Arbuscular mycorrhizal fungi (AMF) are commonly present in most tropical land ecosystems; however, high rates of fertilization can disrupt their life cycle. The AMF occurrence in different crop systems compared to native vegetation in tropical conditions is not well understood. With the aim to analyze potentially benefic symbionts for crop production in an experimental farm, the AMF community structure along different crop cultivation was investigated. Soil subsamples were used to inoculate trap cultures using Sorghum sp. as host. The preserved unfertilized area showed a greater diversity of AMF, followed by the grassland and sugarcane. Number of AMF spores was also greater in the preserved area (47 spores per 100g soil) than the numbers estimated for other sites. Nine AMF genera of Glomeromycotina (spores) and 16 AMF species were observed in the native forest. The natural richness appears to be favored when Sugarcane or grasses are planted; however, the preserved native forest showed unique characteristics. Soils under native forest stored higher amount of SOM, presented higher macro-aggregates and preserved more charcoal than the samples of permanent grassland or under maize. The maize associated AMF community diverged from grassland, as described by the principal component analysis. This study showed differences in AMF communities from cultivated plots compared to native vegetation as well as potential for inoculant formulation for economically crops.

Keywords: Arbuscular mycorrhizal fungi; Atlantic Forest; Sugarcane; Maize; Grassland
where plant species from Atlantic Forest and Cerrado Brazilian biomes, both considered as hotspots of biodiversity [14], occur. Atlantic forest total sampled area represents simply 0.01 % of the remaining biome, Minas Gerais state being one of the better studied [15]. However, most landscapes save for conservation purposes (marginal agricultural lands) can hold high species diversity providing biodiversity conservation services [16]. Thus, to improve the conservation of anthropogenic landscapes in Brazilian tropical forest is urgently needed [17].

With the aim to analyze AMF potentially benefic for crop production in experimental farms, we show in the present study, results of estimates of AMF communities’ structure along one year of crop cultivation. The Experimental farm in Pitangui (an area of 460ha) includes preserved riparian areas between São João and Pará Rivers, localized in the São Francisco River basin, a protected Reserve (its vegetation is primarily Atlantic Forest), forestry, and corn/sugarcane fields. Experimental farm belongs to EPAMIG ITAC Company (Empresa de Pesquisa Agropecuária de Minas Gerais), located in the center-west region of the state of Minas Gerais and dedicated to research with regional cultivars potentially important for intensive systems of production or screening programs. Plant crops are rice, corn, beans, soybean, to name just a few.

Four different areas were studied from maize and sugarcane monocropping, grassland to environmentally protected Native Forest. Among cultivated pastures, Brachiaria grasslands (B. decumbens and B. brizantha) are the most important agronomic species in Brazil, also used for Integrated Crop-Livestock-Forest systems [18]. The grassland for pasture (Brachiaria brizantha), the continuous maize monocropping with high input of fertilizers and pesticides, and sugarcane monoculture (Saccharum officinarum) fertilized only with compost were selected.

We hypothesized that soil grassland could contain high AMF spore numbers, macroaggregates and charcoal, but the native forest can present the highest values. Furthermore, we hypothesized that changes in land-use affect the fertility and structure of the soil matrix so that the overall. We predict monocultures to impact the AMF community in this study. Studies investigating AMF diversity in tropical native forest, grassland and crop fields are scarce, but no available studies have been targeted to the anthropogenic disturbances that drive AMF community shifts together with soil aggregation and charcoal content.

The aims of this study were: (1) to evaluate the communities of AMF in different ecosystems of land use intensity from undisturbed preserved and permanent grassland to monocultures, (2) to associate the AMF communities with soil characteristics and (3) to evaluate the effect of inoculation of soil containing arbuscular mycorrhizal propagules on plant growth, as part of a greenhouse experiment. Since we wanted to focus on the simple on farm inoculation (mixed AMF spores in soil) alone, we chose to compare different non-sterilized soils.

