

Unveiling Microbial Oxidizers in Chihuahuan Desert Rock Varnish (Mexico): A Pioneer Study

Desvelando los oxidantes microbianos en barnices del desierto de Chihuahua (México): un estudio pionero

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ABSTRACT

Arid and semi-arid desert areas characterized by extreme conditions, including northern Mexico, have rocks with micrometric coatings. These coatings are known as desert varnish, an often dark-red microlayer deposited over millennia and developed on different rock types. These microlaminations, rich in clay minerals and Fe- /Mn-oxides, can serve as habitats for various microorganisms, mainly fungi and bacteria. However, the role of microorganisms in these varnish formations is still under debate. In the Chihuahuan desert (Mexico), particularly in the Samalayuca ridge, extensive areas are covered by rocks with desert varnish. Here we present a research effort focused on isolating Fe- and Mn-oxidising microorganisms in different culture media, along with directed sequencing. Using a benzidine spot test to detect oxidant species revealed the presence of bacteria and fungi inhabiting these layers. Likewise, we contribute to a better understanding of the formation of desert varnish, representing the first report of specific microorganisms isolated from rock varnish of the Chihuahuan desert.

Keywords: desert varnish, oxidising microorganisms, Chihuahuan desert, benzidine test.

RESUMEN

Zonas desérticas áridas y semi-áridas caracterizadas por condiciones extremas, incluido el norte de México, presentan rocas con recubrimientos micrométricos. Estos recubrimientos se conocen como barniz del desierto, una microcapa, a menudo de color rojo-oscuro, que se deposita a lo largo de milenios y se desarrolla sobre diferentes tipos de rocas. Estas microlaminaciones, ricas en minerales arcillosos, y óxidos de Fe y Mn, pueden servir como hábitat para diversos microorganismos, principalmente hongos y bacterias. Sin embargo, el papel de los microorganismos en estas formaciones de barniz se encuentra en debate. En el desierto de Chihuahua (México), particularmente en la cordillera de Samalayuca, extensas áreas están cubiertas por rocas con barniz del desierto. Aquí presentamos un esfuerzo de investigación centrado en el aislamiento de microorganismos oxidantes de Fe y Mn en diferentes medios de cultivo, junto con secuenciación dirigida. El uso de una prueba puntual de bencidina para identificar especies oxidantes reveló la presencia de bacterias y hongos que habitan en estas capas. Asimismo, contribuimos a una mejor comprensión de la formación del barniz del desierto, presentando el primer reporte de microorganismos específicos aislados a partir de barnices del desierto de Chihuahua.

Palabras clave: Barniz del desierto, microorganismos oxidantes, desierto de Chihuahua, prueba de bencidina.

1. Introduction

Rock patina or rock varnish is a micrometric layer (50–200 μm thick) that forms on rock surfaces with a slow formation rate of around 1 to 40 μm every 1000 years (Liu and Broecker, 2000). Mineralogically, it is composed of almost 60% clay minerals, ~30% Fe- /Mn-oxides, and ~10% of other elements such as Si, Al, Mg, Ca, Ba, and Ce, among others (Potter and Rossman 1977; 1979; Dorn and Oberlander, 1981; Krinsley *et al.*, 2012; Municchia *et al.*, 2016; Lebedeva *et al.*, 2019). Patinas are common in arid and desert environments (Potter and Rossman, 1977; Jones, 1991; Cremaschi, 1996; Lebedeva *et al.*, 2019), although they can also be present in rocks of other ecosystem types (Krinsley *et al.*, 2012; Lozano and Rossi, 2012; Goossens *et al.*, 2015; Xu *et al.*, 2018).

Desert Varnish can form because of either abiotic or biotic processes. The abiotic one involves varying cycles of deposition, accumulation, and enrichment of minerals with high amounts of manganese (Dickerson, 2011; Otter *et al.*, 2020), weathering and diagenesis of aeolian dust and underlying rock (Municchia *et al.*, 2016; Lebedeva *et al.*, 2019; Andreae *et al.*, 2020), and other physicochemical processes such as photooxidation, humidity, variation in pH and silica gelation (Perry *et al.*, 2006; Dorn, 2007a; 2007b; Xu *et al.*, 2019). On the other hand, biotic processes imply that varnishes are formed by biochemical processes mediated or induced by microorganisms (Krumbein and Jens, 1981; DiGregorio, 2005; Kuhlman *et al.*, 2008), mainly those that mineralize Fe and Mn and deposit these oxidised and/or reduced elements, creating microlaminations.

Finally, some polygenic models imply a complex mixture between both processes (biotic and abiotic) where microorganisms (such as budding bacteria and fungi), physical-chemical and environmental processes all work together as an ecophysiological microsystem (Potter and Rossman, 1979; Dorn and Krinsley, 2011; 2019; Krinsley *et al.*, 2017; Lingappa *et al.*, 2021; Chaddha *et al.*, 2024).

