

Exploring genetic diversity, population structure, and phylogeography in *Paracoccidioides* species using AFLP markers

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Abstract: Paracoccidioidomycosis (PCM) is a life-threatening systemic fungal infection acquired after inhalation of *Paracoccidioides* propagules from the environment. The main agents include members of the *P. brasiliensis* complex (phylogenetically-defined species S1, PS2, PS3, and PS4) and *P. lutzii*. DNA-sequencing of protein-coding loci (e.g., *GP43*, *ARF*, and *TUB1*) is the reference method for recognizing *Paracoccidioides* species due to a lack of robust phenotypic markers. Thus, developing new molecular markers that are informative and cost-effective is key to providing quality information to explore genetic diversity within *Paracoccidioides*. We report using new amplified fragment length polymorphism (AFLP) markers and mating-type analysis for genotyping *Paracoccidioides* species. The bioinformatic analysis generated 144 *in silico* AFLP profiles, highlighting two discriminatory primer pairs combinations (#1 EcoRI-AC/MseI-CT and #2 EcoRI-AT/MseI-CT). The combinations #1 and #2 were used *in vitro* to genotype 165 *Paracoccidioides* isolates recovered from across a vast area of South America. Considering the overall scored AFLP markers *in vitro* (67–87 fragments), the values of polymorphism information content ($PI_C = 0.3345–0.3456$), marker index ($MI = 0.0018$), effective multiplex ratio ($E = 44.6788–60.3818$), resolving power ($R_p = 22.3152–34.3152$), discriminating power ($D = 0.5183–0.5553$), expected heterozygosity ($H = 0.4247–0.4443$), and mean heterozygosity ($H_{avp} = 0.00002–0.00004$), demonstrated the utility of AFLP markers to speciate *Paracoccidioides* and to dissect both deep and fine-scale genetic structures. Analysis of molecular variance (AMOVA) revealed that the total genetic variance (65–66 %) was due to variability among *P. brasiliensis* complex and *P. lutzii* ($\Phi_{IPT} = 0.651–0.658$, $P < 0.0001$), supporting a highly structured population. Heterothallism was the exclusive mating strategy, and the distributions of *MAT1-1* or *MAT1-2* idiomorphs were not significantly skewed (1:1 ratio) for *P. brasiliensis* s. str. ($\chi^2 = 1.025$; $P = 0.3113$), *P. venezuelensis* ($\chi^2 = 0.692$; $P = 0.4054$), and *P. lutzii* ($\chi^2 = 0.027$; $P = 0.8694$), supporting random mating within each species. In contrast, skewed distributions were found for *P. americana* ($\chi^2 = 8.909$; $P = 0.0028$) and *P. restrepiensis* ($\chi^2 = 4.571$; $P = 0.0325$) with a preponderance of *MAT1-1*. Geographical distributions confirmed that *P. americana*, *P. restrepiensis*, and *P. lutzii* are more widespread than previously thought. *P. brasiliensis* s. str. is by far the most widely occurring lineage in Latin America countries, occurring in all regions of Brazil. Our new DNA fingerprint assay proved to be rapid, reproducible, and highly discriminatory, to give insights into the taxonomy, ecology, and epidemiology of *Paracoccidioides* species, guiding disease-control strategies to mitigate PCM.

Key words: AFLP, AMOVA, Endemic mycosis, Genetic diversity, Mating-type, *Paracoccidioides*, Paracoccidioidomycosis.

Available online 30 November 2021; <https://doi.org/10.1016/j.simyco.2021.100131>.

INTRODUCTION

Paracoccidioidomycosis (PCM) is a life-threatening systemic fungal infection first described in Brazil by Lutz and Splendore in 1908 (Lutz 1908). Its description was shortly followed by reported infections throughout South America (Ferguson & Upton 1947, Nino 1950) and, later, Latin America (Gonzalez Ochoa & Esquivel 1950). Following inhalation of soil-born *Paracoccidioides* propagules, patients may develop primary pulmonary foci, subsequently disseminating to other host organs and systems (Brunner *et al.* 1993, Hahn *et al.* 2014) by hematogenic or lymphatic pathway (Restrepo *et al.* 2008). The classical clinical forms of PCM-disease are divided into two groups (Franco *et al.* 1987). The first group includes an acute or sub-acute form ("juvenile"), predominant in children, adolescents, and young adults, and is depicted by tropism of the fungus to the monocyte-phagocyte system. The second and significant group corresponds to the chronic form, found in 80 to 95 % of the total

cases, and men between 30 and 50 years of age are the most affected patients (Nery *et al.* 2021a, 2021b). Brazil accounts for up to 80 % of cases of PCM in Latin America, with an incidence of 7.99 cases per 1000 hospitalizations based on the overall hospital admissions notified to the Ministry of Health in the year 2011 (Giacomazzi *et al.* 2016). The incidence and severity of PCM increase with the progression of human immunodeficiency virus (HIV) infection and reduction in CD4 counts. Therefore, the course of the disease in the HIV-infected patient is similar to those observed in the acute presentation of endemic PCM as it tends to be disseminated and more rapidly progressive (Almeida *et al.* 2017, de Almeida *et al.* 2018, Macedo *et al.* 2018).

PCM is caused by the thermophilic fungi classified in the order Onygenales, family Ajellomycetaceae and genus *Paracoccidioides* (Bocca *et al.* 2013). The etiological agent was first described in 1912 as *Zymonema brasiliensis* and later named *Paracoccidioides brasiliensis* as asserted by Floriano P. de Almeida (Almeida 1930). Historically the taxonomy of

Paracoccidioides is inconsistent with the description of several names that were found to be invalid. A few examples include *Paracoccidioides antarcticus* (Gezuele 1989), *Paracoccidioides cerebriiformis* (Moore 1935), and *Paracoccidioides tenuis* (Moore 1938). All these changes were reduced or described as a synonym of *P. brasiliensis* (Splendore) de Almeida, aiming to describe species names that reflect a natural classification system and a real need for communication among scientists (Almeida 1930, Del Negro et al. 1993, Garcia et al. 1993).

For decades *Paracoccidioides* was considered to be a monotypic taxon. Phylogenetic relationships in *Paracoccidioides* based on the internal transcribed spacer (ITS) region, and partial regions of the mitochondrial genome (COB2, ATP6, COX3, RNS, RNL), or nuclear genome including coding (GP43, PRP8, CHS2, ARF, CDC42, FKS) and non-coding regions (ORN1, 11b12b, 15b16b, AB, KL, MN, R56, TUB, III-IV, and XI-XII) revealed a great deal of diversity among clinical and environmental isolates, supporting the existence of other species beyond *P. brasiliensis* (Feitosa et al. 2003, Carrero et al. 2008, Salgado-Salazar et al. 2010). Using multilocus sequence analysis (MLSA), two distinct biological species *P. brasiliensis sensu lato* (containing at least four cryptic phylogenetic species: S1, PS2, PS3, and PS4) and *P. lutzii* (originally named Pb01-like) were identified (Matute et al. 2006, Teixeira et al. 2009, Theodoro et al. 2012, Turissini et al. 2017). The phylogeographical distribution is diverse in *Paracoccidioides*, and species 1 (S1) is the predominant agent of human PCM recovered from Argentina, Brazil, Paraguay, Peru, and Venezuela. Phylogenetic species 2 (PS2) was found in Brazil, Venezuela, and Uruguay, while the remaining phylogenetic species PS3 and PS4 appear to be restricted to Colombia and Venezuela, respectively (Theodoro et al. 2012). *Paracoccidioides lutzii* (formerly Pb01-like) is prevalent in central-west Brazil, mainly in Mato Grosso state, with scattered cases outside this area (Nery et al. 2021b).

Fungal taxonomy has undergone a significant transformation in recent decades as a method for inferring evolutionary relationships and defining species boundaries, especially with the introduction of molecular data in phylogenetic studies (Lücking et al. 2021). Recently, five species were proposed within *Paracoccidioides*. The *P. brasiliensis* complex includes the classical agent *P. brasiliensis sensu stricto* (formerly S1) in addition to the newly described *P. americana* (formerly PS2); *P. restrepiensis* (formerly PS3), and *P. venezuelensis* (formerly PS4) (Turissini et al. 2017, Teixeira et al. 2020). *Paracoccidioides lutzii* (formerly Pb01-like) is presented as a monophyletic group in phylogenetic analyses (Teixeira et al. 2014c). Nevertheless, there is no consensus among genetic, morphological, and clinical data (Shikanai-Yasuda et al. 2017, de Macedo et al. 2019, Hahn et al. 2019). The morphological markers for the recognition of different *Paracoccidioides* are scarce due to the overlapping phenotypic features. In this diverging scenario, whole-genome sequencing appears as an essential tool for elucidating relationships and resolving *Paracoccidioides* taxonomy (Muñoz et al. 2016).

Judging from a clinical perspective, preliminary studies found no significant clinical differences between the disease caused by members of the *P. brasiliensis* complex (de Macedo et al. 2019) or even between *P. brasiliensis s.l.* and *P. lutzii* (Hahn et al. 2019, Pereira et al. 2020). On the one hand, from a molecular epidemiological perspective, we can benefit from recognizing different genotypes of *Paracoccidioides* (Pinheiro et al. 2020, 2021). Thus, to explore intraspecific variation, it is necessary to

develop and apply molecular tools that are highly discriminatory at an affordable price. To meet this need, amplified fragment length polymorphisms (AFLP) can recognize genetic variations between any two *Paracoccidioides* genomes using a combination of restriction enzyme digestion of DNA, PCR amplification, and separation by capillary electrophoresis (Vos et al. 1995). AFLP has already been used successfully to study genetic variability in fungi, such as *Aspergillus fumigatus* (Warris et al. 2003), *Candida* spp. (Borst et al. 2003), *Coccidioides* species (Duarte-Escalante et al. 2013), *Cryptococcus* spp. (Hagen et al. 2015), *Fonsecaea* spp. (Najafzadeh et al. 2011), *Histoplasma* spp. (Rodrigues et al. 2020a), and *Sporothrix* spp. (de Carvalho et al. 2020).

We took advantage of the whole-genome sequences now available for *Paracoccidioides* in GenBank and conducted extensive bioinformatic analyses to screen for markers that were appropriate to address questions about the epidemiology and genetic diversity. These markers were subsequently evaluated *in vitro* to explore a vast collection of *Paracoccidioides* samples. Here, we report the AFLP primer combinations as an essential step to characterize specimens, species, and genotypes to complement PCM epidemiology with quality data.

MATERIAL AND METHODS

Fungal strains and DNA extraction

This study used one hundred sixty-five clinical and environmental strains of *Paracoccidioides* spp., recovered from Argentina, Brazil, Colombia, Guadeloupe Island, Peru, Uruguay, and Venezuela (Supplementary Table S1). These isolates are deposited in the Laboratory of Emerging Fungal Pathogens culture collection at the Federal University of São Paulo (UNIFESP), São Paulo, Brazil. Yeast cells were grown on Fava-Netto agar at 37 °C and co-cultured every seven days (Fava-Netto 1961, Fava-Netto et al. 1969). DNA extraction was performed from a 14-d-old yeast culture using the FastDNA kit (MP Bio-medicals, Solon, OH, USA) as previously described (Rodrigues et al. 2014). The genomic DNA concentration and purity (A260/A280 nm > 1.8) were analysed by spectrophotometry (NanoDrop 2000; Thermo Fisher Scientific, Waltham, MA, USA), and samples were stored at -20 °C.

Identifying *Paracoccidioides* by TUB1-RFLP

The PCR-restriction fragment length polymorphism (RFLP) of the α -tubulin gene (TUB1-RFLP) was performed using the protocol previously described (Roberto et al. 2016). Briefly, the α -tubulin gene was amplified from genomic DNA with the primers α -TubF and α -TubR (Table 1) (Kasuga et al. 2002). The PCR was incubated in a Mastercycler (Eppendorf, Hamburg, Germany) with an initial denaturation step of 5 min at 95 °C, followed by 35 cycles of 1 min at 94 °C, 45 s at 48 °C, 1 min at 68 °C, and a final extension of 10 min at 68 °C. For RFLP analysis, 3 μ L of the TUB1-PCR product were digested with 2 μ L 10 \times fast digest buffer, 1 μ L BclI endonuclease (10 U/ μ L; Thermo Fisher Scientific), 1 μ L MspI endonuclease (10 U/ μ L; Thermo Fisher Scientific) and ultrapure water to a final volume of 20 μ L. Tubes were incubated at 37 °C for 2 h, and the double-digested products were analysed by electrophoresis at 100 V on 2.5 % (w/v) agarose gels for

Table 1. Primers used in this study for generic amplification, sequencing, and genotyping.

