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Mycological studies as a tool to improve the control of building materials biodeterioration

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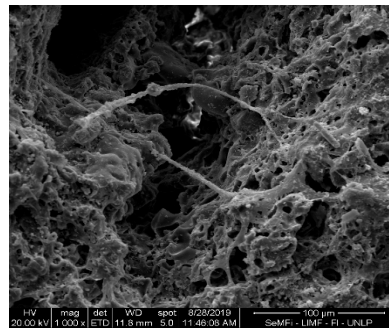
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Fungal
Isolation

Purification and taxonomic identification

Ecological category and biodeterioration assays

C. sphaerospermum
MN371394

A. spectabile
MT071822

A. niger
MN371276

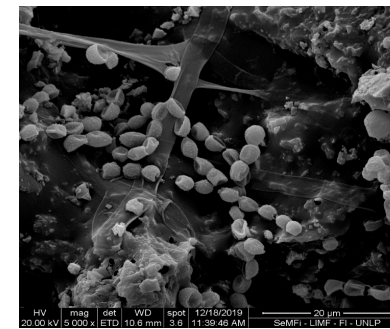*P. commune*
MN371392

L. theobromae
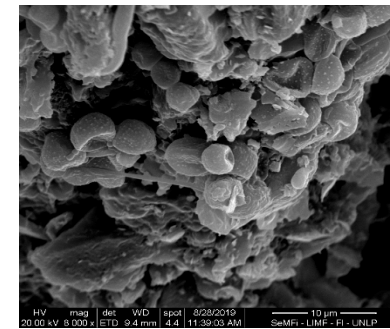MN371283
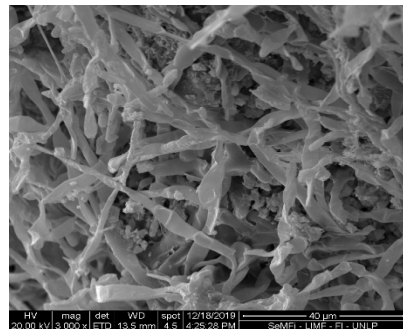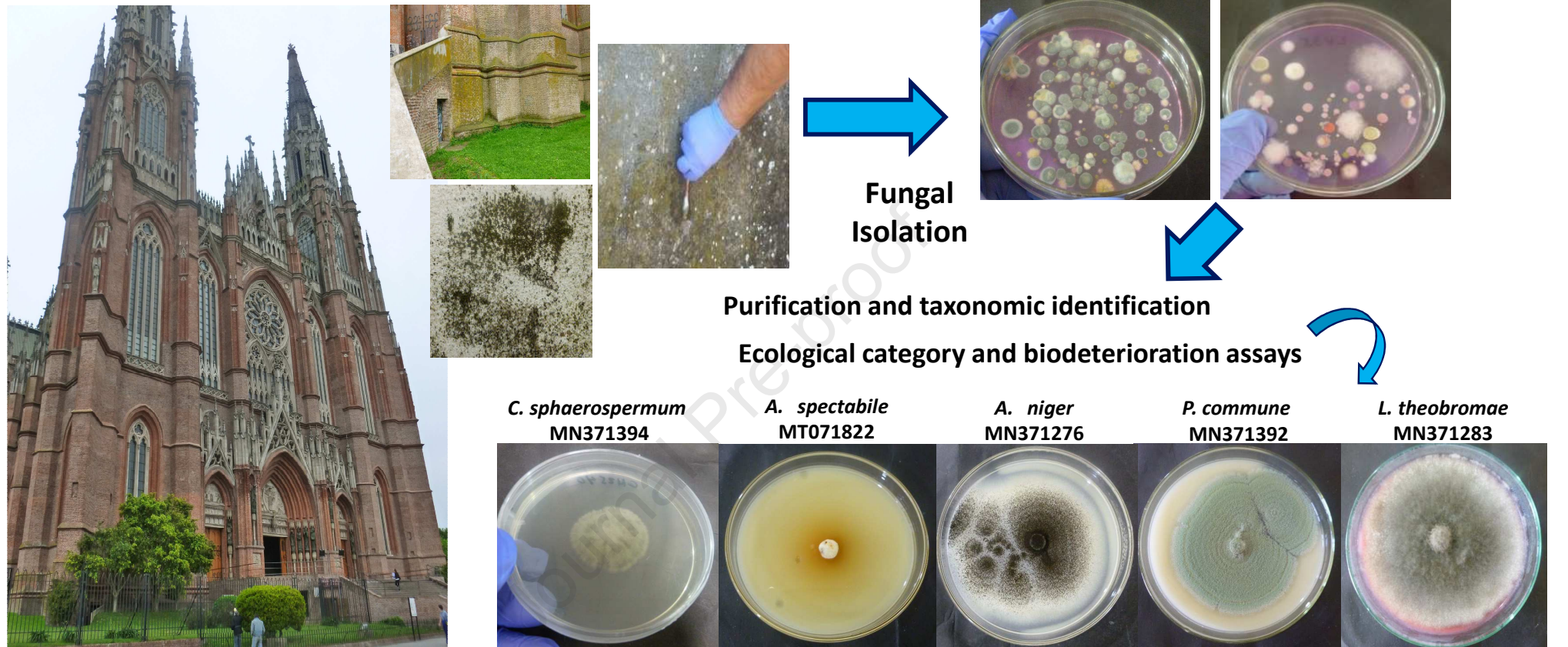
C. sphaerospermum

A. spectabile
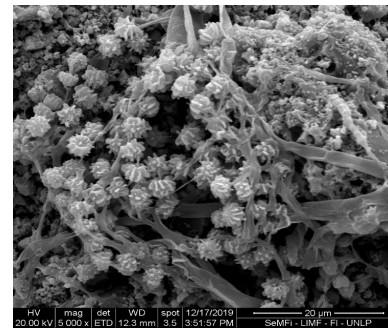
A. niger

P. commune

L. theobromae

Mycological studies as a tool to improve the control of building materials biodeterioration

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Abstract

Fungal deterioration of building materials causes economic losses due to restoration processes and because it affects the health of the people exposed. Therefore, it is necessary to obtain highly deteriorating fungal strains to carry out studies focused on biodeterioration control. The aim of this research was to study the mycobiota involved in deterioration of facades through the isolation and characterization of fungal strains. In this sense, this paper characterizes the mycobiota that contributes to the deterioration of brick and concrete present in those places. Samples were taken from bricks and concrete with visible biodeterioration of the Cathedral of La Plata and the Experimental Station of the Centro de Investigación y Desarrollo en Tecnología de Pinturas. Relative density, apparition relative frequency, and ecological category of fungi isolates were determined. The deteriorating attributes were determined by carbonate dissolution test, pigment secretion assay, acid production test and brick weight variation. *Cladosporium* sp., *Penicillium* sp. and *Aspergillus* sp. were the taxa with the highest apparition relative frequency and abundance. *Aspergillus niger* MN371276, *Lasiodiplodia theobromae* MN371283, *Cladosporium sphaerospermum* MN371394, *Penicillium commune* MN371392 and *Arthothelium spectabile* MT071822 were the strains with the highest biodeteriorant activity. These strains can be used in studies on biodeterioration control of buildings materials.

Keywords: building materials, biodeterioration, filamentous fungi.

1. Introduction

Fungi are ubiquitous heterotrophic organisms able to endure extreme environmental conditions and to adopt several structural, morphological and metabolic strategies to survive [1]. Filamentous fungi are frequently described as deterioration agents of building materials in both indoor and outdoor environments [2,3]. In the built environment, filamentous fungi are associated with physical and aesthetic damages and human health issues such as allergies and toxic reactions. Related problems such as asthma, cough, wheezing, and upper respiratory infections are also described [4]. Fungi are microorganisms with increased survival capacity on surfaces with low water content and can tolerate intense ultraviolet radiation. Solid substrates exposed outdoors are subject to weathering and also to biological growth on their surfaces, which causes undesirable alterations known as biodeterioration [5,6]. The main biological damage is caused by aesthetic biodeterioration due to the production of organic pigments and coverage of the surface by colored biofilms; physical biodeterioration resulting from mechanical forces produced by mycelium within pores that cause the material breakup, and chemical biodeterioration due to the direct action of the metabolic products excreted [7,8].