The presence of microorganisms in rock varnishes has been demonstrated in several studies using techniques such as fluorescence microscopy, electron microscopy, DNA sequencing, and culturing (Taylor-George *et al.*, 1983; Lang-Yona *et al.*, 2018; Esposito *et al.*, 2019; Lingappa *et al.* 2021). These studies have revealed a diverse community of microorganisms, including bacteria, archaea, and fungi associated with the rock varnish (Esposito *et al.*, 2015; Lang-Yona *et al.*, 2018). These microorganisms are thought to play a role in the formation and alteration of the varnish by contributing to the accumulation of minerals and organic matter, such as varnish-associated photosynthetic bacteria, like *Chroococcidiopsis* sp., which accumulates higher concentrations of Mn in its cytoplasm (Lingappa *et al.*, 2021). Also, Mn-oxidiser bacteria such as *Hydrogenophaga* sp. and *Pedobacter* sp., or in consortia, like *Nevskia-Rhizobium* spp., induce individual extracellular mineralization (Sjöberg *et al.*, 2021).

Today, it is well known that the varnishes of the great deserts of America, such as the Sonoran and Mojave, have associated microorganisms (Taylor-George *et al.*, 1983; Perry *et al.*, 2003; Schelble *et al.*, 2005; Kuhlman *et al.*, 2006; Martínez-Paballo *et al.*, 2021). Because of the extreme conditions in which the varnish is found, the inhabiting microorganisms can be considered extremophiles (exposed to high doses of radiation, extreme aridity, and temperature) (Coleine *et al.*, 2021). Here, we present the first demonstration of morphological micro-biodiversity and the presence of Fe- and Mn-oxidising microorganisms in varnish samples from the Samalayuca Desert.

2. Study area

The Chihuahuan Desert is one of the largest desert areas in the American continent, with a quarter of its size in the US states of Arizona, New Mexico, and Texas, and the rest in Mexico states of Chihuahua, Coahuila, Nuevo León, Durango, and Zacatecas (Figure 1A-B). The municipality

of Juárez is in the northern state of Chihuahua, Mexico in the middle of the Chihuahuan Desert (Figure 1A-B), where the locality Samalayuca stands out by its amazing biodiversity and landscape (Hernández *et al.*, 2004). In this area, large amounts of rock varnishes can be found (Figures 1C and 1D), rich in Fe- /Mn-oxides (Martínez-Pabello *et al.*, 2022) and “Los Médanos de Samalayuca”, which has been declared a Protected Natural Area, due to its landscapes, archaeological relevance, and biodiversity (Gatica-Colima *et al.*, 2023). With an annual rainfall of 220 mm and a temperature variation of 23°C (CONANP, 2013), the region has been classified as a cold desert in the BWKx’ (e’) class of the Köppen’s climate classification (Peel *et al.*, 2007). The current climate in the area (Schmidt, 1979) corresponds to an arid highland desert with hot summers, reaching temperatures of up to 41°C, and cold winters, with minimum temperatures dropping to -16°C.

The annual average temperature ranges between 12–18°C. The prevailing winds, which make up 45% of the wind patterns, come from the west at speeds of approximately 20–40 km/h, although the fastest winds, reaching around 50 km/h, are from the southeast. More information on regional settings can be found in Martínez-Pabello *et al.*, 2022.

3. Materials and methods

3.1. COLLECTION OF SAMPLES

Desert varnish was collected in the Sierra de Samalayuca, municipality of Juárez, northeast of the Chihuahuan desert, Mexico (31° 20’ 25.1124”, 106° 30’ 21.0924’; Figure 1A to C). Sample selection was based on the appearance of rock varnish, where the samples with greater thickness, homo-

