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Coprophilous Mucorales (ex Zygomycota) from three areas in the semi-arid of Pernambuco, Brazil

Carlos Alberto Fragoso de Souza\textsuperscript{a}, Diogo Xavier Lima\textsuperscript{a}, Luciana M.S. Gurgel\textsuperscript{b}, André Luiz Cabral Monteiro de Azevedo Santiago\textsuperscript{a,*}

\textsuperscript{a} Programa de Pós-graduação em Biologia de Fungos. Universidade Federal de Pernambuco, Departamento de Micologia, Recife, PE, Brasil
\textsuperscript{b} Instituto Agronômico de Pernambuco (IPA), Recife, PE, Brasil

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\textbf{A B S T R A C T}

Mucorales comprises fungi commonly isolated as saprobes from soil, dung, stored grains and plants. Although these fungi have been studied in several countries, there are relatively a few reports of them in semi-arid areas. Therefore, the aims of the present study were to assess and compare the Mucorales communities in dung from different species and breeds of herbivores in the semi-arid of Pernambuco, based on the frequency of occurrence and species richness of these fungi. Samples of dung collected in the cities of Arcoverde, Serra Talhada and Sertânia were incubated in moist chambers in triplicate. Altogether, 24 taxa of Mucorales distributed in the genera Absidia, Cincinella, Cunninghamamella, Lichtheimia, Mucor, Pilobolus, Rhizopus and Syncephalastrum were identified. The highest species richness was found in sheep excrement. Mucor circinelloides f. grise-cyanus was the most common taxon, followed by \textit{M. ramosissimus}. The similarity of the composition of Mucorales species was greatest between the excrements of Guzerá and Sindi breeds (bovine). All mucoralean species isolated are being cited for the first time from animal dung found in Caatinga and a new species of \textit{Mucor} was recorded. An identification key for species of Mucorales from dung in the semi-arid region of Brazil is provided.

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\section*{Introduction}

\textit{Mucoromycotina} Benny, one of the four subphyla proposed to accommodate certain species of traditional \textit{Zygomycota} C. Moreau (phylum no longer accepted in the new classification due to its polyphyletic nature),\textsuperscript{1,2} covers most saprobes fungi characterized by the production of thick-walled spores of sexual origin, known as zygosporangia. The order \textit{Mucorales} Fr., the largest in number of species within the \textit{Mucoromycotina}, includes fungi forming coenocytic hyphae with septa either at the base of reproductive structures or irregularly distributed in older cultures. Species of this order produce asexual structures such as sporangiophores, sporangia, sporangiospores,
These fungi are commonly isolated from soil, stored grains, plants, and animal excrement, especially that of herbivores and rodents. 

Coprophilous fungi are those that live in or are associated with fecal material, including soil contaminated with feces. These microorganisms are essential for the maintenance of ecosystems and are directly involved in the decomposition of fecal waste, participating in carbon, nitrogen and energy cycles. According to Dix and Webster, fungi may occur in excrement as obligate and optional coprophilous. The obligate coprophilous have spores that require the action of gastric enzymes to break their dormancy, while secondary or facultative coprophilous spores do not need to go through the digestive tract of animals to germinate. Among the Mucorales, only Pilobolus species are considered obligate coprophilous, although other taxa from this order are common in dung.

The Brazilian semiarid region occupies an area of approximately 969,589.4 km², comprising eight northeastern states: Alagoas, Bahia, Ceará, Paraíba, Pernambuco, Piauí, Rio Grande do Norte, Sergipe and part of the state of Minas Gerais. Semi-arid regions are formed, mostly by typical vegetation of Caatinga, an exclusively Brazilian domain that consists of heterogeneous phytophysionomic systems characteristically xerophilic, with vegetation ranging from tree to shrub. The diversity of fungi in Semi-arid ecosystems is admittedly higher than previously thought. However, it is estimated that 41.1% of Caatinga regions have not yet been inventoried. Consequently, data related to the fungal community and the ecological relationship between microorganisms and substrates in these ecosystems are still scarce.

Concerning Mucorales in Brazil, 74 taxa, belonging to 20 genera, have been reported. Of these, 42 were isolated from herbivore dung, which is a highly favorable substrate for the growth of these fungi. Although a number of authors have documented the occurrence of coprophilous Mucorales in Brazil, there are no reports addressing the diversity and ecology of these fungi on dung in semi-arid areas. Therefore, the aims of the present study were to assess and compare the Mucorales communities in dung from different species and breeds of herbivores in the semi-arid region of Pernambuco, based on the frequency of occurrence and species richness of these fungi.