Locus/Region	Primer	Primer sequence 5' to 3'	Tm (°C)	Sense	Amplicon (bp)	Reference
TUB1	α-TubF	CTGGGAGGTATGATAACACTGC	48 °C	Forward	263	Kasuga <i>et al.</i> 2002
	α-TubR	CGTCGGGCTATTGAGATTTAAG	48 °C	Reverse		Kasuga <i>et al.</i> 2002
ITS	ITS1	TCCGTAGGTGAACCTTGCGG	52 °C	Forward	620	White <i>et al.</i> 1990
	ITS4	TCCTCCGCTTATTGATATGC	52 °C	Reverse		White <i>et al.</i> 1990
MAT1-1	GMAT1-1 F	GCAATTGTCTATTCCATCAGT	56 °C	Forward	1 455	Torres <i>et al.</i> 2010
	GMAT1-1 R	CTAGATGTCAAGGTACTCGGTA	56 °C	Reverse		Torres <i>et al.</i> 2010
MAT1-1	MAT1-1 EST2-1 F	GGCATTTAACAAATCTTTACG	52 °C	Forward	400	Torres <i>et al.</i> 2010
	MAT1-1 EST2-1 R	CCCAGTTTGTAGCAATGAGT	52 °C	Reverse		Torres <i>et al.</i> 2010
MAT1-2	GMAT1-2 F	TTCGACCGTCCACGCCTATCTC	56 °C	Forward	1 208	Torres <i>et al.</i> 2010
	GMAT1-2 R	TCATTGCGAAAAGGTGTCAAG	56 °C	Reverse		Torres <i>et al.</i> 2010
MAT1-2	MAT1-2 EST G-F	CATGTCTCTGTCATTGTTCCA	52 °C	Forward	1 000	Torres <i>et al.</i> 2010
	MAT1-2 EST G-R	GGAACAAGGAGGTTGAAGTT	52 °C	Reverse		Torres <i>et al.</i> 2010

120 min in the presence of GelRed (Biotium, Fremont, CA, USA). The 50-bp DNA Step Ladder (Promega, Madison, WI, USA) was used as a size marker. The fragments were visualized using the L-Pix Touch imaging system under UV illumination (Roberto *et al.* 2016).

In silico AFLP analyses

Whole-genome sequence of nine *Paracoccidioides* isolates covering all the phylogenetic species described so far (Table 2) were *in silico* analysed to predict AFLP markers in the range of 50–500 bp. *In silico* AFLP analysis was performed using the software AFLPinSilico v. 2 (Rombauts *et al.* 2003) (available at <http://bioinformatics.psb.ugent.be/webtools/aflpinsilico/>). Briefly, *Paracoccidioides* genomes were retrieved from GenBank and *in silico* digested with EcoRI (a six-base cutter) and MseI (a four-base cutter) restriction enzymes. Afterward, a total of 16 combinations containing two selective bases (EcoRI+2 and MseI+2) were used to mine a subset of fragments. Combinations were selected based on the AFLP Microbial Fingerprinting kit (Applied Biosystems, Foster City, CA, USA). Finally, to accurately simulate the AFLP technique, we determined the length of all fingerprints with the addition of the adaptor and primer lengths. The diversity of the fragments was used to create a matrix of

amplicons, and data were visualized using the software Heat-mapper (Babicki *et al.* 2016). Hierarchical cluster analysis based on average linkage and Euclidean distance was applied to each row cluster.

AFLP fingerprinting

The AFLP fingerprinting analysis was conducted in duplicate with the AFLP Microbial Fingerprinting kit (Applied Biosystems). Digestion of *Paracoccidioides* DNA, adapter ligation, non-selective and selective amplification was performed following the manufacturer's recommendations, with minor modifications (Najafzadeh *et al.* 2011). Briefly, *Paracoccidioides* genomic DNA (200 ng) was digested *in vitro* using EcoRI (GAATTC) and MseI (TTAA) restriction enzymes (New England Biolabs, Ipswich, MA) and ligated to EcoRI and MseI adapters simultaneously. A pre-selective PCR was performed with EcoRI+0 and MseI+0 primers (Vos *et al.* 1995). Fluorescent AFLP was performed with 6-carboxyfluorescein (FAM) or NED fluorescent dye-labelled EcoRI primer with two bases selection (5'-GAC TGC GTA CCA ATT CNN-3') and unlabelled MseI primer with two bases selection (5'-GAT GAG TCC TGA GTA ACT -3'). Two different combinations were chosen to evaluate the potential for genetic characterization of *Paracoccidioides* isolates (combination #1

Table 2. Genomes of *Paracoccidioides* species retrieved from NCBI Genome database (<https://www.ncbi.nlm.nih.gov/genome>) for *in silico* analysis.

Strain	Species	Source	Origin	INSDC ¹ (WGS)	Total length (Mb)	BioProjects	Reference
T16B1	<i>P. brasiliensis</i>	<i>Dasypus novemcinctus</i>	Brazil	SRR4024730	29.1	PRJNA322632	Muñoz <i>et al.</i> 2016
Pb18		Chronic PCM	Brazil	ABKI02	29.5	PRJNA28733	Desjardins <i>et al.</i> 2011
Pb03	<i>P. americana</i>	Chronic PCM	Brazil	ABHV02	28.8	PRJNA27779	Desjardins <i>et al.</i> 2011
Pb262		Dog food	Brazil	SRR4024732	28.9	PRJNA322632	Muñoz <i>et al.</i> 2016
Pb339	<i>P. restrepiensis</i>	PCM	Brazil	SRR4024750	28.7	PRJNA322632	Muñoz <i>et al.</i> 2016
CNH		Chronic PCM	Colombia	LYUC01	29.4	PRJNA288047	Muñoz <i>et al.</i> 2016
Pb300	<i>P. venezuelensis</i>	Soil	Venezuela	LZY001	29.4	PRJNA287815	Muñoz <i>et al.</i> 2016
Pb01	<i>P. lutzii</i>	PCM	Brazil	ABKH02	32.6	PRJNA28731	Desjardins <i>et al.</i> 2011
PIEE		PCM	Brazil	SRR4024735	32.3	PRJNA322632	Muñoz <i>et al.</i> 2016

¹ International Nucleotide Sequence Database Collaboration (INSDC; <http://www.insdc.org/>)

FAM-EcoRI-AC/MseI-CT or #2 NED-EcoRI-AT/MseI-CT). All oligonucleotides were provided by Applied Biosystems in the AFLP Microbial Fingerprinting kit. AFLP fragments were determined by capillary electrophoresis with an ABI3730 Genetic Analyzer alongside a GeneScan 500 ROX internal size standard (35–500 bp; Applied Biosystems) at the Human Genome and Stem Cell Research Centre Core Facility (University of São Paulo, São Paulo, Brazil) under previously described conditions (de Carvalho *et al.* 2020). Electropherograms are representative of two independent experiments.

The selection of amplicons was automated, and only robust and high-quality amplicons were considered. Each electropherogram was carefully inspected to exclude doubtful peaks, setting the minimum threshold at 100 relative fluorescence units (RFU) and considering only peaks with sizes in the range of 50 and 500 base pairs. The size and diversity of the AFLP fragments were determined with BioNumerics v. 7.6 software (Applied Maths, Sint-Martens-Latem, Belgium). AFLP fragments were converted to the dominant presence (1) or absence (0) at probable fragment positions.

Pairwise genetic distances were calculated using the band-based Dice similarity coefficient (Dice 1945) combined with a "Fuzzy logic" option. Dendrograms were built using the unweighted pair group mean arithmetic method (UPGMA). To assess the consistency of a given cluster, we calculated the cophenetic correlation coefficient and its standard deviation, which determines the linear correlation coefficient between the cophenetic distances obtained from the tree and the dendrogram-derived similarities. Therefore, it is a measure of how accurately the AFLP-dendrogram represents the similarities among observations.

To estimate the existence of topological congruence between AFLP dendrograms and their associated confidence level, we determined the congruence index (I_{cong}) (de Vienne *et al.* 2007), based on maximum agreement subtrees (MAST). The correlation between experiments was calculated using the Pearson product-moment correlation coefficient (Pearson correlation) (Schober *et al.* 2018). A scatter plot was used to plot each pair of similarity values as one dot in a similarity plot between two experiment types. Especially for extensive data sets resulting in dense scatter plots, we used a histogram displaying a multi-colour scale ranging from white over blue, green, yellow, orange, and red to black.

Minimum spanning trees (MSTs) were calculated to explore the evolutionary relationships among all the observed genotypes of *Paracoccidioides*. MSTs characterize a set of edges (connections) that connect nodes (isolates) so that the summed distance of all branches is the shortest possible (Vauterin & Vauterin 2006). All figures were exported and treated using Corel Draw X8 (Corel, Ottawa, Canada).

Genetic diversity analysis

To calculate the potential of the two selective primer combinations evaluated here, the following polymorphism indices for dominant markers were calculated: polymorphic information content (PIC) (Botstein *et al.* 1980), expected heterozygosity (H) (Liu 1998), effective multiplex ratio (E) (Powell *et al.* 1996), arithmetic mean heterozygosity (H_{avg}) (Powell *et al.* 1996), marker index (MI) (Powell *et al.* 1996, Varshney *et al.* 2007), discriminating power (D) (Tessier *et al.* 1999), and resolving power (R_p) (Prevost & Wilkinson 1999).

Dimensioning analysis

Alternative grouping approaches such as principal component analysis (PCA) and multidimensional scaling (MDS) were employed to create three-dimensional plots according to their similarity. The optimization and position tolerances for choosing fragments were set to 0.10 %, and automated fragment matching was performed with a minimum profiling of 5 %. Default settings were applied for PCA and MDS, subtracting the average for characters. In addition, the Self-Organizing Map (SOM), a robust artificial neural network algorithm in the unsupervised learning category, was employed to classify AFLP entries in a two-dimensional space (map) according to their likeness (Kohonen 2001). The Kohonen map size was set to 100 (i.e., the number of neural network nodes in each direction). All figures were exported and treated using Corel Draw X8.

Structure analysis

Analysis of AFLP data in STRUCTURE (v. 2.3.4) (Pritchard *et al.* 2000) was performed using the admixture model, allowing alpha to be inferred and assuming correlated allele frequencies, using a burn-in period of 10 000 Markov chain Monte Carlo (MCMC) replications followed by 10 000 sampling replications, with 20 independent runs performed for K values one to twenty. All data were analysed using the method of Evanno and colleagues as implemented in StructureHARVESTER (v. 0.6.94) (Evanno *et al.* 2005, Earl & vonHoldt 2012) to determine the optimal number of clusters (K). Consensus population distributions were obtained with CLUMPP (v. 1.1.2) (Jakobsson & Rosenberg 2007), using the full search for the AFLP data. Final plots were generated using ggplot2 (Wickham 2016) in R (The R Core Team 2014).

Recombination analysis

To explore the relationships among *Paracoccidioides* species, a split network (Neighbor-Net) was constructed using the software SplitsTree v. 5.0.0 alpha (Huson & Bryant 2006) on AFLP profiles. For the construction of networks, we used the Hamming distances method (Hamming 1950) with the Neighbor-Net algorithm (Bryant & Moulton 2004) adapted for binary sequences (Huson & Kloepper 2005).

Analysis of molecular variance (AMOVA)

The AFLP data was transformed into a binary matrix of the presence/absence of each allele for each individual (Peakall & Smouse 2006, 2012). The genetic differentiation among populations was determined using PhiPT (Φ_{PT} , an analogue of F_{ST}). This measure allows intra-individual variation to be suppressed and is therefore ideal for comparing binary data with 9 999 permutations (Teixeira *et al.* 2014a). Analysis of molecular variance among and within populations was performed using GenAlex v. 6.5 (Excoffier *et al.* 1992).

Statistical analysis

We calculated Cohen's kappa coefficient (κ) and its 95 % confidence interval (CI) to determine the degree of concordance between AFLP typing and *TUB1*-RFLP (Roberto *et al.* 2016). Kappa values were read as follows: 0.00–0.20, poor agreement;

0.21–0.40, fair agreement; 0.41–0.60, moderate agreement; 0.61–0.80, good agreement; 0.81–1.00, very good agreement (Altman 1991). A P -value ≤ 0.05 was considered significant. All statistical calculations were performed with MedCalc Statistical Software v. 20.013 (MedCalc Software, Ostend, Belgium; <http://www.medcalc.org>; 2021). We calculated Simpson's diversity (Simpson 1949) and Shannon's diversity (Shannon 1948) for each organism/genetic group with the relative abundances estimated with frequency data.

Characterization of the mating-type idiomorphs

PCR primers targeting the *MAT1-1* or the *MAT1-2* region were used to determine the mating-types idiomorphs, as described before (Torres *et al.* 2010). Approximately 50 ng of genomic DNA was used for PCR with two sets of oligonucleotide primers: GMAT1-1 F and GMAT1-1 R, which amplify a 1 455 bp fragment from the α box region of the *MAT1-1* idiomorph, and GMAT1-2 F and GMAT1-2 R, which amplify a 1 208 bp fragment from the HMG domain gene, present in the *MAT1-2* idiomorph (Torres *et al.* 2010) (Table 1). PCRs were performed with PCR Master Mix buffer (Promega) as described above under the following conditions: 4 min at 95 °C; followed by 35 cycles of 1 min at 94 °C, 1 min at 56 °C, and 1 min at 72 °C; and a final step of 10 min at 72 °C. Samples were visualized on 1.2 % agarose gels as described above.

RESULTS

TUB1-RFLP

We conducted a retrospective molecular epidemiological study using the largest collection of *Paracoccidioides* strains from Latin

America ($n = 165$), preserved in our institution for more than 50 years (1970–2021). The *TUB1* gene was amplified followed by double digestion using the Bcl and MspI endonucleases, which produced four different electrophoretic patterns corresponding to 92 *P. brasiliensis* s. str. (S1; fragments of 155 bp and 108 bp), 22 *P. americana* (PS2; fragments of 62 bp, 93 bp, and 108 bp), 14 *P. restrepiensis* (PS3; amplicon remained intact with 263 bp) and 37 *P. lutzii* (Pb01-like; fragments of 62 bp and 204 bp). *TUB1*-RFLP did not allowed the recognition of *P. venezuelensis* (PS4) (Supplementary Table S1).

