Population density of *Trichoderma* fungi in natural environments and agrosystems of a Cerrado area

João Batista Tavares da Silva¹, Eder Marques², José Eustáquio Menezes¹, Joseane Padilha da Silva¹ & Sueli Corrêa Marques de Mello¹*

¹Empresa Brasileira de Pesquisa Agropecuária Recursos Genéticos e Biotecnologia, Brasília, DF, Brasil.
²Upis Faculdades Integradas, Brasília, DF, Brasil.

*Corresponding author: Sueli Corrêa Marques de Mello, e-mail: sueli.mello@embrapa.br

**SILVA, J.B.T., MARQUES, E., MENEZES, J.E., SILVA, J.P., MELLO, S.C.M. Population density of *Trichoderma* fungi in natural environments and agrosystems of a Cerrado area.** Biota Neotropica 20(4): e20201048. https://doi.org/10.1590/1676-0611-BN-2020-1048

**Abstract:** Soil microorganisms present a great diversity, involving taxonomically distinct groups that play a role in the decomposition of organic matter, nutrient cycling, soil aggregation, among others. In this diversity, the fungi of the genus *Trichoderma* have been successful plant pathogen biocontrol agents, as plant growth promoters and as inducers of plant resistance to diseases. In addition, they are important in the sustainability of natural ecosystems. Aiming to verify the population density of *Trichoderma* fungi in natural environments and agroecosystems, in Cerrado area, samples of soils and roots from native vegetation and agroecological production system were collected in the Federal District, Brazil. The collection points were randomly selected, and each soil or root sample was individually wrapped. The soil adhered to the roots was removed for evaluations. Serial sample dilutions and number of Colony Forming Units (CFUs) of *Trichoderma* isolates were performed. The results showed that the number of CFU varied depending on the plant and location evaluated. The replacement of native vegetation by organic farming systems did not result in a significant reduction in this number.

**Keywords:** soil microflora; biocontrol agent; microbial population.

Densidade populacional de fungos do gênero *Trichoderma* em ambientais naturais e agrossistemas de uma área de Cerrado

**Resumo:** Os microrganismos de solo apresentam uma grande diversidade, envolvendo grupos taxonomicamente distintos que desempenham papel na decomposição da matéria orgânica, ciclagem de nutrientes, agregação dos solos, dentre outros. Nesta diversidade, os fungos do gênero *Trichoderma* têm apresentado sucesso como agentes de biocontrole de fitopatógenos, como promotores de crescimento de plantas e, ainda, como induutores de resistência de plantas a doenças. Além disso, são importantes na sustentabilidade dos ecossistemas naturais. Com o objetivo de verificar a densidade populacional de fungos do gênero *Trichoderma* em ambientes naturais e agroecossistemas, em área de Cerrado, amostras de solos e raízes oriundas de vegetação nativa e de sistema de produção agroecológica foram coletadas na região do Distrito Federal, Brasil. Os pontos de coleta foram selecionados aleatoriamente, e cada amostra de solo ou raiz foi acondicionada individualmente. O solo aderido às raízes foi removido para as avaliações. Foram realizadas diluições seriadas das amostras e contagem do número de Unidades Formadoras de Colônias (UFCs) de isolados de *Trichoderma*. Os resultados mostraram que o número de UFC variou dependendo da planta e da localidade avaliada. A substituição da vegetação nativa por sistemas de cultivo orgânicos não resultou em importante redução neste número.

**Palavras-chave:** microflora do solo; agente de biocontrole; população microbiana.
Introduction

Agroecology is an important instrument for the sustainability of small-scale agricultural activities or family farming, mainly due to the low dependence on external inputs from the recommended production systems (Aquino & Assis 2007). Several studies have shown the importance of this farm model in maintaining soil quality and biological activity, in contrast to conventional agriculture (Crowder et al. 2010). In this sense, agricultural practices adopted in agroecological systems are considered strategic to reduce the impact of agricultural expansion on biodiversity in the edaphic environment (Hol et al. 2005). By prioritizing the use of inputs produced on the property, this production model emphasizes the interrelation of the chemical, physical and biological components of the agroecosystem, promoting the conservation of biodiversity, which is important in soil formation (Vandermer 1995). A challenge for scaling up agroecology lies in translating agroecological principles into practical strategies for soil, water and biodiversity management to increase yield and resilience (Nicholls & Altieri 2018).