### Materials and methods

#### Study areas

Samples of herbivore dung were collected at the Instituto Agronômico de Pernambuco (IPA) in Sertanía-PE (8°03′38″S, 37°13′32″W) [animals: caprine (Capra hircus L., breeds Anglo-Nubiano and Moxotó), and ovine (Ovis aries L., breeds Santa Inês and Morada Nova], and Arcoverde (8°25′00″S, 37°04′00″W) [animal: bovine (Bos taurus L., breeds Holandês, Girolando and Sindi]. Bos taurus L. dung samples (Breed Guzerá Leiteiro) were collected at the IPA in the city of Serra Talhada (7°95′67″S, 38°29′71″W).

### Table 1 – Food supplied to herbivores from Arcoverde, Sertanía and Serra Talhada, PE.

| Herbivore (breed) | Food                  |
|------------------|-----------------------|
|                  | Cane silage | Corn silage | Mineral salt | Graze |
| Anglo-Nubiano    | –          | +           | –            | –     |
| Moxotó           | –          | +           | –            | –     |
| Santa Inês       | –          | +           | –            | –     |
| Morada Nova      | –          | +           | –            | –     |
| Holandês         | +          | –           | +            | –     |
| Sindi            | +          | –           | +            | –     |
| Girolando        | +          | –           | –            | +     |
| Guzerá           | –          | –           | –            | +     |

#### Dung samples

The samples were collected monthly, from September 2013 to April 2014, using sterilized spatulas. They were placed in celophane autoclaved paper bags and kept in polystyrene boxes with ice until they arrived in the laboratory. All samples were collected in the morning, usually after the animals’ first meal of the day.

#### Food supplied to herbivores

The composition of food supplied to herbivores from Arcoverde, Sertanía and Serra Talhada is provided in Table 1.

#### Isolation, purification and identification

Dung samples from each animal were incubated in moist chambers at 28±2 °C for 15 days under alternating light and dark periods, during which time mycelial growth was observed. Fragments of the grown colonies were transferred to malt extract agar (MEA) medium (Merck – EMB), supplemented with chloramphenicol (100 mg L⁻¹). After growth, the fungi were transferred to test tubes containing the same culture medium without antibiotic. The specimens were identified by observing their macroscopic (color, appearance and diameter of colony) and microscopic (microstructures) characteristics as described by Hesselteine and Ellis, Schipper, Hesselteine and Fennel, Zheng and Chen, Hoffmann et al., Zheng et al., and Santiago et al.

#### Molecular analysis

Culture grown in test tubes containing malt extract were incubated at 28 °C for 6 to obtain fungal biomass. The material was transferred to 2 mL microtubes with screw caps. Subsequently, 0.5 g acid-washed glass beads of two different diameters (150–212 μm and 425–600 μm, 1:1; Sigma, USA) were added to each tube. The material was crushed by stirring at high speed in a FastPrep homogenizer. The genomic DNA extraction procedure was conducted as described by Góes-Neto et al. The mycelium was washed with chloroform:isoamyl alcohol (24:1) and then homogenized in 2% cetyltrimethylammonium bromide buffer. The DNA was precipitated in isopropanol, washed with 70% ethanol, and resuspended in 50 μL ultrapure water.
The primer pairs LR1/LSU2 were used for the amplification of the large subunit (LSU) of nuclear ribosomal DNA (rDNA).\textsuperscript{23,25} The polymerase chain reactions were carried out as described by Oliveira et al.\textsuperscript{26} The newly obtained sequence was deposited in the National Center for Biotechnology Information GenBank database (accession number KX133009).

**Frequency of occurrence (FO)**

FO was calculated using the following equation: $\text{FO} = j_i/k$, where FO is the frequency of occurrence of the species $i$, $j_i$ is the number of samples in which the species $i$ has occurred and $k$ is the total number of soil samples.\textsuperscript{27}

**Statistical analysis**

Differences in the associations of Mucorales occurring among different herbivores dung were determined using Similarity Analysis (ANOSIM Primer v.6), in which the matrix of Bray–Curtis similarity was plotted as described by Clarke and Gorley.\textsuperscript{28} Differences between the number of species of Mucorales associated with different breeds of animals and the different months of the year were assessed using the Chi-Square ($\chi^2$) test (adjustment of compliance with expected equality proportions), according to the following formula: $\chi^2 = \sum(0 - e)^2/l$, where $o$ is the observed frequency for each class and $e$ is the expected frequency for that class.\textsuperscript{29} The significance level was set at 0.05 in the analysis.

**Results**

Twenty-four taxa within the genera Absidia, Circinella, Cunninghamella, Lichtheimia, Mucor, Pilobolus, Rhizopus and Syncephalastrum were identified in bovine, caprine and ovine dung found in the semi-arid region of Pernambuco. The highest number of taxa was observed in samples from Morada Nova (14 taxa) and Santa Inês (12) (ovine), followed by Holandês (10) and Girolando (8) (bovine). The lowest species richness was observed in samples from Guzerá (bovine) (3) (Table 2).