Development of AFLP markers for *Paracoccidioides*

The first step in our approach involved the *in silico* characterization of nine *Paracoccidioides* genomes retrieved from NCBI, comprising all medically relevant members described so far. AFLPinSilico was used to inspect restriction spots for EcoRI (GAATTC) and MseI (TTAA). Subsequently, a group of modified genomic fragments was generated by adding adaptor sequences, and an enriched group of modified genetic fragments was chosen based on two selective bases for EcoRI+2 and MseI+2 primers. Thus, we produced a matrix of 144 *in silico* AFLP profiles, which are shown as a heatmap in Fig. 1. A significant diversity of fragments was generated, ranging from 14–87 in 16 combinations evaluated in AFLPinSilico (Fig. 2). *Paracoccidioides lutzii* presented the largest genome core (~32.6 Mb), and we observed the most significant number of AFLP markers in all combinations, supported by a strong positive correlation (Pearson correlation = 0.926, $r^2 = 0.9623$, $P = 0.000337$) (Fig. 2).

We highlighted two combinations (#1 EcoRI-AC/MseI-CT or #2 EcoRI-AT/MseI-CT) to be evaluated *in vitro*, which revealed

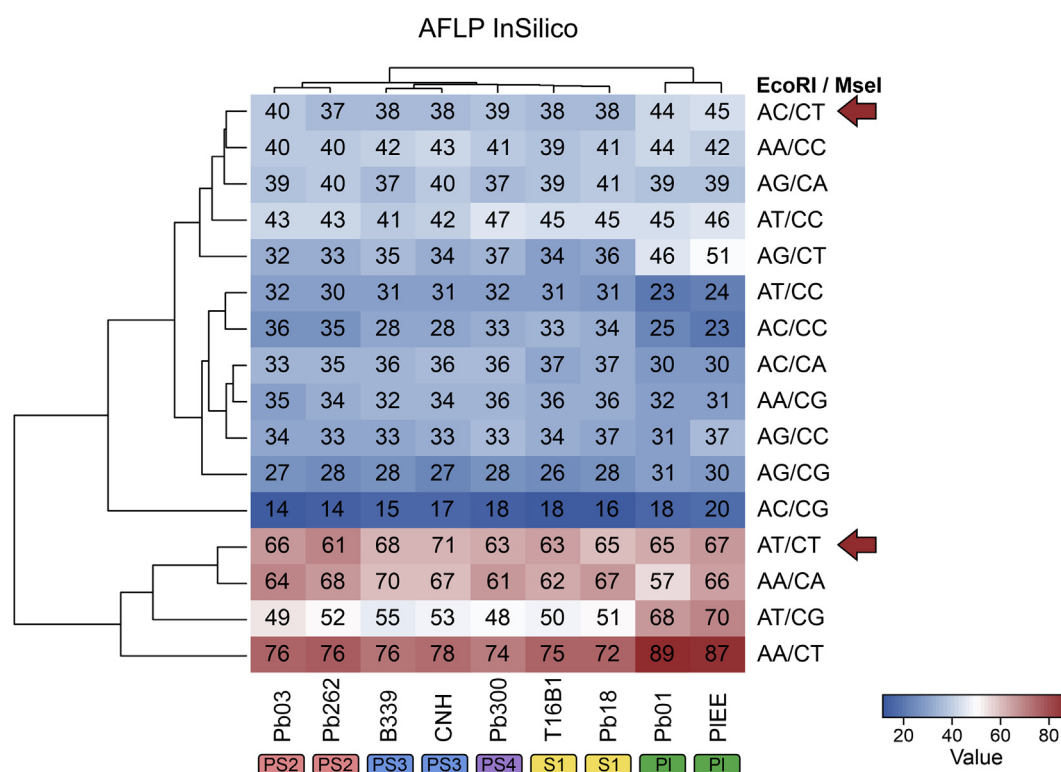


Fig. 1. High-resolution maps based on 16 AFLP fingerprints simulated using nine up-to-date sequenced *Paracoccidioides* genomes available at GenBank. The numbers inside the squares represent the expected numbers of amplicons after *in silico* digestion with EcoRI and MseI endonucleases following ligation of adaptors and selective amplification using selective primer EcoRI (5'-GAC TGC GTA CCA ATT C \overline{N} N-3') and MseI (5'-GAT GAG TCC TGA GTA A \overline{N} N-3'), as indicated.

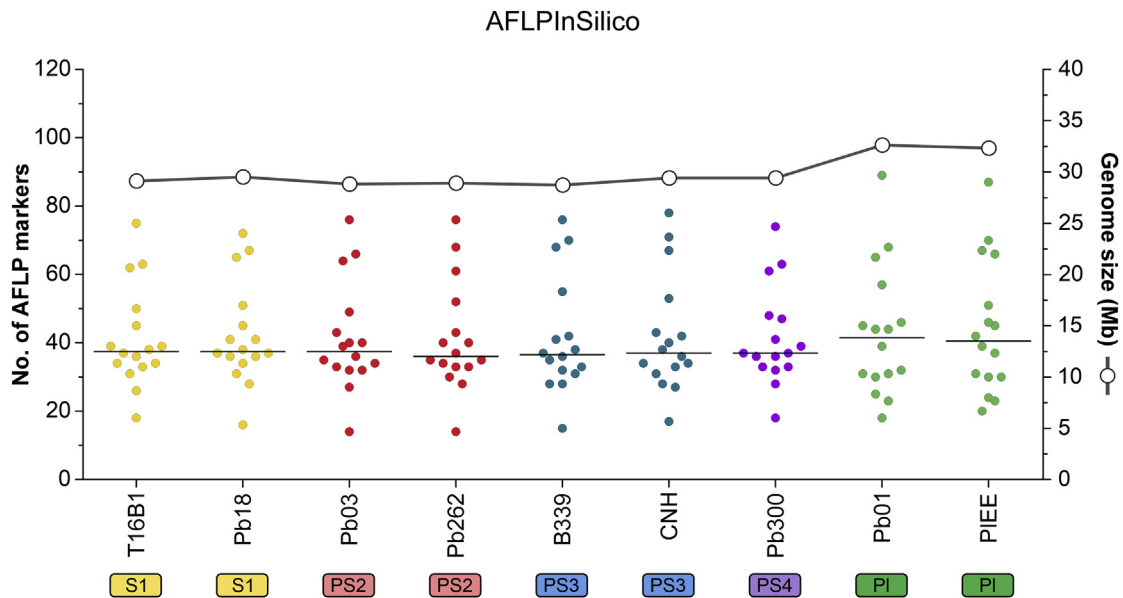


Fig. 2. A total of 16 combinations of selective EcoRI+2 and MseI+2 primer pairs were employed to generate 144 virtual AFLP profiles AFLPinSilico. The dots located on the left X-axis represented the number of fragments generated for each combination and were colour-coded according to their genetic groups. The bold bar represents the average of AFLP markers obtained for all combinations. The white dots located on the right X-axis represent the genome size, estimated by whole-genome sequencing.

the highest number of polymorphic markers (i.e., number and size) with the potential to speciate *Paracoccidioides* (Supplementary Table S2). A total of 154 polymorphic fragments were amplified *in vitro* using the selective primers EcoRI+2 and MseI+2, among them 67 and 87 loci, for combinations #1 and #2, respectively.

The dendrograms generated based on Dice's similarity coefficient are depicted in Figs 3 and 4. Clustering analysis shows five well-supported clades with a global similarity level ranging between 55.84 % \pm 3.53 % and 66.86 % \pm 1.49 %. The global cophenetic correlation coefficient between the dendrogram and the original similarity matrix was significant (96–97 %) for both markers supporting a reasonable degree of confidence in the association obtained for 165 isolates of *Paracoccidioides* (Supplementary Table S3). This AFLP clustering profile agrees with the broadly applied GP43-based classification (Morais et al. 2000).

The AFLP fingerprints revealed that 128 out of 165 isolates were embedded within the *P. brasiliensis* complex (cophenetic correlation values: #1 87 % and #2 78 %), with 79 isolates (48 %) classified as *P. brasiliensis* s. str. (AFLP S1), 22 isolates (13 %) as *P. americana* (AFLP PS2), 14 isolates (9 %) as *P. restrepiensis* (AFLP PS3), and 13 isolates (8 %) as *P. venezuelensis* (AFLP PS4). The second significant genetic cluster refers to 37 out of 165 isolates (22 %) which were classified as *P. lutzii* (cophenetic correlation values: #1 81 %, and #2 74 %) (Figs 3 and 4). The AFLP clusters classification was confirmed by *TUB1*-RFLP genotyping. To determine the level of concordance of the results of the *TUB1*-RFLP and any AFLP assay, we calculated the kappa statistic and its 95 % confidence interval (CI). A very good agreement was observed for *P. brasiliensis* ($\kappa = 0.843 \pm 0.041$, 95 % CI 0.762–0.923), *P. americana* ($\kappa = 1.0$, 95 % CI 1.000–1.000), *P. restrepiensis* ($\kappa = 1.0$, 95 % CI 1.000–1.000) and *P. lutzii* ($\kappa = 1.0$, 95 % CI 1.000–1.000), but poor agreement for *P. venezuelensis* ($\kappa = 0.0$, 95 % CI -2.8859×10^{-8} – 2.8859×10^{-8}). Although *TUB1*-RFLP could not distinguish *P. venezuelensis* (PS4), the AFLP fingerprinting could cluster the isolates into this group, considered

closely related to *P. brasiliensis* s. str. and *P. americana* under both markers (#1 and #2). In this case, the AFLP PS4 group was identified based on the reference strains EPM67 (Pb300/V1) and EPM73 that were characterized as *P. venezuelensis* in previous studies (Salgado-Salazar et al. 2010, Muñoz et al. 2014, Turissini et al. 2017, Pinheiro et al. 2021).

To assess the existence of topological correspondence between the two AFLP dendrograms, we used the congruence index (I_{cong}) (de Vienne et al. 2007), and the Pearson product-moment correlation coefficient (Pearson correlation). A comparable and constant clustering signature was noted in pairwise comparisons, as demonstrated by the great congruence index value and their significant associated *P*-value ($I_{cong} = 2.16$; $P = 3.93 \times 10^{-14}$), as well as a strong positive correlation for the Pearson product-moment correlation coefficient ($r = 87.018$ %, $P < 0.00001$) (Fig. 5). Thus, our AFLP dendrograms are more congruent than expected by chance, supporting the use of new AFLP markers to speciate *Paracoccidioides* and to explore both deep and fine-scale genetic structures.

The averages of fragments for combination #1 (EcoRI-AC / MseI-CT) varied per species between 37–49 for *P. brasiliensis* s. str. (Median = 44; CV = 5.79 %); 44–47 for *P. americana* (Median = 46; CV = 1.90 %); 37–41 for *P. restrepiensis* (Median = 40; CV = 2.38 %); 41–46 for *P. venezuelensis* (Median = 43; CV = 3.35 %); and 44–50 for *P. lutzii* (Median = 49; CV = 4.18 %). A greater number of fragments was observed for combination #2 (EcoRI-AT / MseI-CT) varying between 43–69 for *P. brasiliensis* s. str. (Median = 58; CV = 10.61 %); 62–68 for *P. americana* (Median = 66; CV = 2.79 %); 61–67 for *P. restrepiensis* (Median = 63.5; CV = 2.41 %); 63–69 for *P. venezuelensis* (Median = 67; CV = 2.79 %); and 41–66 for *P. lutzii* (Median = 58; CV = 11.06 %) (Supplementary Table S3). Table 3 shows the features of marker attributes for AFLP primer combinations #1 and #2.

The PIC established for each primer pair was dissimilar among species but comparable between markers. In general, PIC values varied from low polymorphism in *P. americana*