According to Altieri & Nicholls (2000), it is the interactions between the various biotic components of the agroecosystem that will contribute to biological pest control, nutrient recycling, water conservation, soil conservation and / or regeneration, and increased agricultural productivity in a sustainable way. In this regard, microorganisms have played a major role in the sustainability of agroecosystems. Some of the beneficial microorganisms often used in agriculture worldwide include the genera Bacillus, Azospirillum, Trichoderma, Rhizobium, Mycorrhizae, Pseudomonas, Streptomyces and many other groups (Gupta 2012). As an example, Trichoderma strains have been successfully used as biological control agents of various plant pathogens, being one of the most studied and known microorganisms in the world (Verma et al. 2007). But initially, this biopesticide activity was considered as the only benefit to be considered. Subsequently, it was demonstrated that species of this genus could also be used as biofertilizers, biostimulants, among others (Lorito et al. 2010, Woo & Peppe 2018), being used as inoculant in several agricultural crops.

Therefore, any change that may cause a loss in environmental diversity, influenced by agricultural use or the absence / presence of rainfall, for example, may modify biological diversity in the edaphic environment (Laconi et al. 2013). Several methods have been used as indicators of changes in the soil microbial community. The isolation, cultivation and evaluation of microbial density in samples collected in this environment is the most widely used, due to its ease of execution (Antonioli et al. 2010), although techniques based on the use of molecular markers may be more conclusive about the different groups of microorganisms, organisms and their ecology. These include the latest, based on extraction of microbial DNA directly from the soil (McPherson et al. 2018). However, there is little information on surveys and evaluation of the effects of environmental factors on the composition of beneficial fungal populations and plant pathogen antagonists in Brazilian soils.

Trichoderma fungi are often found in soil and organic matter in free-living form, adapt to different ecological conditions and colonize a multitude of substrates, as well as capable of more intimate associations with plant root systems. (Harman et al. 2004). As a constituent of the rhizospheric microbiota, Trichoderma acts on the translocation of minerals, solubilization and availability of nutrients to plants and the production of plant hormones. Increase in productivity is related to the ability to colonize roots, while its action as a biocontroller has been attributed to the mechanisms of antibiosis, hyperparasitism, induction of resistance, favoring the plant in tolerance to biotic and abiotic stresses, solubilization and nutrient sequestration. in addition to inactivation of pathogen-linked pathogen enzymes. However, the functional variability between isolates of the same species in relation to their biocontrol and plant growth promotion activities is a well-proven fact (Martinez et al. 2013, Munir et al. 2014). Given the agricultural importance of Trichoderma, this work was conducted to verify the population density of this fungus in natural ecosystems and agroecosystems (organic production) of the Cerrado biome.

