Conditioning and coating of *Urochloa brizantha* seeds associated with inoculation of *Bacillus subtilis*¹

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INTRODUCTION

In Brazil, the area cultivated with forages occupies about 150 million hectares (IBGE 2017), where the genus *Urochloa* spp. (*Brachiaria*) is present. However, agricultural practices for improving the quantity and quality of forages are necessary to increase the animal productivity (Silva et al. 2008). Thus, grasses seeds should exhibit a high germination, what results in a vigorous seedling growth, and, in turn, this provides a suitable soil cover and inhibits weed species.

Adding value to *Urochloa* spp. seeds by using technology for seed processing, such as seed coating and conditioning, can provide a unique selling point in an increasingly competitive market. Seed coating is the deposition of inert and adhesive materials on the seed surface, with or without modification of

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the seed shape and size (Silva et al. 2002), while in conditioning seeds are soaked with a chemical or biological solution for a certain period of time (Batista et al. 2016). Both techniques may be used to introduce chemical or biological materials that can benefit seeds (Santos et al. 2010). The use of coated or conditioned seeds facilitates sowing by increasing the sanity of the seeds, what, in turn, leads to an improved seedling establishment, allowing the incorporation of nutrients, growth regulators and other inputs (Silva et al. 2002, Bonome et al. 2017). However, Santos et al. (2011) state that techniques such as seed coating, despite being a method used to increase the pasture productivity, may cause inhibition in the germination of Urochloa brizantha.

The plant growth-promoting rhizobacteria that inhabit the soil are isolated from the rhizosphere of several cultivated plants. Among the most studied genera are Bacillus, Pseudomonas, Azospirillum and Rhizobium. The effects of these microorganisms on the plant development are broad and include beneficial effects on seed germination, seedling emergence and plant growth (Figueiredo et al. 2010).

There are few studies about the inoculation of Brachiaria seeds with plant growth-promoting rhizobacteria. For instance, Araujo et al. (2012a) concluded that the inoculation of Bacillus sp. in seeds of U. brizantha promotes the growth of plants. In a study carried out with other grasses, the production of hormones by rhizobacteria changed the root morphology and increased the biomass, thus increasing the soil exploration capacity (Malik et al. 1997). Araujo (2008) evaluated the inoculation of Bacillus subtilis in different plant species and showed that the inoculation significantly increased the nitrogen content in maize leaf tissue, even though the introduced bacterial species is not known as diazotrophic. In addition to effects on plant growth, other benefits related to the inoculation of B. subtilis may be found in disease management, such as nematode control (Morgado et al. 2015).

When inoculating microorganisms in seeds, it is important to ensure that they have enough bacteria for a greater process efficiency. Following this, a study was conducted, wherein several techniques for bacterial inoculation in seeds were evaluated, such as the use of suspensions, pastes, peat or encapsulation (Walker et al. 2004). Seed coating or conditioning techniques can ensure the highest concentrations of bacteria in the seed. Applying beneficial microorganisms to the seed through either conditioning or coating may further enhance the establishment of the crop, particularly if the applied microorganisms settle in the rhizosphere, and subsequently contribute to the long-term plant health or to the promotion of plant growth (Bennett & Whipps 2008).

Seed storage is a very important stage in the seed production process and, in the case of U. brizantha, other aspects should also be explored, such as dormancy and seed viability (Novembre et al. 2006). In addition, it is necessary to evaluate if the introduction of chemical and biological agents may affect the germination and emergence speed of the U. brizantha seedlings (Santos et al. 2010).

Thus, this study aimed to evaluate the effect of the inoculation of B. subtilis on U. brizantha seeds using either coating or osmotic conditioning, as well as the effects of the inoculation on germination, plant development and seed storage.

MATERIAL AND METHODS

The experiments were conducted from September 2016 to June 2017. Non-scarified seeds of U. brizantha (Hochst ex A. Rich.) Stapf. ‘Marandu’ were used from two lots (1 and 2) with lower and higher viability, respectively, based on a tetrazolium test carried out previously (Novembre 2006). Two strains of B. subtilis (PRBS-1 and AP-3) isolated from an agricultural soil and described in Araujo et al. (2005) were used in this study.