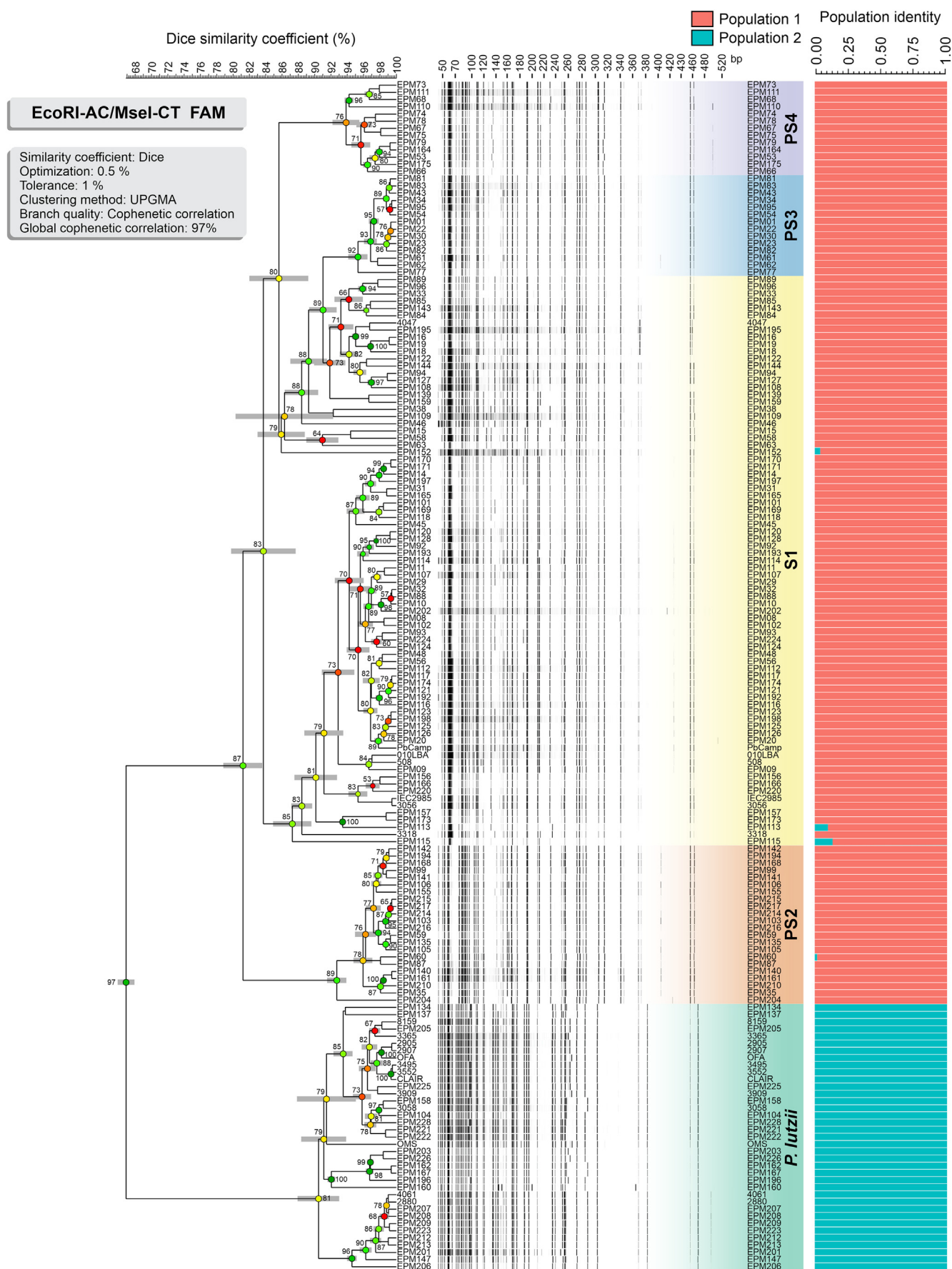


Fig. 3. The UPGMA dendrogram, based on AFLP fingerprint, generated with a total of four selective bases (FAM-EcoRI-AC/MseI-CT) for 165 *Paracoccidioides* spp. originated from Latin America. The dendrogram shows cophenetic correlation values (circles, which are represented by colour ranges between green-yellow-orange-red according to decreasing cophenetic correlation) for a given clade and its standard deviation (grey bar). For pairwise genetic distances calculation, the Dice coefficient was used. The cophenetic correlation of the dendrogram is 97 %. Bayesian cluster analyses with STRUCTURE ($k = 2$) of 165 *Paracoccidioides* spp. based on AFLP. Each vertical bar represents one individual and its probabilities of being assigned to clusters. Further information about isolate sources can be found in [Supplementary Table S1](#).

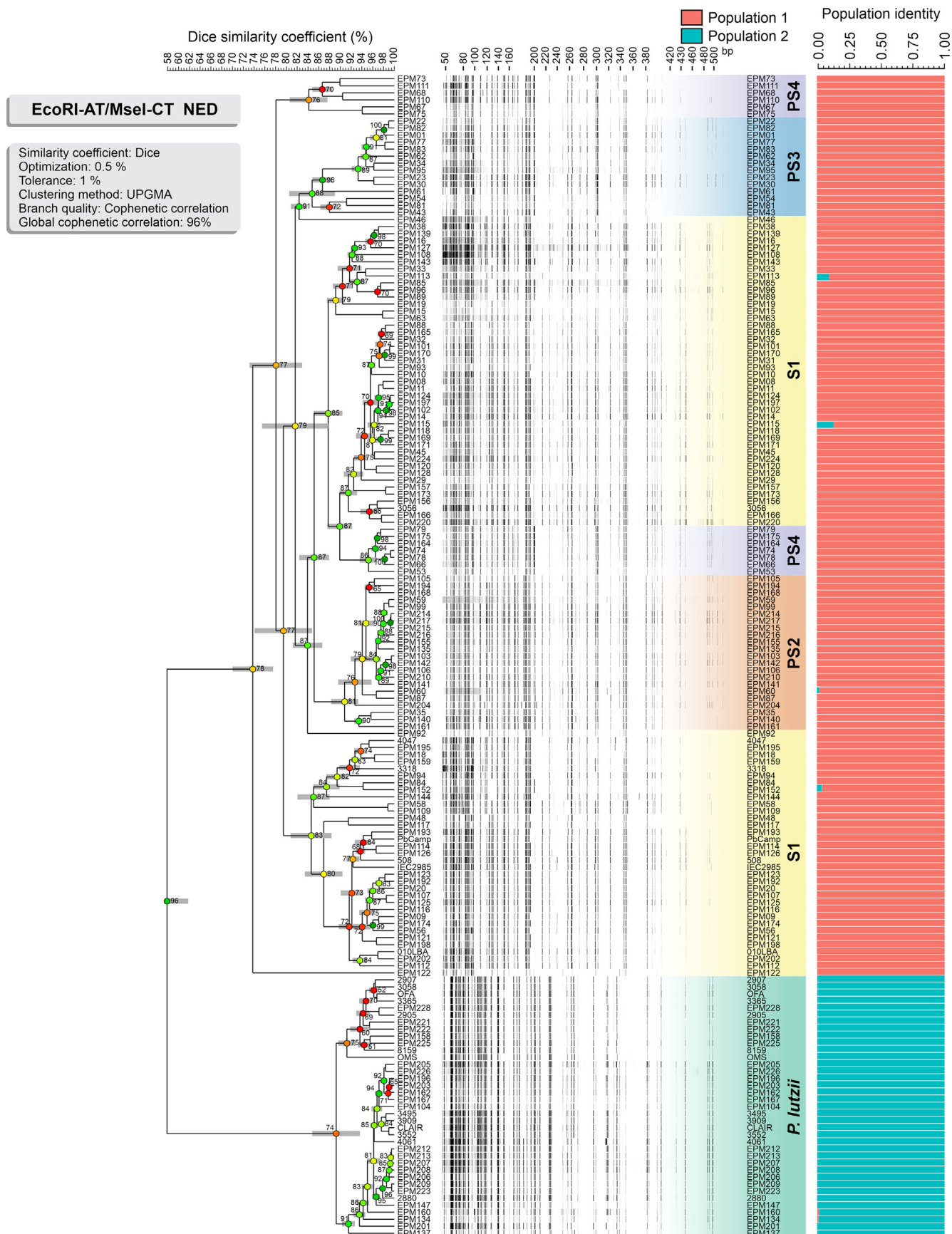


Fig. 4. The UPGMA dendrogram, based on AFLP fingerprint, generated with a total of four selective bases (NED-EcoRI-AT/MseI-CT) for 165 *Paracoccidioides* spp. originated from Latin America. The dendrogram shows cophenetic correlation values (circles, which are represented by colour ranges between green-yellow-orange-red according to decreasing cophenetic correlation) for a given clade and its standard deviation (grey bar). For pairwise genetic distances calculation, the Dice coefficient was used. The cophenetic correlation of the dendrogram is 96 %. Bayesian cluster analyses with STRUCTURE ($k = 2$) of 165 *Paracoccidioides* spp. based on AFLP. Each vertical bar represents one individual and its probabilities of being assigned to clusters. Further information about isolate sources can be found in [Supplementary Table S1](#).

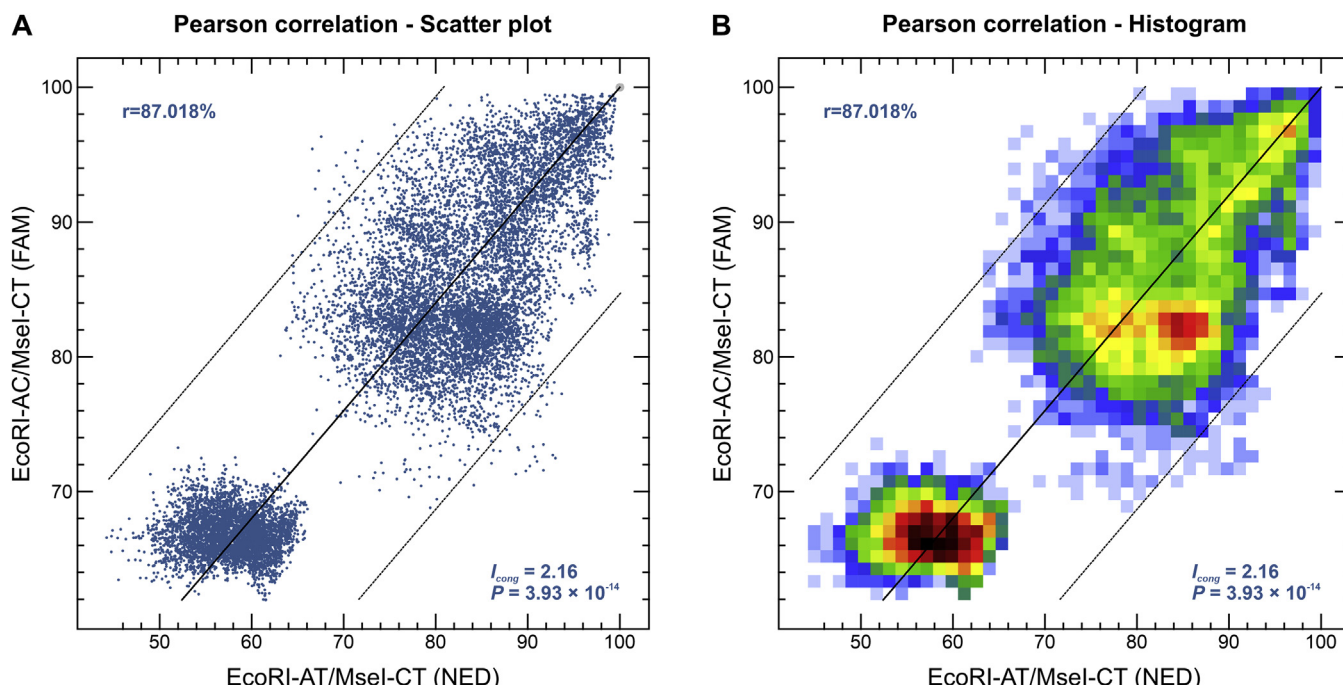


Fig. 5. The correlation between AFLP experiments evaluated for 165 *Paracoccidioides* isolates. A similarity plot for two experiments EcoRI-AC/MseI-CT and EcoRI-AT/MseI-CT was assessed using (A) the Pearson correlation coefficient (scatter plot) to plot each pair of similarity values as one dot, and (B) the Pearson correlation coefficient (histogram) representing the average the number of dots in each area. A multi-colour scale ranges continuously from white over blue, green, yellow, orange, and red to black.

($PIC = 0.1355-0.1364$), *P. restrepiensis* ($PIC = 0.0680-0.1878$), and *P. venezuelensis* ($PIC = 0.1671-0.2008$) to average polymorphism in *P. brasiliensis* s. str. ($PIC = 0.2447-0.2925$), and *P. lutzii* ($PIC = 0.2086-0.2036$), and both markers presented high discriminating power ($D = 0.5183-0.5553$). The highest overall PIC value was observed for primer combination #1 ($PIC = 0.3456$), and the lowest was noted for primer combination #2 ($PIC = 0.3345$), supporting good diversity among the studied *Paracoccidioides*. Overall, *P. brasiliensis* s. str. and *P. lutzii*

showed slightly higher PIC values than the remaining phylogenetic species (Table 3).

The global usefulness of each marker system was estimated using the marker index (MI), which was obtained as a product of polymorphic information content and effective multiplex ratio. Equal overall MI values ($MI = 0.0018$) were obtained for both combinations. A moderate positive correlation was observed between MI and PIC values (combination #1 Pearson correlation = 0.6797, $r^2 = 0.462$, $P = 0.206831$; combination #2 Pearson

Table 3. Summary of polymorphism statistics calculated for two different pairs of selective primers (EcoRI+2 and MseI+2) for *Paracoccidioides* species.

#1 EcoRI-AC/MseI-CT								
Species	Scored bands	H	PIC	E	H_{avp}	MI	D	R_p
S1 (n = 79)	53	0.2854	0.2447	43.8608	0.0001	0.0030	0.3152	10.6076
PS2 (n = 22)	50	0.1472	0.1364	46.0000	0.0001	0.0062	0.1537	2.3636
PS3 (n = 14)	41	0.0705	0.0680	39.5000	0.0001	0.0049	0.0719	1.5714
PS4 (n = 13)	48	0.1841	0.1671	43.0769	0.0003	0.0127	0.1948	4.1538
<i>P. lutzii</i> (n = 37)	56	0.2366	0.2086	48.3243	0.0001	0.0055	0.2554	7.1892
Overall (n = 165)	67	0.4443	0.3456	44.6788	0.00004	0.0018	0.5553	22.3152
#2 EcoRI-AT/MseI-CT								
Species	Scored bands	H	PIC	E	H_{avp}	MI	D	R_p
S1 (n = 79)	77	0.3558	0.2925	59.1772	0.0001	0.0035	0.4094	22.0253
PS2 (n = 22)	71	0.1462	0.1355	65.3636	0.0001	0.0061	0.1525	6.7273
PS3 (n = 14)	63	0.2098	0.1878	55.5000	0.0002	0.0132	0.2240	11.8571
PS4 (n = 13)	65	0.2265	0.2008	56.5385	0.0003	0.0152	0.2435	15.8462
<i>P. lutzii</i> (n = 37)	74	0.2300	0.2036	64.1892	0.0001	0.0054	0.2476	12.3784
Overall (n = 165)	87	0.4247	0.3345	60.3818	0.00002	0.0018	0.5183	34.3152

D = discriminating power; E = effective multiplex ratio; H = expected heterozygosity; H_{avp} = mean heterozygosity; MI = marker index; PIC = polymorphism information content; R_p = resolving power.

correlation = 0.6896, $r^2 = 0.4755$, $P = 0.197644$). The resolving power (R_p), which is the ability of each primer combination to detect level of variation among individuals was found to be higher in primer combination #2 ($R_p = 34.3152$) and lower for primer combination #1 ($R_p = 22.3152$) (Table 3). The R_p values were not correlated with MI for combination #1 (Pearson correlation = 0.4111, $r^2 = 0.169$, $P = 0.491713$), but for combination #2 (Pearson correlation = 0.9336, $r^2 = 0.8716$, $P = 0.020334$).