Mucor exhibited the highest number of species (9), followed by Pilobolus (5) (Table 2).

Mucor circinelloides f. griseo-cyanus occurred in high frequency (FO = 59.35%) in dung from the bovines, caprine and ovine examined, followed by M. ramossissimus (FO = 32.8%) and M. circinelloides f. circinelloides (FO = 31.25%). Cunninghamella echinulata var. echinulata, Mucor sp., P. minutus and R. arrhizus var. arrhizus were the least common (FO = 1.56%) (Table 2).

The highest number of taxa was observed in dung samples from Morada Nova (14) and Santa Inês sheep breeds (12)

| Table 2 - Richness and frequency of occurrence (FO) of Mucorales in herbivore dung from Arcoverde, Sertânia and Serra Talhada, PE. |
|----------------------------------|----------------|----------------|----------------|----------------|----------------|----------------|
| Mucorales                        | Excrement      | FO             |                 |                 |                 |                 |
|                                  | Bovine         | Caprine        | Ovine           |                 |                 |                 |
|                                  | Girolando      | Guzerá         | Holandês        | Anglo-Nubiano   | Moxotó         | Morada Nova    |
|                                  |                |                |                 |                 |                 | Santa Inês     |
| Absidia cylindrospora var. cylindrospora Hagem | –              | –              | +               | –               | –              | –              | 6.25%          |
| Circinella muscae (Sorokin) Berl. & De Toni | +              | –              | +               | +               | +              | +              | 18.74%         |
| Cunninghamella echinulata var. echinulata (Thaxt.) Thaxt. ex Blakeslee | –              | –              | –               | –               | –              | –              | 1.56%          |
| Lichtheimia brasiliensis A.L. Santiago, Lima & Oliveira | –              | –              | –               | –               | –              | +              | 3.12%          |
| L. ramosa (Zopf) Vuill. | –              | –              | –               | +               | –              | +              | 4.68%          |
| Mucor circinelloides f. circinelloides Tiegh. | –              | +              | –               | +               | +              | +              | 31.25%         |
| M. circinelloides f. griseo-cyanus (Hagem) | +              | +              | +               | –               | +              | +              | 59.35%         |
| Schipper                         | –              | –              | –               | +               | –              | –              | 6.24%          |
| M. circinelloides f. janssenii (Lendn.) Schipper | +              | –              | –               | +               | +              | –              | 3.12%          |
| M. lusitanicus Bruderl. | +              | –              | –               | +               | –              | +              | 4.68%          |
| M. hiemalis Wehmer | –              | –              | –               | –               | +              | –              | 3.12%          |
| M. indicus Lendn. | –              | –              | –               | –               | +              | +              | 3.12%          |
| Mucor sp. | –              | +              | –               | –               | –              | –              | 1.56%          |
| M. luteus Linnem. | –              | –              | +               | +               | –              | +              | 4.68%          |
| M. racemosus f. racemosus Fresen. | –              | +              | –               | +               | –              | +              | 3.12%          |
| M. ramossissimus Samouts. | +              | +              | –               | +               | +              | –              | 32.8%          |
| M. variosporus Schipper | –              | –              | –               | –               | +              | –              | 4.68%          |
| Pilobolus crystallinus Tode | +              | +              | –               | +               | +              | –              | 20.31%         |
| P. kleini Tiegh. | +              | +              | –               | +               | –              | +              | 7.81%          |
| P. longipes Tiegh. | –              | –              | +               | +               | –              | +              | 9.37%          |
| P. minutus R.Y. Zheng & G.Q. Chen | +              | –              | –               | –               | –              | –              | 1.56%          |
| P. oedipus Mont. | +              | +              | –               | +               | –              | +              | 14.06%         |
| Rhizopus arrhizus var. arrhizus A. Fisch. | –              | –              | –               | –               | –              | +              | 1.56%          |
| R. stolonifer (Ehrenb.) Vuill. | –              | –              | –               | –               | –              | +              | 7.81%          |
| Syncephalastrum racemosum Cohn ex J. Schröt | –              | –              | +               | +               | –              | +              | 12.50%         |
| Total taxa                      | 8              | 3              | 10              | 5               | 6              | 7              | 14             | 12             |
Fig. 1 – Dendrogram of Bray–Curtis similarity for species composition of Mucorales from the herbivore dung of different animals. Species composition was more similar between Guzerá and Sindi dung, followed by Girolando and Morada Nova dung. Excrements of Anglo-Nubiano and Holandés were less similar.

(Table 2). However, according to the $\chi^2$ test, there were no significant differences in the species richness of Mucorales in dung from the different breeds of animals ($p = 0.0926$), but differences in the number of species isolated across sampling per months of sampling were significant ($p = 0.0458$).

