

Selection of somaclonal variants of the cultivar ‘Prata-Anã’ for resistance to *Fusarium oxysporum* f. sp. *ubense* race 1

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Abstract -The banana tree is one of the most cultivated fruit globally; however, some diseases significantly affect its production, such as Fusarium wilt. The most appropriate measure for controlling this disease in areas with inoculum pressure is the use of resistant cultivars. Therefore, this study aimed to generate banana somaclones of the cultivar ‘Prata-Anã’ resistant to Fusarium wilt by inducing somaclonal variation. ‘Prata-Anã’ stem apexes were established *in vitro* in MS culture medium and, on a monthly basis, subcultivated in AIA and adenine sulfate supplemented MS medium with added plant regulators: 6-benzylaminopurine (BAP, 4 ml L⁻¹), Thidiazuron (TDZ, 1 ml L⁻¹), and Paclobutrazol (PBZ, 10 ml L⁻¹). The treatments were: T0: no regulator, T1: BAP, T2: TDZ, T3: PBZ, T4: BAP + TDZ, T5: BAP + PBZ, T6: TDZ + PBZ, and T7: BAP + TDZ + PBZ. After the twelfth subculture, the regenerated plants were planted in boxes containing sterile soil infected with *Fusarium oxysporum* f. sp. *ubense*, and evaluated after 90 days for resistance to the pathogen. Somaclonal variants T2-1 and T2-2, generated in Treatment 2, with TDZ, were selected as resistant. This result is promising for the launch of a new Fusarium race 1-resistant banana variety.

Index terms: Biotechnology; Tissue culture; *Musa* sp.; Somaclonal variations; Fusarium wilt.

Seleção de variantes somaclonais da cultivar ‘Prata-Anã’ para resistência à *Fusarium oxysporum* f. sp. *ubense* raça 1

Resumo -A bananeira é uma das frutíferas mais cultivadas no mundo; entretanto, algumas doenças têm afetado sua produção, como a murcha de Fusarium. Em áreas onde há pressão de inóculo, a medida mais adequada no controle da doença é a utilização de cultivares resistentes. Diante disso, o objetivo deste trabalho foi obter somaclones de bananeira da cultivar ‘Prata-Anã’ resistentes à murcha de Fusarium pela indução de variação somaclonal. Para isso, ápices caulinares de bananeira ‘Prata-Anã’ foram estabelecidos *in vitro*, em meio de cultura MS. Em seguida, foram transferidos e subcultivados mensalmente para meio MS, suplementado com AIA e sulfato de adenina, acrescidos de diferentes combinações de reguladores: 6-benzilaminopurina BAP (4 ml.L⁻¹), Thidiazuron TDZ (1 ml.L⁻¹) e Paclobutrazol PBZ (10 ml.L⁻¹). Os tratamentos foram: T0: sem regulador; T1: BAP; T2: TDZ; T3: PBZ; T4: BAP + TDZ; T5: BAP + PBZ; T6: TDZ + PBZ; 7: BAP + TDZ + PBZ. As plantas regeneradas, após o décimo segundo subcultivo, foram posteriormente plantadas em caixas d’água contendo solo estéril e infestado com *Foc*, e aos 90 dias foram avaliadas quanto à resistência ao patógeno. Foram selecionados os variantes somaclonais T2-1 e T2-2, resultantes do tratamento 2 com TDZ, como resistentes. Esse resultado é promissor para o lançamento de uma nova variedade de bananeira resistente à murcha de Fusarium raça 1.

Termos para indexação: Biotecnologia; Cultura de tecidos; *Musa* sp; Variações somaclonais; Murcha de Fusarium.

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Received: March 13, 2020
Accepted: August 20, 2020

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Introduction

Bananas are a source of nutrients and income for thousands of families in several countries around the world, such as India, China, Indonesia, Brazil, and some countries of Africa. Brazil is the fourth largest banana producer, accounting for approximately seven million tons of fruit (FAOSTAT, 2018). Prata subgroup cultivars, such as 'Prata-Anã', and derivative selections such as 'Gorutuba' and 'Catarina', form the basis of Brazilian production, representing approximately 70% of the area in the country cultivated with this fruit (IBGE, 2016; COLTRO; KARASKI, 2019). Banana production in Brazil is characterized by small producers and a domestic consumption of 65 to 70% of total production, demonstrating its social importance in terms of food security, job creation, and subsistence (COLTRO; KARASKI, 2019).

Both the production of bananas for subsistence, and for the local and export fruit trade are threatened by the occurrence of Fusarium wilt, caused by the fungus *Fusarium oxysporum* f. sp. *cubense* (*Foc*) (PLOETZ, 2015; DITA et al., 2018). In addition to concerns about managing the pandemic caused by tropical race 4 (TR4) of the pathogen, some countries are searching for strategies to cope with race 1 which is endemic in the main banana producing regions of the world (ARINAITWE et al., 2019; GONÇALVEZ et al., 2019). In Brazil, there are no records of TR4; however, *Foc* race 1 has had a great impact, either by reducing the productivity of banana plantations or by making the soil unsuitable for planting susceptible cultivars, especially in fields under irrigation which constitute a large proportion of banana production areas (REBOUÇAS et al., 2018; GONÇALVEZ et al., 2019).

