.11 - 4.86)					
10	T012	29.31 (12.61 - 46.01)		R041	134.08 (107.10 - 161.05)		R110	29.78 (25.45 - 34.12)		T012	4.49 (4.11 - 4.86)					
11	T054	28.58 (11.89 - 45.28)		S031	125.96 (98.99 - 152.94)		X072	29.69 (25.36 - 34.03)		R124	4.08 (3.70 - 4.45)					
12	X030	25.68 (8.98 - 42.38)		X072	120.46 (93.49 - 147.44)		B006	28.67 (24.34 - 33.00)		Mar	4.08 (3.70 - 4.45)					
13	X072	25.68 (8.98 - 42.38)		T054	119.48 (92.51 - 146.46)		T026	28.47 (24.14 - 32.81)		X072	4.08 (3.70 - 4.45)					
14	R126	23.99 (7.29 - 40.68)		C001	118.56 (91.58 - 145.53)		R126	27.77 (23.43 - 32.10)		B006	3.67 (3.29 - 4.04)					
15	T026	23.50 (6.81 - 40.20)		B006	104.99 (78.02 - 131.97)		X030	27.15 (22.82 - 31.48)		C001	3.67 (3.29 - 4.04)					
16	R120	22.78 (6.08 - 39.48)		X030	101.35 (74.37 - 128.32)		Y021	25.11 (20.78 - 29.44)		S031	3.67 (3.29 - 4.04)					
17	R110	21.81 (5.11 - 38.51)		R124	100.09 (73.12 - 127.07)		R124	24.24 (19.91 - 28.58)		X030	3.67 (3.29 - 4.04)					
18	R087	17.71 (-1.64 - 37.06)		Y021	081.41 (54.44 - 108.39)		T054	24.15 (19.82 - 28.48)		Y021	3.26 (2.88 - 3.63)					
19	R181	16.73 (0.03 - 16.00)														
	GS (%)	30%	20%	10%	GS (%)	30%	20%	10%	GS (%)	30%	20%	10%	GS (%)	30%	20%	10%
		73	114	174		23	26	37		14	16	22		19	22	29

LL: Lower limit of the withdrawal period; UL: Upper limit of the withdrawal period.

and the broad-sense heritability (h^2mc) were of high magnitude (Table 2). For PS, gains with selection (GS%) at 30, 20 and 10% of selection intensity were respectively 149, 221 and 324%; and for ES, it was 23, 27 and 34%, compared to population means (Table 4).

Weight of pure seeds (PS) of the hybrids was, in general, of low magnitude than that of control (Table 3). Three factors may have contributed to this: the high abortion rate in function of hybridization, as suggested by Lutts et al. (1991); the interference by spittlebugs infestation; and the possibility of being a characteristic of the genotype, producing only vegetative tillers, and only under extreme stress conditions it would produce reproductive tillers.

Non-viability of hybrid seeds may be caused by environmental interference, affecting the viability of pollen. Also, it could happen for the hybrid's genotype present gametophytic incompatibility allele, and thus blocking endosperm formation, and consequently seed filling.

Mateus et al. (2015) evaluated *B. decumbens* intraspecific hybrids, including R033, R126, R181, S036, T026 and X117, and also found high nymphal survival of spittlebugs, exceeding 60%. It is noteworthy that all hybrids tested in this study were considered susceptible or mid-resistant in previous works, with spittlebug nymphal survival ranging from 44 to 90%. For values above 40% nymphal survival, genotypes are not subjected to screening for genetic resistance.