The spoilage of structural materials by filamentous fungi causes negative effects on the preservation of historic-cultural and work buildings. An important example of this is the Cathedral of the Immaculate Conception, also known as the Cathedral of La Plata (CP). The CP is affected by growth of lichens, green algae, cyanobacteria, mosses and higher plants [9,10]. However, the biodeterioration by filamentous fungi has not been studied. Regarding the biodeterioration of cultural heritage, it is necessary to evaluate levels of fungal contamination in urban monuments and fungi resistance to certain materials or conservation treatments [11].

In work buildings, it is also important to monitor the deterioration of structural materials by fungi, since they can cause indoor and outdoor air pollution. The mycobiota of indoor environments contains about 150 species and it is considered that 1000 CFU/m^{-3} is a high level of exposure for an indoor occupant of a building [12] . The fungal genera most frequently found in indoor

environments are *Cladosporium*, *Penicillium*, *Aspergillus*, and *Stachybotrys* [13].

The aim of this research was to study the mycobiota involved in deterioration of facades through the isolation and characterization of fungal strains. First, the present research work was focused on the study of the mycobiota involved in the deterioration of brick and concrete in facades by the determination of relative density, apparition relative frequency and ecological category of the fungal isolates. Then, the other goal was to assess the most deteriorating isolated strains by different tests such as calcium carbonate dissolution test, acid production test, pigment secretion assay and brick weight variation in humidity chamber. Molecular analysis of the most deteriorating fungal strains was performed. These strains will be used in further research to study new bioactive materials in the control of building biodeterioration.

2. Materials and methods

2.1 Sampling sites and fungal isolation

Microbiological sampling was carried out during October 2017 (spring time) at the Cathedral of La Plata (CP) (34º55´S, 57º57´O) and at the Centro de Investigación y Desarrollo en Tecnología de Pinturas (CIDEPINT) Experimental Station (CES) (34º50´S, 57º53´O) (Fig. 1). The CP is the main Catholic temple in La Plata city (capital of Buenos Aires province), Argentina and one of the largest churches in the world. It is a neogothic building with an area of 7000 m^2 built with exposed brick and mortar, giving it a very special characteristic appearance. White cement mortar was used in the pinnacles and supporting arches of the vaults and ornaments. CIDEPINT is a center dedicated to obtaining new technological developments related to paints and protective coatings. It has an outdoor experimental station where natural service tests are carried out.

Samples were taken using the swab technique from bricks and concrete with significant biodeterioration at 0.5 and 1.5 meters high with respect to the ground [10,13]. Subsequently, the swabs were introduced into tubes containing sterile physiological solution (0.85 g of NaCl (Anedra, Argentina) /100 mL of sterile distilled water). Serial dilutions (1:10-1:1000) were carried out, and 0.1 mL of

the solution was inoculated on Petri dishes with Potato Glucose Agar (Britania, Argentina), streptomycin (30 mg/100 mL) and Rose Bengal (Cicarelli, Argentina) (5 mg/100 mL). The plates were maintained at 28°C for 15 days and, after close observation, filamentous fungi were isolated at 2, 7 and 15 days following conventional microbiological techniques.

2.2 Taxonomic identification and ecological category of isolates

For the taxonomic analyses, all strains were grown on Potato Dextrose Agar (Britania, Argentina) and 39 g of distilled water per liter were used. The isolates were identified to the lowest possible taxon based on their reproductive, somatic structures and colony cultural characteristics using standard taxonomic keys [14,15]. In some cases, the microculture technique was used. In others, parts of mycelium were taken from mature colonies and observed under light microscope. In order to better observe the conidiophores alcohol was added to the highly sporulated preparations.

Relative density (RD) [16] and apparition relative frequency (RF) were calculated using formula (1) and formula (2), respectively:

RD(%)= (a/b) x 100   (1)

where a is Number of colonies of the taxon, b is Total of colonies of taxa

RF(%)= (c/d) x 100   (2)

where c is Times a taxon is detected, d is Total number of sampling realized

From RF values of the fungi, the following ecological categories were given: abundant (81-100 %); common (61-80 %); moderate (41-60 %); occasional (21-40 %); and rare (0.1-20 %), taking into account what was proposed by Esquivel et al. [17].

2.3 Biodeterioration assays

2.3.1 Inoculum preparation

The spore suspension was carried out from mature colonies. First, physiological solution (0.85 g of NaCl/100 mL of sterile DW) was added to the colonies and finally, it was carefully removed by inoculation loop so as not to drag mycelium. The inoculum concentration was adjusted to 10^5 spores/mL employing a Neubauer chamber.

2.3.2 Calcium carbonate (calcite) dissolution test

The procedure followed was similar to that proposed by Unković et al. [18]. The potential ability of the tested fungi to degrade calcite was screened using Petri dishes with culture medium of the following composition, per liter of distilled water: 5 g of calcium carbonate (Anedra, Argentina); 30 g of malt extract (Circe, Argentina); 3 g of mycological peptone (Oxoid, United Kingdom), and 15 g agar (Parafarm, Argentina). The Petri dishes with $CaCO_3$ malt extract agar were inoculated with agarized 6 mm diameter mycelium disc and incubated for 14 days at 28°C. Positive strains displayed a clear zone around the colony, thus confirming calcium carbonate dissolution. The test was performed in triplicate.

2.3.3 Acid production test

The strains that resulted positive to the carbonate dissolution test were selected to perform the acid production test. To determine the capacity of fungi affecting the pH value of the substrate on which they grow, isolates were cultivated in a Liquid Minimal Medium (LMM) supplemented with glucose 1% and phenol red 0.001% of the following composition, per liter of distilled water: sodium nitrate, 2 g; potassium phosphate, 1g; magnesium sulfate, 0.5 g; ferrous sulfate, 0.01 g and potassium chloride, 0.5 g. The brand of the previous reagents was Anedra, Argentina with the exception of potassium chloride (Cicarelli, Argentina). Before adding the inoculum, the pH was adjusted to 7 and the control was test tubes with spore-free liquid medium. Test tubes with phenol red LMM were inoculated with a spore solution (10^5 spores/ mL) and incubated for 3 days at 28 °C when phenol red pH-indicator showed a change in color [19]. The test was performed in triplicate.

2.3.4 Pigment secretion

The potential ability of the isolates to secrete organic pigments under nutrient-limited conditions (present on the surface of structural materials), and consequently to induce alterations in the original coloration was screened using Petri dishes with culture medium similar to those used in the calcium carbonate dissolution test. The Petri dishes were incubated for 7 days at 28°C. Secretion of fungal pigments was confirmed by changes in color of medium through direct observation. The assay was performed in triplicate.

2.3.5 Exposure studies in Humidity Chamber

The test was carried out according to the procedure of George et al. [20]. The criteria used to select the strains in this assay were the capacity to secrete pigments and the high growth speed. Bricks of 19 ± 1.0 cm^3 were introduced in Petri dishes, each one was inoculated with 200 µL of spore solution (10^5 spores/ mL). Two sets of three bricks were inoculated by each of the strains studied in order to remove the panels after 6 and 12 months. The brick samples were placed in a Humidity chamber at 100% relative humidity. Room temperature of 25°C was maintained inside the chamber. After removal, the bricks were weighted with and without biofilm.

For the fungal biofilm elimination, the bricks were immersed in sodium hypochlorite (10%) solution and dried to constant weight at 60°C. Control bricks without inoculation were used.