The fungus *Foc* is a soil-dwelling pathogen that penetrates the roots of the banana tree and colonizes the tissue, affecting xylem vessels and leading to symptoms such as progressive leaf yellowing, pseudostem cracks, leaf wilt, and plant death (DITA et al., 2018). In addition, the pathogen produces resistant structures called chlamydospores which enable it to survive in the soil for many years (PLOETZ, 2015). Cultural and biological disease control practices are not efficient for controlling *Foc*, especially with high inoculum pressure in very susceptible cultivars, but have shown promising results when associated with less susceptible cultivars with quantitative resistance (HADDAD et al., 2018). So far, the results clearly indicate that the most viable disease management strategy is the development of resistant cultivars (REBOUÇAS et al., 2018; ARINAITWE et al., 2019; GONÇALVEZ et al., 2019). However, most cultivars of the genus *Musa* have high levels of sterility and low seed production, which makes conventional breeding a significant challenge, despite the development of hybrids resistant to *Foc* race 1 by some breeding

programs, such as the Honduran Agricultural Research Foundation (FHIA), and the Brazilian Agricultural Research Corporation (Embrapa) (AMORIM et al., 2011; KHAYAT et al., 2011).

The exploration of somaclonal variations that occur during the micropropagation process is an alternative strategy to progeny breeding and selection programs (CHEN et al., 2013; GHAG et al., 2014). This approach is related to the occurrence of genetic changes induced by *in vitro* cultivation, which can cause genetic or epigenetic variation (PENNA et al., 2019; ANIL et al., 2018). A clear demonstration of success with somaclonal variants resistant to TR4 occurred in Taiwan using the cultivar 'Giant Cavendish' (HWANG; KO, 2004). Another study, using somatic embryos kept in tissue culture for an extended period to generate somaclones which were then challenged with *Foc* race 1, found that four of the 26 banana cv. 'Rasthali' somaclones presented greater *Foc* resistance. In addition, cDNA-RAPD analysis identified at least one differentially expressed gene (lipoxygenase) in these somaclones (GHAG et al., 2014).

In the present study, 'Prata-Anã' stem apices were subcultured in culture medium with different growth regulator combinations and successive cultivations to generate and select somaclonal variants resistant to *Foc* race 1.

Material and methods

Plant material, tissue culture and plant regeneration

Banana plants derived from cv. 'Prata-Anã' (*Musa* spp. AAB) were obtained from the Embrapa Mandioca e Fruticultura. The stem apices were disinfected and established *in vitro* in a culture medium containing MS (Murashige & Skoog, 1962) salts and vitamins, and cultured for 15 days in the dark and 15 days in a growth room under a 16-hour photoperiod and photon flux density of $30 \mu\text{E m}^{-2} \text{s}^{-1}$ at $27 \pm 2 \text{ }^\circ\text{C}$. Thereafter, on a monthly basis, they were transferred and subcultured in MS medium supplemented with AIA and adenine sulfate to which was added different combinations of the plant regulators: 6-benzylaminopurine (BAP, 4 ml.L⁻¹), Thidiazuron (TDZ, 1 ml L⁻¹), and Paclobutrazol (PBZ, 10 ml L⁻¹), as follows: Treatment 1 (T1) BAP, Treatment 2 (T2) TDZ, Treatment 3 (T3) PBZ, Treatment 4 (T4) BAP + TDZ, Treatment 5 (T5) BAP + PBZ, Treatment 6 (T6) TDZ + PBZ, and Treatment 7 (T7) BAP + TDZ + PBZ (Table 1).

Table 1. Plant regulators used in the micropropagation process to generate 'Prata-Anã' somaclonal variants resistant to *Fusarium* wilt race 1.

Sign	Treatments
T0	Without regulator
T1	6-benzilaminopurina BAP (4 ml.L ⁻¹)
T2	Thidiazuron TDZ (1 ml.L ⁻¹)
T3	Paclobutrazol PBZ (10 ml.L ⁻¹)
T4	6-benzilaminopurina BAP (4 ml.L ⁻¹) + Thidiazuron TDZ (1 ml.L ⁻¹)
T5	6-benzilaminopurina BAP (4 ml.L ⁻¹) + Paclobutrazol PBZ (10 ml.L ⁻¹)
T6	Thidiazuron TDZ (1 ml.L ⁻¹) + Paclobutrazol PBZ (10 ml.L ⁻¹)
T7	6-benzilaminopurina BAP (4 ml.L ⁻¹) + Thidiazuron TDZ (1 ml.L ⁻¹) + Paclobutrazol PBZ (10 ml.L ⁻¹)

A total of 12 subcultures were conducted at 30- to 40-day intervals. At the end of the subculture process, the sprouts were individualized and converted into plants, then transferred to MS medium without regulator for rooting. After 30 days, the rooted plants were transferred to tubes containing substrate and acclimated in a greenhouse for 30 to 45 days, until they reached a height of approximately 15 cm.

