BRIEF REPORT

REVISED Xylariales: First results of mycological exploration in the Sangay and Llanganates National Parks, Ecuador [version 2; peer review: 1 approved, 2 approved with reservations]

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Abstract

Fungal samples were collected in the Sangay (SP) and Llanganates (LP) National Parks in Ecuador. Sequences of the internal transcribed spacer regions (ITS1-5.8S-ITS2) of the ribosomal DNA of the samples were analyzed. Taxonomic identification of fungi of the order Xylariales was done through phylogenetic analysis using a Maximum Likelihood method. All analyzed collections presented here belong to the genus Xylaria, of these eight belong to PL and two to SP. Four samples were not identified at the species level, suggesting it could be a new species. This data contributes with base information on the biodiversity of the Parks, necessary to design and implement measures for the conservation of fungi in Ecuador.

Keywords

Diversity, ITS, Llanganates, National Park, Sangay, Xylarial

This article is included in the **Ecology and Global Change** gateway.

Open Peer Review

Approval Status

	1	2	3
version 2			
(revision)			
17 May 2018	view	view	view
version 1			
23 Feb 2018	view	view	

1. **Francisco J. Flores** , Universidad de las Fuerzas Armadas-ESPE, Sangolquí, Ecuador
2. **D. Jean Lodge**, USDA Forest Service, Luquillo, USA
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Any reports and responses or comments on the article can be found at the end of the article.

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Author roles: **Guevara MF:** Conceptualization, Investigation, Methodology, Writing – Original Draft Preparation; **Mátyás B:** Project Administration, Writing – Review & Editing; **Ordoñez ME:** Conceptualization, Investigation, Methodology, Project Administration, Writing – Original Draft Preparation

Competing interests: No competing interests were disclosed.

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REVISED Amendments from Version 1

According to the referee's suggestions we have made the following changes in the manuscript:

- The abstract, introduction and methods were re-written for ease of comprehension.
- The results are explained objectively
- The table and figure legends are also modified for ease of comprehension of the content.
- In addition, we rephrased 2 sentences in Conclusions.
- We have removed one of the authors from the second version of the article as an error was made when she was included in the moment of submitting the first version of the article. We were working on several articles at the same time and because of confusion, her name was included in this article. Paula Salazar is a valuable person for the Universidad Politécnica Salesiana and she has collaborated in several projects, but unfortunately she did not collaborate in this research done in association with Pontificia Universidad Católica.

See referee reports

Introduction

Sangay (SP) and Llanganates (LP) National Parks in Ecuador are considered as high priority conservation units in the Tropical Andes, due to their high biodiversity and high endemism^{1,2}. However, their mycological diversity is still unknown. This

study aims to contribute to the conservation of fungi, showing the results of their diversity, based on molecular taxonomy, by analyzing the ITS (internal transcribed spacer) regions. ITS is the accepted as primary fungal barcode marker for fungi^{3,4}. For this, the DNA sequence of specimens of exploratory fungal collections were analyzed within the aforementioned parks. Here we present results exclusively for the Xylariales order, other fungal orders were also collected but are not shown here.

Methods

Sequencing and molecular identification

Sample collection was carried out during the months of January and February 2015. The fruiting bodies collected were deposited in the QCAM Fungarium (Catholic University Mycology Collection, Quito). **Table 1** displays the collection codes, as stored at the QCAM. The ITS1-5.8SITS2 region was amplified by PCR with primers ITS1F (5'-CTTGGTCATTTAGAGGAAGTAA-3')⁵ and ITS4 (5'-TCCTCCGCTTATTGATATGC-3')⁶. The amplified fragments were sent for sequencing to Macrogen Inc. (Seoul, South Korea). The forward and reverse sequences obtained for each isolate were assembled in Geneious R8 (Biomatter Ltd. 2005–2012), and submitted to GenBank. Sequence data were analyzed by comparison with sequences available in GenBank. An assignment to the lower taxonomic level was made by direct homology of the consensus sequences with the search results in BLASTn (NCBI) optimized for highly similar sequences (megablast). Alignments that presented 100% coverage and at least 99% identity with a sequence

Table 1. Fungi of the order Xylariales collected in the Sangay and Llanganates National Parks in Ecuador. *SNPs: Single nucleotide polymorphisms found between the sample analyzed and the closest hit on the BLASTn search.