For RT, cv. *B. brizantha* was ranked in the fifth position, differing from genotypes C001, R091 and Y021, taking into account the confidence interval. Gains with selection (GS%) at 30, 20 and 10% intensity were, respectively, 73, 114 and 174% (Table 5). The high potential for production of reproductive tillers has heritability of high magnitude of 0.88 and 0.97 in individual plots (h^2g) and in the broad-sense heritability (h^2mc), respectively. Comparing the BLUP of the morphological characters, cv. Marandu was ranked in the second, first and twelfth positions in number of seeds per inflorescence (NSI), number of seeds per raceme (NSR), and number of racemes (NN), respectively. However, the cultivar did not present better performance than R120 genotype for NSI. R120, R120, R126, R181, R091, T026, T054, R110, R041, S036, T012, and R124 genotypes showed higher BLUP in relation to the control, presenting gain in relation to the

Table 6 Predicted genotypic values (BLUP) and gain with (GS% at 30, 20 and 10% selection intensity) and length of racemes (LR) and length of inflorescence (LI) compared to cv. Marandu (Mar) *B. decumbens* for hybrids (Hib) evaluated in the beds in the second harvest

	LR			LI				
	Treatments	BLUP (LL-UL)			Treatments	BLUP (LL-UL)		
1	Mar	14.87 (13.38 - 16.36)			Mar	17.63 (15.24 - 20.02)		
2	R181	6.88 (5.39 - 8.37)			R120	14.63 (12.24 - 17.02)		
3	X030	6.57 (5.08 - 8.06)			T026	13.93 (11.55 - 16.32)		
4	S036	6.50 (5.01 - 7.99)			T054	13.77 (11.39 - 16.16)		
5	S031	6.18 (4.69 - 7.67)			R126	13.03 (10.65 - 15.42)		
6	C001	5.98 (4.49 - 7.47)			S036	11.77 (9.38 - 14.15)		
7	T026	5.89 (4.40 - 7.38)			X072	11.58 (9.19 - 13.96)		
8	T012	5.83 (4.34 - 7.32)			R041	11.57 (9.18 - 13.95)		
9	X072	5.83 (4.34 - 7.32)			R110	11.33 (8.95 - 13.72)		
10	R120	5.79 (4.30 - 7.28)			S031	11.14 (8.76 - 13.53)		
11	R041	5.67 (4.18 - 7.16)			R091	10.99 (8.60 - 13.38)		
12	T054	5.55 (4.06 - 7.04)			R124	10.42 (8.04 - 12.81)		
13	R110	5.48 (3.99 - 6.97)			T012	10.25 (7.86 - 12.64)		
14	B006	5.45 (3.96 - 6.94)			C001	10.18 (7.80 - 12.57)		
15	R124	5.45 (3.96 - 6.94)			B006	9.52 (7.13 - 11.90)		
16	R126	5.19 (3.70 - 6.68)			X030	8.09 (5.70 - 10.47)		
17	Y021	4.66 (3.17 - 6.15)			R181	7.88 (5.50 - 10.27)		
18	R091	4.42 (2.93 - 5.91)			Y021	6.38 (3.99 - 8.76)		
	GS (%)	30%	20%	10%	GS (%)	30%	20%	10%
		12	14	17		22	26	30

LL: Lower limit of the withdrawal period; UL: Upper limit of the withdrawal period.

number of racemes (NR). R120 genotype stood out both for number of seeds per inflorescence (NSI) and number of racemes (NR), taking the first position in the two variables of seed production. This fact may be a promising indicative for its production potential. Although the genotypes present genotypic values of high magnitude when compared with the control in NSI and NR, they had heritability in individual plots (h^2g) inferior to 0.50.

Cultivar Marandu, the control, was ranked in the first position for both length of racemes (LR) and length of inflorescences (LI). However, it appears that a character does not influence the other, since R120 genotype takes the second position in length of inflorescences, and its racemes have the tenth shorter length (Table 6).

CONCLUSIONS

The number of reproductive tillers presents high heritability both in the broad sense (h^2mc) and in individual plots (h^2g), and can be a parameter to predict seed production potential. However, it is necessary further studies on which components really influence seed production of forage plants.

The genotypes that had better performance for production of reproductive tillers (RT) and weight of pure seeds (PS) were C001 and R091.

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