Fungus culture and inoculum preparation

This study used the CNPMF 218A isolate from *Foc* race 1, obtained from the biological collection of *F. oxysporum* f. sp. *cubense* from the Embrapa Laboratory of Phytopathology. This isolate was selected for its virulence and aggressive infection of banana cultivars (HADDAD et al., 2011; COSTA et al., 2015; ROCHA et al., 2020).

The *Foc* 1 isolate was grown on potato dextrose agar at 25 °C under a 12-hour photoperiod. After colony growth, a suspension of conidia was prepared and approximately 20 mL of this deposited in 1 kg of sterilized rice. Subsequently, the culture medium was incubated at 25 °C under a 12-hour photoperiod. After 20 days, colony-forming units (CFU) were counted using a serial dilution to verify the spore concentration and viability. The CFU were counted using a Neubauer camera. The concentration used for inoculations was 10⁶ CFU g⁻¹ of substrate (MADIGAN et al., 2016).

Preliminary selection of *Fusarium* wilt resistant somaclones

The experiment was conducted in a greenhouse using 310 L polyethylene boxes filled with autoclaved soil which was subsequently infected with 1 kg of the *Foc* inoculum. 'Prata-Anã' banana plants, approximately 15 cm high, were planted in the *Foc* infected soil at a density of 30 seedlings per box. Non-treated 'Prata-Anã' seedlings

were planted in each box as a positive control. A total of 2,520 plants subjected to seven treatments (Table 1) were planted, with 360 plants evaluated per treatment.

After 90 days or plant death, internal symptoms of rhizome discoloration were evaluated using the grading scale (ranging from 1 to 5) proposed by Dita et al. (2014), where 1: absence of symptoms, 2: rhizome with initial discoloration, 3: rhizome discoloration throughout the vascular system, 4: rhizome with most internal tissues presenting necrosis, and 5: fully necrotic rhizome.

Plants with no symptoms of disease or with the lowest disease severity score were designated as putative *Fusarium* wilt race 1 resistant somaclones. These seedlings were used as the source material for further tissue culture, where they were introduced *in vitro* and micro-propagated for a second experiment to verify their *Foc* resistance.

Confirmation of resistance in selected somaclones

The micro-propagated seedlings of the somaclones supposedly resistant to *Fusarium* wilt (identified in the preliminary selection) were replanted in the infected soil, in a completely randomized design with ten replicates for each somaclone.

In this second experiment, the plants were evaluated for external leaf yellowing symptoms using the grading scale of Dita et al. (2014), where 1: no symptoms, 2: initial yellowing in old leaves, 3: yellowing in old leaves and initial discoloration in new leaves, 4: intense yellowing in all leaves, and 5: plant death. The symptoms were recorded from onset up to 60 days, at seven-day intervals. External leaf symptom indexes were determined from the scores and used to calculate the area under the disease progress curve (AUDPC) using the formula proposed by Shaner and Finney (1977).

Internal symptoms were analyzed using transverse rhizome sections of the plants 60 days after inoculation or after plant death, applying the same internal symptom grading scale as used in the preliminary experiment (DITA et al., 2014). The scores obtained were used to calculate internal indexes ID (%) values of *Fusarium* wilt symptoms.

Evaluation of fungal structures in the root tissue of the somaclones tested for *Fusarium* wilt resistance

Histological evaluation was performed for the putative somaclones identified in the preliminary test and for those used in the resistance confirmation experiment. In both cases, root fragments were collected after the final internal symptom evaluation during which the rhizomes were sectioned. The Phillips and Haymann (1970) protocol was used for root staining. The root fragments were soaked for 1 h in a 10% potassium hydroxide (KOH) solution in a water bath at 90 °C. Thereafter, the solution was discarded and the biological material washed in water to remove all the KOH. The roots were then transferred to a 1% HCl solution for a 5 min period. The HCl solution was discarded and the roots were stained using Trypan Blue in a 0.05% lactoglycerol solution (2:1:1, lactic acid: glycerin: water) and boiled in a microwave for 40 sec after which the dye was discarded. The roots were then immersed in lactoglycerol solution (2:1:1; lactic acid: glycerin: water) to remove excess dye. The root fragments were visualized and microphotographed under a light microscope.

Statistical analysis

Regarding the preliminary test internal symptom evaluation scores, the frequency of each score per plant was calculated and converted to a percentage. To study the treatments, a cluster analysis using color was also performed, based on a heat map prepared by transforming the internal symptom scores into a binary matrix, where 0 was assigned to plants without symptoms and 1 to plants with disease symptoms. The data for each plant was represented by a color, where green was associated with resistance and red with susceptibility. The ID symptoms and AUDPC data obtained from the second experiment were subjected to analysis of variance, and the means were grouped by the Scott-Knott test at 5% significance. The analyses were performed using the R software.