The weight loss was calculated from:

$$Dw = W_1 - W_2 \quad (3)$$

where W1 is the weight of the brick before introducing it into the chamber and W2 the weight after treatment with sodium hypochlorite.

After one year, the bricks that lost the most weight were observed by scanning electron microscopy (Philips FEI Quanta 200) and the working condition was high vacuum (10^{-6} torr).

2.4 Molecular analysis of the most deteriorating fungal strains

Taxonomic identification was confirmed by molecular methods based on the analysis of sequences coding for ITS (internal transcribed spacer) rRNA region only in the most deteriorating fungal isolates. The ITS sequences were amplified through PCR (polymerase chain reaction) and then, they were sequenced and analyzed [21]. The primers used were ITS1 (5´-TCC GAT GGT GAA CCT GCG G-3´) and ITS4 (5´-TCC TCC GCT TAT TGA TAT GC-3´) [22,23]. This procedure was carried out by Macrogen (www.macrogenlatam.com.ar). The sequences obtained were analyzed by means of the genetic data bank (www.ncbi.nlm.nih.gov/blast/Blast.cgi) using a

search and sequence analysis software to find the closest known analogues with 99% similarity values. In addition, evolutionary history was inferred using the Unweighted Pair Group Method using Arithmetic Averages (UPGMA). This tree-making method assumes that the rate of evolution has remained constant throughout the evolutionary history of the included taxa [24]. The evolutionary distances were computed using the p-distance method and were in the units of the number of base differences per site. This analysis involved 5 nucleotide sequences. Codon positions included were 1st+2nd+3rd+Noncoding. All ambiguous positions were removed for each sequence pair (pairwise deletion option). Evolutionary analyses were conducted in MEGA 10.0.5® [25].

2.5 Statistical analysis

Statistical Analysis of brick weight variation was performed with the RStudio 1.1.463® program. All assays were performed in triplicate and the probability distribution was analyzed by Normal distribution test. A two-way ANOVA analysis was performed. The included factors were time and treatment. The former had two levels: 6 and 12 months and the latter had 8 levels, which is the number of strains tested. Then, Tukey test was applied. Both tests were applied using the multcomp package. Similarly, this program was used to analyze the association between certain variables and the mycobiota present in the sampling sites using Principal Component Analysis (PCA). Specifically for these analysis packages, MASS, ggplot2, scales and factoextra were used.

3. Results and Discussion

3.1 Fungal isolation and ecological category

The swab technique and conventional microbiological techniques allowed the isolation and identification of 43 filamentous fungi. Table 1 presents a list of the isolates belonging to phyla Mucoromycota, Ascomycota or Basidiomycota. These fungal isolates were recorded for the first time on brick and concrete in CP and CES. Fig. 2 presents the RD and the RF from CP and CES. In Fig. 2A, it can be seen that the genera with the highest relative frequency in CP were *Cladosporium* sp., *Penicillium* sp. and *Aspergillus* sp. and together with *Alternaria alternata* were considered abundant. On the other hand, the appearance *of Epicoccum nigrum, Arthothelium spectabile* and *Lasiodiplodia*

theobromae was common in these materials. However, in CES the fungal isolates were classified as abundant, common, moderate and occasional (Fig. 2B). In this sampling place, DUSM 1 and *Chrysonilia* sp. were the strains with the highest relative frequency. In addition, YUSM (yellow unsporulated septate mycelium) and *C. cladosporioides* presented the highest relative density in CP and CES, respectively. Similar fungal strains were reported by Mallo et al. [26] for the city of La Plata in a study related to the presence of airborne fungi detected by various sampling methods. At the same time, Negrin et al. [27] reported that *Cladosporium* sp. was predominant in all the samples collected, followed by genera *Alternaria*, *Penicillium*, *Aspergillus* and *Epicoccum* in bioaerosols in La Plata . In the same way, filamentous fungi were reported in a study that evaluated the development of bioincrustations on marble and mortar surfaces of several historically and architecturally valuable crypts at the cemetery of La Plata [28]. Some of the isolates may be potentially pathogenic for the visitors of such locations. As mentioned in the literature review, filamentous fungi are associated with human health issues such as allergies, toxic reactions and upper respiratory infections [4].

Some of these species are involved in the biodeterioration of building materials worldwide (stone, mortars, glass, marble, etc) for example, *Aspergillus, Penicillium* and *Cladosporium* [2,29,30]. In addition, *L. theobromae, A. spectabile* and *Moniliella* sp. were detected for the first time on concrete and bricks.

There are variations between the number of taxa and the ecological categories for the sampling sites (Fig. 2). These results may be due to parameters related to outdoor climate, petrochemical pollution and differences in the chemical composition of the materials. In this sense, near La Plata city there are Petrochemical Plants located approximately 2.5 and 5 km away from CES and CP, respectively. In a work related to petrochemical pollution [31], different pollutant levels were studied. The authors observed that the concentrations of hexane, cyclohexane, benzene, and toluene were particularly high in the industrial region. In this sense, the presence of these pollutants could influence the mycobiota found in the sampling sites, with greater impact on CES due to the higher proximity to Petrochemical Plants. On the other hand, Diulio et al.

[32] reported, according to the hourly weather data obtained from the Astronomical Observatory of La Plata for 2017, that spring and fall are transition seasons, featuring warm daytime temperatures and cool nighttime temperatures. These temperatures can significantly vary, some days registering temperatures above 32°C and below 0°C. Owing to its riverside location, the city is fairly humid, and its average monthly humidity is higher than 75%. La Plata receives 1092 mm of precipitation annually, with winters being the driest months, and summer the wettest. These climatic conditions may also influence the presence of these strains at the sampling locations. For example, depending on the water activity of the material, certain species prevail. In this sense, secondary (e.g. *Cladosporium*, *Phoma*, *Ulocladium* and *Alternaria*) and tertiary (e.g. *Chaetomium*, *Trichoderma*, *Auraeobasidium* and *Stachybotrys*) colonizers require higher water activity to grow [33]. Taking into account the aforementioned data related to La Plata city which prove high humidity levels throughout the year, it can be seen a correlation with the results obtained since the genus *Cladosporium* classified as secondary colonizer was predominant in all the samples followed by other two with the same classification (*Alternaria* and *Epicoccum*). Finally, there are differences between the chemical composition of concrete and brick. Concrete is a composite material made from cement, aggregates and additives [34]. However, brick is a clayey material generally composed of iron oxide, calcium, and magnesium [7]. Thus, differences in structure and composition of the materials may also influence the abundance and relative frequency of the species at the sampling sites.

3.2 Biodeterioration assays

3.2.1 Calcium carbonate dissolution test

Table 1 shows the results of carbonate degradation test and pigment secretion assay. Only *Penicillium* sp. 1, *Penicillium* sp. 2, *Penicillium commune*, *Penicillium* sp. 3 and *Aspergillus niger* were the strains with capacity to degrade $CaCO_3$, and this can be seen in Fig. 3. It is known that mainly micromycetes from *Aspergillus* and *Penicillium* genera have calcite dissolution activity [18]. It is important to take into consideration that calcite degradation depends on various factors, primarily, on available sources of carbon and nitrogen, and thus, we cannot predict the activity of isolated *in vivo* species [35].

3.2.2 Acid production test

The acid production test was exclusively performed on the strains that resulted positive to the carbonate dissolution test. At the end of this test, the highest decrease in the pH related to control tubes (pH=7.13 ±0.01) was caused by *A. niger* followed by *P. commune* and *Penicillium* sp. 3 showing average pHs with standard deviation equal to 3.27 ±0.06, 5.42 ±0.02 and 5.85 ±0.07, respectively. On the other hand, *Penicillium* sp. 1 and 2 presented a lower pH modification (7.00±0.05) with no difference between them. The pH decrease in the inoculated MML may well be related to the secretion of organic acids by the fungi into the surrounding medium. This would be in agreement with bibliographic data that point out that the secretion of a great variety of organic acids (oxalic, oxalacetic, citric, gluconic, oxoacetic and fumaric) by different fungi makes them highly deteriorating agents for minerals and rocks [6,36]. Organic acids manage to form complexes with calcium, aluminum, iron, manganese, silicon and magnesium ions contributing even more to the degradation of the material [28,37].