**Material and Methods**

**Field analysis**

**Table 1:** Historical characteristics of sites sampled in the experimental farm and adjacent preserved area (Atlantic forest).

| Vegetation/ Standing Crop | Abbreviation | Geographical Coordinates | Altitude (m) | Estimated Age of Forest/Cultivation (year) | Previous Crop / Vegetation | Historic of Fertilizer Use | Historic of Pesticide Use |
|---------------------------|--------------|--------------------------|--------------|------------------------------------------|---------------------------|--------------------------|--------------------------|
| Atlantic Forest           | NF           | 19°42’49.9”S 44°54’07.8”W | 637,05       | 30                                       | —                         | —                        | —                        |
| Grassland                | G            | 19°42’51.5”S 44°53’37.0”W | 628,36       | >25                                      | NA                        | Compost                  | Herbicide VEPAR           |
| Sugarcane                | S            | 19°42’40.9”S 44°53’40.0”W | 642,80       | 2                                        | Bush, grass               | —                        | —                        |
| Maize                    | M            | 19°42’48.1”S 44°53’45.3”W | 632,88       | 10                                       | Maize (high cultivation intensity) | Mineral fertilizer (N:P:K = 8:28:16; 400Kg ha-1); Urea (350Kg ha-1) | LOSBAN, ROUNDUP UpKlorpan 480 CE; Herbicides SANSON 40 SC and DMA 806 BR |

NA: Not available. Grassland: chopped up once a year.

Study sites. The four sites (Table 1) selected for this study are located in the plain of the São Francisco River basin located in Brazil. Four land use types, e.g. maize field (19°42’48.1”S, 44°53’45.3”W), grassland (19°42’51.5”S, 44°53’37.0”W), sugarcane (19°42’40.9”S 44°53’40.0”W) and Atlantic forest (19°42’49.9”S 44°54’07.8”W) (Representative of land use with different intensity in descending order), were selected as sampling sites.

Different areas were sampled:

**Table 2:** List of plant species identified in the permanent forest cover (AF) plots in 2014. Species are ordered by family (in alphabetical order). The characteristic biomes are also reported. Legend: * = risk of extinction.

| Family      | Species            | Biome            | AMF Status    |
|-------------|--------------------|------------------|---------------|
| Areceaceae  | Acrocomia aculeata (Jacq.) Lodd. | Brazilian cerrado | AM (Fonseca et al. 2015) |
| Bignoniaceae| Handroanthus serratifolius | Atlantic forest  | NR            |
| Caricaceae  | Jacaratia heptaphylla | Brazilian cerrado | NR            |
| Caryocaraceae| Caryocar brasiliense Cambess. | Brazilian cerrado | AM (Carneiro et al. 1998) |

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Cecropiaceae  Cecropia pachystachya  Atlantic forest, Brazilian cerrado (Riparian forest)  AM (Pouyú-Rojas and Siqueira 2000)

Fabaceae  Machaerium villosum  Atlantic forest; Brazilian cerrado  NR

Fabaceae  Deguelia costata (Benth.) Az.-Tozzi  Atlantic forest  NR

Fabaceae (Caesalpinioideae)  Copaifera langsordoffii Desf.  Atlantic forest  AM (Siqueira et al. 1998; Zangaro et al. 2003)

Fabaceae Caesalpinioideae  Hymenaea courbaril L.  Atlantic forest  NM (Siqueira et al. 1998) (Pedone-Bonfim et al. 2018)

Fabaceae (Mimosoide)  Inga vera Willd.  Atlantic forest  NR

Fabaceae (Mimosoide)  Styphnodendron adstringens  Brazilian cerrado  NR

Fabaceae (Mimosoide)  Piptadenia gonoacantha  Atlantic forest  NR

Lauraceae  Nectandra oppositifolia  Atlantic forest  NR

Lecythidaceae  Cariniana estrellensis*  Atlantic forest, Brazilian cerrado  NR

Myristicaceae  Virola sebifera Aubl.  Tropical dry forest, Cerradão, Riparian forest border  NR

Monimiaceae  Siparuna guianensis Aublet.  Brazilian cerrado  NR

Myrtaceae  Campomanesia velutina (Cambess) O. Berg, Caatinga  NR

Myristicaceae  Virola sebifera  Brazilian cerrado  NR

Tiliaceae  Luehea grandiflora  Atlantic forest  AM (Siqueira et al. 1998)