Results and discussion

A total of 360 ‘Prata-Anã’ banana seedlings without treatment (susceptible control) and 2,520 somaclonal variants from 12 subcultures with different plant regulator combinations were analyzed for *Foc* race 1 resistance in the preliminary test. Most control plants died within 60 days of planting in *Foc*-infected soil, and only 0.079% of the treatment plants were symptomless, which amounted to 20 plants out of the 2,520 somaclonal variants. The percentage of plants in each treatment (n = 360) with a score of 1 on the internal symptom scale was 2.7% in T2 (TDZ), 0.5% in T4 (BAP + TDZ), 0.8% in T5 (BAP + PBZ), and 1.3% in T6 (TDZ + PBZ) (Figure 1).

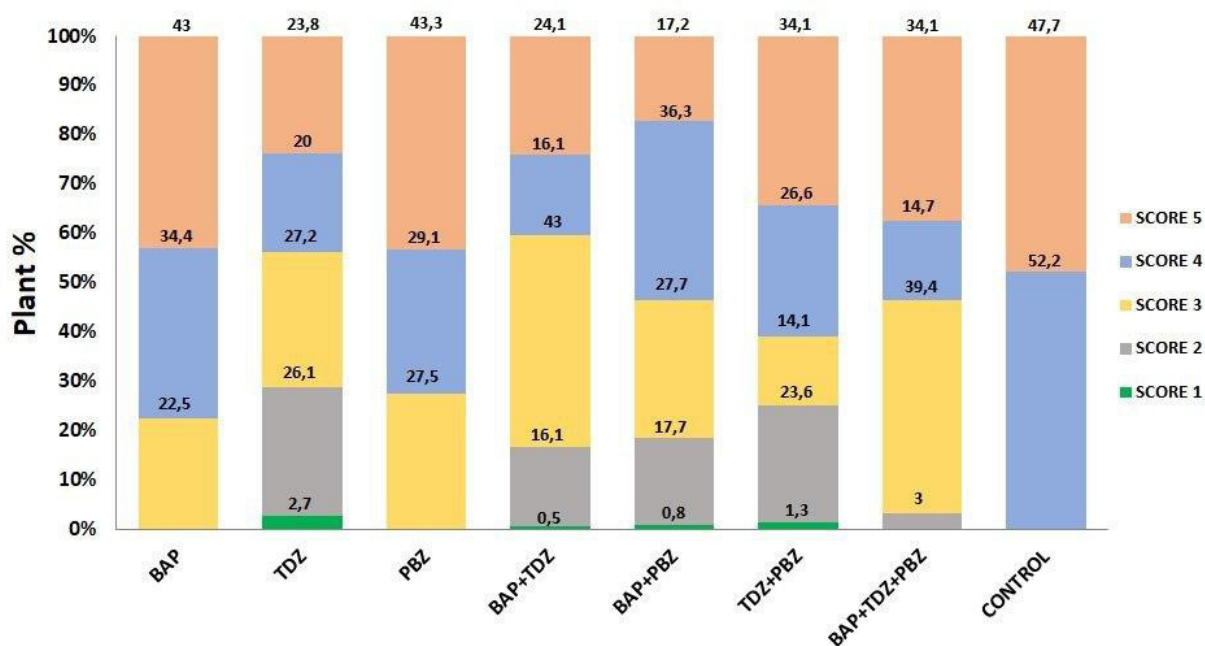


Figure 1. Percentage of plants per grade according to the rhizome internal symptoms scale proposed by Dita et al. (2014) for ‘Prata-Anã’ banana somaclones induced with plant regulators to generate variants resistant to *Fusarium* wilt. A total of 2,520 plants (360 per treatment) and 360 control plants were used. Control: ‘Prata-Anã’ cultivar. Treatments: BAP (6-benzylaminopurine), TDZ (Thidiazuron), PBZ (Paclobutrazol), BAP + TDZ, BAP + PBZ, TDZ + PBZ, BAP + TDZ + PBZ. Internal symptoms scale: 1 = absence of symptoms, 2 = rhizome with initial discoloration, 3 = rhizome discoloration throughout the vascular system, 4 = rhizome with most internal tissues presenting necrosis, and 5 = fully necrotic rhizome.

Owing to contamination problems and losses during the *in vitro* micro-propagation process for the second experiment, only four plants were regenerated and used in the next stage. These plants were designated as putative *Fusarium* wilt race 1 resistant somaclones and named T2-1, T2-2, T4-1, and T5-1. Contamination during the micropropagation process may be related to *Foc* structures in the tissues of plants which, although not manifesting symptoms, may have been infected. Some resistance mechanisms are known to form after infection such that the pathogen enters the tissues, but cannot establish itself because the plant resistance response is faster (LI et al., 2015). The presence of toxins released by the fungus in rhizome tissues is also likely, since *Foc* is a necrotrophic fungus that releases toxins to destroy host tissues (PLOETZ, 2015). Some of the main phytotoxins produced by *Foc* are fusaric acid and beauvericin (PORTAL et al., 2018).