We assessed the expected heterozygosity (H), which is the probability that an individual in the population is heterozygous for the locus. The expected heterozygosity relates to Nei's unbiased gene diversity (H_S), as adapted for dominant markers under the assumptions of Hardy-Weinberg equilibrium and the Lynch-Milligan model (Lynch & Milligan 1994). The overall average expected heterozygosity for *Paracoccidioides* species ranged between 0.4247–0.4443 (Table 3). The high values for expected heterozygosity among *P. brasiliensis* ($H = 0.2854$ – 0.3558) and *P. lutzii* isolates ($H = 0.2300$ – 0.2366) supports high genetic diversity in these groups (Muñoz et al. 2016, Teixeira et al. 2020). The remaining species, such as *P. americana* ($H = 0.1462$ – 0.1472), *P. restrepiensis* ($H = 0.0705$ – 0.2098), and *P. venezuelensis* ($H = 0.1841$ – 0.2265), showed discrete variation, which is in accordance with a prevalently clonal population (Table 3). Moreover, the application of a concordant genotyping method to *Paracoccidioides* species, for which the relationship between the number of AFLP markers used and the estimated genetic diversity converged to the expected variation, further confirms that the approach described here allows assessment of the accuracy of inferences on the genetic diversity of prevalently clonal organisms derived using combinations #1 (EcoRI-AC/MseI-CT) and #2 (EcoRI-AT/MseI-CT) (Arnaud-Haond et al. 2005).

Structure analysis

We found a strong correlation between population structure and *Paracoccidioides* species. The structure analysis indicated two genetic clusters as the most probable number of genetically distinct populations (Figs 3 and 4). The Delta K plot (Fig. 6) showed the highest peak at $K = 2$, supporting the partition into two genetic clusters with no or a very weak signal of admixture (Supplementary Fig. S1). For $K = 2$, members of the *P. brasiliensis* complex clustered with the population 1. On the other hand, the second cluster corresponded to the *P. lutzii* isolates embedded in population 2, originating from endemic areas mainly in mid-west Brazil (Figs 3 and 4).

The AFLP profiles were employed to generate pairwise genetic distance matrices based on Dice's similarity coefficient, which were then analysed using PCA. The PCA plots for combinations #1 and #2 are shown in Fig. 7, and the distribution of 165 *Paracoccidioides* isolates among the three coordinates illustrated a trend similar to cluster analysis. Combination #1 revealed the highest cumulative percentage explained, with 68.2 % of the variation described by the first three components (coordinates X, Y, and Z). PCAs and MDSs analysis indicated considerable intraspecific clustering as well as a large genetic separation between any two taxa (interspecific variation). The structure evidenced by PCA supports the separation of *P. brasiliensis* complex and *P. lutzii*, consistent with the higher level of intraspecific variability shown in dendrogram analysis. Importantly, AFLP results agree with those detected using whole-genome sequencing (Muñoz et al. 2016, Teixeira et al. 2020).

The structure of members of the *P. brasiliensis* complex and *P. lutzii* is confirmed by the AFLP-derived MSTs in Fig. 8, with most isolates having a unique genotype. A few isolates in the

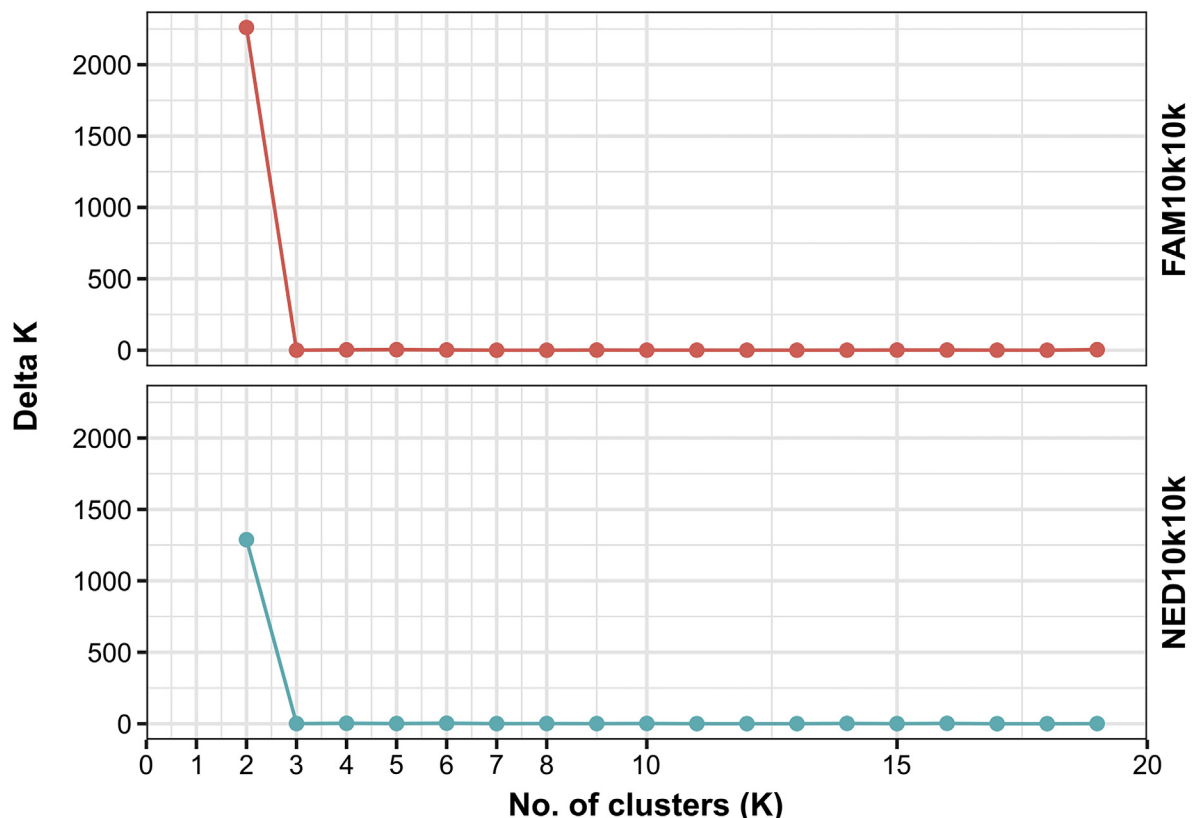


Fig. 6. STRUCTURE Harvester results. The most plausible number of genetic clusters (K) within the complete data set of 165 individuals based on the method depicted by Evanno et al. (2005). Population genetic structure of the estimated ΔK value determined the maximum value at $K = 2$.

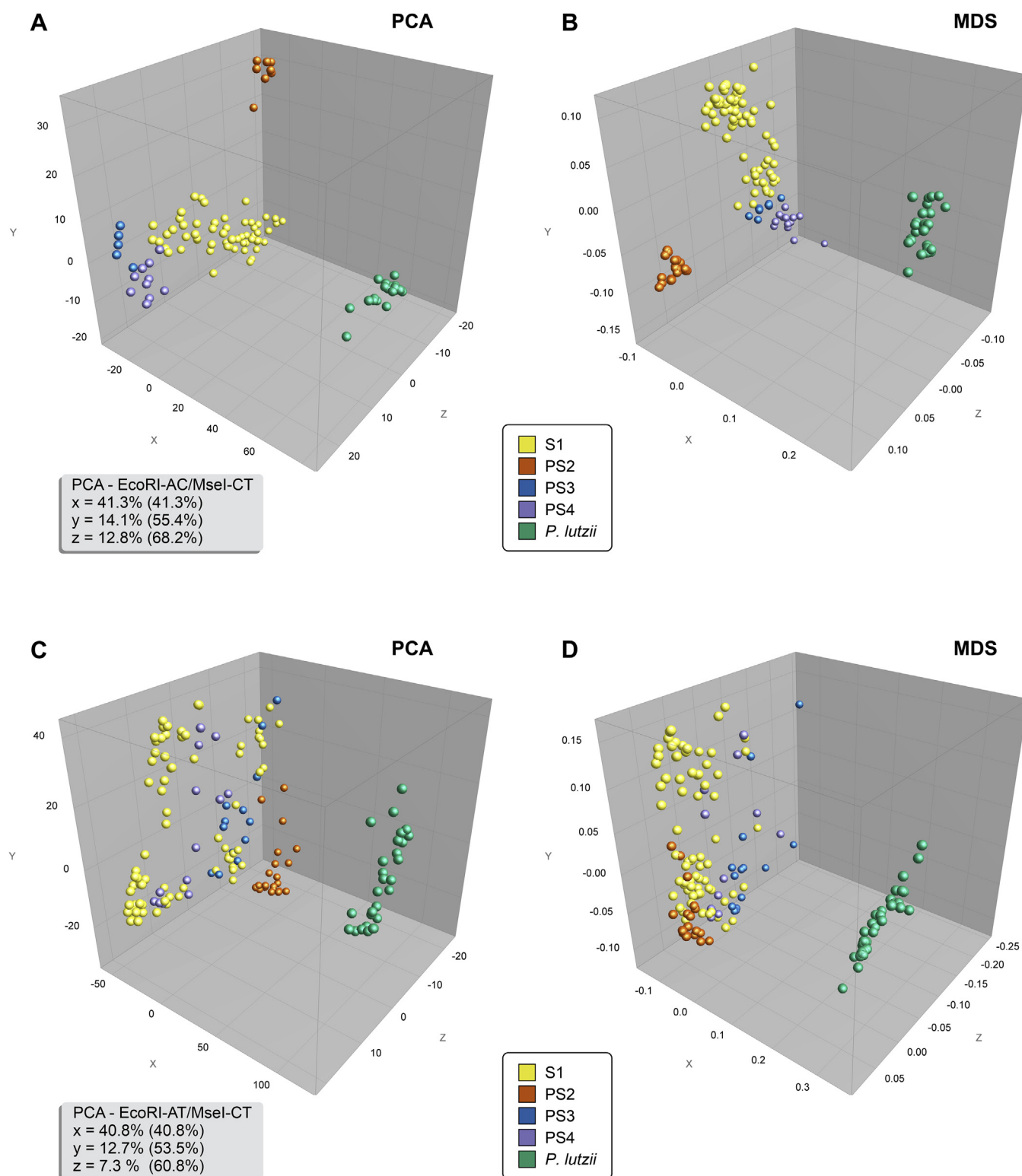
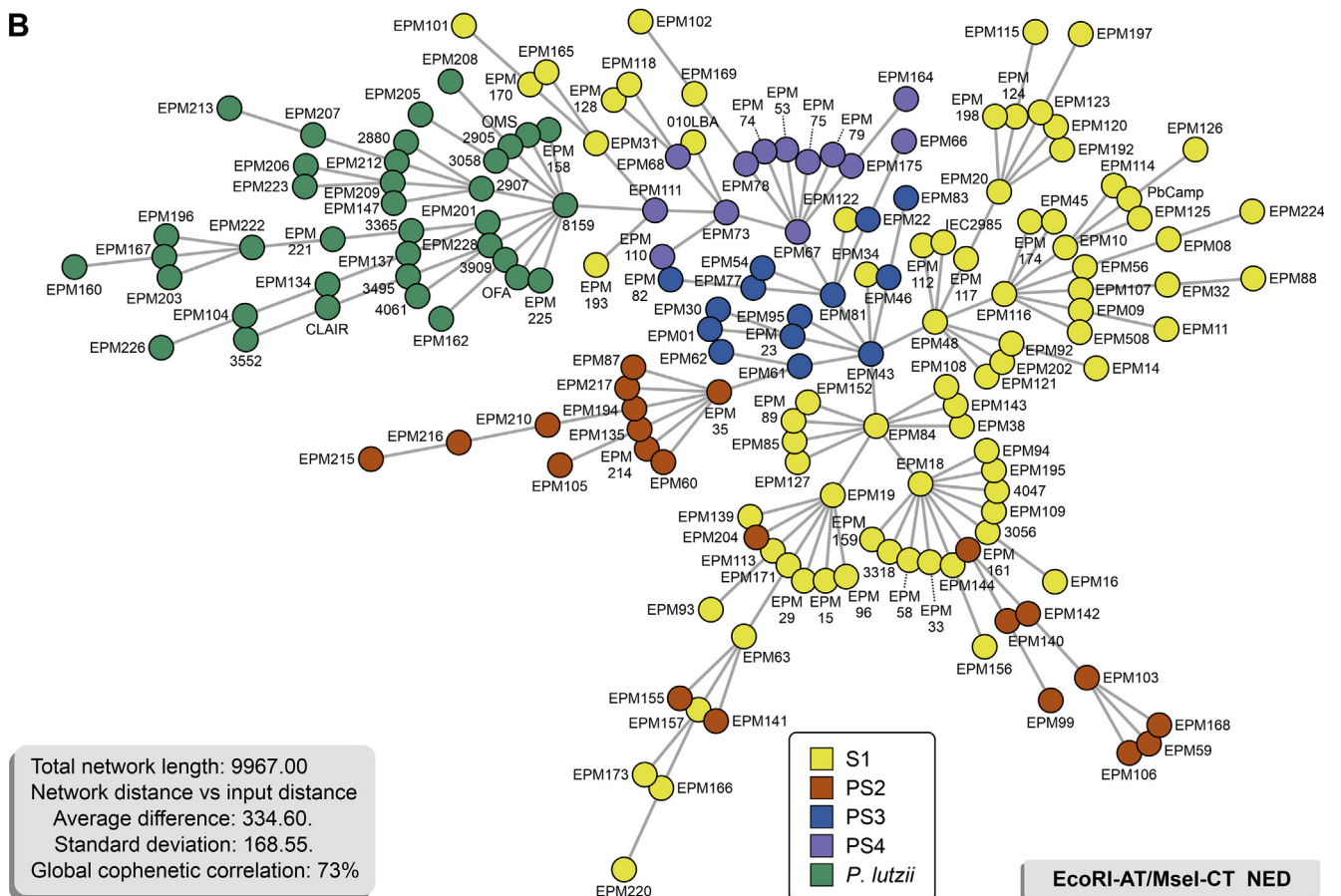
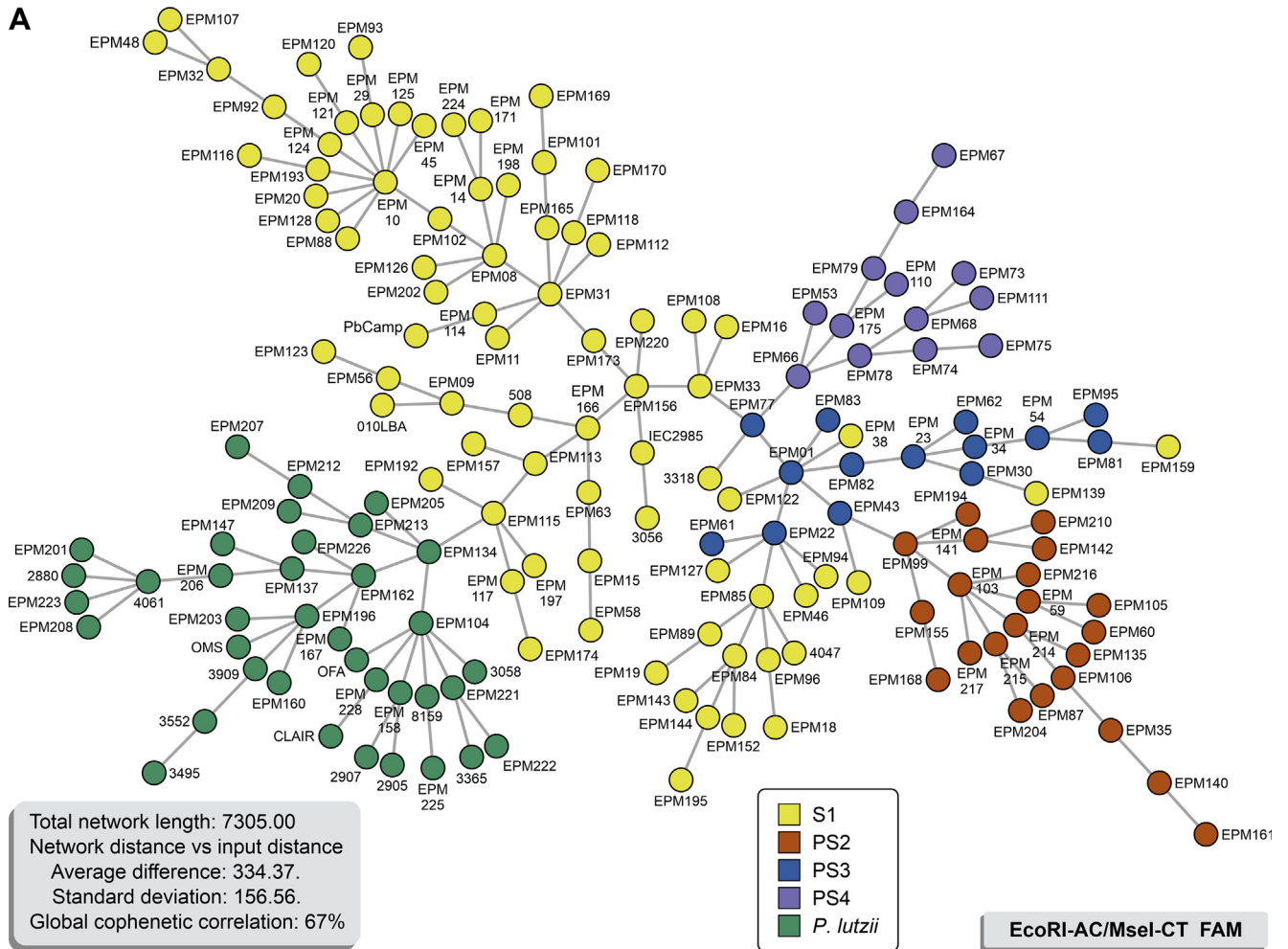