Most Mucorales occurred in the excrement of bovine, caprine and ovine, with the exception of Absidia cylindrospora var. cylindrospora, C. echinulata var. echinulata, Mucor sp. and P. minutus, which were only found in bovine dung, while Lichtheimia brasiliensis, M. variosporus, R. arrhizus var. arrhizus and R. stolonifer were only observed in sheep feces.

Considering that L. ramosa is difficult to be separated from L. corymbifera (Cohn) Vuill. and L. ornata (A.K. Sarbhoj) Alastr.-Izxq. & Walther based exclusively on morphological characters, the LSU rDNA region of our L. ramosa was sequenced. The Blastn analysis showed that our sequence (KX133009) was 99% similar to L. ramosa (CBS 582.65 – NG042518.1), confirming the identity of our specimen.

The species composition was most similar between Guzerá and Sindi (bovine) (75%), followed by Girolando (bovine) and Morada Nova (ovine) (63.63%) and Holandés (bovine) and Anglo-Nubiano (caprine) (60%) (Fig. 1).

**Identification key for species of Mucorales from dung in the semi-arid region of Brazil**

1. Obligatory coprophilous species; sporangiophores bearing subsporangial vesicles and trophocysts
2. Facultative coprophilous species; sporangiophores without subsporangial vesicles and trophocysts
3. Sporangiospores globose, subglobose or ovoid
4. Sporangiospores ellipsoid
5. Subsporangial vesicles ovoid; sporangiogospores globose or subglobose, thin-walled
6. Subsporangial vesicles obovoid; sporangiogospores thick-walled, spherical to broad ovoid
7. Trophocysts short, 180–610 × 125–270 μm; columellae cylindrical; sporangiogospores subglobose, 7–11 μm in diam.
8. Trophocysts long, 410–1800 × 240–420 μm; columellae conical; sporangiogospores subglobose, 10–18.5 μm in diam.
9. Columellae nipple-like; sporangiogospores ellipsoid, pale yellow or hyaline
10.5–17.5 × 5.5–8 μm
11. Sporangiofphores bearing sporangia
12. Sporangiofphores bearing merosporangia or sporangiola

(... continued)
|   | Description                                                                 | Species/Genus                                                                 |
|---|-----------------------------------------------------------------------------|-------------------------------------------------------------------------------|
| 7 | Merospores produced in merosporangia                                         | Syncephalastrum racemosum                                                     |
| 7 | Pedicellate unispored sporangia produced on a fertile vesicle                | Cunninghamella echinulata var. echinulata                                     |
| 8 | Sporangiophores simple or with erect or curved branches; sporangia without sterile spines | Cricinella muscae                                                            |
| 8 | Sporangiophores with circinate branches; sporangia with a sterile spine      | Mucor racemosus                                                               |
| 9 | Sporangia unapophysate; giant cells not produced                            | M. hiemalis                                                                   |
| 9 | Sporangia apophysate; giant cells present or absent                          | M. indicus                                                                    |
| 10| Sporangiospores regular in shape and size                                    | M. luteus                                                                     |
| 10| Sporangiospores with varied shapes and sizes, globose, ovoid, cylindrical or fusiform | Mucor variosporus                                                            |
| 11| Sporangiospores unbranched or slightly branched                              | M. racemosus f. racemosus                                                    |
| 11| Sporangiospores repeatedly branched                                          | M. racemosus f. racemosus                                                    |
| 12| Columellae obovoid; sporangiospores ellipsoidal, plano-convex 2.5–10 × 2–7.5 μm | M. racemosus f. racemosus                                                    |
| 12| Columellae globose; sporangiospores long-elliptical and fusiform 2.5–8.1 (12.5) × 1–5 μm | M. racemosus f. racemosus                                                    |
| 13| Mesophilic species, not growing at 40 ºC                                     | M. racemosus f. racemosus                                                    |
| 13| Thermotolerant, growing at 40 ºC                                             | M. racemosus f. racemosus                                                    |
| 14| Columellae regular in shape; sporangiospores globose, subglobose or ellipsoid | M. racemosus f. racemosus                                                    |
| 14| Columellae of several shapes; sporangiospores subpherical to ellipsoid       | M. racemosus f. racemosus                                                    |
| 15| Sporangiospores with swellings or not; sporangia with short lateral branches; columellae flattened, globose or ovoid; chlamydospores present or not, when present never abundant and never formed in reproductive structures | M. racemosus f. racemosus                                                    |
| 16| Sporangiospores without swellings and with long lateral branches; columellae subglobose, ovoid or ellipsoidal; abundant chlamydospores produced in sporangiophores and in a few columellae | M. racemosus f. racemosus                                                    |
| 16| Colonies high (up to 10 mm); sporangiospores curved or not; swellings under sporangia absent; columellae obovoid, globose or subglobose | M. racemosus f. racemosus                                                    |
| 16| Colonies low (up to 2 mm); sporangiospores curved or not; swellings often viewed under sporangia; columellae flattened | M. racemosus f. racemosus                                                    |
| 17| Colonies initially white, turning gray in older cultures; sporangiophores up to 7 (~10) μm in diam.; sporangia black | M. racemosus f. racemosus                                                    |
| 17| Colonies initially yellow, turning brownish in older cultures; sporangiophores up to 14 (~17) μm in diam.; sporangia gray-brownish | M. racemosus f. racemosus                                                    |
| 18| Sporangiospores sympodially branched; dark sporangia, globose (15–) 20–72.5 (~75) μm; sporangiospores ellipsoid | M. racemosus f. racemosus                                                    |
| 18| Sporangiospores sympodially or monopodially branched (rarely); sporangia dark brown, globose to slightly subglobose, 20–90 μm; sporangiospores globose to slightly subglobose | M. racemosus f. racemosus                                                    |
| 19| Sporangia dark brown and globose, 35–85 μm; columellae obovoid; sporangiospores ellipsoid | M. racemosus f. racemosus                                                    |
| 19| Sporangia brown and globose, 21.5–92.5 μm; columellae globose; sporangiospores ellipsoid ellipsoid and sometimes irregularly shaped | M. racemosus f. racemosus                                                    |
| 20| Sporangiospores arising from stolons, never opposed to rhizoids, spherical sporangia, piriform or subpiriform and apophysid; giant cells present | M. racemosus f. racemosus                                                    |
| 20| Sporangiospores arising from the aerial mycelium and/or stolons, opposed to rhizoids, sporangia apophysid, globose and subglobose; giant cells absent | M. racemosus f. racemosus                                                    |
| 21| Thermophilic species, growing at 40 ºC; subsporangial septum absent or rare; sporangiospores globose, subglobose or ellipsoid; giant cells present | M. racemosus f. racemosus                                                    |
| 21| Mesophilic species, not growing at 40 ºC; subsporangial septum present; sporangiospores cylindrical; giant cells present | M. racemosus f. racemosus                                                    |
| 22| Columellae globose, subglobose and spatulate, often exhibiting projections; giant cells present | M. racemosus f. racemosus                                                    |
| 22| Columellae subglobose and short hemispheric, without projections; giant cells absent | M. racemosus f. racemosus                                                    |
| 23| Rhizoids present, well developed, abundant and rhizopodiform; sporangiophores reaching 3 mm in length; columellae ovoid | M. racemosus f. racemosus                                                    |
| 23| Rhizoids, undeveloped, simple or rarely branched when present; sporangiophores reaching 1.7 mm in length; columellae subglobose, hemiglobose, rarely oblong-ovoid | M. racemosus f. racemosus                                                    |
**Discussion**