Rubiaceae  Rudgea viburnoides (Cham.) Benth.  Brazilian cerrado  NR

Sapindaceae  Cupania vernalis Cambess.  Atlantic forest, Brazilian cerrado  NR

Rutaceae  Zanthoxylum rhoifolium  Brazilian cerrado  NR

1. Native Forest (NF), nature reserve about 25m from São João River, presenting Atlantic forest cover (80ha). The Forest Reserve is a protected area of Atlantic forest with prominent vegetation composed by trees and shrubs having a total density of ~1,029 trees per hectare reaching heights of 10–25m [19]. Dominant species are listed in Table 2. Plant vegetation had been undisturbed for 30 years. Some plant species are at risk of extinction such as *Cariniana legalis*. Exsiccates were deposited in Herbarium PAMG of EPAMIG;

2. The Grassland (G) has gramineous cover with *Urochloa brizantha* syn. Brachiaria brizantha Stapf under no-till cultivation being the dominant grass; (about 1000m from São João River). The G is located in the direct neighborhood of Maize;

3. Maize (M) monocropping (*Zea mays* cultivar AG1051) (conventional hybrid for silage) area, 5ha, fertilized with 400kg 8:28:16 NPK and 350Kg urea; Maize were planted in late January and harvested in December each year from 2004 to 2014. Last maize cultivation was in the year 2014. The maize was manually sowed on December 2014. Assessments of soils were made after the maize harvest, between August 2015 and October 2015. Weeds and pests, when they appeared, were treated with insecticide Klorpan 480 CE (three applications); Herbicides SANSON 40 SC and DMA 806 BR (one application);

4. Sugarcane (S) (0.7ha) *Saccharum officinarum* variety *mellifera melifera* 418, cultivated under undetermined compost, harvested and crushed to feed livestock. This site was only evaluated in the second sample period (Table 2).

Figure 1: Location map showing experimental farm in Brazil. Plot-wise crop variety information. Google Earth image showing the locations of Native forest: Legal Reserve (Brazil’s new Forest Code – Law 12651/2012) and agroecosystems in 2017 (A); Illustrative pictures of the sampling sites: native forest (B); Maize field (C) and grassland (D).
All the sites have the same geological parent material and were classified as clayey Oxisol (Typic Acrustox) according to Soil Taxonomy [USDA 1992]. The cultivated soils are loamy (Figure 1).

The climate of the region is Tropical (Aw) with an annual average temperature of about 23 °C and predominant precipitation in summer and dry winter. This area was characterized by a total solar radiation of 7.2h, an average precipitation of 1337mm per year.

The study area is located in the west of Minas Gerais State, in Brazil, included in the Cerrado- Atlantic Forest region (19°40 ´S 44°53´ W), characterized by 709mm of yearly precipitation concentrated in the spring-summer months as revealed from November to January (Meteorological station Mineração Turmalina, Jaguar Mining Inc. (19°44°36´ S, 44°52°36´ W, 700m), located 6 km south of the town of Pitangui. Near 84% of tropical rainfall falls during the rainy season (October to March). The months of December and January present the most intense precipitation levels. Air humidity, even in the summer, does not exceed 86% on average (Figure 2).

**Brief land-use history**

The original vegetation of the area is derived from Atlantic forest/Brazilian Cerrado ectone, although part of the forest was deforested as early as the 1970s as revealed by interviews with local farmers. The area comprises the São João River with approximately 5m width and 1 to 1.5m deep. The São João basin vegetation was partially degraded and substituted by grasslands or eucalypt plantations.

The experimental farm in Pitangui has been divided into eight sectors: cultivated areas, zootechny in general and Forest reserve, dedicated to research with regional cultivars potentially important for intensive systems of production or screening programs such as rice, corn, beans, soybean, to name just a few. In the 1970s, this area was donated to the Minas Gerais State.

Typical land-use types in the area include forestry, cornfields, sugarcane plantations and grasslands. The agricultural use of the soils includes planted grassland (G); arable lands with moderate-intensity management, such as maize and sugarcane monocropping (M, S) (Table 1). The M site was established in 2004, followed by many years of annual crops. Last maize cultivation was in the year 2014 when it was manually sowed on December. The G site (in the central area, Figure 1) had been cultivated and then abandoned and left for grassland, chopped up once a year.

**Soil sampling and chemical characterization**

Soil samples (three replicate plots per field site; plot sizes, 5 by 20m) were obtained in February (rainy period) and November (after the dry period) 2015 at four sites.