QCAM Fungarium Code	GenBank Accession Number	Identification	Sampling location	
			National Park	Altitude (m)
QCAM4663	MG768840	Xylaria enterogena	Llangantes	1370
QCAM4551	MG768839	Xylaria enterogena	Llangantes	1387
QCAM4537	MG768834	Xylaria fissilis	Llangantes	1377
QCAM4540	MG768836	Xylaria schweinitzii	Llangantes	1377
QCAM4232A	MG768832	Xylaria telfairii	Sangay	2885
QCAM4550	MG768838	Xylaria sp. 1	Llangantes	1387
QCAM4666	MG768841	Xylaria sp. 1	Llangantes	1379
QCAM4306A	MG768833	Xylaria sp. 2	Sangay	2885
QCAM4545	MG768837	Xylaria sp. 3	Llangantes	1373
QCAM4539	MG768835	Xylaria sp. 4	Llangantes	1377

previously reported in GenBank were considered. The results were compared with the previously made morphological identification at the QCAM Fungarium, to check the taxonomic designation.

Phylogenetic analysis

Sequence data were aligned with Geneious R8 and later manually adjusted with Mesquite version 3.04⁷. Public sequences available in GenBank that corresponded to specimens that gave the greatest homology in BLASTn with the sequences of the collected specimens were included. Phylogenetic trees were constructed in Geneious R8 using the PhyML⁸ plugin for Maximum Likelihood (ML) with a custom substitution model (010230), determined by jModelTest 2.1.4.^{9,10}, according to Corrected Akaike Information Criterion (AICc)^{11,12}. A bootstrap of 1000 replica was used.

Results

All the specimens analyzed were of the genus *Xylaria*. The eight specimens from Llanganates National Park were identified as *X. enterogena*, *X. fissilis*, *X. schweinitzii*, *X. telfairii* and three unidentified species. For the two samples from Sangay National Park, one was *X. telfairii* and the other was an unidentified species. Differences in the number of samples found at each park could be due to the sampling effort, and not necessarily to the richness of the Xylariales in the Parks. The unidentified species were different in each park. The analysis shows that there are no shared species of *Xylaria* at the two sampled sites (Table 1), this

is important for conservation decision making. The phylogenetic relationships recovered from the analysis of the ITS sequences (Figure 1) shows two major groups. The first major group, composed by clades A and B, is well supported, it includes specimens from LP and PS. Clade A includes all *X. enterogena* specimens and Clade B includes all *X. telfairii* specimens, and *Xylaria* sp.1 specimens. It is possible that *Xylaria* sp.1 might belong to the *X. telfairii* group, but due to the differences among the sequences it is likely a different species. In the second major group, clade C is sister to clades D, E and F. This major group includes specimens from both LP and SP. Clade C includes all *X. schweinitzii* specimens. Clade D includes *Xylaria* sp. 2. The closest sequence to *Xylaria* sp. 2 from SP was a previously reported collection also from Ecuador¹³ in a cloud forest in the province of Imbabura, that was also identified only at the genus level. Clade E shows *Xylaria* sp. 3, the closest sequence to this individual belongs to the same previously reported study¹³, identified only at the genus level. Clade F includes *Xylaria* fissilis sequences from LP and one from ¹³. Clade F also includes *Xylaria* sp. 4, an unidentified specimen. The number of nucleotide differences (SNPs) between the sample and the closest hit on the BLASTn search suggests that these specimens may belong to new species (Table 1). Additional loci and more detailed morphological analyses are needed to determine this. The genus *Xylaria* is probably the largest in the family Xylariaceae, with 35 estimated genera¹⁴, but the real number remains unknown¹⁵. Studies related to the biological diversity of this order in the

Figure 1. Maximum likelihood phylogenetic tree of specimens of the Xylariales order obtained in Sangay and Llangantes National Parks, based on sequences of the ITS (internal transcribed spacer) region.

National Parks of Ecuador are scarce, more systematic field studies would surely reveal a greater diversity of families, genera and species within the Xylariales in SP and LP, as well as other regions and protected areas of Ecuador, especially if we take into account the cosmopolitan distribution of *Xylaria*¹³. In fact, new fungal species in SP, belonging to the Agaricales, have recently been described¹⁶.