For the seed treatment, initially, B. subtilis was multiplied in a culture medium for five days. Afterwards, the cells were separated by centrifugation (5,000 × g) and resuspended in water. All suspensions were adjusted to 10⁸ colony forming units (CFU) per mL. Initially, the seed coating was carried out using 200 g of U. brizantha seeds and 50 mL of bacterial cell suspension with 0.75 g of methyl cellulose, which were mixed in a rotary drum. During mixing, 200 g of inert talc were added to impart the uniformity in the coating. To perform the osmotic conditioning of the seeds, 200 g of seeds were packed in an Erlenmeyer flask and immersed with 50 mL of bacterial cell suspension with 0.425 g of sodium chloride, which were mixed in a rotary drum. During mixing, 200 g of inert talc were added to impart the uniformity in the coating. To perform the osmotic conditioning of the seeds, 200 g of seeds were packed in an Erlenmeyer flask and immersed with 50 mL of bacterial cell suspension with 0.425 g of sodium chloride. The vials were stirred (120 rpm) for 12 h, at 25 °C. Finally, the seeds were dried for 24 h, at 30 °C. In parallel, treatments with seed coating and conditioning were conducted using sterilized water, instead of bacterial...
suspending, and these were used as control. All treated seeds were stored under a temperature of 15 °C, for 120 days.

The moisture in the seed environment was controlled, and 15% of humidity for conditioning and 12% for coating (Brasil 2009) were maintained. Seeds were collected every 30 days for germination evaluation.

The experiment consisted of a $2 \times 3 \times 5 + 1$ factorial scheme, represented by two seed conditions (coated and conditioned), three inoculants ($B.\ subtilis$ AP-3, $B.\ subtilis$ PRBS-1 and without bacteria), five storage periods (0, 30, 60, 90 and 120 days) and control (bare seeds), in a completely randomized design, with four replicates.

The variables seed germination percentage and germination rate index were evaluated. In the laboratory, for the germination tests, transparent plastic boxes (gerbox type) were used to house four replicates of 100 seeds on blotter paper (moistened with water 2.5 times the weight of the paper). Afterwards, the boxes were transferred to a germination chamber (BOD). Plants were regulated with alternating light and dark periods (8 h and 16 h, respectively) and alternating temperatures of 15 °C and 35 °C. The number of germinated seeds, as well as the number of normal seedlings, were determined for 21 days with daily ratings and used to calculate the germination percentage and the germination rate index (Brasil 2009).

An experiment was also conducted in a greenhouse, for 180 days, to evaluate the $U.\ brizantha$ ‘Marandu’ growth, using the seeds after the treatments previously described, prior to storage. The experimental design was a $2 \times 3 \times 5 + 1$ factorial scheme, represented by two seed conditions (coated and conditioned) and three treatments ($B.\ subtilis$ AP-3, $B.\ subtilis$ PRBS-1 and without bacteria), with five replications.

To fill the pots, 4 kg of a Dystrofic Red Argisol were collected at the 0-20 cm layer. The soil field capacity was determined in the laboratory. Ten seeds per pot were used for sowing. After the emergence of the plants, thinning was done, leaving two plants per pot. The pots were irrigated daily to maintain the soil moisture near the field capacity, using the gravimetric method. The experimental period started with the standardization cut (42 days after sowing).

At the end of the experiment (180 days after sowing), the shoot dry mass was obtained after the final cutting of the plants. Roots were also collected and washed, and both were oven dried at 65 °C, to obtain the dry mass.

Analyses of variance and F-tests at 5% of significance were used to estimate and test the effects of the treatments. The means of the treatments were compared by the Tukey test (5%), and the Dunnett test (5%) was used to compare each one of the treated levels with the control. Polynomial regression was also used for analyzing the response of the treatments over the storage time. All analyses were performed in the Assistat software (Silva 2011).

RESULTS AND DISCUSSION

The evaluation of the seeds of lower viability (lot 1) allowed verifying that the root growth with the conditioning of the seeds seemed promising for dry mass gains, except for those inoculated with $B.\ subtilis$ (PRBS-1). For the shoot dry mass production of $U.\ brizantha$, the use of seed coating, both with and without bacteria, was superior to the bare seeds (control). When the seed conditioning method was used, only those with bacterial inoculation showed gains in this aspect (Table 1).

In the seeds of greater viability (lot 2), it was verified that the conditioning of seeds combined with the use of the $B.\ subtilis$ strain AP-3 was the only treatment that increased the mass of the root system of $U.\ brizantha$, in relation to the control (Table 1).

Table 1. Means1 of root and shoot dry mass of $Urochloa\ brizantha$ (lots 1 and 2) submitted to seed treatments and inoculation with $Bacillus\ subtilis$ (AP-3 and PRBS-1).