At each of the three plots at each field site, five soil core samples were taken up to a depth of 20cm with a mattoc, with the exclusion of the top soil layer (0-5cm), totalizing 30 samples. Samples were stored in plastic bags and transported to the laboratory of EPAMIG. Each composite sample was air dried and carefully conditioned. Each sample was stored in bags, and subsequently brought to the laboratory for analysis. Prior to further processing all visible roots and rocks were removed manually.

**Table 3:** Chemical properties of soils at experimental farm and adjacent Atlantic forest.

| Areas     | Atlantic Forest | Grassland | Cultivated |
|-----------|-----------------|-----------|------------|
| pH (H2O)  | 5.4             | 5.4       | 6.2        |
| OC (dag kg-1) | 4.5       | 2.6       | 1.4        |
| OM (dag kg-1) | 7.7       | 4.4       | 2.4        |
| Ca (cmolc kg-1) | 2         | 1.5       | 4.1        |
| Mg (cmolc kg-1) | 0.8       | 0.9       | 2          |
| Al (cmolc kg-1) | 0.5       | 0.7       | 0.1        |
| K mg kg-1    | 34            | 43        | 230        |
| P mg kg-1    | 1.4           | 1.8       | 5.4        |

About 500g were used to determination of chemical soil parameters, according to EMBRAPA (1997). The rest of soil samples was used for AMF spore extraction and identification. Chemical properties of the soil determined by the traditional methods as described by EMBRAPA (1997). The following chemical analyses were performed: pH in water (1:2.5 soil/water); soil organic carbon (SOC); P, K+, Ca2+, Mg2+ and Al3+ were determined according to EMBRAPA, 1997 (Table 3).

**AMF spore isolation and identification**

Wet sieving and sucrose density gradient centrifugation [20] extracted AMF spores occurring in the original soil samples (50g soil). The resulting supernatant was passed through the 32-μm
sieve, washed with tap water, and transferred to Petri dishes. AMF communities in each area were recorded and their diversity was estimated.

Spores and sporocarps were counted by using a dissecting microscope and mounted on slides with polyvinyl-lactic acid-glucose and mixed 1:1 (vol/vol) with Melzer’s reagent. Only the healthy-looking spores were counted. The spores were examined under a microscope, mounted on slides and identified to the species level or attributed to a specific morphospecies. Identifications were based on current species descriptions.

The old and decaying spores with missing clear features were also counted as an approach for comparison among samples [21]. The results expressed as percentage of unviable AMF spores in the sample. Darkest black color spores or remained damaged were considered as non-viable. Unviable spore counting was done at 40× magnification (Figure 3).

**Greenhouse analysis**

Plant cultures (pots, 20 x 20 x 30cm) were produced for each site. Seeds of Sorghum BR were planted in pots filled with 3 Kg of washed sand, vermiculite and collected soils at each field site: Sugarcane, grassland and Native forest, (1:1:1 [wt/wt]) according to Oehl et al. [10] a methodology denominated Culture Trap, were the host of Fungi symbiont and promoted his growth in soil culture.

The greenhouse experiment was initiated in August 2015 and was maintained in a greenhouse in Pitangui for 3 months under day/light regimes of 12-h: 12-h photoperiod and 28:25 °C temperature, with a mean relative humidity of 65%. Seedlings were irrigated with tap water and plant height were measured at 2 and 3 months after planting (Figure 4).

**Statistical analysis**

The significance of differences between field sites in spore abundance, species numbers, and AMF diversity (Shannon-Weaver index) was tested by using ANOVA one-way. Homoscedasticity and data normality were tested before performing analysis of variance and test of means. Tukey post-hoc tests were used to conduct pairwise tests for significant effects. Associated means for each treatment were reported using the software Minitab 17 [9].

**Results**

**AMF occurrence**

AMF spores were counted and identified for samples taken directly from field sites. Identification of AM fungal species based on morphological features of spores in the root zone of maize, sugarcane, grassland and native forest showed 16 identifiable AMF species. One species of Glomus, one of Sclerocystis, one of Claroideoglomus, one of Ambispora, four of Acaulospora, one of Gigaspora and four of Scutellospora were detected. Four spore types could not be readily distinguished at the species level.