Material and Methods

1. Collection of soil samples

Soil samples from native vegetation and agroecological production system were collected from healthy vegetables in the Cerrado area, in four localities of the Federal District, always in the morning. In each area five subsamples of non-rhizospheric soils (NRhzS) were collected, containing 200g each, at random points, with a distance of 5 cm from the cultivated species (in ridges between ridges) and 0-10 cm deep, making up a sample composed of 1 kg. Similar procedure was adopted in natural vegetation areas, except for the absence of furrows. Roots and root fragments of the plant species were also collected for each rhizospheric soil (RhzS) collection point. Soil and root samples were individually wrapped and, from these, the attached soil was removed (Ethur et al. 2008). The collection sites were: Rajadinha Rural Nucleus II, Planaltina region, in the cultivation of pumpkin (Cucurbita sp.), eggplant (Solanum melongena L.), kale (Brassica oleracea L. var. acephala DC.), cassava (Manihot esculenta Crantz), Mexican Sunflower (Tithonia sp.), maize (Zea mays L.), bell pepper (Capsicum annuum L.), okra (Abelmoschus esculentus L. Moench), cabbage (Brassica oleracea var. capitata L) and tomato (Lycopercis assulentum Mill.) and native plants Cestrum sp. (Solanaceae), Cyathea sp. (Cyatheaeeae), Miconia elegans (Melastomataceae), Tischochima sp. (Melastomataceae); Taguatinga Rural Nucleus, Taguatinga region, for eggplant, coffee (Coffeea arabica L.), persimmon (Diospyros kaki L.), Mexican Sunflower, maize, mucuna (Mucuna pruriens L. DC), tomato, pod (Phaseolus vulgaris L) and native crops Cauthea sp., Cordiera sp. (Rubiaceae) and Trichilia pallida (Meliaeaceae); Laranéo Rural Nucleus, Paraná region, for eggplant, chives (Allium schoenoprasum L.), cabbage, spinach (Spinacia oleracea L.), cassava, Mexican Sunflower, maize, parsley (Petroselinum crispum Mill. Nym.) and native crops Cordiera macrophylla, Miconia elegans, Zanthisoxylum rbofolium (Rutaceaee); Boa Esperança Rural Center, Cuiabá region, in the cultivation of eggplant, coffee, kale, cassava, Mexican Sunflower, maize, mucuna, pepper (Capsicum sp.), cabbage and tomato. In this place there was no native ecosystem area.

2. Fungus isolation

For fungal isolation, 10 g of each soil sample was placed in Erlenmeyer, suspended in 90 mL of sterile water and stirred at 180 rpm at 25 °C for 40 minutes. After the suspensions, the samples were diluted and 100 µL of each concentration were distributed in Petri dishes containing semi-selective Martin medium as described by Mello et al. (2007). For each sample four repetitions were performed.
3. **Morphological identification of fungi**

The plates were incubated at 25 °C in B.O.D. (Biochemical Oxygen Demand) for two days in the dark and for 5-7 days with 12-hour photoperiod exposure. The cultures were evaluated daily until the appearance of a typical *Trichoderma* colony and considered as a colony forming unit (CFU). Slides were made for examination of morphological characteristics under the optical microscope and identification of the fungus at a generic level. The colonies confirmed as belonging to the genus were transferred to purified potato dextrose agar (PDA) medium, used for monosporic cultivation and stored at 4°C.

4. **Statistical analysis**

First, it was verified whether the plant types (explanatory variable), native and cultivated, differed statistically as to the number of isolates of *Trichoderma* (response variable). This analysis was developed under the focus of Generalized Linear Models (GLMs), assigning Poisson distribution to the response variable, a natural choice for variables of this type (discrete counting). When necessary, the heterogeneity factor present in the data (overdispersion) was corrected via the Quasiverosimilitude method. Following this same logic, another model was adjusted to compare the locations within each culture. To obtain the variance analysis table in Poisson distribution GLMs, were used likelihood ratio (LR), which follows approximately Chi-Square distribution, this procedure is known as ANODEV. The analyzes were developed with the R free statistical language program. The adopted significance level was 5% (McCullagh & Nelder 1989).

### Results

From the soil samples from the four properties, 530 isolates of *Trichoderma* were obtained, 361 from agroecosystems and 169 from natural ecosystems.

1. **Analysis of results by location**

1.1. **Rajadinha Rural Center II - Property I**

Vegetation type interfered with the number of *Trichoderma* isolates obtained (LR = 12.4790, df = 1, p-value = 0.0004116), regardless of soil type (NRhzS or RhzS). Native species presented, on average, a larger number of colonies than cultivated ones (Figure 1A).

![Figure 1](https://doi.org/10.1590/1676-0611-BN-2020-1048)

*Figure 1. A) Average number of Trichoderma colony forming units (CFU) + standard deviation according to soil and plant type for property I. B) Average number of Trichoderma CFU obtained from rhizosphere of cultivated species. C) Average number of Trichoderma CFU obtained from rhizosphere of native species.*
Regarding the cultivated species, significant differences were observed (LR = 17.119, df = 9, p-value = 0.04688), detecting two groups: cassava, Mexican Sunflower, eggplant, maize and pumpkin, with the highest number of CFU of *Trichoderma*; cabbage, tomato, kale, bell pepper and okra with the lowest number of CFU (Figure 1B). On the other hand, native species presented, on average, the same number of CFU (Figure 1C), not differing from each other.