A low number of somaclones without symptoms was generated in this study. This result was expected as genetic variation induction is known to occur in low percentages. A similar study in Taiwan induced somaclonal variation in 20,000 Cavendish banana seedlings planted in a field infected with *Foc* TR4 and reported approximately 0.01% of genotypes resistant to *Fusarium* wilt. This particular study registered GCTCV-218, a somaclonal variant of Cavendish resistant to *Fusarium* wilt, for commercial cultivation under the name Formosona, and two other Cavendish somaclonal variants named GCTCV-53 and GCTCV-119 were identified (HWANG; KO, 2004). The findings of the present study indicate that the use of

plant regulators in association with successive *in vitro* multiplication may lead to greater efficiency in generating resistant somaclonal variants because the study conducted in Taiwan showed a lower percentage of resistant variants (0.01%) than were found in the present study (0.079%).

A study by Ghag et al. (2014) regenerated four genotypes of cv. 'Rasthali' banana plants resistant to *Fusarium* wilt using the somaclonal variation tool. In this study, embryos were repeatedly subcultured and maintained for 14 years under tissue culture conditions. This indicates a low frequency of mutations acting on resistance, requiring a significant number of plants or a long period of subcultures to increase the chances of selecting possible somaclonal variants that are resistant to the disease.

Figure 2 shows the incidence of plants with symptoms (highlighted in red) and plants not showing symptoms (highlighted in green) and indicates that a small amount of genetic variability occurred for the resistance factor in the 2,520 somaclonal variants evaluated, considering that most of them showed *Foc* susceptibility (Figure 2). The treatment with the highest number of plants without symptoms was T2 (TDZ), represented by the T2-1 and T2-2 somaclones. This finding indicates that TDZ used on its own and not combined with other plant regulators in the culture medium was most effective in the generation of somaclonal variants resistant to *Foc* race 1. Thidiazuron is a plant regulator of the cytokinin group, responsible for cell division and used in many studies to induce organogenesis and embryogenesis (PELAH et al., 2002; SHEIBANI et al., 2006; AHMAD et al., 2018).

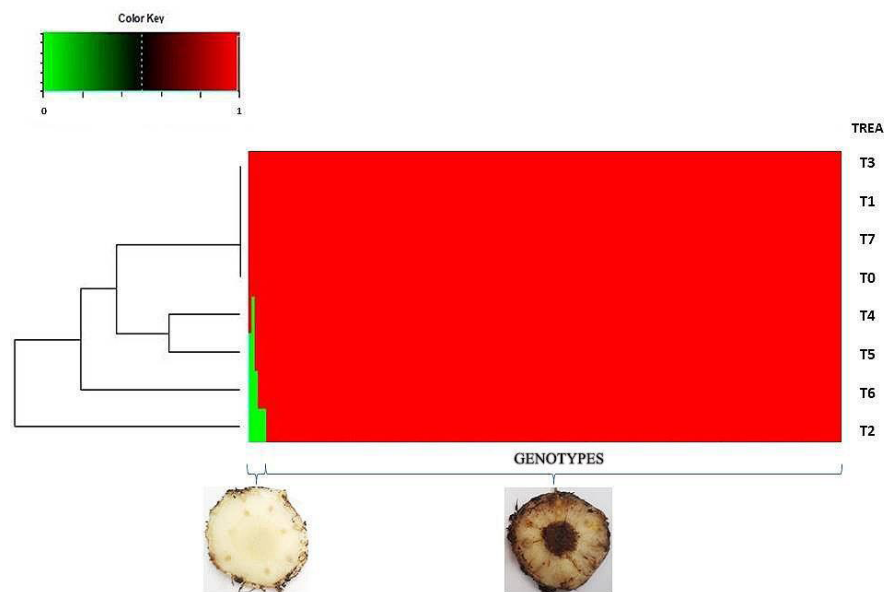


Figure 2. Heat map of the presence (red) and absence (green) of *Fusarium* wilt symptoms in 'Prata-Anã' banana somaclonal variants evaluated in a greenhouse after *Fusarium oxysporum* f. sp. *cubense* race 1 resistance induction with plant regulators. The tree on the left shows the hierarchical grouping of treatments. T0: control ('Prata-Anã'), T1: 6-benzylaminopurine, T2: Thidiazuron, T3: Paclobutrazol, T4: 6-benzylaminopurine + Thidiazuron, T5: 6-benzylaminopurine + Paclobutrazol, T6: Thidiazuron + Paclobutrazol, T7: 6-benzylaminopurine + Thidiazuron + Paclobutrazol, TRAT: treatments.