<table>
<thead>
<tr>
<th>Lot</th>
<th>Treatments</th>
<th>AP-3</th>
<th>PRBS-1</th>
<th>Uninoculated</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Conditioning</td>
<td>28.62*</td>
<td>24.30</td>
<td>28.4*</td>
</tr>
<tr>
<td></td>
<td>Coating</td>
<td>25.54</td>
<td>22.51</td>
<td>25.6</td>
</tr>
<tr>
<td>2</td>
<td>Conditioning</td>
<td>34.42 aA</td>
<td>30.04 aB</td>
<td>30.30 aB</td>
</tr>
<tr>
<td></td>
<td>Coating</td>
<td>33.91 aA</td>
<td>30.56 aA</td>
<td>32.53 A</td>
</tr>
</tbody>
</table>

1Means followed by the same uppercase letter in the rows and lowercase letter in the columns do not differ by the Tukey test at 5% of significance. Means followed by (+) or (-) are respectively higher or lower than the absolute control by the Dunnett test (p < 0.05).
In the shoot mass production, the inoculation of *B. subtilis* strain PRBS-1 provided a significant increase in the root mass, using both the seed coating and conditioning.

The coating and conditioning techniques for *Urochloa* are promising for use in seed processing. In this study, it was observed that, in the less viable lot of seeds, the use of coating and osmotic conditioning techniques with the inoculation of rhizobacteria influenced the growth promotion of the plant shoot, when compared to the bare seeds. Wright et al. (2003) concluded that the simple seed conditioning increases the concentration of bacteria in the seeds, even without inoculation and even after drying beet seeds. Although being a study with a different species, it may suggest that this was the most likely biological phenomenon that happened to explain the growth promotion. This reinforces that the technique is useful for increasing the presence of microorganisms on untreated seeds. Moreover, in this study, it was verified that conditioning and coating, with or without bacterial inoculation, led to significant increases of the biomass of the plant root and shoot in the lot of lower viability, when compared to the bare seeds. This method may be the best technique to treat seeds if they exhibit either a low germination or seed dormancy (Novembre et al. 2006).

The use of seed coatings with rhizobacteria may be considered as a technology with great potential for commercial use (Walker et al. 2004). Studies on beans have shown that coating or conditioning with *B. subtilis* may increase the shoot dry mass production (Junges et al. 2016). The use of *Bacillus* spp. in conditioning cucumber seeds promoted a greater disease control in young plants and increased the final crop yield (Song et al. 2017). However, studies involving the use of coating or conditioning in forage seeds are still scarce. The association of rhizobacteria with the techniques of coating or conditioning on *Brachiaria* seeds has shown to be promising, especially in the promotion of shoot mass in plants of higher viability inoculated with *B. subtilis* (PRBS-1). However, the effect of inoculation of *B. subtilis* (AP-3) via conditioning on root growth in this lot can also be highlighted.

In the germination evaluation during the storage period, a lower germination was observed in low viability seeds (lot 1), when compared to the lot 2 (Figure 1). Coating treatments performed on the seeds of the lot 1 did not promote significant changes for significant regression adjustments over the storage time, what also did not happen with the evaluation of the bare seeds (control). It is observed that, in this lot, germination generally increases at 60-90 days and decreases at 120 days. Analyzing the performance of the treatments, it can also be highlighted that conditioning presented values closer to quadratic adjustments, represented in a significant way by the evaluation of the treatment without the bacteria inoculation.

On the other hand, when analyzing the lot of greater viability, it was observed that the bare seeds presented a significant linear adjustment increase for the germination values. Treatments without inoculation presented quadratic adjustments and maximum germination points close to 60 days, whereas the remaining ones had no significant adjustments.

The germination rate index also showed results with similar adjustments to the germination using regression analysis in both lots (Figure 2).

One of the concerns in the use of seed coating or osmotic conditioning is the physiological disturbances that these techniques can promote over the seed storage time (Santos et al. 2010). These authors concluded that seed coating promotes a reduction in the germination and seedling emergence speed of *U. brizantha* ‘Marandu’ after 12 months of storage. After the evaluation of the two lots in this study, it was determined that these parameters were reduced after 120 days of storage in the treated seeds from the lot of higher viability (Figure 2). The performances of both the coated and conditioned seeds were also observed during the germination of *U. brizantha*, and it was found that conditioned seeds were better for up to 90 days of storage. In conditioning tobacco seeds, the absence of negative effects on germination during 90 days of evaluation was also verified (Caldeira et al. 2014).

Santos et al. (2010) concluded, in a long-term *Urochloa* seed coat study, that this technique promotes a reduction in the seed germination and germination speed index. However, this technique is important because of the other benefits of this practice, such as facilitating sowing and reducing the incidence of phytopathogens. Most likely the seed dormancy was broken with the storage period, what may be expected with *Urochloa*, to which aging decreases the seed vigor. In terms of long-term germination loss, the conditioning of *Brachiaria*
Conditioning and coating of *Urochloa brizantha* seeds associated with inoculation of *Bacillus subtilis* AP-3 and PRBS-1. A: conditioned seeds/lot 1; B: conditioned seeds/lot 2; C: coated seeds/lot 1; D: coated seeds/lot 2.