1.2. Taguatinga Rural Center - Property II

Both soil sample type (NRhzS and RhzS) and vegetation type significantly influenced the average number of *Trichoderma* CFU (LR soil = 15.96, df = 1, p-value < 0.0001; LR plant = 5.1549, df = 1, p-value = 0.02318). There was, on average, more CFU of this fungus in NRhzS than in RhzS, both for native and cultivated species (contrast = 0.39, standard error = 0.18, p-value = 0.0267), and the latter species had lower numbers. average of recovered isolates, in terms of CFU in relation to native vegetation (contrast = 0.74, standard error = 0.18, p-value < 0.0001) - (Figure 2A).

Among the cultivated species there was significant difference (LR = 28.409, df = 9, p-value < 0.0001). Two groups can be established: the first, presenting the highest average number of *Trichoderma* CFU composed of mucuna, pod and persimmon, coffee and Mexican Sunflower and the second group, with less CFU, containing kale, maize, eggplant, tomato and cassava (Figure 2B). Among native species, there were no significant differences in obtaining *Trichoderma* colonies (Figure 2C).

1.3. Lamarão Rural Center - Property III

The soil sample type (NRhzS and RhzS) had no effect on the mean number of CFU, which, however, suffered vegetation type interference (RV = 10.062, df = 1, p-value = 0.001514). Native species presented, on average, more CFU of *Trichoderma* than species of organic cultivation (Figure 3A).

There was a significant difference for at least two cultivated species regarding the average number of *Trichoderma* CFU (LR = 23.291, gl = 9, p-value = 0.005574). Mexican Sunflower and eggplant

![Figure 2](https://www.scielo.br/bn)
presented more CFU than spinach, pumpkin, parsley and tomato (Figure 3B). Regarding native species, there were also differences in the number of CFU (LR = 10,312, df = 2, p-value = 0.005765): *C. macrophylla* had, on average, more CFU than *Z. rhoifolium* and *M. elegans* - (Figure 3C).

1.4. Boa Esperança Rural Center - Property IV

For this property, only isolates from cultivated species were analyzed. As for the type of soil sample, there was no statistical difference regarding the number of CFU (LR = 0.13437, df = 1, p-value = 0.7139) - (Figure 4A).

**Figure 3.** A). Average number of *Trichoderma* CFU + standard deviation according to soil and plant type for property III. B) Average number of *Trichoderma* CFU among cultivated species. C) Average number of *Trichoderma* CFU in native species

**Figure 4.** A) *Trichoderma* CFU mean number + standard deviation according to soil type for property IV. B) Average number of *Trichoderma* CFU among cultivated species.
In this case the average number of CFU in samples from ground cultivated with pepper was higher than in the cases of cabbage, tomato, coffee and kale (Figure 4B).

The locations for each crop were compared for the number of \textit{Trichoderma} CFU obtained from eggplant, cassava, kale, Mexican Sunflower and tomato, species found concurrently in the four studied properties, since maize and mucuna were present in three of the four properties. With eggplant, the lowest average number of \textit{Trichoderma} CFU was found for property II (LR = 10.059, df = 3, p-value = 0.01807). In the case of cassava, this number was higher in property I (LR = 9.5157, df = 3, p-value = 0.02317) and, with kale (LR = 1.2136, df = 3, p-value = 0.7498), Mexican Sunflower (LR = 3.1093, df = 3, p-value = 0.3751) and tomato (LR = 0.73695, df = 3, p-value = 0.8645) there was no significant difference in relation to the sample collection sites (Figure 5).