Altogether, 24 taxa of Mucorales were identified in the dung of herbivores from Arcoverde, Sertânia and Serra Talhada, PE, Brazil. All species identified in the present study are being cited for the first time in the dung of herbivores in Caatinga areas. However, most of the species identified herein have been reported by other authors in excrement from other domains in Brazil, indicating that they are not endemic to the Caatinga.\textsuperscript{14,15,30}

With the exception of *M. indicus* and *Mucor* sp., all taxa of this genus isolated in the present study as well as *Circinella muscae*, *P. crystallinus*, *P. kleinii*, *P. longipes*, *R. arrhizus* var. *arrhizus* and *S. racemosum* were reported by Alves et al.\textsuperscript{14} and by Santiago et al.\textsuperscript{15} in the dung of herbivores in the Atlantic Forest in Recife, PE. *Lichtheimia ramosa* (as *A. ramosa*), *C. muscae*, *M. hiemalis*, *P. crystallinus*, *P. kleinii* and *P. longipes* were described by Trufem\textsuperscript{36} and Trufem and Virlato\textsuperscript{15} in the Atlantic Forest in São Paulo state. Fifteen species reported by the above mentioned authors were observed in the present study, although feces from different herbivores were analyzed in different studies, indicating that the majority of Mucorales do not exhibit specificity for a particular type of animal dung.\textsuperscript{31-33}

A recent phylogenetic study on Mucorales that included critical species of this order\textsuperscript{34} strongly evidenced that *M. ramosissimus* is a synonym of *M. circlinelloides* f. *circlinelloides* or *f. janssennii*. In fact, both species are morphologically very similar to each other. However, considering that our isolates exhibited morphological characteristics, such as small colonies up to 2 mm high, sporangioaphores with a frequent swelling below the sporangia and columellae applanate, that were very similar to those described by Schipper\textsuperscript{35} for *M. ramosissimus*, and considering that *M. ramosissimus* is still a valid species, we prefer to maintain *M. ramosissimus* in our manuscript.