Figure 1. Germination percentage of *Urochloa brizantha* submitted to seed treatments and inoculation with *Bacillus subtilis* AP-3 and PRBS-1. A: conditioned seeds/lot 1; B: conditioned seeds/lot 2; C: coated seeds/lot 1; D: coated seeds/lot 2.

Figure 2. Germination rate index of *Urochloa brizantha* (lot 2) submitted to seed treatments and inoculation with *Bacillus subtilis* AP-3 and PRBS-1. A and C: conditioned seeds; B and D: coated seeds.
seeds showed more promising results than coating for up to 90 days of storage (Figure 1). In order to evaluate the effect of rhizobacteria on bean seeds, El-Mougy & Abdel-Kader (2008) found no loss of seed germination, but observed a loss of rhizobacteria as pathogen antagonists at 90 days.

The presence of rhizobacteria in the rhizosphere of plants can provide numerous benefits to plants and soil. *Urochloa* seeds, in both the evaluated lots, were not affected by the presence of bacteria in the first 90 days, and this may stimulate further studies to evaluate the use of this grass to introduce rhizobacteria like *B. subtilis* in the soil. The introduction of these microorganisms can provide innumerable benefits to the production system, since rhizobacteria are cited as efficient antagonists of pathogens such as fungi (Araujo et al. 2005, Yánez-Mendizábal et al. 2012, Gao et al. 2014). They also provide other indirect benefits such as nutrient solubilization in the soil and reduction of damage caused by abiotic stresses (Ferreira et al. 2018). Bacteria of the genus *Bacillus* can promote plant growth when inoculated previously on *Urochloa* seeds (Araujo et al. 2012a). Pastures may also benefit from the inoculation of *B. subtilis*, as rhizobacteria may confer a greater stress on plants (Gagné-Bourque et al. 2016).

The use of rhizobacteria such as *B. subtilis* may contribute to the sustainability of the system, since these can make the soil suppressive to several pathogens, what may result in an increase in their population over time (Lazzaretti & Bettiol 1997). This bacterial species also has the ability to colonize the roots by forming a thin biofilm layer on them, which then confers benefits to the plant (Beauregard et al. 2013). The treatment of seeds with *B. gaemokensi* to elicit resistance enables the plants to effectively mount long-term resistance responses against plant pathogens and insects (Song et al. 2017).

The efficient colonization of the root system allows a more favorable competitive situation against other species of microorganisms (Kilian et al. 2000). Verma et al. (2018) showed that endophytic microorganisms like *B. subtilis* are responsible for the modulation of the *Urochloa ramosa* growth and the protection of plants against fungal diseases. However, the use of rhizobacteria does not always provide the greatest plant growth (Corrêa et al. 2010): some may be indifferent to the inoculation process, while others may consider the bacteria as pathogens, thus impairing their growth.

Forages are permanent crops, thus requiring a great root development, which aids in the seeking of water and nutrients from the deep layers of the soil and promotes the cycling of nutrients (Santos et al. 2007). Consequently, the use of techniques that allow a greater development of these plants is very important, as plants with a greater root volume have higher production conditions, even in adverse climatic conditions. In addition, a root system with broad growth can improve the soil quality, especially when it comes to the structure and accumulation of organic matter (Salton et al. 2004). Seed coating has been the most widely used enhancement in the forage seed trade, as evidenced by the amount of commercial investment to these processes (Simon et al. 1998).

The results showed potential applications of the techniques of conditioning and coating seeds, with or without bacteria, in the different stages of the forage seed processing. Bonome et al. (2017) reported that coating and conditioning on forage (*U. brizantha*) can also improve sowing operations and serve to introduce other inputs, such as biological agents, that can then associate with the seeds. Most studies on seed conditioning of *Urochloa* has as a main objective the dormancy breaking to obtain a higher germination pattern and vigor and are evaluated in short storage periods (Ribeiro et al. 2019). Based on the results, it can be stated that seed conditioning and coating with *B. subtilis* kept the germination profile similar to the bare seeds for about 90 days, considering the seeds of the most viable lot. These results may stimulate further studies with inoculation of other rhizobacteria using coating or conditioning techniques similar to those in the present study. More studies are needed to explore the ability of the introduced microorganisms to colonize the rhizosphere, as well as tests using other forage species and tests involving longer storage times.

**CONCLUSIONS**

1. Seed physiological conditioning shows a better performance when inoculating *Urochloa brizantha* seeds with *Bacillus subtilis*;
2. Coating and conditioning *U. brizantha* seeds with less viability, both with and without bacteria inoculation, increases the shoot mass in the plants after 180 days of growth;
3. The use of conditioning or coating reduces the germination after 90 days of storage in seeds of greater viability.
REFERENCES