3.2.3 Pigment secretion

E. nigrum, *L. theobromae*, YUSM, *A. spectabile* and OUSM (orange unsporulated septate mycelium) showed the ability to secrete organic pigments in the assay conditions. Most of the isolates produced pigments in various shades of yellow (Fig. 4A, D and E), while red pigments were present in two cultures (Fig. 4B and C). Pigment secretion is a process related to the biology of fungi and it can increase their biodeteriogenic attributes by making them more resistant to certain environmental conditions. Possibly not related to physical or chemical material deterioration, pigment secretion is closely related to aesthetic deterioration because it can change the normal coloration of building materials. In this sense, Trovão et al. [1] showed that *E. nigrum* has pigmented aerial mycelium and reproductive structures and its proliferation can contribute to aesthetic alterations in cathedrals. This damage can also be caused by other dematiaceous fungi, for example *L. theobromae*. However, not only pigment secretion can lead to visual deterioration but also an extensive mycelium development which also causes physical-chemical deterioration. Fungal pigments protect them against many physical and chemical environmental

stressors (high temperatures, DNA damage, accumulation of reactive oxygen species, desiccation and osmotic stresses). For example, the capture of and/or reflected radiation energy by pigments has multiple biological functions ranging from camouflage or makeup to fundamental roles in the maintenance of life including harnessing solar energy for metabolic use and protection against radiation damage [38]. Furthermore, some pigments, such as melanin, can reinforce the cell wall and increase the virulence of pathogenic fungi in plants and animals, reducing their susceptibility to biocides [39]. Another study supports the idea that pigments are then related to biofilm development. Rock-inhabiting Dothideomycetes and Eurotiomycetes have equal number of photoreceptors and the same set of protective pigments. This suggests that photoperception and regulation are important for fungi to receive nutrients through cooperation with phototrophs [36].

3.2.4 Exposure studies in Humidity Chamber

Fig. 5 shows brick samples exposed in the Humidity Chamber for 12 months where it can be seen the deterioration caused by the eight fungal isolates selected. These were selected according to their capacity to secrete pigments (*E. nigrum*, *L. theobromae*, YUSM and *A. spectabile*) and to their growth speed (*Arthrinium* sp. 1, *Arthrinium* sp. 2, DUSM 1 and DUSM 2). Fig. 5A-I shows stereoscopic microscope images obtained at the end of the assay (12 months). Colorless mycelium is observed on the brick surfaces under the assay conditions. Mycelial growth was more invasive in *L. theobromae* and *A. spectabile*, Fig. 5 C and F, respectively. For this reason, SEM was applied to characterize the biofilm formed. In Fig. 5 J, an abundant grouped conidia and vegetative mycelium of *L. theobromae* can be observed. Whereas in Fig. 5K, although only mycelium of *A. spectabile* was observed, the hyphae penetrated the brick pores.

In Fig. 6, it is presented a graph with the brick weight variation calculated by equation (3) and obtained from the inoculated and non-inoculated (control) bricks exposed in the humidity chamber. Bricks inoculated with both *L. theobromae* and *A. spectabile* showed statistically significant difference ($p \leq 0.05$) in weight loss with respect to the rest of the strains. It is worth noting,

that the growth of both strains on the bricks caused greater weight loss at 12 months, but there were no statistically significant differences with the values recorded at 6 months. Even so, *L. theobromae* and *A. spectabile* presented the highest values in brick weight variation *in situ*. The relative humidity is an abiotic factor, which cannot only influence the growth of fungi, but it can also cause modifications in the mechanical properties of bricks [40]. For the first time, these species were found to have biodeteriorating capacity during the course of this study.

On the other hand, the association between certain variables and the mycobiota present in the sampling sites was analyzed with the PCA, whose axes explained 64% of variation using RStudio 1.1.463® program. Fig. 7 showed PCA ordination of fungal isolates according to the variables analyzed. Axis 1 shows relative density, sample place, structural material, relative frequency, carbonate dissolution and acid production with 27.3 % of variation. On the other hand, axis 2 shows that 20.1% corresponded to the variation of brick weight, pigment secretion and the recollection height. As a result, the strains *Cladosporium* sp.1, *Cladosporium* sp.2, *A. alternata* and *Cladosporium* sp. 3 were grouped in points 3, 4, 5 and 6, respectively. Also, the strains belonging to the genus *Penicilium* occurred between points 20 and 24. In this sense, the groups formed by the species can be multifactorial and may lead to similarities in taxonomy and ecology. Especially, if the variables analyzed in this study have sufficient diagnostic characters for these fungi. Thus, isolates that are in the same group can have the same growth and ecological strategies, share the same ecological niches, and use the same substrate. Similarly, Coutinho et al. [41] reported that strains belonging to the *Alternaria, Cladosporium*, *Penicillium* and *Aspergillus* genera were isolated on architectural ceramic material. These last two as primary colonizers of the substrate and r-strategists. On the other hand, the species of *Alternaria* and *Cladosporium* being secondary colonizers of the substrates were catalogued as K-strategists [42]. These characteristics related to the biology of the isolates may be the conditions that favor their participation in building materials biodeterioration. The hyphae can penetrate the pores of these materials and, apart from causing physical damage to the material, they can facilitate the colonization of other species. Then, it is

necessary to protect the monuments from where the species were isolated. This protection can be performed through the application of antifungal coatings on the materials in order to control the fungal deterioration [43].

3.4 Molecular analysis of most deteriorating fungal strains

Taking into account the obtained results and the analyses performed above, molecular analysis was carried out on the following strains: *Cladosporium sphaerospermum* (100 % apparition relative frequency), *Penicillium commune* and *Aspergillus niger* (highest pH variation in LMM) and finally *Lasiodiplodia theobromae* as well as *Arthothelium spectabile* presented the highest brick weight variation values. Accordingly, it was possible to reaffirm the taxonomic identification by molecular techniques. Sequences were uploaded to GenBank database and the access numbers: MN371394, MN371392, MN371276, MN371283 and MT071822 were assigned to *C. sphaerospermum*, *P. commune, A. niger*, *L. theobromae* and *A. spectabile* isolates, respectively.

The clustering analysis shows the relationship between the species with higher biodeteriorating activity (Fig. 8). The values listed from left to right next to each cluster node correspond to the bootstrap values performed using 1000 replications, and the branches shown correspond to values 95%, being fungi grouped according to their Family. In particular, *P. commune* and *A. niger* belong to Aspergillaceae. Nevertheless, *C. sphaerospermum, L. theobromae* and *A. spectabile* are located within Cladosporiaceae, Botryosphaeriaceae and Arthoniaceae, respectively. UPGMA analysis showed consistent clustering of isolates, but the phylogram presented a divergence point where *A. spectabile* and *L. theobromae* did not share a common ancestor with the rest of the species. The optimal tree with the sum of branch length = 1.76794872 is shown. The tree is drawn to scale, with branch lengths in the same units as those of the evolutionary distances used to infer the phylogenetic tree. All positions with less than 100% site coverage were eliminated. There was a total of 195 positions in the final dataset.

The results indicate that the study of the mycobiota, which intervenes in the biodeterioration of building materials based on their ecology and deteriorating attributes, can contribute to subsequently carry out studies focused on the

protection of brick and concrete. Despite these promising results, further research should be undertaken to research on the fungi in consortium and the effect of competition between the strains. In this sense, it would be interesting to study the combined growth of these strains and their effect on bricks or concrete deterioration.