Some of the species isolated in the present study have also been reported in animal dung from other countries. Masunga\textsuperscript{36} isolated *C. elegans* and *P. crystallinus* from the dung of elephants in Africa, while Abdullah\textsuperscript{37} reported *P. kleinii* in donkey, sheep and camel dung collected in Iraq. Studies have shown that this group of fungi has a wide distribution and can adapt to different environmental conditions.\textsuperscript{31,38,39}

*Lichtheimia brasilienensis* and *M. indicus* have been commonly isolated from soils, although they have not as yet been found in dung.\textsuperscript{23,40,41} However, these species are now reported for the first time in the dung of herbivores, thereby expanding our knowledge about the diversity of Mucorales. *Mucor* sp. exhibit morphological and genetic characteristics that differ from other taxa of the genus and will be described and published as a new species in a subsequent paper.

Among the isolated genera, *Mucor* was the most representative in number of taxa, with nine species and four forms, followed by *Pilobolus*, with five species. According to Krug et al.,\textsuperscript{6} *Mucor* is characterized by facultative coprophilous species, while *Pilobolus* is characterized by obligatory coprophilous species. Since the species of the former genus are very common in soil samples,\textsuperscript{42} it is possible that the presence of these taxa in the inventoried excrement was due to the transition of propagules from the soil to the dung, as mentioned by Santiago et al.\textsuperscript{15} All species of Mucorales reported here are being cited for the first time as coprophilous in the Caatinga. In Brazil, the occurrence of *Mucor* species and/or *Pilobolus* in herbivore dung was reported by Batista and Pontual,\textsuperscript{43} Trufem,\textsuperscript{30} Trufem and Virlato,\textsuperscript{15} Richardson,\textsuperscript{34} Alves et al.\textsuperscript{31} and Santiago et al.\textsuperscript{15}. The greatest representation of *Mucor* and *Pilobolus* in relation to the number of taxa in the dung analyzed was confirmed by Santiago et al.\textsuperscript{35}

Our results indicated that Morada Nova dung (ovine) was the richest in terms of the number of taxa, followed by the dung of Santa Inês (ovine) and Holandês (bovine). However, according to the $\chi^2$ test, no significant difference ($p=0.0926$) was found for the species richness of Mucorales in the dung of the different animals analyzed. According to Ebersohn and Eicker\textsuperscript{18} and Santiago et al.\textsuperscript{15} differences in the composition of a community can be correlated with abiotic and biotic factors that influence the mycobiota of the substrate. The fact that dung samples were kept in similar experimental conditions (temperature, light incidence, moisture) and were free of mycophagous insects could explain the similarities found for the species richness of Mucorales in dung. However, statistical analysis revealed significant differences in the number of taxa isolated from dung between the different sampling months ($p=0.0458$), indicating that seasonality influences the number of species, but not the composition of the Mucorales. The opposite result was observed by Santiago et al.,\textsuperscript{15} who reported that the composition of Mucorales species was affected by seasonal changes of the year, differing from the results observed for the number of taxa. Bell\textsuperscript{44} conducted a three-year study on the mycota in possum dung (*Trichosurus Vulpecula* Kerr) in New Zealand and attributed the higher incidence of certain fungal species to winter rains.

Concerning the frequency of occurrence of the isolates, *M. circlinelloides* f. *griseo-cyanus* was the most common taxon (FO = 93.35%), while *C. echinulata* var. *echinulata*, *Mucor* sp. and *R. arrhizus* var. *arrhizus* were the least common taxon (FO = 1.56%). *Mucor* and *Rhizopus* include many species, coprophilous or not,\textsuperscript{6} while *Cunninghamella* species primarily colonize other substrates.\textsuperscript{42} Most species identified in the present study exhibited low frequencies of occurrence, not exceeding 32.8%. Similar behavior was observed by Richardson,\textsuperscript{34} Nyberg and Persson\textsuperscript{32} and Santiago et al.\textsuperscript{15} in studies of coprophilous fungi from herbivorous dung. *Pilobolus kleinii* has been commonly associated with a high frequency in herbivorous animal dung in other domains.\textsuperscript{31,32} However, this species was uncommon in the samples analyzed in the present study (7.81%). It is possible that this taxon is more sensitive to the water and temperature stress that are characteristic of the semi-arid region.

Considering the different breeds of herbivores, species composition was more similar between Guzerá and Sindi (bovines) (75%), which although not sharing the same environment, were exposed to a similar nutritional status and were the only animals kept on pasture (Table 1). According to Santiago et al.,\textsuperscript{15} nutritional differences between the animals can influence the mycota of dung. The fact that these animals spent most of their time grazing in the pasture may have contributed to the high similarity found. Curiously, high similarity was also observed between the Girolando (bovine) and Morada Nova (ovine) species, and between Anglo-Nubiano (caprine) and Holandês (bovine) species, despite the fact that they were grazed in different cities (although in the same domain) and...
have different diets. Studies have shown that although the food given to the animals is a limiting factor for the appearance of certain taxa, other factors, such as the geographic location, humidity and high temperatures may influence the composition of fungal communities in animal dung.\textsuperscript{31,38,45}