For some species (maize and mucuna), the fungus \textit{Trichoderma} was recovered only at three sites (Figure 6). In this case, there was no significant difference between sites with corn crop (LR = 4.1822, df = 2, p-value = 0.1235) and, with mucuna, a higher number of CFUs were recovered in property II (LR = 12.552, df = 2, p-value = 0.001881).

\begin{figure}[h]
\centering
\includegraphics[width=\textwidth]{figure5.png}
\caption{Average number of \textit{Trichoderma} CFU + standard deviation observed at the four sites.}
\end{figure}
Discussion

Only in property II the number of Trichoderma CFUs was higher in NRhzS than in RhzS (Figure 2A), probably due to the incorporation of crop residues or other organic matter in the soil, since in the other properties there were no significant differences in this number (Figures 1A, 3A and 4A). Moreover, according to Ethur et al. (2008), Trichoderma species settle better in soils when they contain vegetable remains and other forms of organic matter. Although in these soils there is greater interaction between microorganisms and plants (Mohammadi et al., 2011), it is also in them that the microbial flora suffers the most competition pressure (Dantas et al. 2009). Kredics et al. (2018) complement that various biotic and abiotic factors affect diversity populations of microbial communities in agroecosystems, including plant species and their growth stage, total microbial competition, pesticide or fertilizer application, as well as geographic region. However, only a few studies address the population, abundance and diversity of the genus Trichoderma in specific fields or agroecosystems. From the results obtained in this work, there is no need to remove parts of plant roots to obtain a representative number of Trichoderma isolates, since the number of Trichoderma isolates recovered from root-attached soil samples was similar to non-rhizospheric soil samples. In areas cultivated with tomato and cucumber in conventional system, Ethur et al. (2008) obtained similar results. It is worth mentioning that, in studies of antagonistic potential, Jash & Pan (2007) found no differences in antagonism against Rhizoctonia solani when testing Trichoderma isolates from RhzS and NRhzS.

The areas of native vegetation, located around the crop, suffer a reduced anthropic effect, so the soil supposedly represents the ecological conditions of environmental stability because it is not influenced by disturbances of preparation and application of inputs, unlike cultivated areas, even treating it itself from organic production. Probably, this fact explains the higher number of CFUs found in native vegetation areas in the three evaluated sites (Figures 1C, 2C and 3C), compared to those of cultivated area (Figures 1B, 2B and 3B). Brouwer & Riezebos (1998) has mentioned that in forest soils, nutrient losses from the ecosystem are lower. This provides better soil cover, higher organic matter content and greater floristic diversity, determining factors for larger soil settlement in number and microbial diversity (Ramos et al. 2012). According to Lourente et al. (2011), the diversity of native vegetation species (quantity and quality) implies the continuous deposition of organic substrates with varied composition, favoring the microbial mass content, but the substitution of native vegetation by cultivation systems may cause important changes in the attributes. soil chemicals in the first year of implementation. There are several reports that disturbances caused by land use and crops may also result in decreased microbial biodiversity (Bending et al. 2004, Mendes et al. 2012). Louzada et al. (2009), who studied the antagonistic action of Trichoderma isolates from various regions of Brazil against plant pathogens Sclerotinia sclerotiorum and Fusarium solani, mention studies by other authors showing that isolates from native areas would result in a higher percentage of potentially active Trichoderma isolates against plant pathogens in in vitro tests, with a mycelial growth inhibition of around 80%. However, in this work, the substitution of native vegetation by organic cultivation systems did not cause a significant reduction in the number of Trichoderma CFUs, confirming postulations made by different authors about the advantages of the organic cultivation system regarding the preservation of soil microflora and stabilization of agroecosystems.

Therefore, the results presented show the effect on the number of CFU of Trichoderma, depending on the culture and location evaluated (Figures 5 and 6). These properties, although located in Cerrado areas, have soils probably subjected to different treatments, which would have interfered with the number of recovered Trichoderma colonies. According to Frazão et al. (2010), soil microbial community is generally influenced by variations in soil temperature, water content and aeration, aggregate disruption, decreased soil cover, nutrient availability and organic substrates. Studies by Saravanakumar et al. (2016) from samples of coastal regions showed that the biodiversity of Trichoderma spp. was influenced by temperature, redox potential and pH. In the set of results, it was found a variation between Trichoderma population levels in the various organic crops evaluated, due to the different factors mentioned, although no correlation was studied or made regarding soil types and characteristics, only with the cropping systems.