Fig. 7. Principal component analysis (PCA) and Multidimensional scaling (MDS) analysis of the combinations #1 EcoRI-AC/MseI-CT (67 loci) and #2 EcoRI-AT/MseI-CT (87 loci) informative AFLP markers plotted in three-dimensional space coloured according to the genetic groups. **(A)** PCA, and **(B)** MDS based on combination #1 EcoRI-AC/MseI-CT ($n = 165$). **(C)** PCA, and **(D)** MDS based on combination #2 EcoRI-AT/MseI-CT ($n = 165$). PCAs and MDS were created in the software BioNumerics v. 7.6.

P. brasiliensis complex were randomly distributed, similar to the dendrogram, particularly for combination #2 (Fig. 8B), suggesting a plausible chain of disease transmission in molecular epidemiology (Salipante & Hall 2011). Many invariable fragments were observed in *P. americana*, *P. restrepiensis*, and *P. venezuelensis* (as evidenced by low *PIC* values in Table 3), and together with the overall high similarity of > 80 % between the fingerprints, agreed with a monophyletic origin of the isolates.

SOMs, an unsupervised artificial neural network, were used to cluster high-dimensional AFLP data by projecting it according to genetic clusters onto a low-dimensional map (Fig. 9). Contrasting to PCA, the distance between entries in the SOMs is not proportional to the taxonomic distance between the entries (Felix et al. 2015). Therefore, in Fig. 9, *Paracoccidioides* strains with low genetic distance form clusters (typified by black blocks). The relative genetic distance between neighbouring groups (black



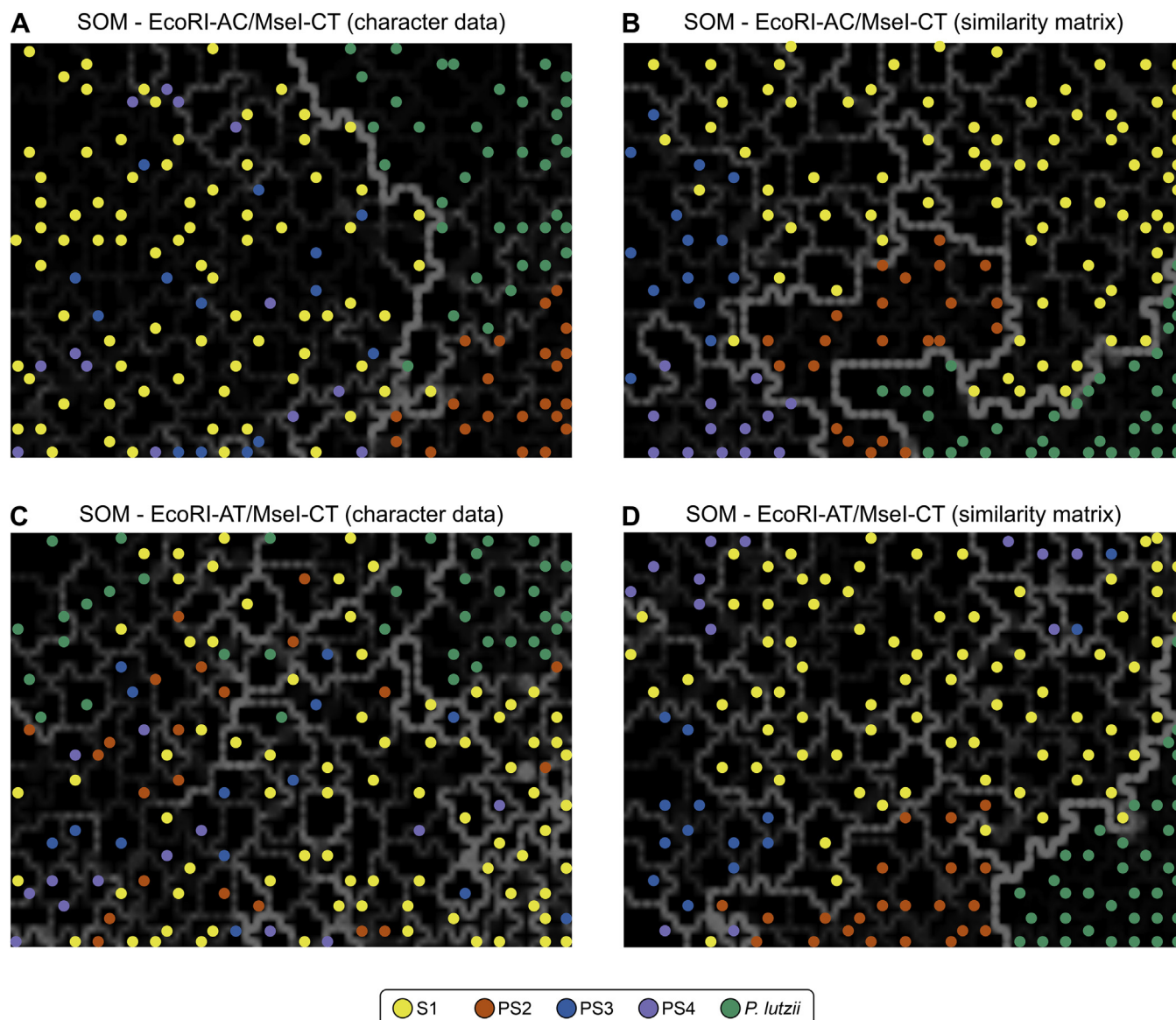


Fig. 9. The distribution of the studied AFLP genotypes of 165 *Paracoccidioides* species originated from Latin America, using self-organizing mapping (SOM). The dimensioning analyses were performed using BioNumerics v. 7.6 to determine the consistency of the differentiation of the populations defined by the cluster analysis. (A) and (B) show the SOM for EcoRI-AC/MseI-CT combination (67 loci) using character data (binary matrix) and similarity matrix, respectively. (C) and (D) show the SOM for EcoRI-AT/MseI-CT combination (87 loci) using character data (binary matrix) and similarity matrix, respectively. The lighter and thicker the line (white, grey) between black blocks, the more distant are those samples contained in the black block from the adjacent black block. Isolates were colour-coded according to their genetic groups.

blocks) is designated by the intensity of white lines separating the clusters, with closely related groups separated by faint dark lines and more distantly related strains separated by increasingly lighter thicker lines. Thus, phylogenetic species displaying slight intraspecific variation, such as *P. americana*, *P. restrepiensis*, and *P. venezuelensis*, tended to remain closer, separated by thinner lines, but bright solid lines were observed separating clusters interspecifically (Fig. 9).

In a phylogenetic network analysis of *Paracoccidioides* AFLP profiles, we found that members of the *P. brasiliensis* complex and *P. lutzii* were the most differentiated from each other in both

markers (Fig. 10). Moreover, in the reconstruction of evolutionary history, phylogenetic networks revealed large sets of parallel edges, suggestive of recombination events.

AMOVA

We used AMOVA to investigate genetic variance among 165 individuals of the two populations in *Paracoccidioides* (*P. brasiliensis* complex, $n = 128$, population 1; *P. lutzii*, $n = 37$, population 2). Table 4 shows the AMOVA findings for the population genetic analysis. AMOVAs performed for marker #1 in

Fig. 8. Minimum Spanning Trees (MSTs) showing the genetic relationship between 165 *Paracoccidioides* genotypes using (A) EcoRI-AC/MseI-CT (Total network length 7 305.00) and (B) EcoRI-AT/MseI-CT (Total network length 9 967.00). Each genotype was considered unique. Isolates were colour-coded according to their genetic groups. The distance between genotypes in the diagram does not reflect any relationship with the genetic distance between genotypes. The annotation of the genetic distance between each edge that connects the nodes is shown in Supplementary Fig. S2 (EcoRI-AC/MseI-CT) and Supplementary Fig. S3 (EcoRI-AT/MseI-CT).