The present work reports 24 taxa of Mucorales in herbivore dung from the semi-arid of Brazil and is a pioneer study for coprophilous Mucorales in the Caatinga. Considering the adverse conditions of temperature and humidity, typical of semi-arid regions, and comparing the results described herein with those of other authors, it is clear that the species richness of Mucorales in the inventoried areas is high. Abdullah\textsuperscript{37} reported only three species of Mucorales in donkey, sheep and camel dung collected in semi-arid regions of southern Iraq. Masunga et al.\textsuperscript{36} reported six species of this group in semi-arid areas of Botswana, Africa. Upon comparison of the results of the present study with previous studies of the diversity of coprophilous Mucorales in Atlantic Forest areas, Santiago et al.\textsuperscript{21,15} reported 39 taxa in this domain in Pernambuco, whereas Trufem and Viriatio\textsuperscript{33} and Viriato and Trufem\textsuperscript{46} together only obtained 23 taxa in the same domain in São Paulo. Thus, it is relevant to continue surveying the fungal diversity in the Caatinga in order to demystify the erroneous idea that the richness of fungal species in this area is low.

Conflicts of interest

The authors declare no conflicts of interest.

Acknowledgments

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REFERENCES

1. Hibbett DS, Binder M, Bischoff JR, et al. A higher-level phylogenetic classification of the Fungi. Mycol Res. 2007;101:509–547.
2. Hoffmann K, Voigt K, Kirk PM. Mortierellomyctota subphyl. nov., based on multi-gene genealogies. Mycotoxnon. 2011;115:353–363.
3. Humber RA. Entomophthoromyctota: a new phylum and reclassification for entomophthorid fungi. Mycotoxnon. 2012;120:477–492.
4. Santiago ALCM de A, dos Santos PJ, Maia LC. Mucorales from the semiarid of Pernambuco, Brazil. Braz J Microbiol. 2013;44(1):299–305.
5. Alexopoulos CJ, Mims CW, Blackwell M. Introductory Mycology. 4th ed. New York: John Wiley & Sons; 1996.
6. Krug JC, Benny GL, Keller HW. Coprophilous fungi. In: Mueller GM, Bills GF, Foster MS, eds. Biodiversity of Fungi. Hardbound: Elsevier Academic Press; 2004:468–499.
7. Richardson MJ. Records of coprophilous fungi from the Lesser Antilles and Puerto Rico. Caribb J Sci. 2008;44(2):206–214.
8. Dix NJ, Webster J. Fungal Ecology. London: Chapman & Hall; 1995.
9. Richardson MJ. Coprophilous fungi. Field Mycol. 2003;4(2):41–43.
10. Drumond MA, Kiill LHF, Nascimento CES. Inventário e sociabilidade de espécies arbóreas e arbustivas da Caatinga na Região de Petrolina, PE. Bras Florest. 2002;74:37–43.
11. Gusmão LFP, Barbosa FR, Barbosa FF. Fungos Conídios. In: Gusmão LFP, Maia LC, eds. Diversidade e caracterização dos fungos no semi-árido. Vol 1. Recife: Associação Plantas do Nordeste; 2006;27–47.
12. Santiago ALCM de A. Mucorales. In: Lista de Espécies da Flora do Brasil. Jardim Botânico do Rio de Janeiro.; 2015. Available from http://floradobrasil.brd.gov.br/jabot/floradobrasil/bf120276.
13. Trufem SFB, Viriato A. Mucorales do Estado de São Paulo. Mucoraceae coprófilas. Rikia. 1985;12:113–123.
14. Alves MH, Trufem SFB, Milanez AI. Táxons de Mucor Fresen. (Zygymycota) em fezes de herbívoros, Recife, PE, Brasil. Rev Bras Bot. 2002;25(2):147–160.
15. Santiago ALCM de A, Trufem SFB, Malosso E, dos Santos PJP, Cavalcanti MA de Q. Zygomycetes from herbivore dung in the ecological reserve of Dois Irmãos, Northeast Brazil. Braz J Microbiol. 2011;42:85–95.
16. Benny GL. The methods used by Dr. R.K. Benjamin, and other Mycologists to isolate Zygomycetes. Aliso. 2008;26:37–61.
17. Hesseltine CW, Ellis JJ. The genus Absidia: Gongonella and cylindrical-spored species of Absidia. Mycologia. 1964;56:659–610.
18. Schipper MA. On certain species of Mucor with a key to all accepted species. Stud Mycol. 1978;17:1–69.