**Table 4:** AMF fungal genera found in the different sampled soils. Atlantic forest (AF), grassland (G) and cultivated (M and S).

| AMF Family         | Genera               | AF | G  | M  | S  |
|--------------------|----------------------|----|----|----|----|
| Acaulosporaceae    | Acaulospora          | 3  | 0  | 1  | 1  |
| Ambisporaceae      | Ambispora            | 1  | 0  | 0  | 1  |
| Claroideogluomerace| Claroideoglomus      | 1  | 0  | 1  | 0  |
| Gigasporaceae      | Gigaspora            | 0  | 0  | 1  | 1  |
| Glomeraceae        | Funneliformes        | 1  | 1  | 1  | 0  |
|                    | Glomus               | 1  | 1  | 0  | 0  |
|                    | Sclerocystis         | 1  | 0  | 0  | 0  |
| Pacisporaceae      | Pacispora            | 1  | 0  | 0  | 0  |
| Scutellosporaceae  | Denticutata          | 1  | 1  | 1  | 1  |
|                    | Racocetra            | 2  | 0  | 1  | 1  |
|                    | Scutellospora        | 1  | 1  | 0  | 0  |
| Total genera       |                      | 13 | 4  | 6  | 5  |

In the soil samples taken from the studied systems, 16 AMF species could be distinguished. The 10 AMF species observed in soil sampled in the dry and rainy season consisted of seven families and ten genera (Table 4). Four species belonged to the genera Acaulospora while the genera Claroideoglomus, Gigaspora, Scutellospora, Denticutata and Ambispora were represented by a single species each. The species Denticutata heterogama, was observed in all the...
areas, in both the dry and rainy season, the last species being the most frequent (100%). The species *Sclerocystis rubiforme, Sclero-
cystis taiwanensis* and *Racocetra verrucosa* were exclusive to the NF
area, whereas *Gigaspora margarita* was observed only in the cultivated/sugarcane areas (Table 4). *Glomus* sp. 3 was unique to NF and
*A. mellea, A. scrobiculata* and *A. appendicula* were shared between
NF and Sugarcane (Figure 5).

**Spore numbers**

AMF spore numbers in the soil samples ranged from 1 to 23
spores per 100g of soil, with an average of 8 to 12 spores, and signif-
nicant differences were detected between sites (Table 5). The spore
numbers in the maize field was significantly lower in the rainy
period than in the NF, G and S. The spore numbers in the NF (dry period), G, S and Maize field (dry period) were not significantly
different.

**Table 5:** Total AMF spore numbers (50g soil) of each family in soils from
Natural forest (NF), grassland (G) and cultivated (M and S) (n = 3).

| AMF Family         | Rainy Period | Dry Period |
|--------------------|--------------|------------|
|                    | NF | G | M | NF | G | M | S |
| Acaulosporaceae    | 6.6 | 0 | 0.3 | 2.6 | 0.6 | 0.3 | 3.3 |
| Ambisporaceae      | 0.3 | 0 | 0 | - | - | - | 2.3 |
| Claroideoglomerace  | 4.6 | 1 | 0.6 | 11 | 0.6 | 1 | 1.3 |
| Gigasporaceae      | - | 1 | - | - | - | - | 0.2 |
| Gomerae             | 8.9 | - | - | 18.6 | - | 0.6 | 0.6 |
| Pachyspora         | - | - | - | 0.67 | 2 | - | 0 |
| Scutellospora       | 2.9 | - | 0.3 | - | 1.6 | - | 1.9 |

**Species richness and diversity**

**Table 6:** Richness (S), Shannon-Weaver (Hs), Simpson (D) and Even-
ness (E) indices calculated from AMF associated with each soil at exper-
imental farm at rainy and dry periods.

| Site             | S  | H   | D  | E  |
|------------------|----|-----|----|----|
| Rainy Dry        |    |     |    |    |
| Atlantic Forest  | 16 | 9   | 2.3| 1.3 |
| Grassland        | 2  | 4   | 0.33| 1.27|
| Maize            | 3  | 3   | 1.04| 1.01|
| Sugarcane        | NE | 7   | NE | 1.66|

Mean values n = 3.

In both seasons, the values of total species richness (S) and the
Shannon diversity index were different among the areas (Table 6),
indicating different composition of AMF communities, ranging from
0.33 to 2.12. In the dry season, the AMF community had greater
similarity between sugarcane and native forest areas, while in the
rainy season, the AMF community of the NF had greater richness.

The species numbers in the grassland was slightly, but signifi-
cantly lower in the rainy than in the dry period despite the fact that
the similar precipitation had been registered, but more humidity
was detected in the rainy period.