4.Conclusions

Through the sampling method and the isolation techniques used, it was possible to study the mycobiota present in facades of two buildings in La Plata (the Cathedral of La Plata and the Experimental Station of the Centro de Investigación y Desarrollo en Tecnología de Pinturas).

The tests performed allowed obtaining the most deteriorating strains: *A. niger* MN371276, *L. theobromae* MN371283, *C. sphaerospermum* MN371394, *P. commune* MN371392 and *A. spectabile* MT071822. Molecular method was used to confirm the strains identification and to analyse their evolutionary relationship. In this sense, it is interesting to highlight that the two strains *L. theobromae* and *A. spectabile,* which resulted older than the rest of the strains from an evolutionary point of view, also produced the highest deteriorating effect on bricks reflected in weight loss after the humidity chamber test.

The research has also shown mycological studies as a tool to obtain information about the culturable species present in the biofilm of building spoilage. In this sense, some of these strains can be used for their adaptability to grow under laboratory conditions in a study that seeks the efficient assessment of new materials with biocidal potentialities, and therefore the improvement of building biodeterioration control.

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Table 1: Pigment secretion and carbonate dissolution of fungi isolated from CP and CES. The isolates are ordered by the place of sampling

No.	Fungal Isolate	Sample place	Structural material	Heigth (m)	Carbonate dissolution	Pigment secretion
1	*Penicillium* sp. Link 1	CP	Brick	1.5	+	-
2	*Epicoccum nigrum* Link.	CP	Brick	1.5	-	+
3	*Cladosporium* sp. Link 1	CP	Brick	1.5	-	-
4	*Cladosporium* sp. Link 2	CP	Brick	1.5	-	-
5	*Alternaria alternata* (Fr.) Keissl.	CP	Brick	1.5	-	-
6	*Cladosporium* sp. Link 3	CP	Brick	1.5	-	-
7	*Penicillium* sp. Link 2	CP	Brick	0.5	+	-
8	*Cladosporium* sp. Link 4	CP	Brick	0.5	-	-
9	Coelomycetes Grove. 1	CP	Brick	0.5	-	-
10	*Cladosporium* sp. Link 5	CP	Brick	0.5	-	-
11	Orbiliomycetes O.E. Erikss. & Baral	CP	Brick	1.5	-	-
12	*Arthrinium* sp.Kunze. 1	CP	Brick	1.5	-	-
13	*Lasiodiplodia theobromae* Griffon & Maubl.	CP	Brick	1.5	-	+
14	*Penicillium commune* Thom	CP	Brick	1.5	+	-
15	*Cladosporium sphaerospermum* Penz.	CP	Brick	0.5	-	-
16	*Penicillium* sp. Link 3	CP	Brick	0.5	+	-
17	*Penicillium* sp. Link 4	CP	Brick	0.5	-	-
18	*Cladosporium* sp. Link 6	CP	Brick	1.5	-	-
19	*Arthrinium* sp.Kunze. 2	CP	Brick	1.5	-	-
20	*Penicillium* sp. Link 5	CP	Concrete	0.5	-	-
21	*Cladosporium cladosporioides* (Fresen.)	CP	Concrete	0.5	-	-
22	Coelomycetes Grove. 2	CP	Concrete	0.5	-	-
23	*Penicillium* sp. Link 6	CP	Concrete	0.5	-	-
24	*Penicillium* sp. Link 7	CP	Concrete	0.5	-	-
25	Coelomycetes Grove. 3	CP	Concrete	0.5	-	-

26	*Aspergillus niger* Tiegh.	CP	Brick and Concrete	0.5	+	-
27	YUSM	CP	Concrete	1.5	-	+
28	*Arthothelium spectabile* A. Massal.	CP	Brick	1.5	-	+
29	*Moniliella* sp. Stolk & Dakin 1	CES	Concrete	0.5	-	-
30	DUSM 1	CES	Concrete	1.5	-	-
31	DUSM 2	CES	Brick	0.5	-	-
32	*Cladosporium cladosporioides* (Fresen.)	CES	Brick	0.5	-	-
33	DUSM 3	CES	Concrete	0.5	-	-
34	*Moniliella* sp. Stolk & Dakin 2	CES	Concrete	0.5	-	-
35	*Chrysonilia* sp. Arx.	CES	Concrete	1.5	-	-
36	DUSM 4	CES	Brick	1.5	-	-
37	*Alternaria* sp. Nees, 1	CES	Concrete	1.5	-	-
38	SUNEM	CES	Brick	1.5	-	-
39	*Moniliella* sp. Stolk & Dakin 3	CES	Concrete	0.5	-	-
40	*Aureobasidium pullulans* (de Bary)	CES	Concrete	0.5	-	-
41	*Geomyces* sp. Traaen	CES	Concrete	1.5	-	-
42	*Mucor* sp. Fresen.	CES	Concrete	1.5	-	-
43	OUSM	CES	Concrete	0.5	-	+