These entire unidentified specimens might represent new species. Additional loci and more detailed morphological analyses are needed to determine this. The genus *Xylaria* is probably the largest in the family Xylariaceae, with 35 estimated genera¹³, but the real number remains unknown¹⁵. Studies in relation to the biological diversity of this order in the National Parks of Ecuador are scarce, more systematic field studies would surely reveal a greater diversity of families, genera and species within the Xylariales in SP and LP, as well as other regions and protected areas of Ecuador, especially if we take into account the cosmopolitan distribution of *Xylaria*¹³. In fact, new fungal species in PS, belonging to the Agaricales, have recently been described¹⁶.

Conclusions

The results obtained allow us to establish a baseline for the biological diversity of the Xylariales in SP and LP, an important step to the conservation of fungi. This is the main contribution of this study. We found four species of *Xylaria*: *X. enterogena*, *X. telfairii*, *X. schweinitzii*, and *X. fissilis*, and four potential new species based on ITS sequences divergence; the species found in LP are different from those found in SP. However, there is much more to discover. A huge and complex task is pending. To advance our understanding of the Kingdom Fungi we must start by deciphering the diversity of species present in these sites.

Data availability

The sequencing data are available on the NCBI Genbank webpage: *Xylaria enterogena*:

<https://www.ncbi.nlm.nih.gov/nuccore/MG768840>

<https://www.ncbi.nlm.nih.gov/nuccore/MG768839>

Xylaria fissilis: <https://www.ncbi.nlm.nih.gov/nuccore/MG768834>

Xylaria schweinitzii: <https://www.ncbi.nlm.nih.gov/nuccore/MG768836>

Xylaria telfairii: <https://www.ncbi.nlm.nih.gov/nuccore/MG768832>

Xylaria sp. 1: <https://www.ncbi.nlm.nih.gov/nuccore/MG768838>

<https://www.ncbi.nlm.nih.gov/nuccore/MG768841>

Xylaria sp. 2: <https://www.ncbi.nlm.nih.gov/nuccore/MG768833>

Xylaria sp. 3: <https://www.ncbi.nlm.nih.gov/nuccore/MG768837>

Xylaria sp. 4: <https://www.ncbi.nlm.nih.gov/nuccore/MG768835>

Competing interests

No competing interests were disclosed.

Grant information

The Secretaria de Educación Superior, Ciencia, Tecnología e Innovación del Ecuador (SENESCYT), Arca de Noé Initiative and the QCAM Fungarium from “Pontificia Universidad Católica del Ecuador” supported the project.

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Current Peer Review Status: ? ? ✓

Version 2

Reviewer Report 16 July 2018

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Juan Enrique Tacoronte Morales

Central University of Ecuador, Quito, Ecuador

I have read this submission proposal and all related information reported in (introduction, methodology, results, discussion and cited references). Considering my expertise in secondary metabolites and bioactive natural products isolated from invertebrates (defensive secretions and applied chemical ecology) and fungi (volatile organic compounds) I believe that I could analyze the reported information data with an appropriate, conceptual and methodological level to confirm that this report devoted to mycological exploration of Ascomycota-Xylariales in tropical Andean region has a quite recognized scientific standards for being published in *F1000Research*.

In this context, please, I have some considerations regarding the submitted article:

- it could be of great significant and relevance to evaluate some biogeochemical parameters of eco-geographical collecting zones for potential determining the zonal distribution of Xylariales in these areas, including any potential correlation with tree diversity.

- It would have been highly didactic, and recommended, the inclusion, in the proposal submitted, of some photographs in real time, taken during the collection phase in the national parks, and some, comparative ones, taken during the preparation phase in the lab (for morphological purposes only!)

Competing Interests: No competing interests were disclosed.

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard.

Reviewer Report 29 May 2018

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Francisco J. Flores

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There are still several issues that need to be addressed before indexing. Please see the file in the provided link for specifics:

[Click here for annotated PDF](#)

Competing Interests: No competing interests were disclosed.

Reviewer Expertise: Phylogenetics

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard, however I have significant reservations, as outlined above.

Reviewer Report 24 May 2018

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D. Jean Lodge

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Abstract:

4th line from top - change 'done' to 'inferred' because you can't be sure of an identification based on ITS blast.