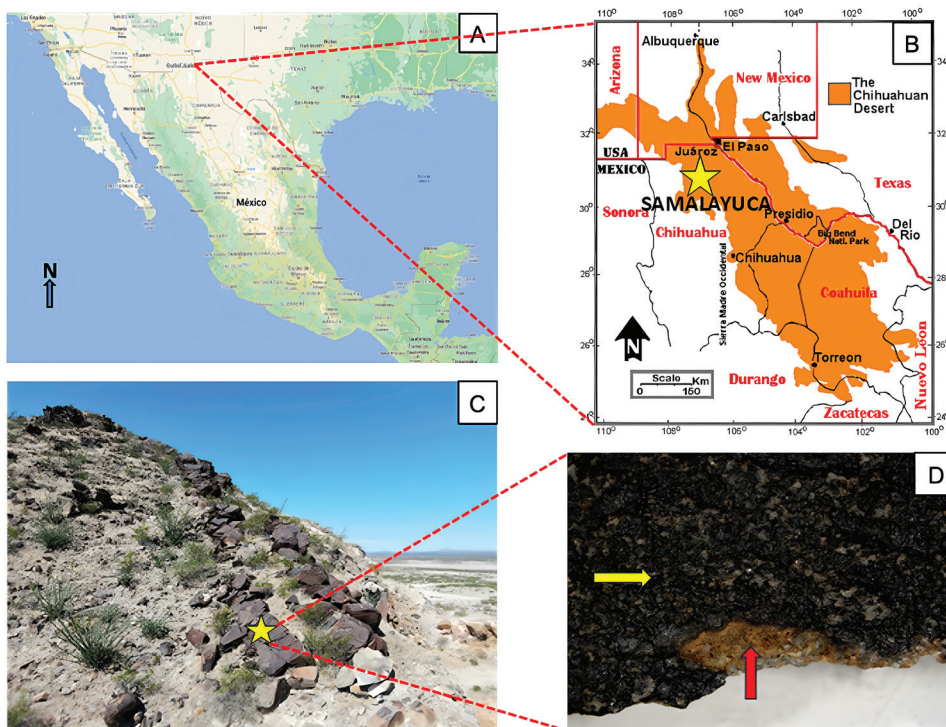


Figure 1 A and B. Location of the Chihuahuan Desert and Samalayuca collecting area (yellow star), Image taken and modified from Rivera-Rivera, 2006; C, Sampling site in Samalayuca. *Prosopis glandulosa* and *Larrea tridentata* are the dominant vegetation; D, Sample kind of collected varnish for this work. Dark varnish enriched with Mn-oxides (yellow arrow) and contact with underlying rock, sandstone (red arrow).

geneity, and dark pigmentation in the varnish were selected. The varnish was extracted with a cleaned geological hammer and stored in sterile conical Corning® tubes (50 mL) at 21°C until use.

3.2. SAMPLE PREPARATION

Since the objective of this study was to provide an initial approach to isolating microorganisms from the varnish, one sample was selected for the analysis. The sample was prepared according to the modified protocol of Taylor-George *et al.* (1983). Briefly, 1 cm² of varnish was separated from the source rock.

A manual grinding machine (Dremel ® 300) was adapted with a previously cleaned (hypochlorite-reagent grade) diamond tip to eliminate as much of the source rock as possible. Under sterile conditions, the varnish was placed in a Corning® conical tube with 40 mL of sodium phosphate buffer (NaCl 8.0 g/L, KCl 0.2 g/L, Na₂HPO₄ 1.44 g/L, and KH₂PO₄ 0.24 g/L; pH 7.4, J.T. Baker®), and vortexed for 10 min to remove traces of surface dust. Subsequently, the varnish was irradiated with UV-AB light for 30 min on each side of the varnish (face/back) to eliminate surficial microbial populations.

Finally, this fragment was pulverized in a porcelain mortar until fragments of ≈ 0.5 mm in length were obtained. The sample was separated into two fractions according to size: A) >0.5 and B) <0.5 mm. Fraction A was used for scanning electron microscopy (SEM), and fraction B was used for microbial cultures.

3.3. SEM

Fraction A consisted of 10 fragments (>0.5mm) that were analysed in a Zeiss Evo MA 10 microscope. Prior to observation, the samples were coated with carbon in a cathode sputter (Denton Vacuum® Desk V) for 10 min. Different areas of each fragment were scanned, looking for biological structures.

3.4. MICROBIAL CULTURES

Four culture solid media were used to cultivate the microorganisms: i) PDA, potato dextrose agar (39 g per litre, BD Bioxon™); ii) AA, water agar (8.5 g agar per litre, BD Bioxon™); iii) CZ (Czapek medium, 30 g sucrose, 2 g NaNO₃, 1 g K₂HPO₄, 0.5 g MgSO₄·7H₂O, 0.5 g KCl, 0.01 g FeSO₄, 8.5 g agar per litre, J.T. Baker®); and iv) TM (Thornton medium, MT, 1 g mannitol, 1 g K₂HPO₄, 0.5 g KNO₃, 0.5 g asparagine, 0.2 g MgSO₄·7H₂O, 0.1 g CaCl₂, 0.1 g NaCl, 0.002 g FeCl₃, 8.5 g agar per litre, J.T. Baker®). All culture media were sterilised 15 min at 121°C, 15 psi.

20 mg of sterile fraction B were suspended in 1.5 mL of sterile water-peptone (0.01%) and vigorously shaken. 10 µL of the suspension was spread with a Drigalski spatula in triplicate in each medium. The cultures were incubated for 22 days at 21° ± 1°C (room temperature seeking lichen growth), 25°C (according to Wei (2020), most mesophilic microorganisms grow mostly at this temperature), and 37°C (standard bacterial growth temperature).

3.5. SEQUENCING ANALYSIS

Two morphologically different microorganisms were chosen to sequence from the isolates: samples from the AA medium with a colony morphology like a bacterium and one from the PDA medium corresponding to a fungus (strain 5-1 and 13-3, respectively). The strains were identified by amplifying the gene 16S rRNA of 5-1 strain, using oligonucleotides 27F (5'-AGA GTT TGA TCC TGG CTC AG-3') and 1492R (5'-GGT TAC CTT GTT ACG ACT T-3') both at 0.3 µM (Lane, 1991) and according to the instructions of the GoTaq® G2 Flexi DNA Polymerase kit (Promega©). The reactions were run in the program: 5' at 95°C; 35 cycles of 1' at 95°C, 30" at 55°C and 1.5' at 72°C; 10 min at 72°C and an indeterminate time of 4°C.