The species richness in the maize fields was slightly, but signifi-
cantly lower in the both sampling periods. This analysis indicated
that the different AMF communities with plant cover of grassland/
cultivated samples did not associate with the native fertile soil (Ta-
ble 7).

**Table 7:** Distribution of AMF species associated with each soil at exper-
imental farm, Minas Gerais state, Brazil (+, presence; -, absence) at both
sampling periods.

![AMF Species](attachment:image)

Data obtained at 0-20cm soil depth. bMaximal AMF richness found in
each sample (n = 6).

**Table 8:** Mean number of unviable AMF spores and macro-charcoal
fragments in soils from Atlantic forest (AF), grassland (G) and cultivated with
sugarcane (S) or maize (M).

| Site | Unviable Spores | Macroscopic Charcoal |
|------|-----------------|----------------------|
| AF   | 4.6±3.7b        | 28.3±13.05ab         |
| G    | 27.6±9.6ab      | 19.6±3.21ab          |
| M    | 62.3±24.7a      | 8.3±5.03bc           |
| S    | 13.6±8.9ab      | 47±15.71a            |

†Average for three replicates plots (25g soil) of field site in November
2015. Data are means ± SE. Mean values with the same uppercase letter
in the treatment do not differ among themselves in the ANOVA Tukey test
(p<0.05).

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The number of viable AMF spores in the soil samples, which was evaluated in the dry period, ranged from ~5 to 62 spores per 100g of soil, with an average of 36.2±15.5 spores, but no significant differences were detected between sites (Table 8).

Greenhouse experiment

Figure 6: Abundance of AMF morphotypes detected in natural and cultivated.

Figure 7: Sorghum height Growth in greenhouse experiment at 2 and 3 months after cultivation with inoculum from NF, G and S. Experiment realized at 2015.

Mean values with the same letter among the inoculum sites do not differ among themselves in the ANOVA Tukey test (p<0.05).

Figure 8: Sorghum height Growth under greenhouse conditions after 2 and 3 months after cultivation with inoculum from NF, G and S. a) Experiment realized at year 2015 and b) Experiment realized at 2018.

The results of the greenhouse experiment showed higher growth of Sorghum inoculated with soil from the native forest and from the grassland sites (Figure 4) than with soil from the cultivated area. Sorghum height in greenhouse experiment at 2 and 3 months after cultivation was significantly higher by the use of inoculum from NF and G sites (Figure 6,7,8 & 9).

Discussion

Our study provide support for the hypothesis that AMF communities are limited in their diversity by monocultures. We found that preserved ecosystem (Atlantic Forest ecotone) was more diverse than agroecosystems, not only in plant diversity but also in AMF diversity and in improved soil OC content and soil aggregation.

The cultivated soils presented high fertility, a higher pH value, and lower organic matter content than native forest soil, which presented moderate fertility and low P and K levels. This shows that agriculture as practiced in tropical agro-systems for Brazil, negatively affects the satisfactory nutrient levels (with excess of Ca, K, P and Mg), increases pH and decrease OM in top soils. Consistent supplies of fertilizers and lime are required in naturally acid soils (high contents of aluminum and iron oxides, low CEC values and OM content) in Cerrado [22].

For P applications in Brazilian crop production systems, soil test P methods indicate that quantity of plant-available soil P [23] and the required fertilizer level is then inferred. However, the loss of fertilizers in agricultural runoff from soils (negative impacts of Brazilian agriculture) can be controlled via use of conservation tillage, cover crops, buffer or riparian zones [23], and conservation/augmentation of biofertilizers/soil conditioners [24].

The proportion of aggregates >2 mm in the investigated soils slightly decreases in the following order: NF and G > crop. However, microaggregates were significantly lower in forest soils than in G and cropland. Similar results from Portella et al. (2012) [8] reported that macroaggregates are sensitive to changes in management practices and, from Blanco-Canqui and Lal (2004) [24] reported some variation in microaggregates. It is also known that water-stable aggregates >0.25 mm together with soil OM content can significantly control erodibility [27]. Moreover, the tillage practices adopted in croplands can disrupt soil aggregates [9], increase soil compaction and disturb plant and animal populations that contribute to soil aggregation.