3rd line from bottom, change sentence beginning "This data" to These data provide baseline information"

The Xylaria sp. 3 is still listed as unidentified. The molecular phylogeny indicates this is likely X. curta. X. curta is expected from that area and has been collected before in that part of Ecuador. It is easily identified in photographs, with confirmation by examining spores and ascus plugs under a microscope. It is not clear why this point was not addressed in revisions, though it would require

renumbering of the unidentified species in tables and figures and text. Also, *Xylaria* sp. 4 appears to be *X. fissilis* based on the phylogeny.

The abstract says there were 4 unidentified *Xylaria* species, but if sp. 3 is *X. curta* and sp. 4 is *X. fissilis*, there are only two suspected undescribed species - sp. 1 and sp. 2.

The conclusions say there were four unnamed species also, but *X. fissilis* is among the named species based on one collection, but sp. 4 has an identical sequence to the named one, so that is also *X. fissilis* and not a new species.

The manuscript should not be indexed until the authors reduce the number of undescribed species from 4 to 2 and make the corresponding changes to the text throughout.

Competing Interests: No competing interests were disclosed.

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard, however I have significant reservations, as outlined above.

Version 1

Reviewer Report 19 April 2018

<https://doi.org/10.5256/f1000research.14800.r32603>

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D. Jean Lodge

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Introduction. Because the aim is to compare these two parks in Ecuador, it is important that the elevation ranges of each are included, as well as the distance between the two parks.

Methods, p. 3, last line seems to be an incomplete sentence. It ends with 'BLASTn' (NCBI) optimized for highly similar sequences (megablast), alignments that presented 100'
It is probably meant to say 'overlapped 100%.'

Results: Text for Clade E does not match the phylogram. The text says that sp. 3 was related to another collection identified only at the genus level. However, the phylogram shows a well-supported clade (98% BS) comprised of sp. 3 and *X. curta* KP133352 ECU. I would expect *X. curta* to be reasonably common there. There is either an error in identification of KP133352 ECU or the text

in results needs to be changed, and the abstract and first line of results and conclusions also need to be changed to include *X. curta*.

That also means that the unidentified species need to be renumbered in the text, table, and phylogram, with sp. 4 becoming sp. 3.

There are likely 3 undescribed species included in this study. The journal allows for photographs, and photos should be added to the manuscript. Even if photos were not taken of the fresh specimens, photos of the dried specimens can be used, and are very helpful. If photos of the asci, especially the ascus plug stained with iodine/Melzer's reagent, and photos of the ascospores showing the germ slits, that would be very helpful also for future work. The ascospore length and width ranges, the shape and extent of the germ slit, and the shape and size of the ascus plug would be helpful additions.

Is the work clearly and accurately presented and does it cite the current literature?

Partly

Is the study design appropriate and is the work technically sound?

Yes

Are sufficient details of methods and analysis provided to allow replication by others?

Yes

If applicable, is the statistical analysis and its interpretation appropriate?

Not applicable

Are all the source data underlying the results available to ensure full reproducibility?

Yes

Are the conclusions drawn adequately supported by the results?

Yes

Competing Interests: No competing interests were disclosed.

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard, however I have significant reservations, as outlined above.

Reviewer Report 01 March 2018

<https://doi.org/10.5256/f1000research.14800.r31232>

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The article "Xylariales: First results of mycological exploration in the Sangay and Llanganates National Park, Ecuador " provides internal transcribed spacer sequences from *Xylaria* isolates collected from Ecuadorian national parks. Information about fungal diversity in Ecuador is still scarce, which makes this manuscript relevant. Nevertheless, I have several concerns, mainly:

- Phylogenetic analysis using only the ITS sequence is very superficial, and, most likely, it won't show an accurate representation of the real evolutionary relationships between isolates.
- There is no mention about the morphological characteristics of isolates. Is the morphology congruent with the genotype?

The manuscript needs major revision to improve clarity.

Several suggestions are made in the file linked below.

https://f1000researchdata.s3.amazonaws.com/linked/196501.Bence_Matyas_rev.pdf

Is the work clearly and accurately presented and does it cite the current literature?

Partly

Is the study design appropriate and is the work technically sound?

Partly

Are sufficient details of methods and analysis provided to allow replication by others?

Yes

If applicable, is the statistical analysis and its interpretation appropriate?

Partly

Are all the source data underlying the results available to ensure full reproducibility?

Yes

Are the conclusions drawn adequately supported by the results?

Partly

Competing Interests: No competing interests were disclosed.

Reviewer Expertise: Phylogenetics

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard, however I have significant reservations, as outlined above.

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