A study by Goelzer et al. (2019) showed that a combination of TDZ and ANA had a negative effect on sprout growth, suggesting that the use of TDZ alone in the culture medium intensified cytokinin effects on cell division, which could cause DNA polymerase errors, consequently generating mutations. This may account for why, in this study, T2 (TDZ) emerged as the best treatment for generating possible genetic variation in comparison with the other treatments that used combinations of plant regulators.

Analysis of the second experiment results confirmed the resistance of the somaclonal variants T2-1 and T2-2. Evaluation of the external symptoms of somaclones T2-1 and T2-2 by AUDPC yielded a value of 250. This was statistically different from the AUDPC values of somaclones T4-1, T5-1, and T0 (control), which had AUDPC values ranging from 600 to 1,500 (Figure 3). As for the internal symptoms evaluated by the ID, the somaclonal variants T2-1 and T2-2 had the lowest

indexes (0% and 5%, respectively). These values were statistically different to the ID values of somaclones T4-1, T5-1, and T0, which ranged from 40 to 70% (Figure 3). The transverse sections of plant rhizomes showed characteristic *Fusarium* wilt symptoms, with reddish brown coloration in the vascular tissues in T4-1, T5-1, and T0 (Figure 3 A, D, E). Through these analyses, somaclones T4-1 and T5-1, which appeared to be *Foc* resistant during the preliminary test, were shown to be susceptible to *Fusarium* wilt during the second experiment.

The histological evaluation at the end of the preliminary test revealed *Foc* structures such as chlamydozoospores and hyphae in the tissues of T0 and somaclonal variant T2-2 only (Figure 4 A and C). At the end of the second experiment to confirm resistance, the histological evaluation showed that somaclonal variants T2-2, T4-1, T5-1 and T0 had *Foc* structures inside the root tissues. However, the T2-1 variant lacked evidence of fungal structures in the tissues, corroborating the preliminary test result for this somaclone (Figure 4 B and G).