+ : Positive; - : Negative

YUSM: yellow unsporulated septate mycelium

DUSM: dematiaceous unsporulated septate mycelium

OUSM: orange unsporulated septate mycelium

SUNEM: salmon unsporulated non-septate mycelium

Figure 1: Mycological sampling sites. A: Cathedral of La Plata. B: Experimental Station of the Centro de Investigación y Desarrollo en Tecnología de Pinturas. Biodeteriorated facades details. C: Highlighted in red Buenos Aires Province, taken from Google maps.
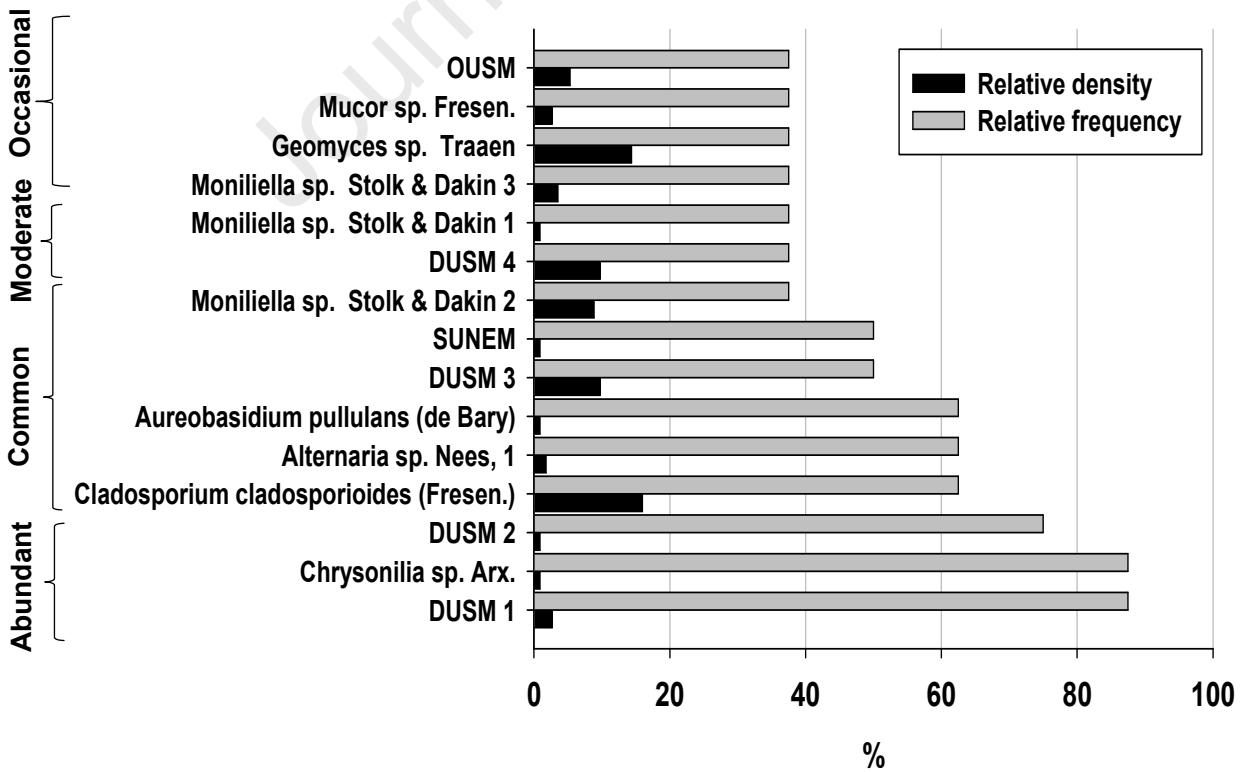
A
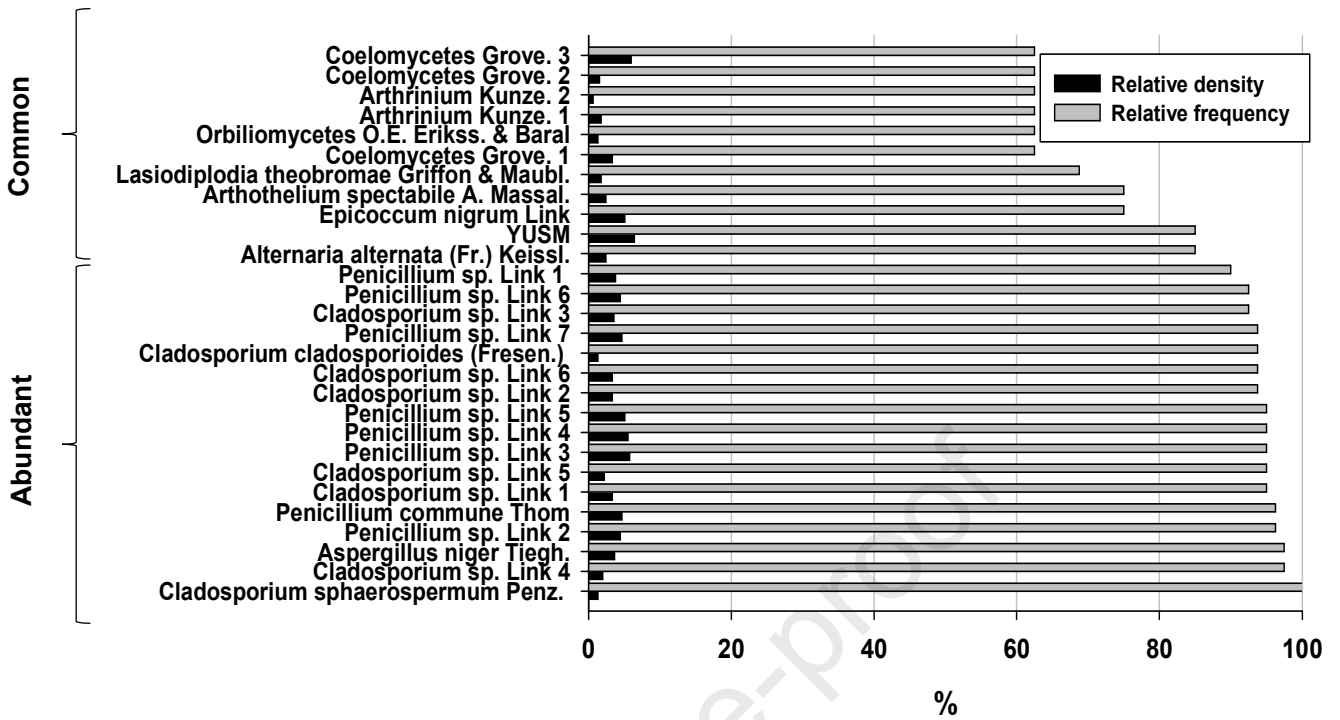
B

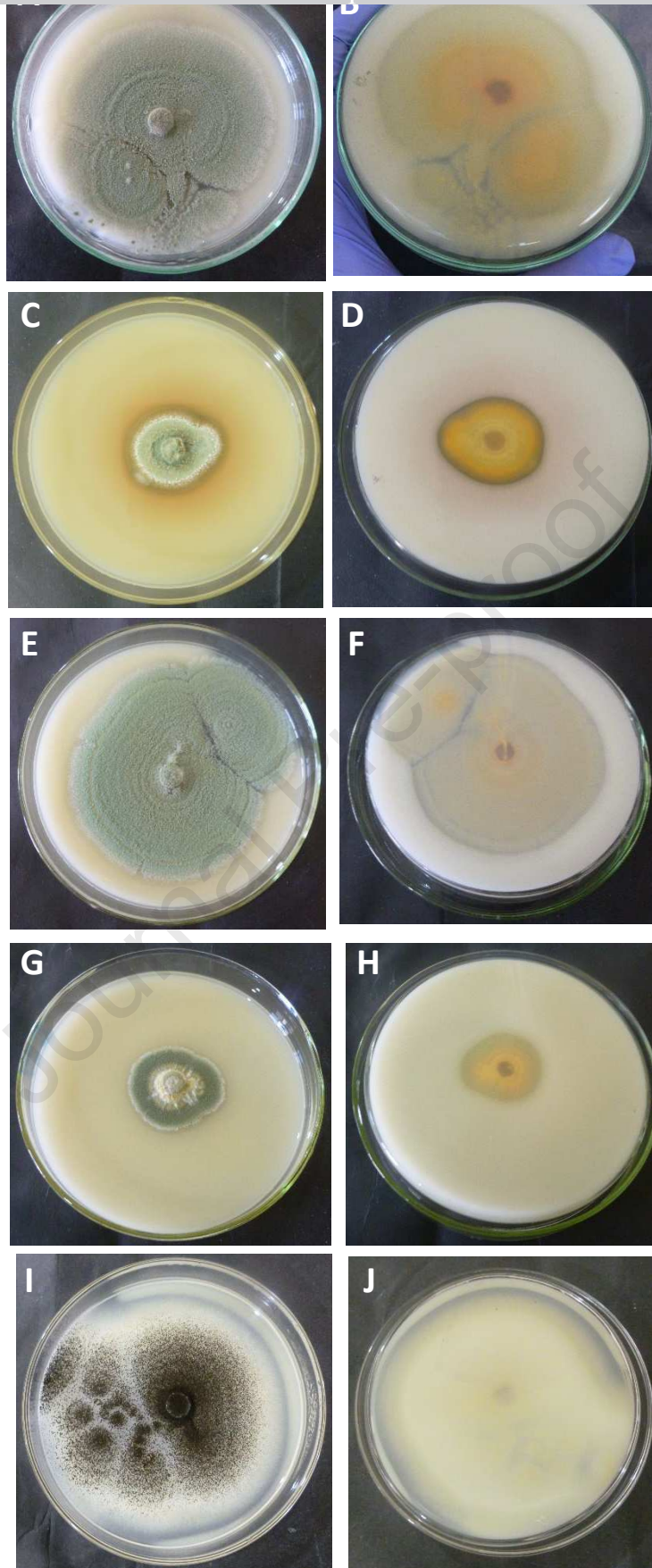Figure 2: Relative density, relative frequency and ecological category of isolated fungi. A: CP; B: CES.