In the case of the 13-3 strain, we used the ITS molecular marker using the oligonucleotides, ITS1-F (5'-CTTGGTCATTTAGAGGAAG-

TAA-3') and ITS4 (5'-TCCTCCGCTTATTGATATGC-3'), both at 0.3 μ M (Gardes and Bruns, 1993). The reactions were run in the program: 5' at 95°C; 35 cycles of 1' at 95°C, 30" at 55°C and 45" at 72°C; 5 min at 72°C and an indeterminate time of 4°C.

A 1% agarose gel in 1X TAE (Tris, glacial acetic acid, and EDTA, Sigma-Aldrich®) was made to verify the amplifications and purification of fragments. The runs were at 100 volts for 40 minutes (ENDURO™, LabNet® chamber). The gels were stained with ethidium bromide at 500 nM for 10 minutes. To identify the size of the amplicons, the 1kb Ladder (Promega®) was used and the purification of the amplicons was carried out with the gel extraction kit (QIAGEN®). The McLab company in San Francisco (USA) sequenced the fragments by the Sanger method with ABI 3730XL sequencers.

3.6. BIOINFORMATICS ANALYSIS

The sequences were cured using BioEdit® (Version 5.09). For bacteria identification, each sequence was subjected to two alignments with BLAST from NCBI®; the first excluded sequences of non-culturable bacteria; the second considered the genus with the highest percentage of identity and coverage from the first alignment was also excluded. Sequences of at least two strains of each species with similarity were downloaded. Finally, the ITS sequence was aligned in the UNITE 8.3 (Kõljalg *et al.*, 2020) database for fungus identification.

For phylogenetic analysis, in both cases, alignments were done using MAFFT® (Version 7), applying the "Q-INS-i" strategy. A suitable evolutionary model was calculated for each group of sequences using JModelTest® (Version 2.1.10). Cladograms were created with Bayesian inference; for this, the MrBayes program (Version 3.2.7) was configured with the evolutionary models and applied at least 1,000,000 generations and sampling frequencies every 1,000 generations. Each generated file was edited with FigTree® (Version 1.4.4).

3.7. MICROSCOPIC ANALYSIS OF CULTURES

The primary cultures were observed at different incubation times in the Leica MZ® 12.5 with camera (DFC300) and the Zeiss® Axio Zoom V16 with camera Axiocam ICC5 microscopes, using transmitted light at 40X. For strain 5-1, the Gram test was performed using crystal violet (5% alcoholic solution, Hycel®), Lugol (5% in water, Hycel®), alcohol-acetone (Hycel®) in a proportion of 1:1, and safranin (1% in water, Hycel®). The preparation was observed under a microscope (Zeiss® PrimoStar) at 100X. The catalase and oxidase tests were performed on this strain, dripping 10 % H₂O₂ or BBL® Oxidase Reagent over isolated colonies.

3.8. BENZIDINE SPOT TEST

Reagent-grade benzidine (1,1'-biphenyl-4,4'-diamine, Sigma-Aldrich®) was used to prepare a 1% solution in acetic acid (7%). After 22 days of incubation, the primary cultures were flooded with 2 mL of the solution to determine the presence of Fe- and Mn-oxidising microorganisms. The colour changes from pale yellow to deep blue were considered positive results. Both chosen strains (5-1 and 13-3) were also tested.

4. Results and discussion

4.1. VARNISH SURFACE ANALYSIS

Ten fragments \geq 0.5 mm were selected to be analysed by SEM to show the micromorphology of the microbiota present in varnish before pulverising. The varnish micrographs show microorganisms embedded in (Figure 2). We observed bacteria biofilm (Figure 2A) coexisting with fungus mycelium and spores (Figure 2B). The distribution of the different types of microorganisms is heterogeneous, frequently accumulated in small grooves, gaps, and microcavities on the surface of the varnish. These biological structures and distribution have also been observed on the surface of varnishes from

the Sonoran Desert (Taylor-George *et al.*, 1983; Martínez-Pabello *et al.*, 2021) and the Mojave Desert (Perry *et al.*, 2003; Kuhlman *et al.*, 2006; Lebedeva *et al.*, 2019), such as the brown filaments typical of Mn-oxidising bacteria described in New Mexico varnishes (Sjöberg *et al.*, 2021).

4.2. CULTIVABLE MICROBIAL VARNISH STRAINS

Viable colonies were recovered from the second fraction (B) of the same varnish that was cultured in four different media at three different temperatures ($21^{\circ} \pm 1^{\circ}\text{C}$, 25°C , and 37°C). After 22 days of incubation, bacterial colonies and mycelial morphologies were identified and photographed at different times (Figure 3): as time passed, the different morphologies appeared. Also, the cultures were incubated at room temperature ($21^{\circ} \pm 1^{\circ}\text{C}$) near a natural light source to promote the growth of photosynthetic microorganisms. However, it

was impossible to ensure the isolation of microorganisms with this capacity. It has been reported that the growth of cyanobacteria associated with lichens is around 25°C (Lange *et al.*, 1998), which was the reason to incubate them at this temperature. Finally, greater growth of mesophilic bacteria was favored by culturing at 37°C .