A higher hyphal length, root colonization by AMF and AMF (spore) species diversity is expected in native forest than cultivated sites, due to a permanent plant cover, higher soil OM content and soil humidity [27]. Thus, the higher macroaggregates content under NF in rainy period can be due to a higher hyphal growth. Soils under permanent grassland can also store higher soil OM in larger aggregates than samples under permanent maize [9].

The observed higher quantity of macroscopic charcoal in sugarcane and NF than in grassland and maize was consistent with results by Schneider et al. (2011) [28] showing unchanged charcoal chemical quality and stocks along 100 years in a tropical ecosystem in western Kenya. The notable increase of charcoal in the sugarcane
site suggesting fire activity maybe by burning management in a previous period.

Indeed, results showed that each site correlate with distinct soil attributes. NF associated with OC, charcoal, and macroaggregates. Thus, soil chemical properties such as Al, Ca, Mg, P and K concentrations of soil C, and soil aggregates are the main soil factors distinguishing sites. This study evaluated the AMF communities in different sampling sites showing that spores belonging to 9 genera of Glomeromycotina were detected. The results showed that preserved ecosystem (Atlantic Forest) was more diverse than agroecosystems, wherein pH is higher, and higher levels of Ca, P and K were detected.

In general, spore-based morphological studies demonstrated that application of fertilizers, pesticides, and fungicides results in significant loss of AMF diversity in many ecosystems [29]. We predicted that spore-based morphological studies could detect the occurrence of Gigasporaceae and Scutelllosporaceae in more preserved areas (two species of *Racotetra* and one of *Scutellospora* were found). The total number of AMF species varied among the evaluated areas being higher in preserved area of Atlantic Forest (NF), observing that number of AMF spores in preserved area ranged from 9 to 47 spores in 50 g soil.

In both seasons, the values of total species richness (S) and the Shannon diversity index were different among the areas, indicating different composition of AMF communities. In dry season, the AMF community had greater similarity between sugarcane and forest areas, while in rainy season, the AMF community of NF had greater richness. This pattern suggests that AMF community is affected not only by the host, but also by climatic conditions, and possibly by soil conditions [30]. Azevedo et al. (2014) [31] also found a high number of AMF species (37) in a small area of sugarcane plantations. Aidar et al. (2004) [32] found that H (from 1.10 to 1.96) late season of AM fungal diversity.

The S, where the plant protection strategy is non-burning management and compost addition, resembled more the G than the conventional M. The successional dynamics described here are in line with Whittaker’s (1972). The species numbers in crop fields and grassland was slightly, but significantly lower in the conventional than in organic treatments despite the fact that same crop rotation had been applied. The species richness in crop fields and grassland was slightly, but significantly lower in the conventional than organic treatments despite the fact that the same crop rotation had been applied. This result was confirmed in the trap cultures.

This study agrees with Oehl et al. [10] who also found a decrease of spore abundance of AMF species not belonging to the genus Glomus in Monocultures. Many of these species appeared to prefer or even to be restricted to forest systems. This was the case for most species of *Acaulospora* identified from field as well as for *Acaulospora* sp. and *Entrophospora infrequens*. A previous study indicated a trend towards an increase in AMF belonging to genera *Acaulospora*, *Scutellospora* and *Entrophospora* under long-term reduced tillage managements [33].

Our study complements this finding, indicating that not only reduced tillage but also organic farming may increase the diversity of AMF species and genera in arable lands. Our findings indicate that some AMF species, especially *Acaulospora* spp., find an ecological niche in soils of organic farming systems, and that this may be connected with characteristic low level of readily available P in these soils [10]. We hypothesize that these AMF species are functionally important to natural ecosystems and low-input sustainable farming; if this is true, their loss under conventional farming would be alarming.

The reduced AM fungal diversity in annual croplands compared to grasslands corresponds with some other studies in tropical regions [10] and supports the reported inverse relationship between agricultural land use intensity and AM fungal diversity [10]. The number of spores in cultivated area was much lower. This spore distribution points to hypothesis of greater sporation of AMF detected in more preserved areas.

Unviable AMF spore number in soil samples ranged from 5 to 62 spores per 100g of soil. The average spores number agree with the proportion of viable spores, in surface soil, from 35 to 60% detected by the MTT test [21,34]. In the present study, the proportion of unviable spores in NF was lower but not significant than in other sites, with a possible stable spore production and loss from spore bank.