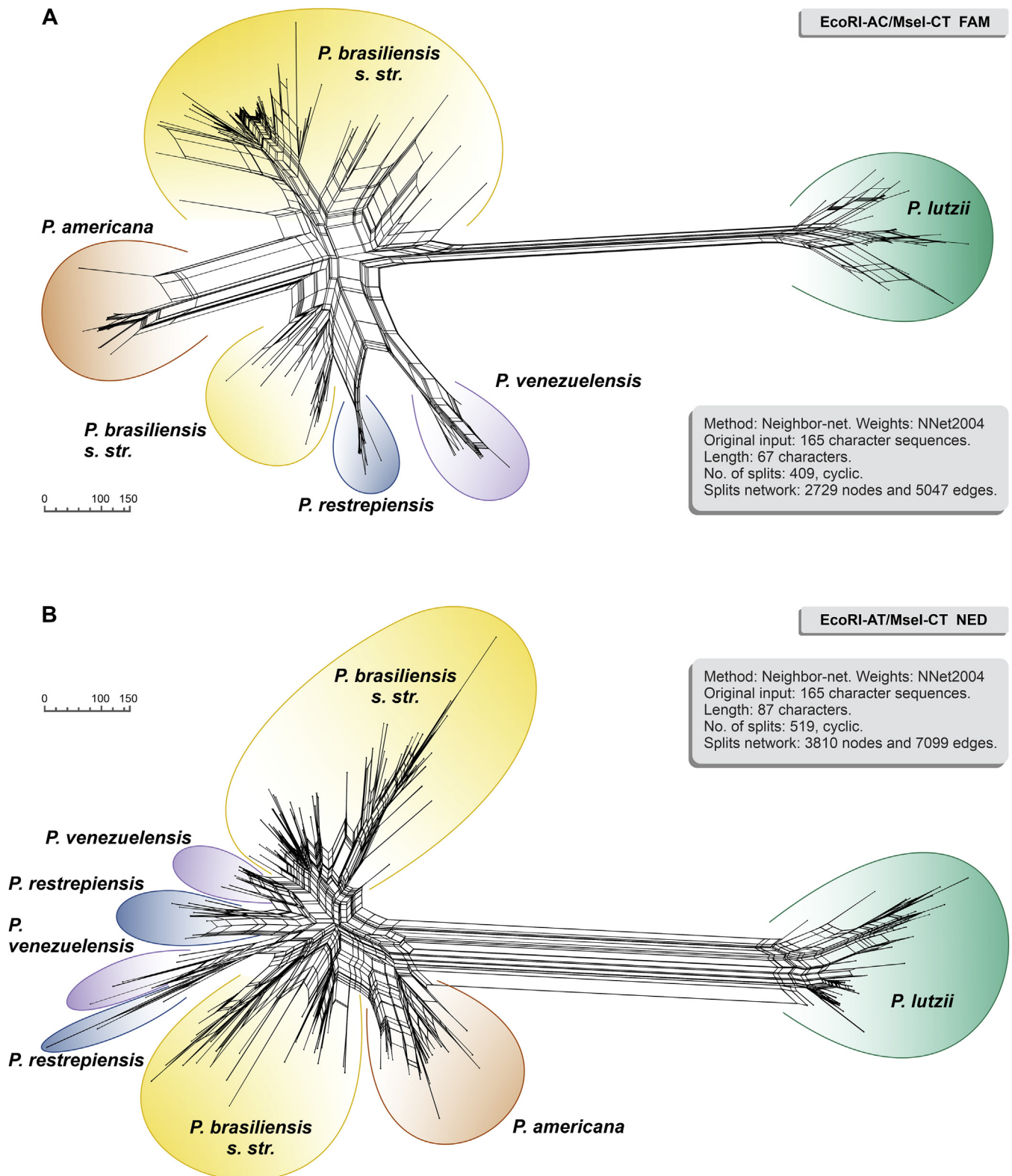


Fig. 10. Neighbor-Net network showing genetic relationships based on AFLPs among *Paracoccidioides* species (scale equals genetic distance). **(A)** EcoRI-AC/MseI-CT and **(B)** EcoRI-AT/MseI-CT split networks. Analysis was performed using SplitsTree v. 5.0.0_alpha (Huson & Bryant 2006) for binary sequences (Huson & Kloepper 2005), and the original input consisted of 165 standard character sequences.

P. brasiliensis complex and *P. lutzii* showed that 66 % of the total genetic variance was triggered by variability among populations, whereas 34 % was driven by variability within populations (PhiPT = 0.658, $P < 0.0001$). A similar trend was observed for marker #2 with 65 % of total variation among populations and 35 % within populations (PhiPT = 0.651, $P < 0.0001$) (Supplementary Fig. S4). The hierarchical analysis of molecular variance leaves a strong differentiation among the groups,

supporting a highly structured population. The results were highly significant ($P < 0.0001$).

Mating-type

A mating type-specific PCR assay was used to amplify the *MAT1-1* or the *MAT1-2* regions among 165 *Paracoccidioides* isolates. The *MAT1-1* region was detected in 88 isolates, while

Table 4. Analysis of molecular variance (AMOVA) shows the partitioning of genetic variation within and between *Paracoccidioides* species populations.

Marker	Source of variation	Df	SS	MS	Est. var.	%	P-value
#1 EcoRI-AC /MseI-CT	Among Population	1	520.417	520.417	8.984	66 %	0.0001
	Within Population	163	760.516	4.666	4.666	34 %	0.0001
#2 EcoRI-AT /MseI-CT	Among Population	1	775.274	775.274	13.380	65 %	0.0001
	Within Population	163	1166.811	7.158	7.158	35 %	0.0001

df = degree of freedom, SS = sum of squares, MS mean squares, Est. var. = estimate of variance, % = percentage of total variation, P-value is based on 9 999 permutations.

the *MAT1-2* region was detected among 77 isolates ($\chi^2 = 0.733$; $P = 0.3918$). Heterothallism (self-sterility) was the universal mating strategy amongst *Paracoccidioides* species. The distribution of each sexual idiomorph within molecular species (S1, PS2, PS3, PS4, and *P. lutzii*) is presented in Table 5. The distributions of *MAT1-1* or *MAT1-2* idiomorph were not significantly skewed (1:1 ratio) for *P. brasiliensis* s. str. ($\chi^2 = 1.025$; $P = 0.3113$), *P. venezuelensis* ($\chi^2 = 0.692$; $P = 0.4054$), and *P. lutzii* ($\chi^2 = 0.027$; $P = 0.8694$), supporting the presence of random mating within each species. However, a biased distribution was found for *P. americana* ($\chi^2 = 8.909$; $P = 0.0028$) and *P. restrepiensis* ($\chi^2 = 4.571$; $P = 0.0325$) with an overwhelming presence of *MAT1-1* idiomorphs.

Phylogenetic trends in *Paracoccidioides*

Distribution patterns were explored combining our dataset (n = 165) with data from 333 *Paracoccidioides* isolates reported in the literature and identified down to species level using molecular methods (e.g., whole-genome sequencing, MLSA, DNA fingerprint) (Matute et al. 2006, Carrero et al. 2008, Teixeira et al. 2009, Salgado-Salazar et al. 2010, Theodoro et al. 2012, Turissini et al. 2017, de Macedo et al. 2019, Hahn et al. 2019, Cocio et al. 2020b, Teixeira et al. 2020, Mattos et al. 2021, Nery et al. 2021b). Phylogenetic trends revealed that the *P. brasiliensis* complex species, including the four cryptic species, are widely distributed among different

countries in Latin America. Most cryptic siblings occur in sympatry, as exemplified by *P. brasiliensis* s. str. and *P. americana*, with a clear overlapping distribution. In contrast, *P. lutzii* is endemic to Brazil (Fig. 11). The source of isolation revealed an overwhelming occurrence of clinical isolates (90.77 %, n = 452 out of 498) followed by animals (e.g., armadillo, dog, and penguin; 7.23 %, n = 36 out of 498), and from the environment (e.g., soil and dog food; 1 %, n = 5 out of 498). The source of isolation was unknown for five strains (1 %). Most molecularly characterized isolates (n = 498) are from Brazil (74.69 %, n = 372), followed by Venezuela (7.83 %, n = 39), Argentina (5.62 %, n = 28) Colombia (5.42 %, n = 27), Peru (2.20 %, n = 11), Paraguay (1.4 %, n = 7), Uruguay (0.8 %, n = 4), Bolivia (0.4 %, n = 2), Ecuador (0.2 %, n = 1) and Guadeloupe Island (0.2 %, n = 1). These data reveal the urgency to increase genetic surveillance in *Paracoccidioides*-affected areas (Fig. 11A).

Paracoccidioides brasiliensis s. str. (S1) is predominantly found in southeastern and southern Brazil, Argentina, Peru, Venezuela, Paraguay, Uruguay, Bolivia, and Guadeloupe Island. *P. americana* has sporadic distribution and is less frequently reported, with human cases described thus far in Brazil, Venezuela, Uruguay, and Argentina. The remaining species, such as *P. restrepiensis* (PS3) and *P. venezuelensis* (PS4), are sporadic PCM agents, and cases have been found in Colombia and Venezuela, respectively. Occasional cases related to *P. restrepiensis* have been found outside Colombia, mainly in Brazil, Argentina, Peru, and Uruguay. *Paracoccidioides lutzii*, on the other hand, comprises a single species and is primarily distributed in the Midwest and Amazon regions of Brazil. A single *P. lutzii* strain was reported from Ecuador (Fig. 11A).

In Brazil, an essential difference in the geographical incidence of each phylogenetic species was noted (Fig. 11B). The south-east region corresponds to the majority of PCM agents and presents the highest levels of diversity (Simpson Index = 0.772; Shannon Index = 0.721), with all species being reported. In the central-west region, PCM cases are mainly due to *P. lutzii*, followed by *P. brasiliensis* s. str. and *P. americana* (Simpson Index = 0.576; Shannon Index = 1.124). *Paracoccidioides americana* was the principal agent in the south region, followed by *P. brasiliensis* s. str. and *P. restrepiensis* (Simpson Index = 0.464; Shannon Index = 1.157). The lowest index of diversity was found for the North region (Simpson Index = 0.500; Shannon Index = 0.811), albeit only six isolates were recovered from this region. We did not detect species diversity in Northeast Brazil, with only six isolates characterized as *P. brasiliensis* s. str. (Fig. 11B).

Table 5. Distribution of mating type alleles determined by PCR with mating-type allele-specific primers in *Paracoccidioides* isolates.

Species	No. of isolates	No. of isolates by mating-type		Chi-square value	P-value
		<i>MAT 1-1</i> (%)	<i>MAT 1-2</i> (%)		
<i>P. brasiliensis</i> (S1)	79	35 (44.30)	44 (55.69)	1.025	0.3113
<i>P. americana</i> (PS2)	22	18 (81.81)	4 (18.18)	8.909	0.0028
<i>P. restrepiensis</i> (PS3)	14	11 (78.57)	3 (21.42)	4.571	0.0325
<i>P. venezuelensis</i> (PS4)	13	5 (38.46)	8 (61.53)	0.692	0.4054
<i>P. brasiliensis</i> complex	128	69 (53.90)	59 (46.09)	0.781	0.3768
<i>P. lutzii</i>	37	19 (51.35)	18 (48.64)	0.027	0.8694
Overall	165	88 (53.33)	77 (46.66)	0.733	0.3918

P. brasiliensis complex = S1, PS2, PS3, and PS4.

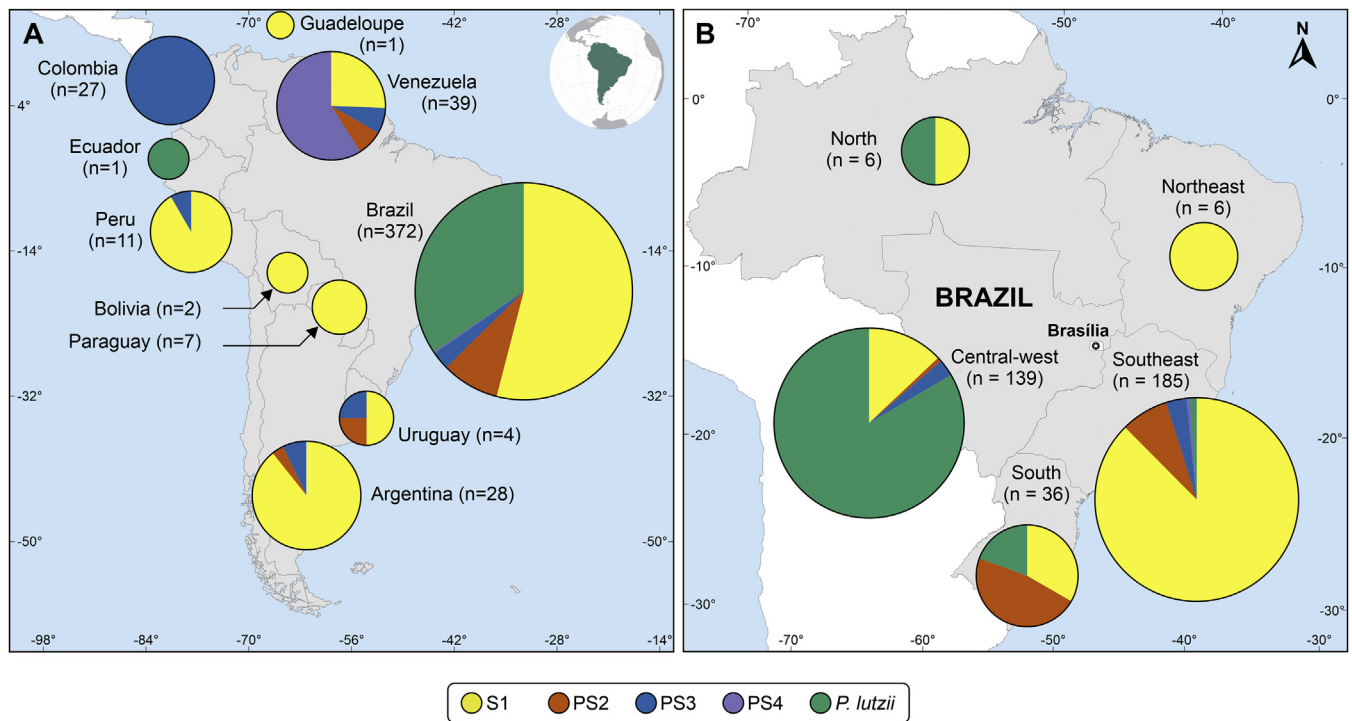


Fig. 11. Distribution patterns of 498 *Paracoccidioides* spp. isolates based on molecular characterization. **(A)** Distribution patterns observed in South America. **(B)** Distribution patterns observed in Brazil ($n = 372$). The sizes of circumferences are roughly proportional to the number of strains included. Codes reported within the pies denote genetic groups. Further information about isolate sources can be found in [Supplementary Table S1](#).