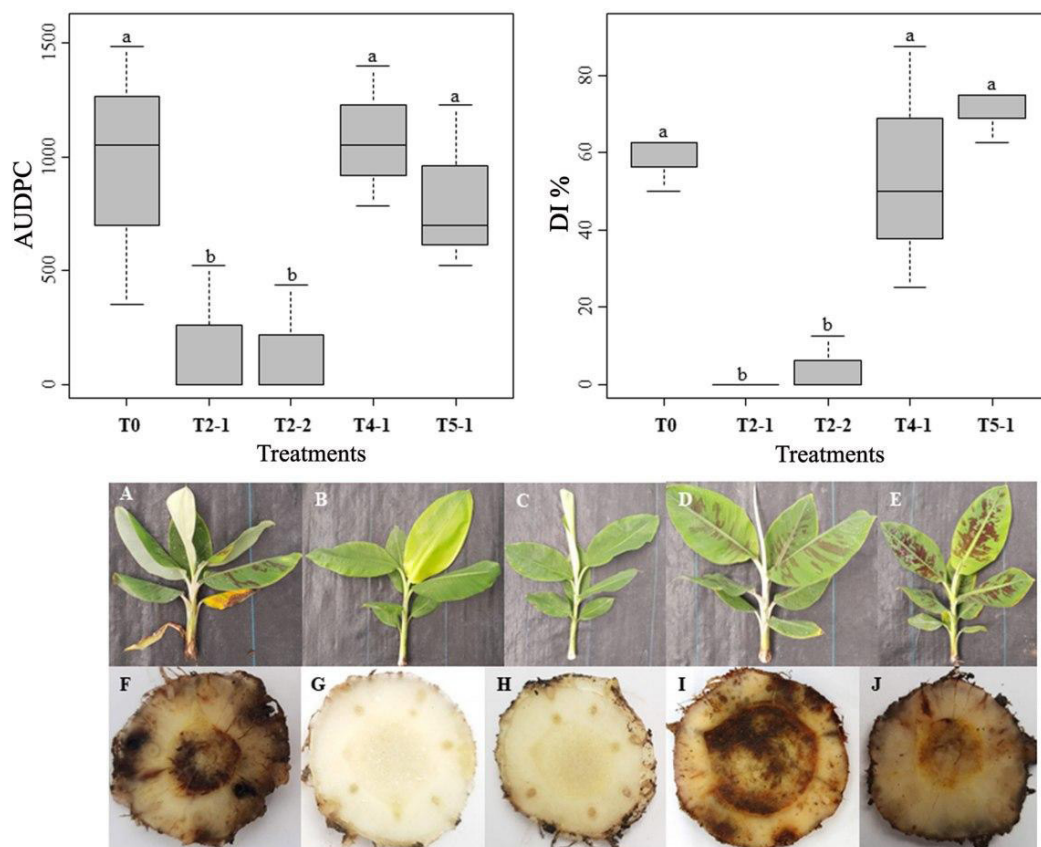


Figure 3. Boxplots of the area under the disease progress curve (AUDPC) and the disease index (ID) of ‘Prata-Anã’ banana plants evaluated in a greenhouse after somaclonal variation induction with plant regulators for *Fusarium* wilt resistance. T0: ‘Prata-Anã’ control without induction, T2-1: variant induced with Thidiazuron, T2-2: variant induced with Thidiazuron, T4-1: variant induced with 6-benzylaminopurine + Thidiazuron, T5-1: variant induced with 6-benzylaminopurine + Paclobutrazol. The bars followed by the same letter do not differ by the Scott-Knott test ($p < 0.05$). Typical external leaf yellowing symptoms are shown from A to E, and transverse sections of the rhizome showing internal symptoms from F to J. The plants were evaluated for symptoms 90 days after inoculation with *Fusarium oxysporum* f. sp. *cabense* race 1 in a greenhouse. ‘Prata-Anã’ (susceptible control) (A, F), T2R1 variant induced with Thidiazuron (B, G), T2R2 variant induced with Thidiazuron (C, H), T4R1 variant induced with 6-benzylaminopurine + Thidiazuron (D, I), T5R1 variant induced with 6-benzylaminopurine + Paclobutrazol (E, J).

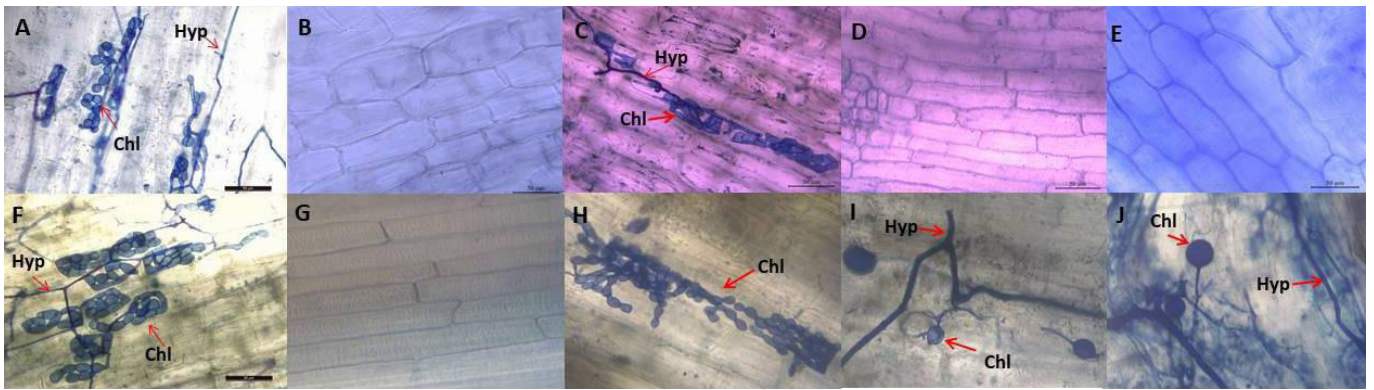


Figure 4. Histological evaluation of internal root tissues of 'Prata-Anã' banana plants induced by somaclonal variation with plant regulators for *Fusarium* wilt resistance caused by *Fusarium oxysporum* f. sp. cubense (Foc). First stage of the study (A, B, C, D, E): Only the 'Prata-Anã' control (A) and the T2-2 somaclonal variant (C) had Foc structures. Second stage of the study (F, G, H, I, J): All somaclonal variants had Foc structures, except T2-1 (G). The roots were clarified, and Foc structures were stained with Trypan Blue. 'Prata-Anã' control without induction showing pathogen colonization and sporulation in the tissue (A and F), T2-1 somaclonal variant induced with Thidiazuron (B and G), T2-2 somaclonal variant induced with Thidiazuron (C and H), T4-1 somaclonal variant induced with 6-benzylaminopurine + Thidiazuron (D and I), T5-1 somaclonal variant induced with 6-benzylaminopurine + Paclobutrazol (E and J). Chl: chlamydospores, and Hyp: hyphae.

Considering that the absence of pathogen structures in root tissues indicates that the pathogen has not successfully penetrated these tissues, the resistant variant T2-1 may have developed physical and chemical barriers to block pathogen penetration (PETIT-HOUDENOT et al., 2017; BANI et al., 2018). As reported in a histological study by Costa (2013), the cultivar 'BRS Platina', a Foc race 1 resistant 'Prata' cultivar (launched by Embrapa in 2011), showed no Foc structures inside the roots, a finding similar to that for somaclonal T2-1. The T2-2 somaclonal variant showed no external or internal *Fusarium* wilt symptoms, indicating Foc resistance; however, it had pathogen structures in the root tissues. These findings suggest that, whereas this somaclone has no resistance to

Foc tissue penetration, it has resistance mechanisms that prevent pathogen infection, possibly because the genetic resistance of plants corresponds not only to the capacity to prevent, but also to delay disease development (VALE et al., 2001; LI et al., 2015).