19. Hesseltine CW, Fennel DI. The genus Cincinella. Mycologia. 1995;47:193–211.
20. Zheng R-y, Chen G-q. A monograph of Cunninghamella. Mycologia. 2001;83:1–75.
21. Hoffmann K, Discher S, Voigt K. Revision of the genus Absidia (Mucorales, Zygomycetes) based on physiological, phylogenetic, and morphological characters; thermostolerant Absidia spp. form a coherent group, Mycocladiae fam. nov. Mycol Res. 2007;111:1169–1183.
22. Zheng R-y, Chen G-q, Huang H, Liu X-y. A monograph a Rhizopus. Sydowia. 2007;59(2):273–372.
23. Santiago ALCM de A, Hoffmann K, Lima DX, et al. A new species of Lichtheimia (Mucoromycotina, Mucorales) isolated from Brazilian soil. Mycol Progr. 2014;13:343–352.
24. Gőes-Neto A, Loguericio-Leite C, Guerrero KT. DNA extraction from frozen field collected and dehydrated herbivoral fungal basidiomata: performance of SDS and CTAB-base methods. Biotemas. 2005;18:19–32.
25. van Tuinen D, Zhao B, Giannazzi-Pearson V. PCR in studies of AM fungi: from primers to application. In: Varma AK, ed. Mycorrhizal Manual. Berlin: Springer; 1998:387–399.
26. Oliveira RJV, Lima TEF, Cunha IB, et al. Cornicularia brasiliensis, a new species of coelomycetes in the rhizosphere of Caesalpinia echinata (Fabaceae, Caesalpinioideae) in Brazil. Phytopath. 2014;138(3):197–204.
27. Brower JC, Zar JH, Von Ende CN. Field and Laboratory Methods for General Ecology. Dubuque: McGraw Hill; 1990.
28. Clarke KR, Gorley RN. Primer v6 User Manual/Tutorial. Plymouth: Primer-E Ltd; 2006.
29. Melo RFR, Bezerra JL, Cavalcanti MA de Q. Diversity of coprophilous ascomycetes from captive wild animals in Dois Irmãos State Park, Brazil. Nova Hedwig. 2012;94(2):153–162.
30. Trufem SFB. Mucorales do Estado de São Paulo. 4. Espécies coprófilas. Rikia. 1984;11:65–75.
31. Richardson MJ. Diversity and occurrence of coprophilous fungi. Mycol Res. 2001;105(4):387–402.
32. Nyberg Å, Persson IL. Habitat differences of coprophilous fungi on moose dung. Mycol Res. 2002;106(11):1360–1366.
33. Delgado-Ávila AE, Urdaneta-García LM, Piñeiro-Chávez AJ. Coprophilous fungi of Zulia state, Venezuela. Divisions: Myxomycota, Zygomycota and Basidiomycota. Rev Cien Fac Cien Vet Univ Zulia. 2005;15(1):5763.
34. Walther G, Pawłowska J, Alastruey-Izquierdo A, et al. DNA barcoding in Mucorales: an inventory of biodiversity. Persoonia. 2013;30:11–47.
35. Schipper MAA. On Mucor circinelloides, Mucor racemosus and related species. Stud Mycol. 1976;12:1–40.
36. Masunga GS, Andresen Ø, Taylor JE, Dhillion SS. Elephant dung decomposition and coprophilous fungi in two habitats of semi-arid Botswana. Mycol Res. 2006;110:1214–1226.
37. Abdullah SK. Coprophilous mycoflora on different dung types in southern desert of Iraq. Sydowia. 1982;35:1–5.
38. Ebersohn C, Eicker A. Determination of the coprophilous fungal fruit body successional phases and the delimitation of species association classes on dung substrates of African game animals. Bot Bull Acad Sin. 1997;38:183–190.
39. Caretta G, Piontelli E, Savino E, Bilgheroni A. Some coprophilous fungi from Kenya. Mycopathologia. 1998;142:25–34.
40. Szwedek-Trzaska A, Glowacka A. Seeking ways to eradicate potentially pathogenic fungi isolated from soil. Pol J Environ Stud. 2011;20(5):1313–1318.
41. Lima DX, Santiago ALCM de A, Souza-Motta CM. Diversity of Mucorales in natural and degraded semi-arid soils. Rev Bras Bot. 2015;38:1–7.
42. Domsch KH, Gams W, Anderson TH. Compendium of Soil Fungi. San Francisco: IHW-Verlag; 2007.
43. Batista AC, Pontual D. Alguns fungos coprófilos de Pernambuco. Bol Secr Agric Ind Comér. 1948;15:27–44.
44. Bell A. Fungal succession on dung of the brush-tailed opossum in New Zealand. N Z J Bot. 1975;13:437–462.
45. Caretta G, Piontelli E. Coprophilous fungi from confined deers in Pavia (Lombardia, Italy). Bol Micol. 1996;11:41–50.
46. Viriato A, Trufem SFB. Mucorales de São Paulo. Espécies Merosporangiadas. Rickía. 1985;12:147–154.