DISCUSSION

We here present the broadest population genetic study of *Paracoccidioides* species to date using isolates recovered from across a vast area of South America. Two sets of highly discriminatory AFLP markers were developed and shown to be a promising tool to dissect both deep and fine-scale genetic structures. The typing method proposed here combines robustness, reproducibility, high discriminatory power, and affordability, which is desirable for an important neglected mycosis such as PCM that is usually associated with poverty.

Pathogens with higher genetic diversity, significant effective population size, a mixed reproduction system, and great mutation rates are assumed to possess the highest evolutionary potential ([Nath et al. 2013](#)). Therefore, information regarding the current *Paracoccidioides* population and its evolutionary potential helps make informed disease-control strategies to mitigate PCM. Our AFLP technique demonstrated polymorphisms among closely related *Paracoccidioides*, which may contribute to resolve local epidemiological patterns as well as broader changes within populations over time and in response to selection pressures imposed by the environment and host resistance ([McDonald & Linde 2002](#)).

The availability of complete genome sequences for *Paracoccidioides* allowed us to predict the DNA fragments that AFLP would generate ([Desjardins et al. 2011](#), [Muñoz et al. 2014, 2016](#), [Teixeira et al. 2020](#)). Here, we demonstrated the best combination of restriction enzymes (EcoRI and MseI) by modelling their performance *in silico* for each species. Our analysis showed that the fragments observed following AFLP with EcoRI-AC / MseI-CT or EcoRI-AT / MseI-CT primers represent the optimal combinations to explore genetic diversity in *Paracoccidioides*. A similar *in silico* framework has been successfully applied for medically relevant *Sporothrix* species ([de](#)

[Carvalho et al. 2020, 2021a](#)), supporting that combining bioinformatics tools and whole-genome sequences can make the AFLP method more predictable instead of, rather, using random combinations of suboptimal endonuclease-combinations ([Rombauts et al. 2003](#), [Paris et al. 2010](#)).

Our AFLP dendrograms for *Paracoccidioides* species complex are compatible with the evolutionary history of the etiological agents of PCM, based on multilocus sequencing of proteins-encoding genes such as *ARF*, *GP43*, *TUB1*, and intein *PRP8*, or phylogenomic analyses ([Morais et al. 2000](#), [Theodoro et al. 2008, 2012](#), [Turissini et al. 2017](#), [Teixeira et al. 2020](#)). Convergence between fingerprints and genomic methods has already been demonstrated for *Candida auris* using AFLP ([Schelenz et al. 2016](#)), short tandem repeat typing ([de Groot et al. 2020](#)), and whole-genome sequencing ([Lockhart et al. 2017](#)). Phylogenetic studies of *Paracoccidioides* suggest that *P. brasiliensis* s. str., *P. americana*, *P. restrepiensis*, and *P. venezuelensis* are closely related taxonomic entities ([Muñoz et al. 2016](#), [Turissini et al. 2017](#)), and this clustering profile was recognized in our AFLP dendrograms.

Our dendrograms were more congruent than expected by chance, supported by the I_{cong} value and a positive Pearson correlation, confirming that different markers reveal congruent evolutionary histories. In each case, *P. lutzii* is basal to members of the *P. brasiliensis* complex, and our AFLP data indicates that the *P. brasiliensis* complex members all share a more recent common ancestor with each other than they do with *P. lutzii*. *P. americana*, *P. restrepiensis*, *P. venezuelensis* and *P. brasiliensis* s. str. remained as sister species, as previously reported ([Theodoro et al. 2012](#), [Muñoz et al. 2016](#), [Turissini et al. 2017](#), [Teixeira et al. 2020](#)).

From the fragment's profiles observed, *P. americana*, *P. restrepiensis*, and *P. venezuelensis* reveal more invariant fragments than *P. brasiliensis* s. str., suggesting a more recent

differentiation and a monophyletic origin of these lineages. Nearly all *P. restrepiensis* and *P. venezuelensis* occur within Colombia and Venezuela, respectively, suggesting that they evolved in these regions. Clusters of strains that presented many invariant fragments were mainly collected at a proximate geographic distance from each other. This finding suggests that vectors of dispersal for *Paracoccidioides* species are slow, leading to detectable regional diversification. Moreover, it may indicate a founder effect, the species being the most recently emerged taxon in *Paracoccidioides*, like the patterns found in *Fonsecaea* species (Najafzadeh *et al.* 2011). In this scenario, cases reported outside these areas may be regarded as imported cases. Contrasting to the above species, *P. brasiliensis* s. str. is by far the most diverse taxon in our dataset. Coding and non-coding nuclear markers also support the reciprocal monophyly in members of the *P. brasiliensis* complex (Turissini *et al.* 2017). These observations match their close arrangement in the PCAs and MDSs plots. MSTs and Neighbor-Net also capture phylogenetic proximity, an association further supported by our Kohonen maps (SOMs).

Currently, *P. lutzii* is described as a new biological species (Teixeira *et al.* 2014c), mainly due to the geographic, antigenic, and genetic differences when compared to the cryptic species of the *P. brasiliensis* complex (S1, PS2, PS3, and PS4) (Rodrigues *et al.* 2020b). Studies of evolutionary history suggest that *P. lutzii* diverged from *P. brasiliensis* around 22.5 million years ago (Muñoz *et al.* 2016); however, divergence times between *Paracoccidioides* species pairs range between 0.03 and 33 million years (Teixeira *et al.* 2020). This genetic distance between *P. lutzii* and the four members of the *P. brasiliensis* complex observed in molecular phylogeny studies was also reflected in our AFLP dendrograms and Neighbor-Net analysis.

The results of our study indicate no or very limited genetic introgression between *P. brasiliensis* complex and *P. lutzii* in South America. The assessment of the genetic structure based on two sets of AFLP markers indicate the coexistence of two genetic clusters with no or minimal admixture. This agrees with the results observed by Teixeira *et al.* which suggests that there is a signature of introgression in only one species pair out of ten possible pairs in *Paracoccidioides* (Teixeira *et al.* 2020). This scenario was confirmed using whole-genome sequencing and structure analysis (Muñoz *et al.* 2016). Further confirmation for this consideration is given by: (1) genetic diversity criterion assessed for the studied *Paracoccidioides* populations. Although they are not measures of genetic variation, they may indicate a genetic distinctiveness between the *P. brasiliensis* complex and *P. lutzii*. Indeed, PCA, MSTs, and SOMs performed for *P. brasiliensis* complex and *P. lutzii* combined clearly indicated different genetic clusters; (2) significant genetic differentiation (PhiPT) between *P. brasiliensis* complex and *P. lutzii*; and (3) slight genetic differentiation among isolates embedded in the *P. brasiliensis* complex in Neighbor-Net analysis.

We found high diversity in the *P. brasiliensis* s. str., suggesting that this lineage has high fitness favouring its dispersion, allowing the survival and adaptation to varied geographic conditions throughout Latin America. Likewise, the epicentre of occurrence for *P. lutzii* lies in the Mato Grosso state, an area characterized by the biogeographic formations of the Cerrado savannas, Pantanal, and the Amazon rainforest (Simoes *et al.* 2020). These biomes may have contributed to the geographic isolation and population structure in *Paracoccidioides*. Currently, with less than 50 % of the native vegetation cover remaining, the deforestation

of the Cerrado surpass those in Amazonia (Grecchi *et al.* 2014, Beuchle *et al.* 2015, Espírito-Santo *et al.* 2016), and along with the occupation of the Cerrado lands for mechanized agricultural production may lead to the emergence and expansion of the area of occurrence of *P. lutzii*. Indeed, *Paracoccidioides* species propagules inhabit a complex environment in the soil with several amoeboid predators that can impose selective pressure, selecting for virulence traits (Albuquerque *et al.* 2019). A hypothesis has been raised in recent years whereby biodiversity loss may increase pathogen transmission and disease incidence, especially if it reduces predation and competition on reservoir hosts, thereby increasing their density (Keesing *et al.* 2010).

Our data shows that *Paracoccidioides* is a heterothallic fungus with a single mating-type locus that produces two alleles, *MAT1-1* and *MAT1-2*, in agreement with a previous report (Torres *et al.* 2010). The initial stages of a sexual cycle in *Paracoccidioides* have been observed under laboratory conditions (Torres *et al.* 2010, Teixeira *et al.* 2013), and along with the recombination events reported in genomic studies (Muñoz *et al.* 2016) could support the hypothesis of a sexual cycle leading to diversification in these pathogens (Teixeira *et al.* 2013). After evolutionary divergence, genetic hybridization may be mainly observed when (i) the ranges of closely related species overlap, or (ii) one species is uncommon, and individuals have to find mates from a closely related species. In the first scenario, this can lead to two species being genetically more related when in parapatry or sympatry than in regions where they are in allopatry (Palme *et al.* 2004, Behm *et al.* 2010, McKinnon *et al.* 2010). On the other hand, asymmetric introgression may occur when one species exists at a low density (Choleva *et al.* 2014). Unbalanced gene flow may also be affected by sex-biased dispersal or philopatry. Therefore, it is tempting to hypothesize that this phenomenon could orchestrate genetic hybridization among members of the *P. brasiliensis* complex, as our results reveal a mixed-mode of reproduction in *Paracoccidioides* that occurs in sympatry. A mating-type idiomorph-biased distribution was not found to be a significant feature in *P. brasiliensis* s. str., *P. venezuelensis*, and *P. lutzii*, but in *P. americana* and *P. restrepiensis*. Skewed *MAT* loci distribution could result from the scarcity of sexual reproduction, strong selection for pleiotropic effects of a mating-type allele (Nieuwenhuis & James 2016), or even a phenomenon of small populations (Valero *et al.* 2018). This paradoxical reproduction system has been observed in *Sporothrix* (Teixeira *et al.* 2015, de Carvalho *et al.* 2021b), *Histoplasma* (Rodrigues *et al.* 2020a), and *Cryptococcus* (Nielsen *et al.* 2005), with species prevalently clonal along with recombinant molecular siblings coexisting in the same geographical range.

Geographical trends observed for *P. brasiliensis* s. str. revealed a widely distributed species throughout Latin American, present in Argentina, Brazil, Bolivia, Guadeloupe Island, Paraguay, Peru, Uruguay, and Venezuela. Our results corroborate the distribution reported previously by other authors (Matute *et al.* 2006, Teixeira *et al.* 2009, Theodoro *et al.* 2012), including Bolivia and Guadeloupe Island areas for lineage S1. *P. americana* geographical distribution agrees to the countries described in the literature for this genetic group, i.e., Argentina, Brazil, Uruguay, and Venezuela (Theodoro *et al.* 2012, Roberto *et al.* 2016).

Initially, it was thought that *P. restrepiensis* was restricted to Colombia, but this phylogenetic species has already been found in Brazil and Venezuela (Roberto *et al.* 2016, Cocio *et al.* 2020a, Mattos *et al.* 2021). Although most isolates originated from Colombia, we found a single isolate occurring in Uruguay.

P. venezuelensis was the last group recognized in the *P. brasiliensis* complex, and until recently, it was thought to be exclusive to Venezuela (Teixeira et al. 2014b, Turissini et al. 2017). However, phylogenetic analysis detected one strain from São Paulo, Brazil, characterizing a new location for this species.

Finally, we confirm that *P. lutzii* has its epicentre in Central-west Brazil with an overwhelming occurrence in Mato Grosso state (Gegembauer et al. 2014, Hahn et al. 2014, Teixeira et al. 2014b, Hahn et al. 2019, Rodrigues et al. 2020b). The distribution patterns in *Paracoccidioides* species have a notable impact on the serological diagnosis of PCM (Rodrigues et al. 2020b). Therefore, our data draws attention to the urgent need to expand the offer of serological diagnostic tests using antigenic preparations from *P. lutzii* (Gegembauer et al. 2014, Queiroz Junior et al. 2014, Maifrede et al. 2021) or the availability of PCR assays for the detection of *P. lutzii* DNA (Pinheiro et al. 2021), which is occurring in areas beyond the known endemic range in Brazil.

CONCLUSION

Our study illustrates the need to improve genetic surveillance in endemic areas for *Paracoccidioides* species to ensure that the results of molecular epidemiological studies are accurate. AFLP analysis identifies *P. brasiliensis* s. str. and *P. lutzii* as the most diverse species in the genus. In contrast, markedly low genetic diversity was noted for *P. americana*, *P. restrepiensis*, and *P. venezuelensis*. This straightforward typing method will enable the cost-effective analysis of more *Paracoccidioides* isolates to improve our understanding of the eco-epidemiology trends in PCM infections, help progress toward a consensus taxonomy, assess species boundaries, and explore the significance of genetic diversity in *Paracoccidioides* species in the clinical scenario.

ACKNOWLEDGMENTS

The authors acknowledge the financial support granted by the São Paulo Research Foundation (FAPESP 2017/27265-5 and FAPESP 2018/21460-3), National Council for Scientific and Technological Development (CNPq 429594/2018-6), and Coordination for the Improvement of Higher Education Personnel (CAPES 88887.177846/2018-00). TNR was a fellow of FAPESP (2013/05405-9). AMR is a CNPq Research Productivity Fellow (CNPq 304902/2020-9). The funders had no role in the study design, data collection, and analysis, decision to publish, or preparation of the manuscript. MAB was funded by the Wellcome Trust (#206194). MCF was funded by the UK Medical Research Council and Wellcome Trust and is a Fellow in the CIFAR 'Fungal Kingdoms' program. For the purpose of Open Access, the author has applied a CC-BY public copyright license to any Author Accepted Manuscript version arising from this submission.

APPENDIX A. SUPPLEMENTARY DATA

Supplementary data to this article can be found online at <https://doi.org/10.1016/j.simyco.2021.100131>.

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