Figure 6. Average number of Trichoderma CFU + standard deviation observed at the three sites.
Regarding diversity, molecular characterization work of *Trichoderma* isolates from target properties should be performed to verify the prevalent species in the soil of organic crops and native vegetation, because the methodology used for sample dilution and calculation of forming units of colonies (CFUs) do not allow the distinction of an introduced strain of *Trichoderma* populations residing in the investigated environment (Kredics et al. 2018). According to Louzada et al. (2009), there are no data in the literature reporting the loss of diversity of *Trichoderma* spp. with the continuous agricultural use of soils or even the possible relationship of such interferences with the reduction of the frequency of antagonism to pathogens. *Trichoderma* spp. are highly successful settlers in their habitats and are able to overcome adversity related to environmental variations around the world (Schuster & Schmoll 2010). Studies of this nature coupled with the knowledge of the real distribution and population dynamics of this fungal genus and its associations with different plant species and soils are crucial to ensure the efficiency and safety of the use of these microorganisms, especially in the biocontrol and promotion of plant growth.

Given the above and based on the results observed in this work, it is concluded that the types of crop and native vegetation influenced the distribution of the population of *Trichoderma* fungi in soils of organic farming system. On the other hand, the substitution of native vegetation by organic cultivation systems did not result in a significant reduction in the number of *Trichoderma* CFUs.

**Author Contributions**

João Batista Tavares da Silva: substantial contribution in the concept and design of the study; contribution to data collection, contribution to data analysis and interpretation, contribution to manuscript preparation, contribution to critical revision and adding intellectual content.

José Eustáquio Menezes: contribution to data collection, contribution to data analysis and interpretation and contribution to manuscript preparation.

Eder Marques: contribution to data analysis and interpretation and contribution to manuscript preparation.

Joséane Padihla da Silva: contribution to data analysis and interpretation.

Sueli Corrêa Marques de Mello: substantial contribution in the concept and design of the study; contribution to data collection, contribution to data analysis and interpretation, contribution to manuscript preparation, contribution to critical revision and adding intellectual content.

**Conflicts of Interest**

The authors declare that they have no conflict of interest related to the publication of this manuscript.

**References**

ALTIERI, M. & NICHOLLS, C.I. Agroecologia: Teoria y práctica para una agricultura sustentable. Série Textos Básicos para la Formación Ambiental. 1ª Edición. México: PNUMA, 2000, 250p. ISBN 968-7913-04-X

ANTONIOLLI, Z.I., SANTOS, L.C., LUPATINI, M., LEAL, I.T., SCHIRMER, G.K. & REDIN, M. 2010. Efeito do cobre na população de bactérias e fungos do solo, na associação micorrízica e no cultivo de mudas de *Eucalyptus grandis* W. Hill ex Maiden, *Pinus elliotti* Engelm e *Peltophorum dubium* (Sprengel) Taubert. Ciência Florestal 20(3):419–428. http://dx.doi.org/10.5902/198050982057

AQUINO, A.M. & ASSIS, R.L. 2007. Agricultura orgânica em áreas urbanas e periurbanas com base na agroecologia. Ambiente & Sociedade 10:137–150. http://dx.doi.org/10.5902/1414-753X200700100009

BENDING, G.D., TURNER, M.K., RAYNS, F., MARX, M.C. & WOOD, M. 2004. Microbial and biochemical soil quality indicators and their potential for differentiating areas under contrasting agricultural management regimes. Soil Biology & Biochemistry 36(11):1785–1792. http://dx.doi.org/10.1016/j.soilbio.2004.04.035

CROWDER, D.W., NORTHELD, T.D., STRAND, M.R. & SNYDER, W.E. 2010. Organic agriculture promotes evenness and natural pest control. Nature 466:109–112. https://doi.org/10.1038/nature09183

DANTAS, J.S., SOUZA, A.P., FARIAS, M.F. & NOGUEIRA, V.E.F. 2009. Interações entre grupos de microorganismos com a zigoafera. Pesquisa Aplicada & Agrotecnologia 2(2):213–218. https://revistas.unicentro.br/index.php/repa/article/download/113/308