Sporulation and dormancy are of prime importance for the long-term persistence and survival of AMF in any habitat. The AMF spore bank in soil (formation of new spores and removal from the bank by death and by germination) will determine the spore numbers in soils [10]. AMF species differ in their sporulation and germination dynamics [10].

Sorghum growth experiment at 2 and 3 months after cultivation was significantly higher by use of inoculum from NF and G sites. This can be due to different propagules that can be present in NF and G such as hyphae and spores, corroborating a great bank of propagules. The finding of higher AMF diversity and plant infection potential in organic as compared to conventional treatments fits well with previous observations of a higher soil aggregate stability. AM inoculation in combination with biochar application may be applicable not only for maize production but also for Phyto stabilization of Cd-contaminated soil [35]. Dai et al. [36] AM fungal resources of soils being resilient to disturbance and that richness of AM fungi under crop management has been maintained, despite evidence of a structural shift imposed by this type of land use. Roadside in prairie are a good repository for conservation of AM fungal diversity.

AMF colonization of sugarcane roots in current study is within the range observed in other studies, i.e. 10 to 89% in sugarcane under different field and greenhouse conditions [37-39]. However, no-burning management resulted in higher root colonization after the first harvest, as compared to pre-harvest burning management. Possible explanations for these results are: (I) an enhanced degradation of the litter driven by the tillage, and (II) more abundant organ mineral interactions preserving SOM from degradation and...
driven by the higher root density under grassland. Lignin balance showed that lignin stocks were more efficiently preserved under ley and permanent grassland than under permanently cropped soils.

AMF spores were counted and identified for sample taken directly from field sites. Identification of AM fungal species basing on morphological features of spores in rhizosphere soil of maize, sugarcane and native and restored forest.

In both seasons, the values of total SR and the Margalef diversity index were very similar among the areas, indicating similar composition of AMF communities, ranging from 50 to 66 %. In the dry season, the AMF community had greater similarity between pasture and forest areas, while in the rainy season, the AMF community of the regeneration area had greater similarity to that of forest fragment. This pattern suggests that AMF community is affected not only by host, but also by climatic conditions, and possibly by soil conditions [30].

The species numbers in crop fields and grassland was slightly, but significantly lower in conventional than in organic treatments despite the fact that same crop rotation had been applied. This result was confirmed in trap cultures. In this way, the present study demonstrated the importance of symbiosis with AMF in preserved areas as potential for the establishment of AMF inoculants for species of economic interest typical of region.

The inoculum potential of soil samples from the different treatments in field trial was determined in AMF trap cultures by measuring the proportion of root length of trap plants after 2 months of culture. It was found to be highest in the decreased samples from in same order as the spore abundances observed.

The benefits of AMF to their host plants could not be explained by improved nutrition alone, since interaction with the remaining soil organisms (i.e., priming effect of mycorrhiza) also differed between inoculated AMF. Due to distinct socialization strategies between inoculated AMF and remainder soil microbes and fauna, inoculation with irregulars and especially with Scutellospora sp. did not overrule the soil’s legacy from maize monocropping (and thus the negative feedbacks), while inoculation with C. claroideum, F. mosseae and Gigaspora sp. Did [40].

Conclusion

The soil management adopted in cultivated areas decreases the density and diversity of AMF spores in relation to adjacent natural vegetation. The present analysis of the surface profile of soils revealed that Sugarcane cultivation leads to significant accumulation of charcoal. Sugarcane showed most benefit than maize cultivation systems. The most accumulation processes occurred in reined soils, which is due to input and mineralization of plant residues and soil amendments [41].

The best strategy for reclaimed the degraded cultivable soils could be to add AMF inoculum better than abandoning cropland for succession, as unviable spores and lower spore abundances are present. Thus, soil content can indicate more closely related to natural soils attributes, when soil. Soil inoculation with propagules of AMF positively influenced the plant height grown in the greenhouse.

These results demonstrated good prospects for study of functional diversity of mycorrhizae to attain greater productivity of regional agricultural systems. Stimulates more effective and ambitious conservation actions in the region to address the increasing human pressure and demand for agricultural land expected in coming decades.

We highlight the need of introduction or maintenance of AMF spore bank in soil for agronomic management practices. Recommendations are made for AM functional groups, such as to maintain preserved forest on rural farm as a source of ecosystem services, which can be the dispersal of AMF propagules and for use in the transition of conventional agriculture to organic.

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Conflict of Interest

No conflict of interest.

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