Slow-growing filamentous colonies were observed in all media and all incubation temperatures. Some of the colonies grew directly from millimetric varnish particles dispersed over the Petri dish (Figure 3A). The greatest number of microorganisms growths was found in the AA media (Figure 3A and 3B), followed by CZ (Figure 3C1 and 3D) and TM media (Figure 3C2). In the PDA media, we observed some colonies on the 9th day of incubation compared with the above other media, where colonies appeared at least after 14 days of incubation.

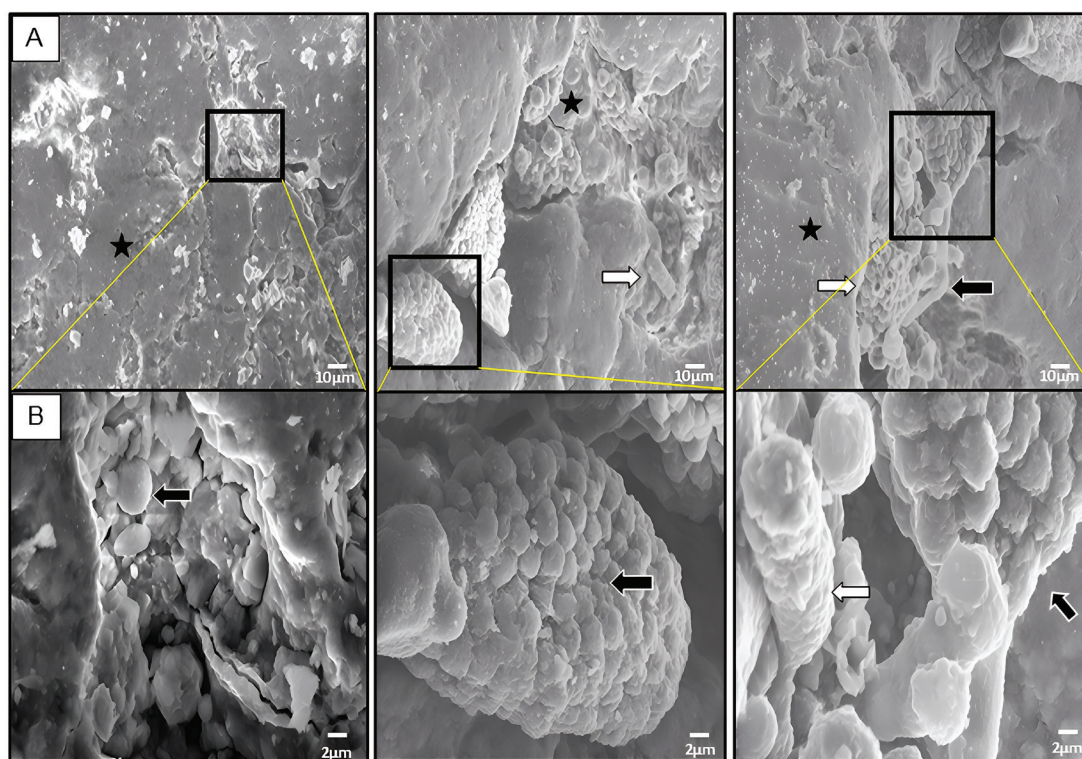


Figure 2 Analysis of varnish surface by SEM. A, Micrographs show many microorganisms morphologies embedded in the varnish such as: botryoidal biofilm (black star), ascomycetes colonies, hyphae and spores (black arrows), and tubular cells (white arrows). B, Magnification areas delimited by black rectangle (in A). Scale is indicated at bottom right by white bar ($10\ \mu\text{m}$ in A and $2\ \mu\text{m}$ in B, respectively).

Fungal structures (spores and hyphae) were also present in all samples. The hyphae were composed of spherical or slightly elongated cells, with approximately 5–10 μm in diameter and thick melanized walls typical of black meristematic fungi. Some colonies resemble Mn-oxidiser fungi or bacteria, such as *Cladosporium* sp. or *Pedobacter* sp., respectively (Sjöberg *et al.*, 2021). Microbial communities appear to be composed mainly of bacteria with few individual coccoid cells (Antonelli *et al.*, 2020).

Although we observed differences in growth between the isolated bacteria and the fungus, their role in varnish colonization and/or mineralization processes is still indeterminate. Further studies are needed to understand and clarify their contributions.

4.3. MANGANESE AND IRON OXIDE PRODUCTION (REDOX)

The cultures were subjected to the benzidine spot test to determine the presence of Fe- and Mn-oxidising microorganisms. This test is based on the oxidation of benzidine in the presence of Fe^{3+} or Mn^{4+} to form a benzidine radical cation (deep blue; do Nascimento *et al.*, 2006). It was confirmed that some of the obtained microorganisms were Fe- and Mn- oxidisers, such as the brown (Figure 4A) and black filamentary colonies (Figure 4B) that were coloured deep blue. Oxidising microorganisms were found mainly in AA and TM media, at 37°C and 25°C, respectively. On the one hand, Fe^{2+} and Mn^{2+} were dissolved in the TM medium, and oxidizer strains could directly mineralize

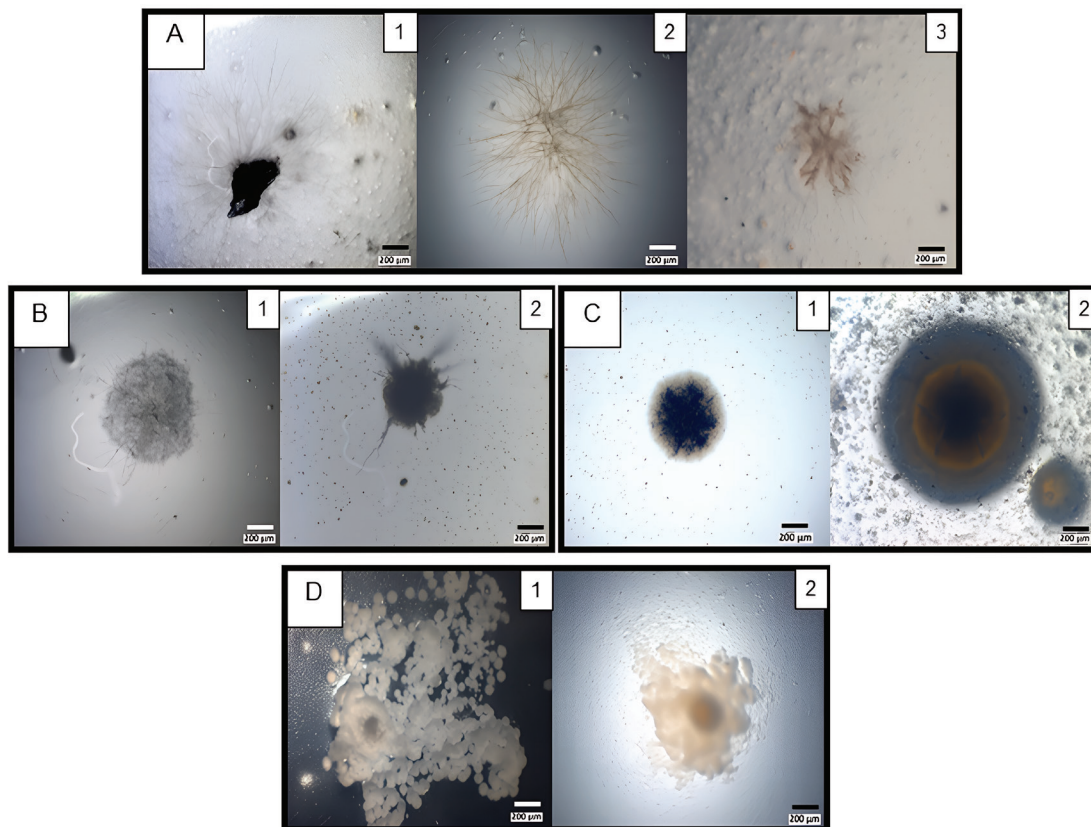


Figure 3 Strain colonial diversity recovered from pulverised varnish. A, Brown mucilage colonies with thin (A1-2) and astral mycelia (A3) morphologies. Micrographs (A1-3) correspond to AA Petri dishes incubated for 9, 10 or 8 days and, at 21°, 25° or 37°C, respectively. B, Mycelium with sporangia morphologies. Micrographs (B1-2) correspond to AA Petri dishes incubated for 9 days at 21°C. C, Rounded colonies. Micrographs correspond to CZ (C1), or TM (C2) Petri dishes incubated for 10 or 9 days and, at 25° or 21°C, respectively. D, Umbonate colonies. Micrographs correspond to CZ Petri dishes incubated for 7 (D1) or 9 (D2) days and, 37° or 21°C, respectively.

them; conversely, in the AA medium, the dark hue comes from dispersed varnish particles.

Manganese and iron oxidation/reduction by microorganisms contribute to desert varnish's complex mineralogy and geochemistry. The relative abundance and distribution of manganese and iron oxides within desert varnishes can provide valuable insights into the microbial communities and environmental conditions that have shaped their deposit over time.

Manganese and iron are two abundant elements in desert varnish, often occurring in the form of oxides. Microorganisms, particularly bacteria and cyanobacteria, play a significant role in the oxidation and reduction of these metals. Manganese-oxidising bacteria, such as *Bacillus*, *Pseudomonas* (Wright *et al.*, 2018), and *Arthrobacter* (Liang

et al., 2017) genders, can oxidise Mn^{2+} to Mn^{4+} ; these facilitate the oxidation of soluble Mn^{2+} ions to insoluble Mn^{4+} oxides resulting in their precipitation. Conversely, manganese-reducing bacteria, like *Geobacter* sp. (Mehta *et al.*, 2005) and *Shewanella* sp. (Wright *et al.*, 2016), and some fungi species can reduce Mn^{4+} to Mn^{2+} , leading to the dissolution of manganese oxides (Wei *et al.*, 2012). Similarly, iron-oxidising bacteria, such as *Leptothrix* sp. and *Gallionella* sp., oxidise Fe(II) to Fe(III), promoting the formation of iron oxides (Eggerichs *et al.*, 2020), while iron-reducing bacteria, such as *Geobacter* sp. (Straub and Schink, 2004) and *Clostridium* sp. (Mishra and Pradhan, 2024), reduce Fe(III) to Fe(II), resulting in the dissolution or alteration of iron oxides.

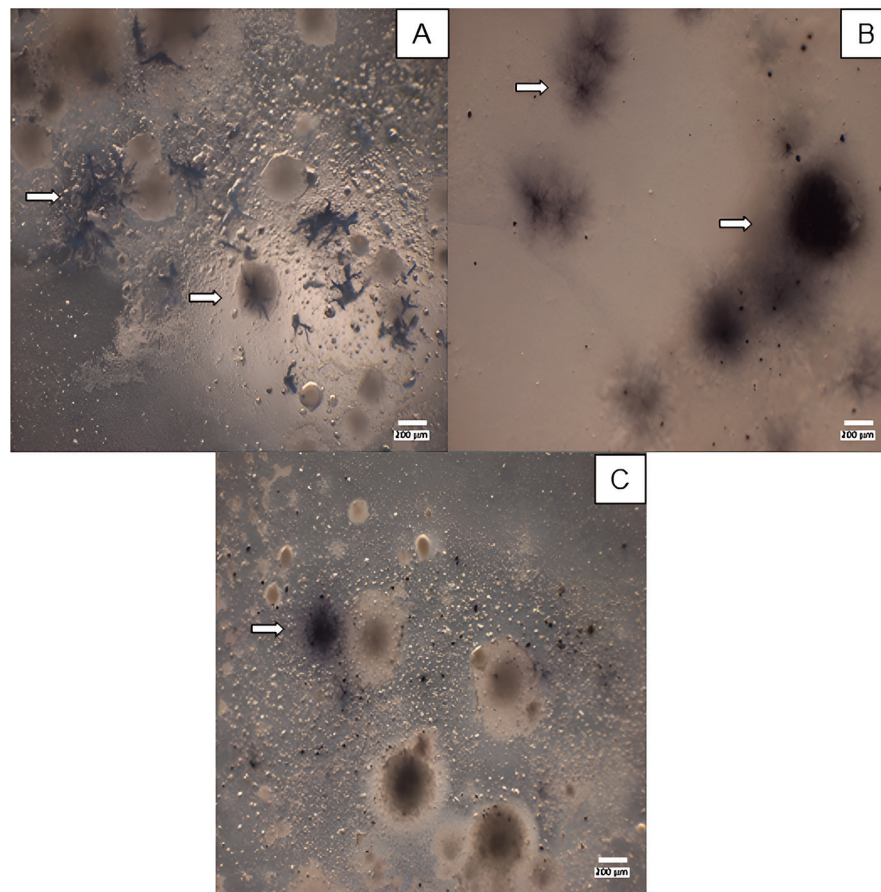


Figure 4 Revealing of oxidising microorganisms. A and B, AA medium culture incubation at 37°C. C, TM medium culture incubation at 25°C. All cultures were revealed in 22nd incubation day. Arrows highlight the positive presence of colonies with different morphologies.

places where desert varnishes have been reported; *P. nigra* has previously been isolated in rocks from this same site (Trujillo *et al.*, 2017); it has also been found that there are genes of the genus in biofilms that rest on sandstone rocks (Duan *et al.*, 2021), the same type of rock on which varnishes grow in the Samalayuca mountain range.

The 13-3 strain was a slow-growing black fungus in PDA media (Figure 6A). It has branched mycelium with thick-walled, cylindrical, and swollen cell hyphens (Figures 6B and 6C). *Spissiomycetes aggregatus* was its close relative (Figure 6D), with whom it shares the same morphological characteristics (Su *et al.*, 2015). The species of this gender are melanin-producing fungi with a rock-inhabiting lifestyle (Ametrano *et al.*, 2019), also reported as Mn-oxidisers (Chaput *et al.*, 2015).