ETHUR, L.Z., BLUME, E., MUNIZ, M.F.B., ANTONIOLLI, Z.I., NICOLINI, C., MILANESI, P. & OLIVEIRA, F. 2008. Presença dos gêneros *Trichoderma* e *Fusarium* em solo rizosférico e não rizosférico cultivado com tomateiro e pepino, em horta e estufa. Ciência Rural 38(1):19–26. https://doi.org/10.1590/S0103-84782008000100004

BROUWER, L.C., RIEZEBOS, H.T. 1998. Nutrient dynamics in intact and logged tropical rain forest in Guyana. In: Schulte A., Ruhiyat D. (eds). Soils of Tropical Forest Ecosystems. Springer, Berlin, Heidelberg. https://doi.org/10.1007/978-3-662-03649-5_7, pp 73–86.

FRAZÃO, L.A., PICCOLO, M.C., FEIGL, B.J., CERRI, C.C. & CERRI, C.E.P. 2010. Inorganic nitrogen, microbial biomass and microbial activity of a sandy Brazilian Cerrado soil under different land uses. Agriculture Ecosystems & Environment 135:161–167. https://doi.org/10.1016/j.agee.2009.09.003

GUPTA, V.V.S.R. Beneficial microorganisms for sustainable agriculture. 2012. Microbiology Australia 33(3):113–115. https://10.5902/198050982057

HARMAN, G.E., HOWELL, C.R., VITERBO, A., CHET, I. & LORITO, M. 2004. *Trichoderma* species – opportunistic, avirulent plant symbionts. Nature Reviews/Microbiology 2:43–56. https://doi.org/10.1038/nmicrobot79

HOLE, D.G., PERKINS, A.J., WILSON, J.D., ALEXANDER, I.H., GRICE, P.V., EVANS, A.D. 2005. Does organic farming benefit biodiversity? Biological Conservation 122:113–130. https://doi.org/10.1016/j.biocon.2004.07.018

JASH, S. & PAN, S. 2007. Variability in antagonistic activity and root colonizing behavior of *Trichoderma* isolates. Journal of Tropical Agriculture 45(1–2):29–35. http://jropag.kau.in/index.php/oj2/article/view/169/169

KREDICS, L., CHEN, L., KDEVES, O., BÜCHNER, O., HATVANI, L., ALAGA, H., NAGY, V.D., KHALED, J.M., ALHARBI, N.S. & VÁGVÖLGYI, C. 2018. Molecular Tools for Monitoring *Trichoderma* in Agricultural Environments. Frontiers in Microbiology 9:1–17. https://doi.org/10.3389/fmicb.2018.01599

LANÇONI, M.D., TAKETANI, R.G., KAVAMURA, V.N. & MELO, I.S. 2013. Microbial community biogeographic patterns in the rhizosphere of two Brazilian semi-arid leguminous trees. World Journal of Microbiology & Biotechnology 29(7):1233–1241. http://dx.doi.org/10.1007/s11274-013-1286-4

LORITO, M., WOO, S.L., HARMAN, G.E. & MONTE, E. 2010. Translational research on *Trichoderma*: from ‘omics to the field. Annual Review Phytopathology 48:395–417. https://doi.org/10.1146/annurev-phyto-073009-114314

LOURENTE, E.R.P., MERCANTE, F.M., ALOVISI, A.M.T., GOMES, C.F., GASPARINI, A.S. & NUNES, C.M. Atributos microbiológicos, químicos e físicos de solo sob diferentes sistemas de manejo e condições de Cerrado. Pesquisa Agropecuária Tropical 41(1):20–28, 2011. https://10.20.96.pat. v411.8459
LOUZADA, G.A.S., CARVALHO, D.D.C., MELLO, S.C.M., LOBO JÚNIOR, M., MARTINS, I. & BRAÚNA, L.M. Antagonist potential of *Trichoderma* spp. from distinct agricultural ecosystems against *Sclerotinia sclerotiorum* and *Fusarium solani*. Biota Neotropica. 9(3):145–149. http://dx.doi.org/10.1590/S1676-06032009000300014

MARTÍNEZ, B., INFANTE, D. & REYES, Y. *Trichoderma* spp. y su función en el control de plagas en los cultivos. Revista de Protección Vegetal 28(1):1–11, 2013. http://scielo.sld.cu/pdf/rpv/v28n1/ rpv01113.pdf

McCULLAGH, P. & NELDER, J.A. Generalized Linear Models. 2.ed. London: Chapman & Hall, 1989. 511p.