The leaves of the somaclonal variants showed morphological differences. Somaclones T4-1 and T5-1 had reddish spots on their leaves, similar to the spots on the T0 leaves (Figure 5). The leaves of somaclones T2-1 and T2-2 had no such reddish spots, suggesting some form of genetic modification in these two variants. Previous studies have reported morphological changes in banana trees in relation to somaclonal variation (ALVARES et al., 2002).

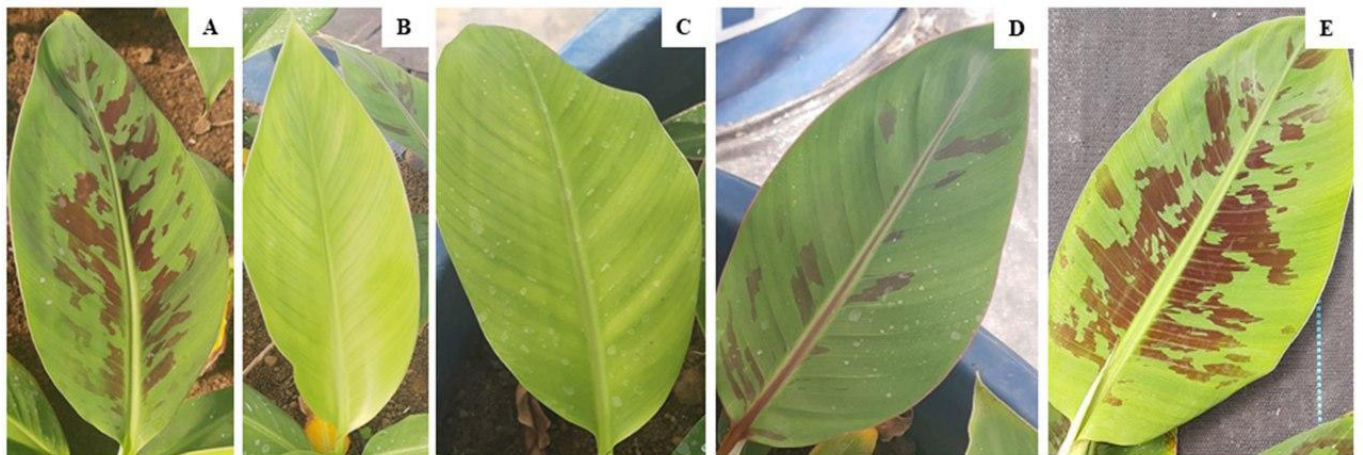


Figure 5. Leaf surface features of 'Prata-Anã' banana plants induced by somaclonal variation with plant regulators for *Fusarium* wilt resistance. 'Prata-Anã' control without induction showing reddish spots on the leaf surface (A), T2-1 variant induced with Thidiazuron with no leaf surface spots (B), T2-2 variant induced with Thidiazuron with no leaf surface spots (C), T4-1 variant induced with 6-benzylaminopurine + Thidiazuron with reddish spots on the leaf surface similar to the control (D), and T5-1 variant induced with 6-benzylaminopurine + Paclobutrazol with reddish spots on the leaf surface similar to the control (E).

The pathogen *Foc* race 1 is present in most banana fields in Brazil. The 'Prata-Anã' cultivar is susceptible to this pathogen and the disease it causes has led to a 60% loss in production in plantations around the country. This fungus produces chlamydospores, resistant structures that can survive long periods in the soil (PLOETZ, 2015; PEGG et al., 2019). Cultural, chemical, and biological control methods have not been effective in controlling the disease (TUSHE-MEREIRWE et al., 2000; KARANGWA et al., 2016), consequently, identification of a 'Prata-Anã' somaclone resistant to *Fusarium* wilt race 1 that maintains other agronomic and sensory characteristics of the original cultivar is essential to protect the cultivar and ensure that it continues to be a source of income for farmers, especially given its preference by Brazilian consumers (IBGE, 2016; MOSTERT et al., 2017; HADDAD et al., 2018).

The findings of this study demonstrate that somaclones T2-1 and T2-2 are promising variants for further studies by the Embrapa banana genetic improvement program. Such studies should include both agronomic characterization to confirm the permanence of desirable characteristics similar to those of the 'Prata-Anã' cultivar, and the elucidation of resistance mechanisms generated in these somaclones. The results of current and future research on the somaclones generated in this study will have a significant impact on the launch of a new *Fusarium* race 1 wilt resistant cultivar.

Conclusion

Induction of somaclonal variations of the 'Prata-Anã' cultivar is viable, as indicated by the generation of the somaclones T2-1 and T2-2, which showed evidence of resistance to *Fusarium* wilt race 1.

Acknowledgment

The authors thank the Coordination for the Improvement of Higher Education Personnel (CAPES) for granting the scholarship, the Breeding Better Bananas/IITA/Bill and Melinda Gates Foundation project, and Embrapa Cassava and Fruit Crops for providing the facilities for conducting the study.

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