Figure 3: Fungi with positive result in carbonate dissolution test, t= 14 days. A and B: *Penicillium* sp. 1. C and D: *Penicillium* sp. 2. E and F: *P. commune*. G and H *Penicillium* sp. 3. I and J: *A. niger*. Colony morphology on $CaCO_3$ malt extract agar A, C, E, G and I: Obverse; B, D, F, H and J: reverse. T= 28°C, t= 14 days.
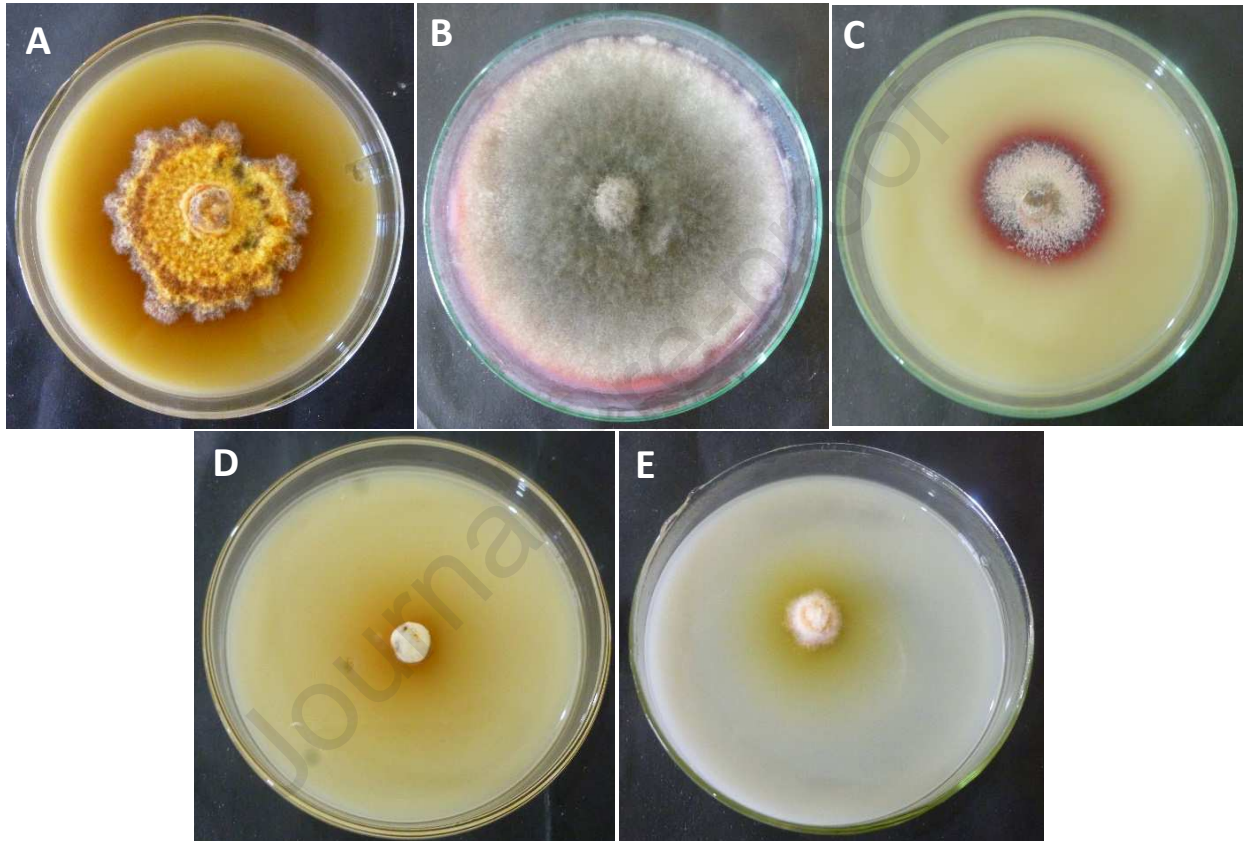
Figure 4: Fungal isolates capable of secreting pigments. A: *E. nigrum.* B: *L. theobromae*. C: YUSM. D: *A. spectabile*. E: OUSM. Obverse colony morphology. T= 28°C, t= 7 days.
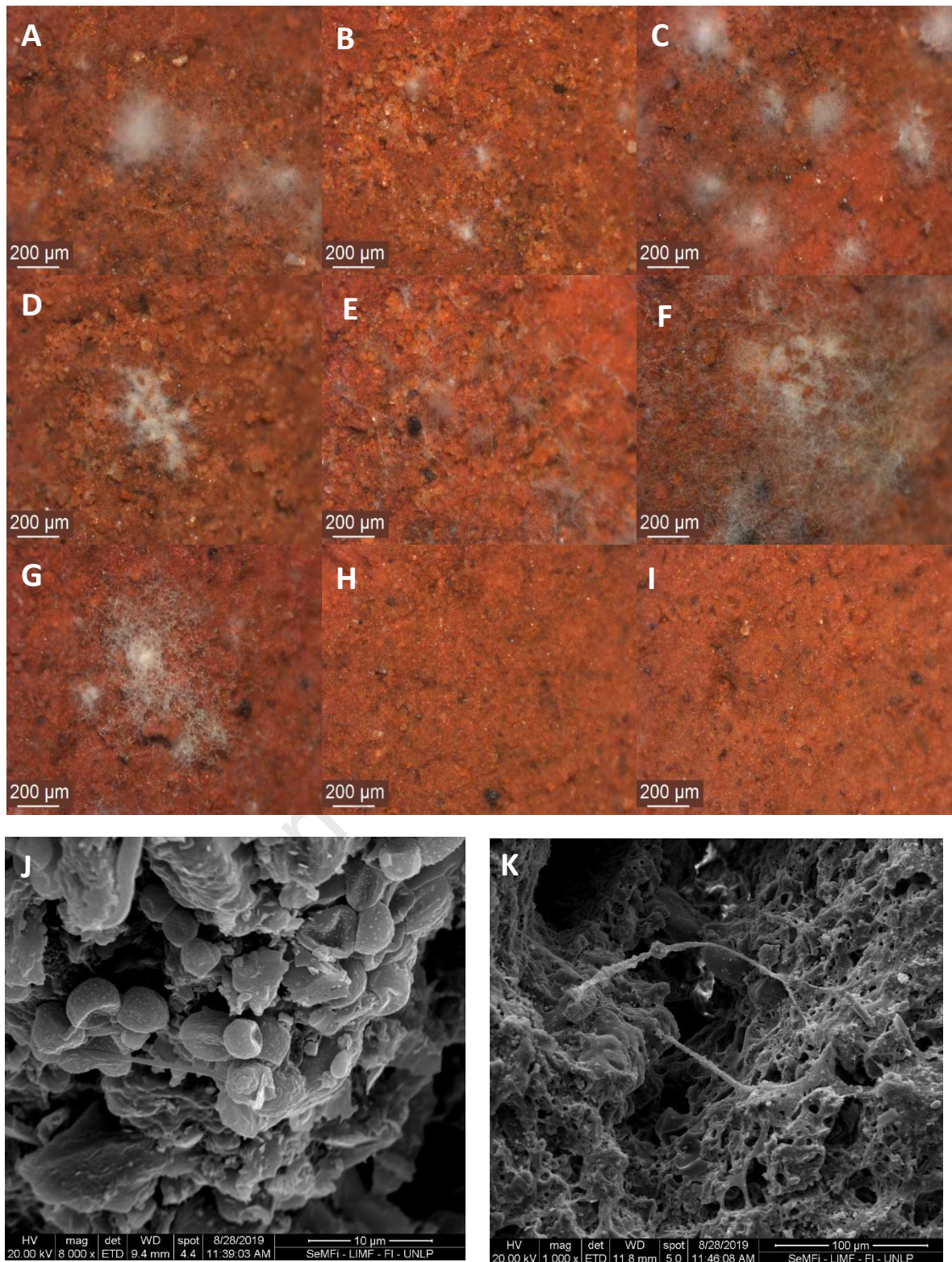
Figure 5: Bricks inoculated and exposed in the Humidity Chamber. A-I: Stereoscopic microscope images. J and K: SEM micrographs (t= 12 months). A: *E. nigrum*, B: *Arthrinium* sp. 1, C: *L. theobromae,* D: *Arthrinium* sp. 2, E: YUSM, F: *A. spectabile*, G: DUSM 1, H: DUSM 2 and I: Uninoculated brick.
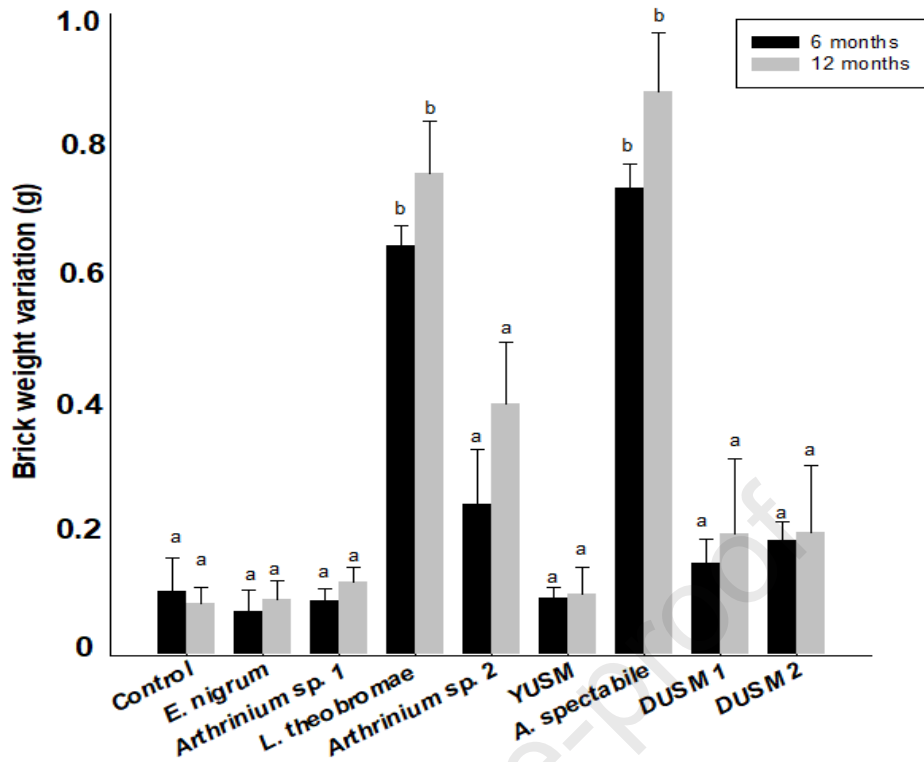
Figure 6: Loss of weights values in the bricks at different times, different letters on bars indicate difference statistically significant (ρ≤0.05) and for Tukey test.
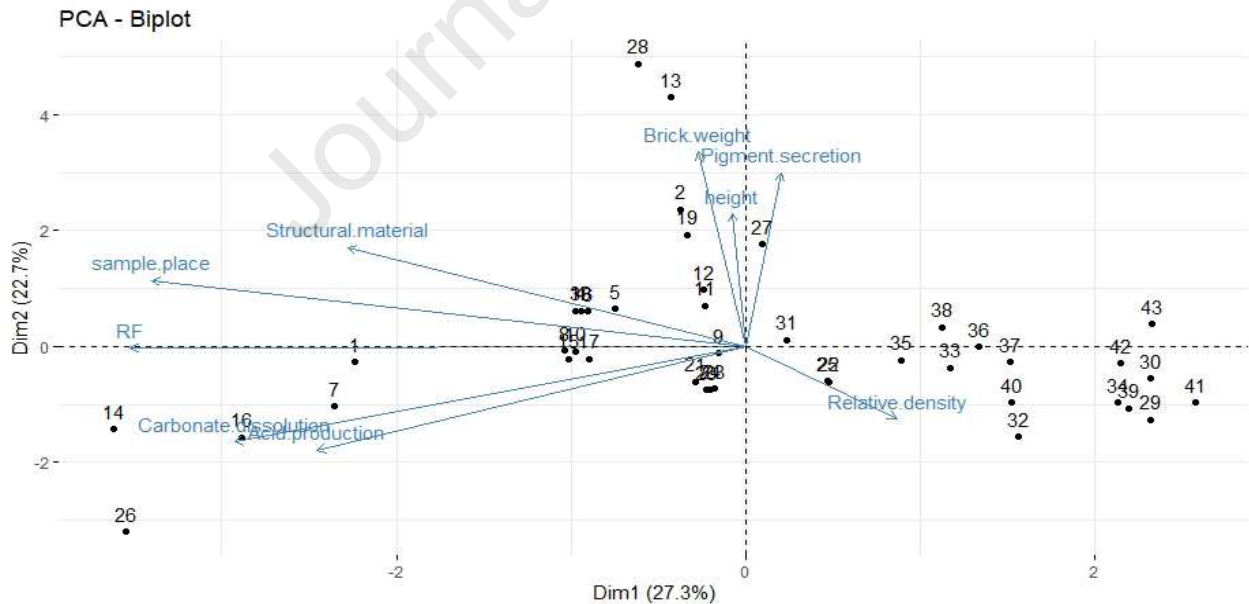