5. Conclusions

This is the first study that explores the microorganism diversity inhabiting the rock varnishes in the Chihuahuan Desert (Mexico). Varnishes of the Samalayuca desert have a rough surface that allows the colonization of various reproductive structures of different microorganisms (spores and conidia). As expected, the Samalayuca desert varnish harbors microbial communities (mainly bacteria and fungi) that oxidise metals such as Mn or Fe. Bacterial colonies mineralise Fe and Mn directly from the varnish in media with any supplementation (*e.g.*, PDA or AA media). These bacteria have rapid growth (~7 days) compared to isolated black fungi (at least ~22 days) in the selection media; this may be due to the metabo-

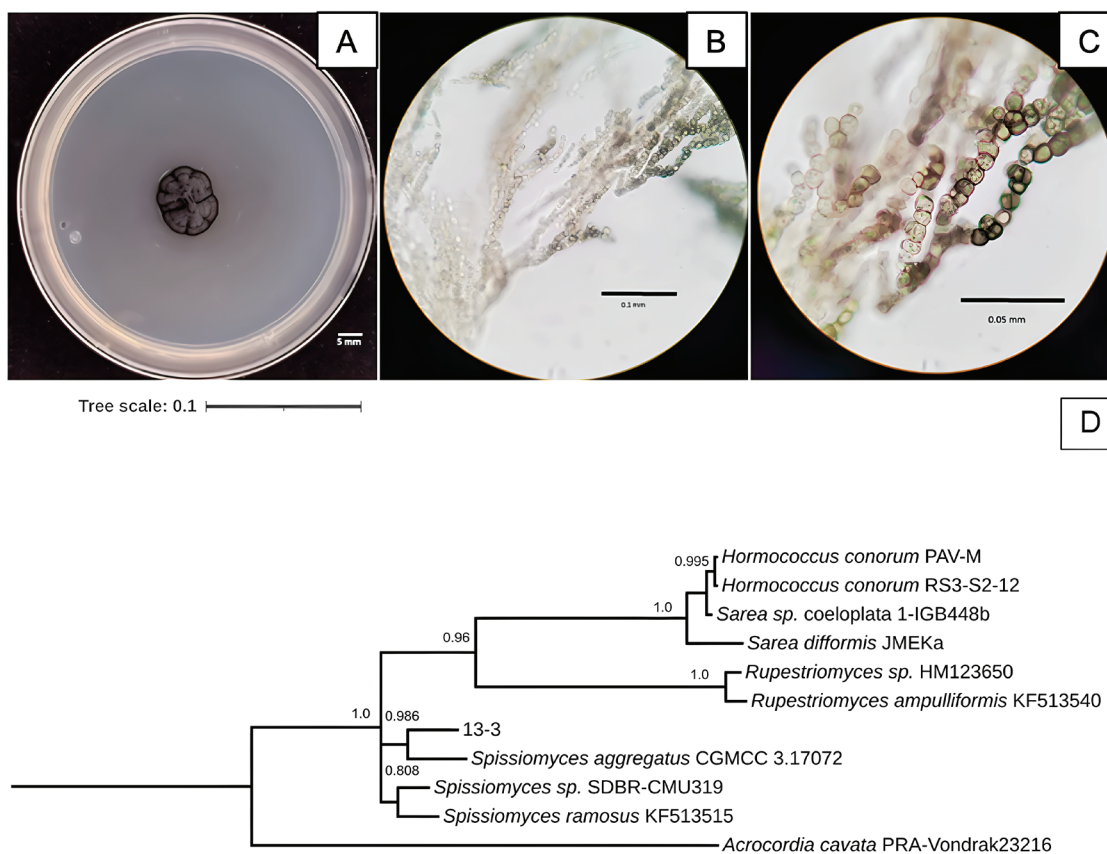


Figure 6 Characteristics of 13-3 isolated fungus strain from Samalayuca desert varnish. A, Colony morphology in PDA medium after 37 incubation days at 25°C; B and C, mycelium micrographs. D, Bayesian inference phylogenetic tree formed with ITS1 sequences. PRA-Vondrak23216 *Acrocordia cavata* was the external group. The probability branch supports values are shown.

lism of each one and the conditions in which it was experimented. However, this can be evaluated in future works. The specific role of those isolated microorganisms in the formation of rock varnishes remains unknown. Still, the positive response of some of them to benzidine is a first step to understanding the complex ecological relationship of varnish microbiota.

Also, future studies on the molecular (metagenomics or metabarcoding analysis) and physiological characterization (*i.e.* mycelial morphology, secondary metabolites production in different culture media) of these isolates could clarify which species are involved with desert varnish formation and which are wind-borne environmental organisms.

Contributions of authors

(1) Conceptualization: MAR, PUMP, MCG; (2) Data analysis or acquisition: MAR, PUMP, MCG, VALR, ISQ, MPOL, PDVP, AIP, HMMP; (3) Methodological/technical development: PUMP, VALR, ISQ, MPOL, PDVP, AIP, HMMP; (4) Writing of the original manuscript: MAR, PUMP, MCG, VALR; (5) Drafting of the corrected and edited manuscript: MAR, PUMP, MCG, VALR, ISQ, MPOL, PDVP, AIP; (6) Graphic design: MAR, PUMP, VALR, ISQ; (7) Fieldwork: VALR, AIP; (8) Interpretation: MAR, PUMP, MCG, PDVP; (9) Financing: MAR, MPOL, MCG.

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Conflicts of interest

The authors declare no conflicts of interest.

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Alexander Correa-Metrio.

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