MCPherson, M.R., Wang, P., Mitchell, R.B. & Schachtman, D.P. Isolation and analysis of microbial communities in soil, rhizosphere, and roots in perennial grass experiments. Journal of Visualized Experiments 24(137):1–11, 2018. https://doi.org/10.3791/57932

Mello, S.C.M., Ávila, Z.R., Braúna, L.M., Pádua, R.R. & Gomes, D. 2007. Cepas de *Trichoderma* para el control biológico de *Sclerotium rolfsii* Sacc. Fitosanidad 1(1):3–9. https://ainfo.cnptia.embrapa.br/digital/bitstream/item/178234/1/209116144001-2-8.pdf

Mendes, I.C., Fernandes, M.F., Chaer, G.M. & Reis Junior, F.B. 2012. Biological functioning of brazilian cerrado soils under different vegetation types. Plant and Soil 352(1):183–195. https://10.0.3.239/s11034-012-1195-6

Munir, S., Jamal, Q., Bano, K., Shewani, S.K., Abbas, M.N., Azam, S., Kan, A., Ali, S. & Anees, M. 2014. *Trichoderma* and biocontrol genes: review. Scientia Agriculturae 2(2):40–45. https://10.59.88/PSCP.SA.2014.1.2.4045

Nicholls, C.I. & Altieri, M.A. 2018. Pathways for the amplification of agroecology. Agroecology and Sustainable Food Systems, 1–24. https://doi.org/10.1080/21683565.2018.1499578

Mohammadi, k. Heidari, G., Khalesro, S., Sohrab, Y. 2011. Soil management, microorganisms and organic matter interactions: A review. African Journal of Biotechnology,10(84):19840–19849. https://doi.org/10.5897/AJBX11.006

Ramos, M.L.G., Meneghin, M.F.S., Pedroso, C., Guimarães, C.M., & Konrad, M.L. 2012. Efeito dos sistemas de manejo e plantio sobre a densidade de grupos funcionais de microrganismos, em solo de Cerrado. Bioscience Journal 28(1):58–68. http://docs.bvsalud.org/biblioref/2018/09/912350/efeito-dos-sistemas-de-manejo-e-plantio-sobre-a-densidade-de-gr_6s2MkPN.pdf

Schuster, A. & Schmoll, M. 2010. Biology and biotechnology of *Trichoderma*. Applied Microbiology and Biotechnology 87(3):787–799. https://doi.org/10.1007%2Fs00253-010-2632-1

Saranakumar, K., Yu, C., Dou, K., Wang, M., Li, Y., Chen, J. 2016. Biodiversidade da comunidade de *Trichoderma* nos planos de marés e zonas úmidas do sudeste da China. PLoS ONE 11(12): e0168020. https://doi.org/10.1371/journal.pone.0168020

Vandermar, J. 1995. The ecological basis of alternative agriculture. Annual Review Ecology Systematics, 26:201–224. https://www.jstor.org/stable/2097205

Verma, M., Brar, S.K., Tyagi, R.D., Surampalli, R.Y. & Valero, J.R. 2007. Antagonistic fungi *Trichoderma* spp.: Panoply of biological control. Biochemical Engineering Journal 37(1):1–20. https://doi.org/10.1016/j.bej.2007.05.012

Woo S.L. & Peppe, O. Microbial consortia: promising probiotics as plant biostimulants for sustainable agriculture. Frontiers in Plant Science 9(4,9):1801. https://doi.org/10.3389/fpls.2018.01801

Received: 08/06/2020
Revised: 24/08/2020
Accepted: 31/08/2020
Published online: 25/09/2020