Figure 7: PCA ordination of fungal isolates (represented as numbers in the figure) according to the variables analyzed.
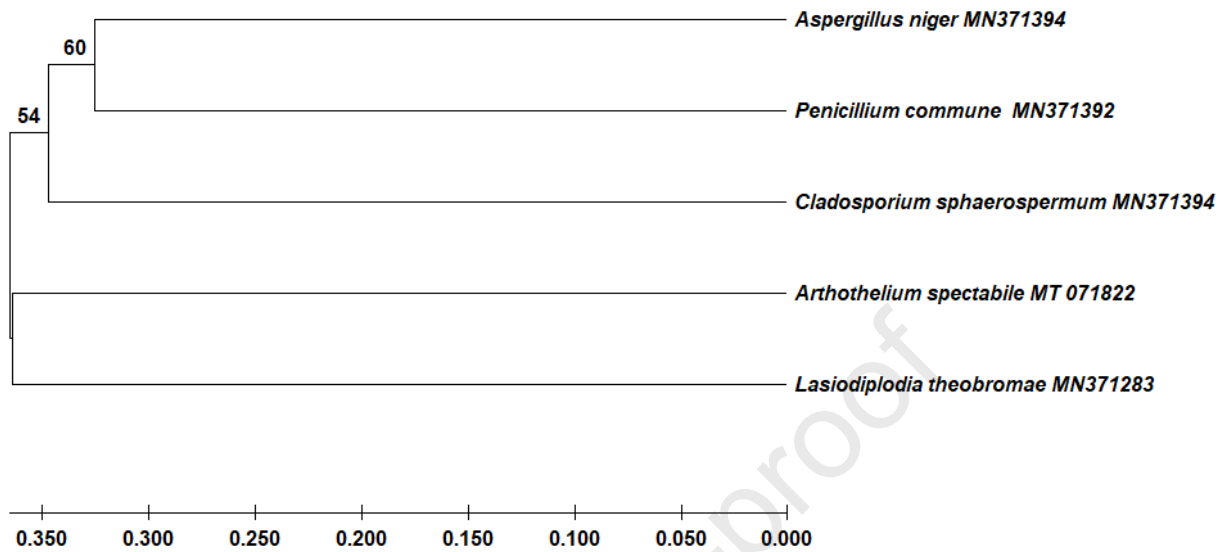

Figure 8: UPGMA cluster analysis of fungi isolated with greater deteriorating potential. The evolutionary distances were calculated using the p-distance model. The Bootstrap numbers shown on the left (95%) were obtained using 1000 replicates. The horizontal scale indicates the level of dissimilarity.

Highlights

- For the first time, mycobiota in facades at the Cathedral of La Plata and the Experimental station of the Centro de Investigation y Desarrollo en Tecnología de Pinturas was studied.
- The strains are useful to the assessment of the antifungal potentialities of new building materials.
- *L. theobromae* and *A. spectabile* were associated with a significant deterioration of a ceramic material for the first time.

Declaration of interests

☒ The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

☐ The authors declare the following financial interests/personal relationships which may be considered as potential competing interests: